BEFORE THE

INDEPENDENT CITIZENS' OVERSIGHT COMMITTEE TO THE CALIFORNIA INSTITUTE FOR REGENERATIVE MEDICINE ORGANIZED PURSUANT TO THE CALIFORNIA STEM CELL RESEARCH AND CURES ACT

REGULAR MEETING

LOCATION: SHERATON GATEWAY LOS ANGELES HOTEL

6101 WEST CENTURY BOULEVARD

LOS ANGELES, CALIFORNIA

DATE: DECEMBER 17, 2015

9 A.M.

REPORTER: BETH C. DRAIN, CSR

CSR. NO. 7152

BRS FILE NO.: 98031

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1	LOS ANGELES, CALIFORNIA
2	THURSDAY, DECEMBER 17, 2015; 9 A.M.
3	
4	CHAIRMAN THOMAS: GOOD MORNING, EVERYBODY,
5	FROM THE SHERATON LAX. WELCOME TO THE DECEMBER
6	BOARD MEETING FOR CIRM. IT HAS BEEN A NUMBER OF
7	MONTHS SINCE WE HAD AN IN-PERSON MEETING. WE'VE
8	HAD, AS YOU KNOW, MONTHLY MEETINGS TELEPHONICALLY,
9	BUT IT'S GREAT TO SEE EVERYBODY AGAIN AND WELCOME.
10	WE COULD START. MARIA, COULD YOU LEAD US IN THE
11	PLEDGE OF ALLEGIANCE.
12	(THE PLEDGE OF ALLEGIANCE.)
13	CHAIRMAN THOMAS: MARIA, COULD YOU PLEASE
14	CALL THE ROLL.
15	MS. BONNEVILLE: DAVID BRENNER. LINDA
16	BOXER.
17	DR. BOXER: PRESENT.
18	MS. BONNEVILLE: KEN BURTIS. ANNE-MARIE
19	DULIEGE.
20	DR. DULIEGE: YES.
21	MS. BONNEVILLE: MICHAEL FRIEDMAN.
22	DR. FRIEDMAN: HERE.
23	MS. BONNEVILLE: JUDY GASSON.
24	DR. GASSON: HERE.
25	MS. BONNEVILLE: SAM HAWGOOD. DAVID
	, a
	4

1	HIGGINS.	
2		DR. HIGGINS: HERE.
3		MS. BONNEVILLE: STEPHEN JUELSGAARD.
4		MR. JUELSGAARD: HERE.
5		MS. BONNEVILLE: SHERRY LANSING.
6		MS. LANSING: HERE.
7		MS. BONNEVILLE: KATHY LAPORTE.
8		DR. LAPORTE: HERE.
9		MS. BONNEVILLE: BERT LUBIN. SHLOMO
10	MELMED.	
11		DR. MELMED: HERE.
12		MS. BONNEVILLE: LAUREN MILLER.
13		MS. MILLER: HERE.
14		MS. BONNEVILLE: ADRIANA PADILLA.
15		DR. PADILLA: HERE.
16		MS. BONNEVILLE: JOE PANETTA.
17		MR. PANETTA: HERE.
18		MS. BONNEVILLE: ROBERT PRICE.
19		DR. PRICE: HERE.
20		MS. BONNEVILLE: FRANCISCO PRIETO.
21		DR. PRIETO: HERE.
22		MS. BONNEVILLE: CARMEN PULIAFITO.
23		DR. PULIAFITO: PRESENT.
24		MS. BONNEVILLE: ROBERT QUINT. AL
25	ROWLETT.	
		5

160 S. OLD SPRINGS ROAD, SUITE 270, ANAHEIM, CALIFORNIA 92808 1-800-622-6092 1-714-444-4100 EMAIL: DEPO@DEPO1.COM

1	MR. ROWLETT: HERE.
2	MS. BONNEVILLE: JEFF SHEEHY.
3	MR. SHEEHY: HERE.
4	MS. BONNEVILLE: OSWALD STEWARD. JONATHAN
5	THOMAS.
6	CHAIRMAN THOMAS: HERE.
7	MS. BONNEVILLE: ART TORRES.
8	MR. TORRES: HERE.
9	MS. BONNEVILLE: KRISTINA VUORI.
10	DR. VUORI: HERE.
11	MS. BONNEVILLE: DIANE WINOKUR.
12	CHAIRMAN THOMAS: THANK YOU VERY MUCH.
13	WE'LL NOW PROCEED TO THE CHAIR'S REPORT. SINCE WE
14	LAST MET IN PERSON, I'VE BEEN TO A NUMBER OF EVENTS
15	OF SOME INTEREST AND NOTE REPRESENTING CIRM, AND I'D
16	LIKE TO GIVE YOU A COUPLE THOUGHTS FROM THOSE IN NOT
17	EXACTLY CHRONOLOGICAL ORDER. WE'RE GOING TO START
18	WITH THE WORLD ALLIANCE FORUM, WHICH IS SOMETHING
19	THAT BRINGS TOGETHER PRINCIPALLY UNITED STATES AND
20	JAPANESE SCIENTISTS IN THE STEM CELL FIELD. IT WAS
21	HELD IN GOLDEN GATE PARK, A COUPLE-DAY AFFAIR.
22	WE HAD A MOST INTERESTING PANEL ON THE
23	SUBJECT, WHICH WAS SORT OF A TOUGH ONE, ENTITLED
24	"REGENERATIVE MEDICINE 2020 AND 2030." AND WE HAD
25	AMONGST OTHERS ON THE PANEL DR. SHINYA YAMANAKA,
	6
	U U

6

1	WHO, AS YOU KNOW, IS THE CREATOR OF THE IPS
2	TECHNOLOGY FOR WHICH HE RECEIVED THE NOBEL PRIZE AND
3	WHICH FORMS THE UNDERPINNINGS FOR A GREAT DEAL OF
4	EXCITING RESEARCH GOING ON IN THE FIELD.
5	IN CONNECTION WITH THAT PANEL, I DID A
6	LITTLE SAMPLING OF PEOPLE'S OPINIONS ON THE SUBJECT
7	OF REGENERATIVE MEDICINE 2020 AND 2030. AND I HAVE
8	A COUPLE RESPONSES HERE WHICH I THOUGHT YOU MIGHT
9	FIND INTERESTING FROM PEOPLE THAT I SPOKE TO. AND
10	IF YOU WILL BEAR WITH ME, I JUST WANT TO READ A
11	COUPLE OF E-MAILS. ONE IS FROM DOUG MELTON, WHO YOU
12	WILL RECOGNIZE AS ONE OF THE PREEMINENT STEM CELL
13	SCIENTISTS IN THE TYPE 1 DIABETES FIELD. AND SO
14	THIS WAS DOUG'S RESPONSE ON WHAT TO EXPECT IN THE
15	NEXT FIVE TO FIFTEEN YEARS.
16	"HI, JON. THIS SOUNDS LIKE A FUN PUZZLE,
17	AND YOU'RE KIND TO ASK MY OPINION. FACT IS I'VE NOT
18	GIVEN IT ENOUGH THOUGHT, BUT I IMAGINE WE'LL SEE A
19	DIFFERENT CONNECTION BETWEEN PATIENTS AND THEIR
20	HEALTHCARE PROVIDERS (HOSPITALS). I THINK HOSPITALS
21	AND DOCTORS WILL HAVE TO MOVE FROM TREATING PATIENTS
22	WHEN THEY ARE SICK TO AN EARLIER INTERACTION, VERY
23	EARLY, IN FACT. MAYBE A GOOD WAY TO DESCRIBE IT
24	WOULD BE TO REIMAGINE THE PATIENT'S MEDICAL HISTORY.
25	"AT PRESENT THAT HISTORY IS TAKEN ON
	7

1	ADMISSION IN MOST CASES. IMAGINE INSTEAD THAT
2	HOSPITALS FORM A LIFELONG RELATION WITH THE FAMILY,
3	NOT UNLIKE A CHURCH OR SYNAGOGUE, FROM BIRTH TO
4	DEATH. WHEN THE BABY IS BORN, DNA SEQUENCE IS
5	DETERMINED AND COMPARED TO THE PARENTS' DNA AS WELL
6	AS SOME CELLS TO MAKE IPS CELLS FOR LATER USE AND
7	ANALYSIS. MAKING PATIENT-SPECIFIC CELLS AND TISSUES
8	WILL BE EASIER BY THEN. SO, WHEN NEEDED, THEY'LL BE
9	AT HAND, AT HAND FOR CELL TRANSPLANTATION, FOR DRUG
10	TESTING PATIENT-SPECIFIC TOXICITY, AND
11	SUSCEPTIBILITY OR INCLINATION TO WHAT WE NOW CALL A
12	DISEASE.
13	"AS THE JUSTIFIABLE EXCITEMENT IN CANCER
14	IMMUNOTHERAPY SHOWS US NOW, WE SHOULD IMAGINE A TIME
15	WHEN WE MAKE A PATIENT'S THYMUS AND USE THAT TISSUE
16	IN CONJUNCTION WITH THEIR HEMOPOIETIC STEM CELLS TO
17	MAKE IMMUNE CELLS OF CHOICE. AMONG THE MANY USES
18	WOULD BE IMMUNE CELLS TO SPEED HEALING AND/OR
19	ELIMINATE INFECTIONS AND CANCER. THERE WILL BE A
20	NEW MEANING TO IMMUNIZATIONS. LOOKING A BIT FARTHER
21	AHEAD, I WON'T BE SURPRISED TO SEE THE PRODUCTION OF
22	MINI BRAINS, LITTLE CUBES OF NEURONS THAT CAN GIVE
23	CLUES ABOUT MENTAL HEALTH BASED ON THE PATIENT'S
24	PREDISPOSITIONS. IT WILL BE EXCITING TIMES."
25	THAT'S FROM DOUG MELTON.

8

1	CLOSER TO HOME, I ASKED OUR OWN DR. PAT
2	OLSON WHAT SHE THOUGHT ON THE SUBJECT. SHE WAS KIND
3	ENOUGH TO RESPOND. SO HERE IS PAT'S COMMENT.
4	"ADVANCES IN TECHNOLOGY MANUFACTURING AND
5	REGULATION WILL RESULT IN ON-DEMAND, PERSONALIZED
6	AUTOLOGOUS STEM CELL-DRIVEN REPLACEMENT THERAPIES TO
7	TREAT, FOR EXAMPLE, HEART FAILURE, LIVER, LUNG, AND
8	NEUROLOGIC DISEASE AND INJURY. SINGLE-CELL GENETIC
9	DISEASES WILL BE A THING OF THE PAST DUE TO GENE
10	CORRECTION IN STEM PROGENITOR CELLS. THIS WOULD
11	NEED TO BE DONE DURING EARLY DEVELOPMENT.
12	"THE BEST DRUG FOR YOU AND YOUR DISEASE
13	WILL BE DEVELOPED ON DEMAND USING AN ASSAY SYSTEM
14	DEVELOPED FROM YOUR REPROGRAMMED CELLS, A TAILORED
15	COMPOUND LIBRARY GENERATED FROM AN APP OF COMPOUNDS
16	SHOWN TO HAVE PROMISE FOR YOUR DISEASE AND 3D
17	PRINTED, AND ONCE THE BEST COMPOUND FOR YOU IS
18	IDENTIFIED, MANUFACTURED ON YOUR 3D PRINTER.
19	"BANKS WILL EXIST, INCLUDING CELL SAMPLES
20	FROM ALL NEWBORNS (WON'T HAVE THE MUTATION LOAD OF
21	ADULTS) FOR FUTURE REPROGRAMMED STEM PROGENITOR CELL
22	THERAPY DEVELOPMENT, SCREENING FOR PERSONALIZED
23	DRUGS, AND FOR ORGAN OR ORGANOID GENERATION FOR
24	ORGAN REPLACEMENT."
25	THANK YOU, PAT.

1	THAT'S A LOOK. I THOUGHT YOU FOLKS WOULD
2	APPRECIATE A LOOK INTO THE FUTURE OF REGENERATIVE
3	MEDICINE 2020 AND 2030.
4	WE JUST GOT BACK FROM, A NUMBER OF US WENT
5	TO WORLD STEM CELL SUMMIT IN ATLANTA, WHICH IS AN
6	EVENT THAT'S HELD AROUND THE COUNTRY EVERY YEAR THAT
7	IS DIFFERENT THAN MOST OF THE EVENTS WHICH TEND TO
8	BE ENTIRELY SCIENTIFIC IN NATURE. THIS EVENT ALWAYS
9	HAS A NUMBER OF PATIENTS AND PATIENT ADVOCATES. SO
10	THERE IS A GENERAL ATTEMPT TO HAVE PANELS AND
11	PRESENTATIONS THAT ARE A BIT MORE UNDERSTANDABLE
12	THAN THOSE MEETINGS THAT ARE ENTIRELY IN SCIENCE.
13	I MODERATED A PANEL THERE. I SHOULD SAY,
14	BY THE WAY, RANDY AND I BOTH WENT TO THIS. AND FOR
15	THE LAST TWO YEARS, THEY'VE HAD RANDY AND I DO
16	DUELING PANELS AT THE SAME TIME EVEN THOUGH IT'S A
17	FOUR-DAY CONFERENCE. SO RANDY AND I DISCUSSED HOW
18	OURS WENT. MINE, FOR YOUR INTEREST, HAPPENED TO BE
19	ON THE SUBJECT OF INTERNAL AND EXTERNAL
20	COLLABORATIONS AND HOW YOU CAN LEVERAGE WHAT YOU ARE
21	DOING WITH OTHER FOLKS. IT WAS QUITE INTERESTING.
22	THE MILKEN INSTITUTE, AS YOU KNOW, PART OF
23	THAT IS THE FASTER CURES ORGANIZATION IN WASHINGTON
24	WHICH LOBBIES CONGRESS ON THE IMPORTANCE OF FUNDING
25	FOR MEDICAL RESEARCH. THEY HAVE A SPECIFIC MEETING
	10

1	EVERY NOVEMBER IN NEW YORK CALLED "PARTNERING FOR
2	CURES" AT WHICH YOU GET A HOST OF RESEARCHERS,
3	FOUNDATIONS, PATIENTS, INVESTORS, ETC. AMY LEWIS,
4	NEAL LITTMAN, AND I WENT ON BEHALF OF CIRM TO THAT
5	THIS YEAR AND HAD A NUMBER OF MEETINGS WITH PEOPLE
6	FROM DIFFERENT DISEASE FOUNDATIONS TOWARDS THE END
7	OF GETTING THEM INTERESTED IN SPECIFIC PROJECTS THAT
8	WE HAVE; FOR EXAMPLE, THE FOUNDATION FOR FIGHTING
9	BLINDNESS, THE TYPE 1 DIABETES EXCHANGE, AND THE
10	AMERICAN HEART ASSOCIATION, ALL OF WHICH WE'RE
11	LOOKING TO BRING IN TO IN SOME CAPACITY HELP US WITH
12	PROJECTS WE HAVE IN THOSE FIELDS.
13	MILKEN JUST HAD HIS SUMMIT ON CALIFORNIA
14	WHERE YOU HAVE LEADERS FROM AROUND THE STATE OF ALL
15	DIFFERENT INDUSTRIES COME AND SPEAK. SENATOR TORRES
16	DID A TERRIFIC JOB ON A PANEL AT THAT CONFERENCE.
17	AND WE WERE ABLE TO MEET WITH A NUMBER OF THOUGHT
18	LEADERS THROUGHOUT THE STATE AND GIVE THEM UPDATES
19	ON WHAT'S GOING ON WITH CIRM, WHICH CONTINUES TO BE
20	A MATTER OF GREAT INTEREST.
21	WE HAD THE STEM CELL MEETING ON THE MESA,
22	WHICH IS A TWO-DAY EVENT DOWN IN LA JOLLA, WHICH
23	BRINGS INDUSTRY AND INVESTORS TOGETHER, AND THIS
24	YEAR HAD ALMOST AN OVERFLOW CROWD REFLECTING THE
25	DEVELOPMENT OF THE FIELD AND THE LEVEL OF INTEREST

1	THAT YOU SEE IN IT FROM ALL PARTICIPANTS.
2	WE HAD OUR ANNUAL MEETING WITH THE STATE
3	CONTROLLER, WHO, AS YOU MIGHT RECALL, IS THE
4	CONSTITUTIONAL OFFICER WHO HAS OVERSIGHT OVER CIRM.
5	THE BATON WAS PASSED SINCE OUR LAST MEETING FROM NOW
6	STATE TREASURER JOHN CHIANG TO NOW STATE CONTROLLER
7	BETTY YEE. WE WENT AND HAD, I THINK, A VERY
8	PRODUCTIVE MEETING WITH HER AND HER STAFF, WHICH
9	WAS, I WOULD SAY, ART, A HUNDRED PERCENT UPBEAT,
10	HIGHLY SUPPORTIVE OF WHAT EVERYBODY IS DOING, HIGHLY
11	SUPPORTIVE OF THE STATE OF PLAY HERE AT CIRM, WHICH
12	IS A GOOD THING. SO WE WANT TO MAKE SURE THAT WE
13	HAVE GREAT RELATIONS WITH THE FOLKS IN SACRAMENTO TO
14	SHOW THEM WHAT GREAT STUFF WE'RE ALL DOING.
15	TOWARDS THAT END, I HAD A SEPARATE MEETING
16	WITH THE STATE TREASURER, ABOUT AN HOUR AND A HALF
17	WORTH, GAVE HIM CHAPTER AND VERSE ON ALL OF RANDY'S
18	NEW PROGRAMS AND HOW THAT'S ADVANCED THINGS HERE TO
19	EVEN GREATER HEIGHTS. HE WAS VERY ENTHUSIASTIC
20	ABOUT THAT.
21	WE HAD, OF COURSE, THE ANNUAL BRIDGES AND
22	CREATIVITY MEETINGS. THOSE ARE AMONGST ALL OF OUR
23	FAVORITE EVENTS BECAUSE YOU GET THESE KIDS FROM HIGH
24	SCHOOL ALL THE WAY UP TO POST DOCS WHO ARE
25	UNBELIEVABLY IMPRESSIVE. YOU TALK TO THESE PEOPLE

1	AND YOU THINK THEY'VE BEEN IN THIS FOR YEARS AND
2	YEARS, EVEN THE HIGH SCHOOLERS. HAD A FRIEND, JUST
3	THE CALIBER OF PEOPLE THAT GET TO PARTICIPATE IN
4	SOME OF THESE PROGRAMS. HAD A COUPLE OF PEOPLE THAT
5	I RECOGNIZED FROM LOS ANGELES AND GOT TO LISTEN TO
6	THEIR POSTERS. WE HAD ONE GREAT ONE, LAUREN, ON
7	ALZHEIMER'S. THIS KID HAD BEEN THERE FOR THREE
8	MONTHS AND THIS POSTER, AND IT SOUNDED LIKE A
9	POST-DOC. IT WAS JUST EXTRAORDINARY, HAVING GONE
10	INTO IT WITH NO KNOWLEDGE OF THE VERNACULAR, THE
11	SUBJECT, OR ANYTHING ELSE. SO IT'S A VERY
12	SUCCESSFUL PROGRAM.
13	WE HAD SOME MEETINGS WITH BIG PHARMA
14	TOWARDS FORGING A STRATEGIC ALLIANCE ON A NUMBER OF
15	OUR PROJECTS GOING FORWARD WHICH ARE IN PROCESS AND
16	WE'LL BE ABLE TO REPORT ON MORE AT A LATER DATE.
17	LOTS OF ACTIVITIES SURROUNDING WHAT YOU'LL
18	BE HEARING LATER IN THE SESSION, OUR ACCELERATED
19	THERAPIES PUBLIC PRIVATE PARTNERSHIP CONCEPT, A
20	GREAT DEAL OF WORK ON THAT.
21	I WOULD LIKE TO MENTION I DON'T KNOW IF
22	HE'S HERE, BUT IT WAS A REALLY NICE EVENT WE HAD IN
23	THE OFFICE DON REED. IS DON HERE? I DON'T SEE
24	DON. DON, AS YOU KNOW, IS A LONGTIME, ARDENT
25	SUPPORTER OF CIRM, WROTE A BOOK ON PROP 71. AND WE

1	HAD AN EVENT AT CIRM WHERE HE GOT UP AND TALKED
2	ABOUT THE BOOK AND WHAT WENT INTO IT. AND IT WAS
3	VERY HEARTFELT AND A WONDERFUL RECOUNTING OF
4	EVERYTHING THAT LED UP TO THE PASSAGE AND WHAT'S
5	HAPPENED AT CIRM SINCE. HE HAS COPIES OF THE BOOK,
6	AND WE'LL SEE IF WE CAN GET COPIES FOR ALL OF YOU.
7	I THINK IT'S SOMETHING THAT'S NICE TO HAVE GIVEN THE
8	AMOUNT OF TIME YOU'VE PUT INTO THIS.
9	MS. LANSING: CAN WE BUY THE BOOK?
10	CHAIRMAN THOMAS: YES.
11	MS. LANSING: LIKE ON AMAZON?
12	CHAIRMAN THOMAS: WE WILL GET YOU
13	INFORMATION ON THAT. IT'S REALLY IT'S A VERY
14	EASY READ, AND HE PUT A TON OF TIME INTO THIS. AND,
15	AGAIN, IT'S SOMETHING I THINK YOU'D ALL ENJOY
16	HAVING.
17	SO ANOTHER THING, YOU MAY HAVE HEARD OUR
18	TEN-YEAR FREE LEASE RAN OUT IN NOVEMBER. WE SPENT A
19	LOT OF TIME WORKING ON THE SUBJECT OF WHERE WE WOULD
20	PUT OUR NEW OFFICES BECAUSE THE SPOT WE WERE AT DOWN
21	BY THE BALLPARK IS A WHITE HOT AREA THAT'S OFF THE
22	CHARTS EXTRAORDINARILY EXPENSIVE. WE COULD NOT STAY
23	THERE. LOOKED AT A NUMBER OF PLACES IN SAN
24	FRANCISCO. THE CITY, THE REAL ESTATE MARKET IS
25	HUMMING, AND IT JUST UNFORTUNATELY MADE IT SO THAT

1	WE COULD NOT AS A PUBLIC AGENCY AFFORD TO STAY IN
2	THE CITY. SO AFTER A GREAT DEAL OF TIME AND EFFORT,
3	WE CHOSE A SPOT IN OAKLAND AT LAKE MERRITT.
4	WOULD LIKE TO SINGLE OUT SENATOR TORRES
5	FOR HIS EXTRAORDINARY EFFORT THROUGHOUT THE PROCESS
6	OF LOOKING FOR A NEW SPOT. IN OAKLAND HE WAS KEY TO
7	INTRODUCING CIRM TO THE MAYOR, WHO WAS VERY HELPFUL
8	IN A NUMBER OF WAYS IN CONNECTION WITH OUR NEW
9	SPACE, INCLUDING PARKING AND OTHER THINGS. AND,
10	ART, I KNOW THIS TOOK A LOT OF WORK ON YOUR PART.
11	IN HOUSE PRESIDENT MILLS HAS INSTITUTED
12	WHAT HE CALLS THE GAME BALL IDEA FOR SOMEONE WHO PUT
13	IN ABOVE AND BEYOND THE CALL OF DUTY ON SOMETHING.
14	SO I WOULD LIKE, ON BEHALF OF CIRM AND THE BOARD, TO
15	PRESENT TO ART A CIRM GAME BALL.
16	(APPLAUSE.)
17	MR. TORRES: OH, THAT'S GREAT. THANK YOU.
18	CHAIRMAN THOMAS: THANK YOU.
19	SO THE FIELD IS MAKING GREAT PROGRESS. WE
20	HAD A PRESS RELEASE BY UCLA A FEW WEEKS AGO ON A
21	PROJECT THAT EARLY CIRM FUNDING HAD HELPED TO HAVE
22	AN IMPACT ON THAT'S IN THE FIELD OF SEVERE COMBINED
23	IMMUNODEFICIENCY DISEASE OR SCID, BETTER KNOWN
24	COLLOQUIALLY AS BUBBLE BABY DISEASE, WHERE DON KOHN,
25	WHOM YOU'RE FAMILIAR WITH, HAD A PROCEDURE WHICH
	15

1	COMBINED STEM CELL AND GENE THERAPY FOR KIDS WITH
2	THIS CONDITION, WHICH IS A TERRIBLE CONDITION. AND
3	BASICALLY, WITHOUT GETTING INTO TOO MUCH DETAIL, THE
4	COMBINED THERAPEUTIC TREATMENT A YEAR AGO RESULTED
5	IN 18 KIDS WHO HAVE THIS CONDITION NO LONGER BEING
6	ON ANY MEDICATION, NO LONGER BEING QUARANTINED, AND,
7	IN FACT, BEING IN SCHOOL WITH KIDS JUST LIKE NORMAL
8	CHILDREN AND WERE A YEAR AGO THOUGHT TO BE
9	FUNCTIONALLY CURED.
10	SO A COUPLE OF WEEKS AGO THEY HAD ANOTHER
11	PRESS RELEASE, MARKING THE ONE-YEAR ANNIVERSARY OF
12	THE FIRST PRESS RELEASE AND THAT NOTED THEY'D HAD AN
13	ADDITIONAL FIVE KIDS WHO WERE TREATED, WERE
14	SIMILARLY DOING WELL. AND THEY LOOKED BACK AT THE
15	FIRST 18, AND IN COMBINATION, ALL 23 OF THESE KIDS
16	ARE FINE. AND IT LOOKS LIKE THEY ARE FUNCTIONALLY
17	CURED USING STEM CELL AND GENE THERAPY.
18	I MENTION THIS FOR SEVERAL REASONS. ONE
19	IS THAT IT'S EVIDENCE OF THE EXCITING DEVELOPMENTS
20	IN THE FIELD. THIS IS A TRUE WIN.
21	SECONDLY, DON KOHN YOU MAY RECOGNIZE AS
22	THE INVESTIGATOR IN OUR SICKLE CELL DISEASE TEAM
23	PROJECT IN WHICH HE'S USING BASICALLY THE SAME IDEA
24	TO TREAT THAT DISEASE. AND WE ARE, OF COURSE, MOST
25	HOPEFUL THAT IT ACHIEVES SIMILAR RESULT.
	16

SO I THOUGHT THAT THE BOARD WOULD ENJOY
SEEING A SHORT PIECE, VERY SHORT, CBS NEWS ON DON'S
WORK THAT WENT PUBLIC NATIONALLY. AND WITHOUT
FURTHER ADO, AMY, IF YOU COULD JUST SHOW THAT.
(VIDEO WAS SHOWN, BUT NOT REPORTED
NOR HEREIN TRANSCRIBED.)
CHAIRMAN THOMAS: THANKS, AMY. SO I'D
LIKE JUST TO CONCLUDE MY CHAIR'S REPORT WITH A
SPECIAL SHOUT OUT TO MY DAUGHTER LIZZY WHO TURNS 20
TODAY. SO HAPPY BIRTHDAY TO LIZZY.
(APPLAUSE.)
CHAIRMAN THOMAS: ON NOW TO THE
PRESIDENT'S REPORT.
YES, ANNE-MARIE.
DR. DULIEGE: FIRST OF ALL, AMAZING REPORT
ON THESE CHILDREN WITH SEVERE COMBINED
IMMUNODEFICIENCY. AS A PEDIATRIC IMMUNOLOGIST,
HAVING IN THE PAST SEEN KIDS AND TAKEN CARE OF KIDS
IN THEIR BUBBLE, THIS IS ABSOLUTELY AMAZING.
WERE THERE ANY COLLABORATION WITH CIRM?
DID CIRM FUND PART OF THIS PROJECT, OR IS IT JUST A
MAJOR MILESTONE IN THE FIELD?
CHAIRMAN THOMAS: WE HAD VARIOUS EARLY
GRANTS THAT WE HAD THAT IMPACTED WHAT THEY ARE DOING
THROUGH TRAINING, THROUGH SHARED LABS, AND SOME
17

1	OTHER THINGS. I BELIEVE THEY HAD NIH FUNDING FOR
2	THE ACTUAL CLINICAL TRIAL ITSELF.
3	DR. DULIEGE: BUT CIRM CAN CLAIM TO HAVE
4	CONTRIBUTED PARTIALLY TO THIS SUCCESS?
5	CHAIRMAN THOMAS: YES.
6	DR. DULIEGE: IT'S PHENOMENAL.
7	AND THE SECOND COMMENT IS ACTUALLY I WANT
8	TO SAY WE REALLY APPRECIATE, AT LEAST I DO
9	APPRECIATE, RECEIVING FROM CIRM ON A REGULAR BASIS
10	SOME NEWS LIKE THIS AND WE'RE ON THE DISTRIBUTION
11	LIST, WHICH IS GREAT. I WAS WONDERING IF WE COULD
12	ALSO BE INFORMED ON SOME OF THE EVENTS THAT YOU
13	MENTIONED; FOR INSTANCE, DON REED COMING TO CIRM,
14	BECAUSE FOR THOSE OF US WHO ARE NEARBY, MAYBE WE
15	COULD, IF WE CAN, ATTEND THOSE AS WELL. SO JUST LET
16	US KNOW WHEN THAT HAPPENS WITH ADVANCE NOTICE.
17	CHAIRMAN THOMAS: THANK YOU.
18	MS. LANSING: SO THIS IS NOT TO BE VIEWED
19	AS A CRITICISM. SO, PLEASE, I JUST WANT TO TAKE IT
20	AS HOW CAN I AND OTHERS WHO MAYBE HAVE CONTACT IN
21	THE MEDIA BE HELPFUL BECAUSE THIS IS WHAT WE'VE BEEN
22	TALKING ABOUT, THESE BIG WINS THAT WE NEED TO GET TO
23	THE PUBLIC. THEY'RE ALWAYS ON OUR SIDE, BUT LET'S
24	SAY EVEN MORE ON OUR SIDE BECAUSE I WASN'T REALLY,
25	AND I KIND OF FOLLOW THIS, I WASN'T REALLY AWARE OF
	10

1	THE EXTRAORDINARY IMPACT OF THIS. I HEARD A LITTLE
2	BIT ABOUT IT. I DID NOT SEE THE NEWS PIECE, AND
3	THAT'S PROBABLY MY FAULT. I'M SURE YOU SENT US
4	SOMETHING TO LET US KNOW. BUT THIS SHOULD BE HUGE,
5	JUST HUGE.
6	AGAIN, THIS IS NOT A CRITICISM. I'M SURE
7	EVERYTHING HAS BEEN DONE. IT'S NOT TOO LATE, I
8	GUESS, IS WHAT I'M SAYING. WHEN YOU HAVE SOMETHING
9	LIKE THIS, LET'S SEE HOW WE CAN REALLY MAKE IT THE
10	FRONT PAGE OF EVERY BLOG, EVERY NEW YORK TIMES, MORE
11	NEWS PEOPLE CARRYING IT BECAUSE THIS IS WHAT WE
12	NEED. IF WE WERE A SMALL PART OF IT, THAT'S FINE.
13	THE POINT IS STEM CELLS ARE WORKING, AND WE HAVE TO
14	MAKE THE PUBLIC UNDERSTAND THAT SO WE CAN CONTINUE
15	TO GET THEIR SUPPORT.
16	THERE WAS A MOVIE DONE ON THE BUBBLE BABY
17	WITH JOHN TRAVOLTA. THAT'S HOW OLD I AM THAT I
18	REMEMBER THIS, BUT I NEVER FORGOT IT. I NEVER
19	FORGOT IT. MOST OF THE JOURNALISTS WILL HAVE SEEN
20	IT TOO. SO THIS IS INCREDIBLE.
21	CHAIRMAN THOMAS: THANKS, SHERRY. POINT
22	WELL TAKEN. THE PRESS EFFORT ON THIS WAS, AS YOU
23	WOULD IMAGINE, SPEARHEADED BY UCLA, BUT WE ALSO,
24	YOU'RE RIGHT, COULD HELP IN THAT REGARD. AND
25	TOWARDS THAT END, WE HAVE TWO AUGUST MEMBERS OF THE
	10

1	FOURTH ESTATE HERE IN THE AUDIENCE WHO ARE COVERING
2	AND I THINK WOULD BE MORE THAN HAPPY TO TAKE NOTE OF
3	THIS. AS I SAY, WE'RE VERY HOPEFUL THAT THE DISEASE
4	TEAM THAT WE'RE FUNDING WITH DR. KOHN ON SICKLE CELL
5	USING THE SAME TECHNOLOGY WILL BE ABLE TO REPORT
6	BACK DOWN THE ROAD THAT THEY'RE GETTING VERY GOOD
7	RESULTS AS WELL.
8	MS. LANSING: THIS IS NOT ABOUT CIRM
9	GETTING CREDIT. THIS IS ABOUT, AS RANDY OFTEN SAYS,
10	SAVING LIVES AND PROVING THAT THE WORK THAT IS GOING
11	ON REALLY IS CHANGING THE WORLD. AND I HOPE THAT
12	THERE WILL BE A LOT WRITTEN ABOUT IT.
13	CHAIRMAN THOMAS: THANK YOU. ANY OTHER
14	COMMENTS FROM MEMBERS OF THE BOARD? DR. MILLS,
15	PRESIDENT'S REPORT.
16	DR. MILLS: THANK YOU, CHAIRMAN THOMAS,
17	MEMBERS OF THE BOARD. I WILL ATTEMPT TO KEEP MY
18	PRESIDENT'S REPORT BRIEF TODAY BECAUSE I WANT TO
19	RAMBLE ON ABOUT THE STRATEGIC PLAN WHEN I GET A
20	CHANCE TO DO THAT. SO I DON'T WANT YOU TO GET SICK
21	OF ME TOO SOON.
22	AS ALWAYS, WE'LL GO OVER THE MISSION. I
23	WANT TO GIVE YOU AN UPDATE ON THE FISCAL YEAR FIRST
24	QUARTER FINANCIAL UPDATE, WHICH, REMINDING FOR US,
25	OUR FISCAL YEAR FOR 2016 BEGAN JULY 1ST. THEN TALK

1	A LITTLE BIT ABOUT OUR NEW CLINICAL STAGE PROGRAM
2	REVIEW PROCESS THAT WE IMPLEMENTED UNDER CIRM 2.0
3	ALMOST A YEAR AGO. ACTUALLY WE ADOPTED A YEAR AGO
4	AT THE LAST DECEMBER BOARD MEETING. AND THEN,
5	LASTLY, JUST END ON A NOTE OF OUR NEW HOME.
6	SO, AS ALWAYS, I THINK IT'S IMPORTANT THAT
7	WE NEVER LOSE SIGHT OF WHY WE'RE HERE. WE ARE AN
8	AGENCY THAT WAS CREATED TO HELP PATIENTS, AND OUR
9	MISSION IS TO ACCELERATE STEM CELL TREATMENTS TO
10	THOSE PATIENTS WITH UNMET MEDICAL NEEDS.
11	NOW MOVING TO THE FINANCIAL UPDATE. SO I
12	LIKE TO TALK ABOUT CIRM'S FUNDING IN TWO SEPARATE
13	BUCKETS. WE HAVE A BIG BUCKET WHICH HOLDS THE MONEY
14	THAT WE DISTRIBUTE OUT IN AWARDS. THAT IS INITIALLY
15	A \$2.75 BILLION BUCKET. AND WE HAVE THE SMALLER
16	ADMINISTRATIVE BUCKET WHICH IS CAPPED AT \$180
17	MILLION FOR THE LIFE OF CIRM, WHICH AT THAT TIME WAS
18	THOUGHT TO BE IN THE BALLPARK OF AROUND TEN YEARS.
19	SO FAR WE'VE SPENT 105 MILLION OF THAT.
20	WE HAVE 75 MILLION OF THAT REMAINING, WHICH GIVES US
21	OUT OF THIS SMALL BUCKET, AT THAT CURRENT SPEND
22	RATE, THAT WOULD GIVE US APPROXIMATELY FIVE YEARS OF
23	RUNWAY. WE OBVIOUSLY HAVE PLANS AND THE ABILITY TO
24	SCALE THAT ONE WAY OR THE OTHER AS NECESSARY AS
25	EVENTS UNFOLD.
	21

1	WITH REGARDS TO THE AWARD BUCKET, THAT WAS
2	THE MUCH LARGER OF THE TWO, 2.75 BILLION TO START
3	WITH IN THAT. WE HAVE TWO BILLION THAT'S EITHER
4	BEEN COMMITTED, WHAT WE CALL AWARDED, OR THAT HAVE
5	ALREADY BEEN SPENT. WE HAVE ANOTHER 759 MILLION
6	THAT IS CURRENTLY UNCOMMITTED. SO WITH A PLANNED
7	SPEND RATE OF ABOUT 190 MILLION IN NEW AWARDS, WE
8	WOULD EXPECT A NET SPEND RATE OF ABOUT 170. AND
9	THAT'S BECAUSE JUST BECAUSE WE COMMIT FUNDS WHEN WE
10	MAKE AN AWARD DOESN'T MEAN ALL OF THOSE FUNDS GET
11	USED. WHEN PROGRAMS DON'T WORK, THE AWARDS ARE
12	TERMINATED, THE FUNDS COME BACK TO CIRM SO THEY CAN
13	BE REISSUED.
14	I'VE SHOWN THIS SLIDE BEFORE, BUT THERE'S
15	A POINT HERE THAT I WANT TO MAKE ON WHAT'S GOING ON.
16	SO IN THE FIRST QUARTER OF 2016, SO EARLIER THIS
17	YEAR, JULY, WE COMMITTED AN ADDITIONAL 38 MILLION IN
18	NEW AWARDS. SO GWG MADE RECOMMENDATIONS AND THIS
19	DOADD ADDDOVED \$20 MTH TON THE NEW AWARDS DUT TH
	BOARD APPROVED \$38 MILLION IN NEW AWARDS. BUT IN
20	THAT SAME PERIOD, WE HAD AWARD REDUCTIONS OF 16
20 21	
	THAT SAME PERIOD, WE HAD AWARD REDUCTIONS OF 16
21 22	THAT SAME PERIOD, WE HAD AWARD REDUCTIONS OF 16 MILLION, AND WE HAD AWARD REPAYMENTS OF AN
21	THAT SAME PERIOD, WE HAD AWARD REDUCTIONS OF 16 MILLION, AND WE HAD AWARD REPAYMENTS OF AN ADDITIONAL 5 MILLION. SO OUR ACTUAL NET COMMITMENT
21 22 23	THAT SAME PERIOD, WE HAD AWARD REDUCTIONS OF 16 MILLION, AND WE HAD AWARD REPAYMENTS OF AN ADDITIONAL 5 MILLION. SO OUR ACTUAL NET COMMITMENT RATE FOR THE QUARTER WAS ONLY \$16 MILLION.

1	DID WAS WE WENT BACK AND LOOKED AT HOW DID THE FULL
2	YEAR FOR 2015'S FY GO. SO THIS IS THE WHOLE YEAR.
3	WE MADE \$130 MILLION IN NEW AWARDS, AND WE HAD 29
4	MILLION COME BACK, WHICH IS AN AWARD RETURN RATE OF
5	ABOUT 22 PERCENT.
6	THE ONLY REASON I BRING THIS UP IS THE
7	STRATEGIC PLAN THAT I'M GOING TO TALK ABOUT LATER
8	ASSUMED A RETURN RATE OF ONLY 10 PERCENT. IF THE
9	ACTUAL RETURN RATE ENDED UP BEING SOMETHING CLOSER
10	TO 22 PERCENT, THAT WOULD MEAN WE WOULD HAVE AN
11	ADDITIONAL \$150 MILLION IN AWARD MONEY THAT WOULD BE
12	AVAILABLE TO BE RECOMMITTED UNDER NEW AWARDS. THE
13	PROBLEM SO THAT SEEMS LIKE GOOD NEWS. THE
14	PROBLEM WITH THAT IS WE DON'T GET ANY MORE
15	CORRESPONDING ADMINISTRATIVE MONEY TO ADMINISTER
16	THAT \$150 MILLION IN NEW AWARDS.
17	OBVIOUSLY CHAIRMAN THOMAS IS GOING TO TALK
18	MORE ABOUT THIS AS THE MEETING CONCLUDES. BUT IT'S
19	SOMETHING THAT WE'LL JUST HAVE TO WATCH AND SEE
20	UNFOLD, BUT SO FAR RETURN RATE IS HIGHER THAN OUR
21	CURRENT EXPECTATIONS. I THINK THAT'S ALSO A GOOD
22	SIGN BECAUSE WE KNOW BIOTECH DOESN'T WORK AT A
23	HUNDRED PERCENT, AND WE WOULD EXPECT PROGRAMS TO RUN
24	INTO CHALLENGES AND SOME THINGS TO JUST NOT WORK.
25	AND I THINK WE'RE BEING VERY GOOD STEWARDS OF THE
	22

1	TAXPAYERS' MONEY WHEN WE PROSPECTIVELY SET UP
2	PROGRAMS AND WE HAVE AGREEMENTS GOING INTO THOSE
3	THAT SAY IF THIS DOESN'T WORK, WE'RE NOT JUST GOING
4	TO KEEP THROWING MONEY AT IT. WE'LL TERMINATE THE
5	PROGRAM AND ALLOW THOSE REMAINING FUNDS TO BE
6	REINVESTED.
7	I JUST WANT TO MAKE THE BOARD AWARE THAT
8	RETURN RATE IS HIGHER THAN WE CALCULATED IN THE
9	STRATEGIC PLAN. THAT COULD HAVE SOME UP SIDE TO IT.
10	IT ALSO CREATES SOME FUNDING CHALLENGES FOR
11	SUBSEQUENT ADMINISTRATION.
12	DOES ANYONE HAVE QUESTIONS ON THAT?
13	MR. PANETTA: THANKS. SO, RANDY, WHAT
14	YOU'RE SAYING IS YOU'VE GOT FIVE YEARS OF
15	ADMINISTRATIVE FUNDING LEFT, BUT YOU COULD
16	POTENTIALLY HAVE IN EXCESS OF FIVE YEARS OF GRANT
17	FUNDING. SO YOU COULD GET TO FIVE YEARS FROM NOW
18	AND HAVE GRANT FUNDING, BUT NOT THE ABILITY TO
19	ADMINISTER THE GRANT FUNDING?
20	DR. MILLS: THEORETICALLY, YES. AGAIN,
21	THE FIVE YEARS OF ADMINISTRATIVE FUNDING DOESN'T
22	QUITE WORK THAT WAY BECAUSE OUR ASSUMPTION IS WE'LL
23	ACTUALLY GET DONE MAKING AWARDS SOONER THAN THAT BY
24	ABOUT FOUR YEARS, AND THE EXPENSE LEVEL TAILS OFF.
25	WE WOULDN'T KEEP BURNING OUT OF THAT ADMINISTRATIVE

1	BUCKET. SO REALISTICALLY WE THINK WE'LL PROBABLY GO
2	SEVEN YEARS ON THE ADMINISTRATIVE BUCKET AND FOUR
3	AND A HALF ON AWARDS. BUT WITH \$150 MILLION NEW
4	THROWN INTO THAT, IT WOULD CHANGE THE GAME.
5	OKAY. THE NEXT THING I'D LIKE TO TALK
6	ABOUT IS JUST GIVE YOU A BRIEF, AND THIS IS A BRIEF,
7	UPDATE ON OUR CLINICAL STAGE PORTFOLIO AND THEN TALK
8	ABOUT HOW WE'RE GOING TO BE UPDATING OUR
9	CLINICAL PORTFOLIO GOING FORWARD.
10	SO FIRST OF ALL, I THINK IT'S VERY
11	IMPORTANT CIRM 2.0 FOR THE CLINICAL STAGE
12	PROGRAMS WE PUT IN PLACE A YEAR AGO, SO I THINK IT'S
13	IMPORTANT FOR US TO TAKE A ONE-YEAR SNAPSHOT AND
14	LOOK AT THAT AND SEE HOW THAT'S BEEN GOING AND IF
15	IT'S BEEN DOING WHAT WE INTENDED IT TO DO.
16	SO I THINK IN A NUTSHELL I CAN SAY WE'RE
17	VERY, VERY PLEASED WITH HOW THE PROGRAM IS GOING.
18	WE RECEIVED SO FAR ACTUALLY THIS YEAR 28
19	APPLICATIONS. NINETEEN OF THOSE APPLICATIONS WENT
20	ON TO PASS ELIGIBILITY. SIXTEEN OF THOSE 19 HAVE
21	BEEN GIVEN FINAL DISPOSITIONS BY THE GRANTS WORKING
22	GROUP. THREE ARE CURRENTLY UNDER REVIEW. AND THE
23	APPLICATIONS THAT HAVE BEEN RECOMMENDED FOR FUNDING
24	AND SUBSEQUENTLY FUNDED BY THIS BOARD IS SIX OR 38
25	PERCENT OF ELIGIBLE APPLICATIONS.

1	SO JUST FOR COMPARISON, ALL OF LAST YEAR
2	WE HAD THREE APPLICATIONS, AND NONE OF THEM ENDED UP
3	BEING ELIGIBLE AND WE MADE NO NEW AWARDS. SO THE
4	AMOUNT OF INTEREST WE'VE SEEN IN CIRM'S CLINICAL
5	STAGE PROGRAM HAS SHOT UP DRAMATICALLY. AND I THINK
6	IT'S BECAUSE OF THE TREMENDOUS WORK OF THE GWG,
7	WHICH INCLUDES MANY MEMBERS FROM THIS BOARD ON IT,
8	WILLING TO MEET ON A MONTHLY BASIS, AND THEN THE
9	CIRM TEAM WILLING NO. WILLING KIND OF LIKE COME
10	ON, GUYS, YOU KNOW WE'RE GOING TO DO THIS WILLING,
11	BUT WILLING AND, MOST IMPORTANTLY, ABLE TO CONDUCT
12	MONTHLY REVIEWS. SO OUR CYCLE TIME HAS BEEN REDUCED
13	PRACTICALLY FROM 22 MONTHS TO FOUR MONTHS FROM THE
14	TIME THAT PEOPLE ACTUALLY HAVE THEIR FUNDING, AND
15	IT'S 61 DAYS UNTIL THE TIME THEY KNOW THEY HAVE
16	THEIR DECISION. AND SO THAT HAS BEEN A REAL GAME
17	CHANGER FOR INTEREST INTO CIRM FOR OUR CLINICAL
18	STAGE PROGRAMS, AND IT'S SHOWING UP IN THE VOLUME
19	THAT WE'RE SEEING.
20	JUST TAKING A LOOK AT THE SCORING BECAUSE
21	I THINK WE INTRODUCED SOME PRETTY INNOVATIVE THINGS
22	HERE TOO. AS YOU WILL RECALL, WE USED TO DO SCORING
23	1 TO A 100, AND WE WOULD BASICALLY APPROVE ANYTHING
24	GREATER THAN 75 AND REJECT ANYTHING LESS THAN 65 AND
25	SORT OF THROW IT TO THE BOARD ON WHAT YOU WANTED TO

1	DO IN BETWEEN. WE CAME UP WITH THIS NEW SCORING
2	SYSTEM THAT SAID WE'RE GOING TO EVALUATE THESE
3	CLINICAL TRIALS. AND BECAUSE WE'RE HOLDING MONTHLY
4	REVIEWS, WE DON'T HAVE TO BASICALLY DO THIS PUNT
5	IT. IF IT'S NOT A GREAT APPLICATION, WE DON'T HAVE
6	TO GIVE IT TO THE BOARD AND SAY FIGURE OUT WHAT YOU
7	WANT TO DO. INSTEAD, WE CAME UP WITH AN OPPORTUNITY
8	WHERE WE WOULD ALLOW THE APPLICANTS TO LISTEN TO THE
9	FEEDBACK FROM THE GRANTS WORKING GROUP AND SEE IF
10	THEY WOULD MODIFY THEIR APPLICATIONS TO IMPROVE
11	THEM.
12	AGAIN, THE GOAL HERE WASN'T JUST TO GET
13	MORE THINGS THROUGH FASTER, BUT IT WAS ALSO TO GET
14	THINGS THROUGH BETTER. WE DIDN'T WANT TO PUT MORE
15	75S THROUGH. WE WANTED TO PUT 95S THROUGH. SO
16	HERE'S HOW THIS WORKS.
17	SO ON INITIAL SCORING OUT OF ALL OF THE
18	THINGS THAT HAVE BEEN SCORED BY THE GWG, THREE WENT
19	THROUGH FIRST TIME AS TIER I. THEY WERE GREAT
20	PROGRAMS AND SHOULD BE FUNDED. EIGHT WENT THROUGH
21	AS TIER III, WHICH IS THEY'RE FLAWED AND THE GWG
22	BELIEVES THE FLAW IS BASICALLY A FATAL FLAW TO THE
23	PROGRAM AND NOT REMEDIABLE. BUT SEVEN OF THOSE WERE
24	GIVEN A TWO, WHICH IS HERE ARE OUR COMMENTS, SEE IF
25	YOU CAN MAKE YOUR APPLICATION BETTER. TWO OF THOSE

1	ARE STILL PENDING. SO IF YOU TAKE THEM OUT, THAT
2	MEANS WE HAD FIVE APPLICATIONS THAT HAD THE
3	OPPORTUNITY TO AMEND THEIR AWARDS AND SUBMIT. ONE
4	REFUSED TO AMEND THEIR AWARD, AND SO IT DIDN'T GO
5	ANY FURTHER. BUT OF THE FOUR THAT DID FIX THEIR
6	APPLICATION, THREE OF THOSE WENT ON TO A 1, AND ONE
7	OF THEM WENT ON TO A 3.
8	AND SO I THINK THE IDEA OF INSTEAD OF
9	PUTTING YOU GUYS IN AN UNCOMFORTABLE POSITION ON
10	WHAT DO YOU DO WITH A MEDIOCRE AWARD, THIS PROCESS
11	WAS ABLE TO TAKE THAT AND AMEND THOSE. AND BY THE
12	WAY, THE THREE WENT ON TO GO FROM 2S TO 1S, THAT
13	WHOLE THING WAS DONE WITH ONLY 30 DAYS OF EXTRA
14	WORK, INCLUDING OUR FEEDBACK TO THEM AND THEIR
15	REVISION, BACK TO THE GWG AND THE GWG REREVIEW ALL
16	TAKING PLACE WITHIN 30 DAYS. SO I THINK THIS
17	PARTICULAR SCORING ASPECT OF IT HAS WORKED PRETTY
18	WELL.
19	AND THEN THE LAST THING I JUST WANTED TO
20	SHOW HERE IS WHAT WE'RE FUNDING BECAUSE IT'S A
21	LITTLE DIFFERENT THAN WHAT IT USED TO BE. SO COMING
22	INTO THIS, WE DIDN'T HAVE ANY PHASE III PROGRAMS,
23	AND I THINK WE ONLY HAD ONE PHASE II PROGRAM IN OUR
24	CLINICAL PORTFOLIO. SO TWO OF THE SIX THINGS THAT
25	WE'VE FUNDED HAVE BEEN IN PHASE III, ONE IN PHASE

1	II, ONE IN PHASE I, AND THEN TWO THAT ARE IN THE
2	IND-ENABLING PHASE OF CLINICAL RESEARCH GETTING
3	READY TO START.
4	SO THAT GIVES US A CLINICAL PROGRAM THAT
5	LOOKS LIKE THIS TODAY. SOME OF THIS STUFF YOU CAN
6	SEE, BUT IT'S BROKEN UP BY MAJOR DISEASE CATEGORIES.
7	YOU CAN SEE NEUROLOGICAL IS THE LARGEST AT 31
8	PERCENT OF OUR FUNDING FOLLOWED BY CANCER AND
9	CARDIOVASCULAR. AND THEN YOU CAN SEE ON THE LEFT
10	THAT'S JUST A LIST OF THE DIFFERENT CLINICAL TRIALS
11	THAT WE CURRENTLY HAVE ACTIVE. AND I WANT TO SAY
12	THESE ARE CLINICAL TRIALS THAT WE HAVE DIRECTLY,
13	ONE, OPINED ON AS A GWG AND A BOARD, AND, TWO, ARE
14	DIRECTLY FUNDING. SO WE HAVE SOMETHING LIKE ANOTHER
15	13 TO 15 TRIALS THAT IN VARIOUS WAYS WE HAVE
16	SUPPORTED PREVIOUSLY, BUT THESE ARE THE ONES THAT WE
17	ARE ACTUALLY ACTIVELY FUNDING RIGHT NOW.
18	SO THE LAST THING IN THIS SECTION THAT I
19	WANTED TO TALK ABOUT WAS A PROCESS FOR UPDATING.
20	ONE OF THE THINGS THAT WE DO WITH THE GWG IS, EVEN
21	THOUGH WE MEET MONTHLY TELEPHONICALLY, TWICE A YEAR
22	WE GET THE GRANTS WORKING GROUP TOGETHER FOR LONGER
23	SESSIONS SO WE CAN SYNC UP AND SO WE CAN HEAR IDEAS
24	AND CHALLENGES AND HOW WE CAN MAKE THE PROCESS
25	BETTER. AND OUT OF THAT LAST MEETING, ONE OF THE
	20

1	THINGS THAT CAME UP WAS HOW DO WE KEEP THE GWG
2	INFORMED AND HOW DO WE BEST UTILIZE THE GWG FOR
3	ACTIVE PROGRAMS IN OUR PORTFOLIO.
4	SO RIGHT NOW WE HAVE 26 CLINICAL STAGED
5	PROGRAMS IN OUR PORTFOLIO. WHAT I MEAN IS WE HAVE
6	26 PROGRAMS THAT AT THE IND THAT HAVE ALREADY HAD
7	THEIR PRE-IND MEETING ONWARD. EITHER THEY HAVE
8	THEIR IND OR THEY'RE IN PHASE I, THEY'RE IN PHASE
9	II, THEY'RE IN PHASE III. SO WE HAVE 26 OF THOSE
10	PROGRAMS. WHAT WE'RE DOING IS WE'RE BREAKING THOSE
11	DOWN, THEN, BY DISEASE AREA AND WE'RE CREATING HIGH
12	LEVEL SUMMARIES. WE WILL NOW BE REVIEWING, AND
13	WE'RE GOING TO TEST THIS OUT COMING UP NEXT WEEK, WE
14	WILL NOW BE REVIEWING ON A MONTHLY BASIS THESE
15	PROGRAMS WITH THE GRANTS WORKING GROUP. AND WE WILL
16	BRING TO THEM BECAUSE WE COULDN'T GO OVER ALL 26 IN
17	A GWG MEETING AND STILL ACTUALLY CONDUCT OUR REVIEW
18	BUSINESS, BUT INSTEAD BRING TO THEM ANY MATERIAL
19	CHANGES THAT WE HAVE WITHIN THE LAST MONTH OR ANY
20	PROBLEMS OR QUESTIONS THAT WE THINK MIGHT BE ON THE
21	HORIZON FOR ANY PARTICULAR PROGRAMS, AND THEN
22	OBVIOUSLY ANSWER ANY QUESTIONS ANYONE ELSE HAS.
23	SO BY HAVING MONTHLY REVIEW, WE'LL HAVE
24	THE GWG MORE ON TOP OF IT. AND THEN WHERE
25	APPROPRIATE WE'LL HAVE THE GWG MAKE RECOMMENDATIONS
	20

1	BACK TO CIRM ON WHAT WE SHOULD, COULD, MIGHT DO IN
2	ORDER TO EITHER MAKE THE PROGRAM BETTER OR TO
3	PERHAPS TERMINATE THE PROGRAM, WHATEVER IT MIGHT BE.
4	AND THEN WE WILL TAKE ALL OF THAT AND THEN ON A
5	QUARTERLY BASIS BRING THAT BACK TO THE BOARD AND
6	CONDUCT QUARTERLY REVIEWS IN PERSON TO THE BOARD OF
7	THAT PORTFOLIO.
8	SO WE'RE GOING TO TEST THIS OUT. IF YOU
9	HAVE FEEDBACK ABOUT THAT, IF YOU HAVE COMMENTS, YOU
10	WANT MORE OR LESS INFORMATION, LET US KNOW.
11	OBVIOUSLY WE WANT TO HAVE THIS BE AS, ONE, AS
12	HELPFUL A PROGRAM AS WE CAN TO OUR AWARDEES SO WE
13	CAN GET THEM THE BEST INFORMATION THEY CAN TO MAKE
14	THEIR PROGRAM SUCCESSFUL. AND THEN OBVIOUSLY WE
15	WANT THIS PROGRAM TO BE AS INFORMATIVE FOR YOU ALL
16	SO THAT YOU CAN MAKE THE BEST DECISIONS GOING
17	FORWARD.
18	SO ANY QUESTIONS ON THAT?
19	DR. DULIEGE: ACTUALLY, FIRST OF ALL,
20	CONGRATULATIONS. GREAT. IT'S EXACTLY THE MISSION.
21	IT'S INSPIRING. AND IT'S THE RIGHT LEVEL OF
22	INFORMATION THAT PERSONALLY I'M HAPPY WITH.
23	ONE QUESTION, YOU MAY HAVE SAID IT, YOU
24	MENTIONED TWO PHASE III TRIALS. WHAT ARE THESE
25	PHASE III TRIALS?

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1	DR. MILLS: WHAT ARE THEY? WE HAVE ONE
2	IN THEY WERE BOTH APPROVED THIS YEAR. ONE OF
3	THEM IS IN GLIOBLASTOMA AND THE OTHER ONE IS IN
4	MALIGNANT METASTATIC MELANOMA.
5	DR. DULIEGE: PHENOMENAL. THANKS.
6	DR. MILLS: OTHER QUESTIONS?
7	THEN, LASTLY, I JUST WANT TO SAY A WORD
8	ABOUT OUR NEW HOME. OVER THE THANKSGIVING WEEKEND,
9	CIRM SUCCESSFULLY COMPLETED OUR MOVE FROM SAN
10	FRANCISCO TO OAKLAND. WE'RE UP AND RUNNING. IT
11	WORKED. THERE ARE A NUMBER OF PEOPLE THAT DID A
12	PHENOMENAL JOB. AND THEN THERE WAS AMANDA MORA WHO
13	BASICALLY HAS LIVED THIS MOVE AND HAS TAKEN
14	OWNERSHIP OF MAKING IT BE SUCCESSFUL AND HAS DONE A
15	PHENOMENAL JOB. AND I THANK HER FOR HER DEDICATION
16	AND REALLY OWNERSHIP OF THAT PROGRAM.
17	WE ARE THERE. I WANT TO TELL THE BOARD
18	WE'RE THERE AND WE'RE UP AND WE'RE RUNNING, BUT
19	THIS, IN CONJUNCTION WITH YOU HAVE A NEW LEADER AND
20	WE HAVE A NEW STRATEGIC PLAN AND WE HAVE A REFINED
21	MISSION AND WE HAVE A NEW CULTURE, IS A TREMENDOUS
22	AMOUNT OF CHANGE THAT THE ORGANIZATION IS DIGESTING.
23	AND SO YOU'VE HEARD WHO MOVED MY CHEESE. WE DIDN'T
24	MOVE THEIR CHEESE. WE PUNTED THEIR CHEESE OUT THE
25	WINDOW AND ALL THE WAY ACROSS THE BAY. AND I JUST
	22

1	WANT TO TELL THE BOARD HOW SINCERELY I APPRECIATE
2	THE FLEXIBILITY AND THE COMMITMENT AND THE
3	DETERMINATION OF THIS GROUP OF PROFESSIONALS BEHIND
4	ME AND THE REMAINDER OF THE GROUP THAT STAYED IN SAN
5	FRANCISCO AND THEIR WILLINGNESS TO WALK WITH US ON
6	THIS JOURNEY AND GO THROUGH THESE KINDS OF CHANGES.
7	IT IS NOT EASY. IT'S TAXING. THERE'S ANXIETY
8	ASSOCIATED WITH IT. AND I VERY MUCH WANT YOU TO
9	KNOW HOW MUCH I APPRECIATE ALL THEY'RE DOING AND
10	THEIR CONTINUED OUTSTANDING EFFORT AS WE'VE GONE
11	THROUGH THIS. SO THANK YOU.
12	CHAIRMAN THOMAS: THANK YOU, DR. MILLS.
13	WE'RE GOING TO MOVE ON. WE HAVE THE
14	CONSENT CALENDAR HERE. IS THERE ANYTHING ON THIS
15	CONSENT ITEM THAT ANYBODY ON THE BOARD WOULD LIKE
16	REMOVED? HEARING NOBODY, DO I HAVE A MOTION TO
17	APPROVE?
18	MR. TORRES: MOVE TO ADOPT.
19	MS. LANSING: SECOND.
20	CHAIRMAN THOMAS: MOVED BY SENATOR TORRES.
21	SECONDED BY MS. LANSING. ALL THOSE IN FAVOR PLEASE
22	SAY AYE. THOSE ON THE PHONE?
23	MS. BONNEVILLE: LINDA BOXER.
24	DR. BOXER: YES.
25	MS. BONNEVILLE: KATHY LAPORTE.
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1	MS. LAPORTE: YES.
2	CHAIRMAN THOMAS: THANK YOU. WE'RE
3	APPROVED.
4	ON TO THE ACTION ITEMS. THE FIRST ITEM,
5	DR. MILLS, YOU'RE BACK UP WITH THE DISCUSSION OF THE
6	CIRM STRATEGIC PLAN.
7	DR. MILLS: THANK YOU, CHAIRMAN THOMAS AND
8	THE BOARD. IT'S BEEN AWHILE SINCE WE'VE SPOKEN. SO
9	TODAY I WOULD LIKE TO TAKE YOU AND OBVIOUSLY MEMBERS
10	OF THE PUBLIC THROUGH OUR PROPOSED AND WHAT WE HOPE
11	IS THE FINAL VERSION OF OUR STRATEGIC PLAN. WE'VE
12	GONE THROUGH OBVIOUSLY SEVERAL ITERATIONS AND
13	REFINING AND DRIVING. WE'VE HAD COMMENT FROM THE
14	PUBLIC; WE'VE HAD COMMENT FROM BOARD. WE HAD A
15	DRAFT VERSION OBVIOUSLY THAT WE PUT OUT TO SCIENCE
16	SUBCOMMITTEE AND EVERYONE ELSE, AND THEN WE'VE
17	GOTTEN SOME GREAT COMMENTS AND GREAT FEEDBACK. I AM
18	VERY EXCITED ABOUT THIS PLAN, AND I THINK WHAT WE
19	WILL BE ABLE AS AN AGENCY TO DO GOING FORWARD, IF WE
20	SO CHOOSE, YOU SO CHOOSE, TO ADOPT IT.
21	TO START PERHAPS, I COULD ASK JEFF SHEEHY,
22	WHO IS CHAIR OF THE SCIENCE SUBCOMMITTEE, TO MAYBE
23	GIVE A BRIEF INTRODUCTION AS THE PLAN WENT TO THE
24	SCIENCE SUBCOMMITTEE FOR APPROVAL BEFORE COMING TO
25	THE FULL BOARD.
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1	MR. SHEEHY: THANK YOU, RANDY. AND I ALSO
2	JUST WANT TO FOLLOW UP ON YOUR REMARKS ABOUT THE
3	INCREDIBLE EFFORTS OF THE TEAM OVER THE LAST YEAR.
4	THEY REALLY HAVE GONE THROUGH A LOT OF CHANGES, AND
5	WE REALLY OWE A LOT TO THESE INDIVIDUALS WHO HAVE
6	STAYED WITH US AND HAVE CREATED SUCH I MEAN THE
7	LEVEL OF INNOVATION WE'VE SEEN OVER THE LAST YEAR
8	HAS JUST BEEN ASTONISHING FROM MY PERSPECTIVE. IT'S
9	TAKEN A LOT OF REALLY, REALLY HARD WORK AND
10	DEDICATION AND PASSION. AND HAVING SEEN A LOT OF
11	THAT TAKE PLACE, I AM SO PROUD TO BE A MEMBER OF
12	THIS ORGANIZATION.
13	AND THIS STRATEGIC PLAN, I THINK, IS JUST
14	ONE MORE OUTCOME OF THIS INCREDIBLE WORK THAT WE'RE
15	SEEING DONE BY OUR TEAM.
16	THE SCIENCE SUBCOMMITTEE LOOKED AT THIS
17	EXHAUSTIVELY, ASKED VERY, VERY TOUGH QUESTIONS, AND
18	I THINK THE RECEPTION WAS INCREDIBLY POSITIVE. AND,
19	AGAIN, TALKING ABOUT THE TEAM AND DR. MILLS, THE
20	INNOVATION THAT'S INCLUDED IN THIS PLAN IS SOMETHING
21	I DON'T THINK WE'VE SEEN BEFORE OR A PUBLIC FUNDING
22	AGENCY HASN'T DONE BEFORE. AND I THINK IT'S HIGH
23	RISK, BUT IT'S ALSO HIGH REWARD. WE USE THOSE TERMS
24	A LOT, BUT THIS REALLY IS TRUE IN THIS INSTANCE. I
25	THINK IT HAS THE POTENTIAL TO REALLY, REALLY

1	TRANSFORM THE WHOLE FIELD IF IT WORKS OUT.
2	BUT I ALSO I THINK ALL OF US ARE VERY
3	REALISTIC THAT, LIKE EVERYTHING WE'VE DONE, IT'S
4	ALWAYS A WORK IN PROGRESS. NOTHING SET IN STONE.
5	AND I'M SURE THAT THEY'LL INNOVATE AS WE GO ALONG
6	BASED ON THE OUTCOMES THAT THEY GET FROM THE
7	IMPLEMENTATION OF THE PLAN.
8	I DON'T KNOW IF DR. STEWARD HAS ANYTHING
9	OR ANY OTHER MEMBER OF THE SCIENCE SUBCOMMITTEE, BUT
10	IT WAS TRULY EXCITING TO SEE THE PRODUCT THAT'S COME
11	OUT. I REALLY PERSONALLY WANT TO SAY HOW MUCH I
12	APPRECIATE THE METRICS THAT ARE INCLUDED IN THIS
13	PLAN. THERE'S SOME VERY DEFINITIVE OUTCOME METRICS,
14	IF YOU LOOK IN THE BACK, I THINK IT'S IN THE
15	APPENDIX, THAT ARE VERY CLEAR. AND I ALWAYS
16	APPRECIATE HAVING CLEAR METRICS. I THINK IF YOU
17	CAN'T MEASURE IT, YOU DON'T KNOW IF YOU'RE DOING IT.
18	WHAT I ALSO THOUGHT WAS VERY INTERESTING
19	IS THERE'S ACTUALLY PROCESS METRICS FOR EACH TEAM
20	WITHIN THE LARGER CIRM TEAM THAT, AS I UNDERSTAND
21	IT, THE TEAMS COLLABORATIVELY CAME UP WITH THESE
22	METRICS IN ORDER TO MEASURE THEIR OWN PROGRESS
23	TOWARDS ADVANCING THIS TOWARDS ACHIEVING THE LARGER
24	OUTCOME METRICS OF THE STRATEGIC PLAN. I THINK
25	THAT'S ALSO INCREDIBLY INNOVATIVE AND VERY HELPFUL

IN TERMS OF GETTING TO ACHIEVE THE KIND OF SUCCESS
WE HOPE TO ACHIEVE WITH THIS PLAN.
DR. MILLS: THANK YOU, JEFF.
MR. HIGGINS: I WOULD JUST LIKE TO ADD A
COUPLE COMMENTS TO JEFF'S COMMENTS. EVERYBODY KNOWS
THAT STRAT PLANS ARE BORING AND THAT THEY USUALLY
GET DONE UNDER THE RECOMMENDATION OF A CONSULTANT
AND THEY GO IN A DESK DRAWER AS SOON AS THE
CONSULTANT LEAVES TOWN WITH THE CHECK. I WANT TO
SAY THAT THIS STRAT PLAN IS NONE OF THAT. THIS IS
AN EXCITING DOCUMENT. IT'S GOING TO BE A LIVING
DOCUMENT THAT WE CAN FOLLOW AND USE FOR GUIDANCE.
AND I WOULD ENCOURAGE EVERYONE TO STUDY IT, LOOK AT
IT, AND APPRECIATE THE DIFFERENCE FROM ANY STRAT
PLAN YOU'VE EVER BEEN INVOLVED IN.
SO IF YOU'VE BEEN INVOLVED IN STRAT PLAN
DEVELOPMENT IN YOUR PAST LIVES OR YOUR CURRENT
LIVES, THIS IS NOT THAT. THIS IS SOMETHING TOTALLY
DIFFERENT. AND I THINK THE BOARD CANNOT JUST
APPROVE THIS. I THINK THE BOARD CAN GET BEHIND IT
WITH EXCITEMENT.
DR. MILLS: THANK YOU. SO LET'S GET INTO
THIS. WE'VE SET THE BAR HIGH NOW, AND I FEEL A
LITTLE INTIMIDATED TO DELIVER, BUT WE'RE GOING TO
TRY.
37

1	THE FIRST THING THAT I'D LIKE TO SAY IS AS
2	WE GO THROUGH THE STRAT PLAN, YOU'LL SEE THERE'S A
3	LOT OF USE OF METAPHOR, AND THERE'S A LOT OF
4	DESCRIPTIONS AND CARTOONS AND TRUCKS AND THINGS LIKE
5	THAT. AND THE PURPOSE OF THAT REALLY IS TO DRIVE
6	HOME CLARITY OF WHAT WE'RE TRYING TO ACCOMPLISH AND
7	HAVE THAT DIGESTIBLE. AND I WANT TO RECOGNIZE ONE
8	BOARD MEMBER WHO REALLY TOOK US TO TASK ON THAT AND
9	PUSHED US TO DO THAT. AND THAT IS ACTUALLY LAUREN
10	MILLER, WHO WE SAT DOWN WITH AND SAID, "LOOK, YOU'RE
11	NOT EFFECTIVELY COMMUNICATING ANYTHING IF EVERYONE
12	CAN'T UNDERSTAND IT. AND WE NEED TO HAVE CLEAR
13	COMMUNICATION THAT ALL OF OUR STAKEHOLDERS CAN
14	UNDERSTAND AND PULL OUT THE COMPLEXITY." SO WE
15	REALLY TRIED TO DO THAT.
16	I WAS ASKED ABOUT THE STRATEGIC PLAN. SO
17	WHAT'S DIFFERENT ABOUT THIS STRATEGIC PLAN THAN THE
18	LAST STRATEGIC PLAN? I SAID WHAT ARE THE THREE MOST
19	IMPORTANT THINGS FROM THE LAST STRATEGIC PLAN? THE
20	RESPONSE WAS I DON'T KNOW. I SAID THAT'S HOPEFULLY
21	GOING TO BE THE DIFFERENCE. I DO HOPE AT THE END OF
22	THIS YOU WILL KNOW AT LEAST THE THREE MOST IMPORTANT
23	THINGS THAT WE'RE TRYING TO DO.
24	SO TO GET ON WITH IT, THIS IS THE TABLE OF
25	CONTENTS FOR THE STRATEGIC PLAN. IT ENDED UP BEING
	20

1	A LITTLE LONGER THAN WE WANTED, ABOUT 50 PAGES. WE
2	WERE AIMING FOR 30, BUT IT JUST HAPPENED THAT WAY.
3	AND I'M GOING TO GO THROUGH THESE VARIOUS ELEMENTS
4	TODAY. THE PLAN ITSELF REALLY CENTERS FROM PAGE 4
5	OR SECTION 4 TO SECTION 10 WHERE WE TALK GOING
6	FORWARD ABOUT WHAT WE'RE GOING TO DO TO ACCOMPLISH
7	OUR MISSION.
8	SO, AGAIN, A LITTLE BIT ABOUT THE
9	STRATEGIC PLANNING PROCESS THAT WE EMPLOYED. NO
10	CONSULTANTS. WE DIDN'T MAKE IT OVERLY COMPLEX.
11	REALLY SUCCESSFUL STRATEGIC PLANNING CAN BE
12	ACCOMPLISHED IF YOU DO JUST THREE THINGS. ONE IS
13	VERY, VERY HONESTLY ASSESS WHERE YOU ARE NOW, WHERE
14	THE ENVIRONMENT IS NOW, ASK THE RIGHT QUESTIONS, ASK
15	AS MANY PEOPLE AS YOU CAN, AND GET A GOOD SENSE OF
16	WHERE YOU ARE.
17	THEN, TWO, FIGURE OUT ASPIRATIONALLY WHERE
18	IT IS YOU WANT TO GO. WHAT DOES GOOD LOOK LIKE
19	SOMETIMES I'LL REFER TO IT AS. AND THEN SIMPLY COME
20	UP WITH A STRATEGY THAT CONNECTS THOSE DOTS.
21	STRATEGIC PLANNING DOESN'T HAVE TO BE MORE
22	COMPLICATED THAN THAT.
23	IF WE ARE SUCCESSFUL, WE SHOULD BE ABLE TO
24	GET THESE THREE THINGS OUT OF THE STRATEGIC PLAN.
25	FIRST IS SITUATIONAL AWARENESS. IT'S NOT A GOOD
	20

1	IDEA TO STRATEGIC PLAN CONTINUOUSLY BECAUSE YOU
2	DON'T WANT TO KEEP CHANGING WHERE THE GOAL LINE IS.
3	BUT EVERY ONCE IN A WHILE, IT IS A GOOD IDEA TO COME
4	UP AND REASSESS THE WORLD. ONE OF THE THINGS THAT
5	WE SAW IN THIS STRATEGIC PLAN WAS HOW MUCH THE WORLD
6	HAS CHANGED SINCE 2004. AND I THINK HOW PROUD CIRM
7	CAN BE FOR HAVING DRIVEN SUCH A SIGNIFICANT CHANGE
8	IN THE WORLD AROUND STEM CELLS. THIS PLAN WOULD NOT
9	HAVE TO BE SO DIFFERENT IF CIRM HAD NOT BEEN SO
10	SUCCESSFUL IN ITS FIRST TEN YEARS OF LIFE.
11	SECONDLY, WHAT WE WERE TRYING TO DO WITH
12	THIS PLAN IS CREATE ORGANIZATIONAL CLARITY. SO THE
13	BOARD, THE LEADERSHIP TEAM, THE TEAM AT CIRM, CIRM
14	STAKEHOLDERS, EVERYONE HAVING A CLEAR UNDERSTANDING
15	AND BEING COMPLETELY ALIGNED ON WHAT WE'RE TRYING TO
16	DO AND HOW WE'RE TRYING TO DO IT.
17	LASTLY, AS JEFF SAID, A STRATEGIC PLAN
18	WITHOUT MEASURABLE GOALS MIGHT AS WELL BE PUT IN THE
19	DESK DRAWER AND FORGOTTEN ABOUT BECAUSE YOU CAN'T
20	COME BACK IN FIVE YEARS AND SAY I WONDER HOW THAT
21	ALL WORKED OUT. THIS IS A DIFFICULT PLAN, AND IT IS
22	GOING TO TAKE A TREMENDOUS COMMITMENT FOR US TO
23	ACCOMPLISH. NOTHING IN THIS PLAN REQUIRES US TO
24	BEND THIS TIME SPACE CONTINUUM. BUT IF WE DO NOT
25	WORK VERY HARD, WE CERTAINLY WILL NOT ACCOMPLISH IT.

1	THESE GOALS ARE NOT EASY-TO-ACCOMPLISH GOALS. EASY
2	WAS NOT ALLOWED TO BE BROUGHT UP AS A GOAL OF THIS
3	STRATEGIC PLAN. INSTEAD, WHAT WE WANTED TO FIGURE
4	OUT WAS HOW COULD WE HAVE THE BIGGEST IMPACT
5	POSSIBLE WITH OUR REMAINING FUNDS AND OUR REMAINING
6	LIFE IN A WAY, AS JEFF SAID, THAT WOULD LITERALLY
7	TRANSFORM REGENERATIVE MEDICINE OVER THE NEXT FIVE
8	YEARS.
9	EVERYTHING BETWEEN US, WHERE WE WERE TODAY
10	IN OUR GOAL, HAVING THESE THERAPIES AVAILABLE TO
11	PATIENTS IN NEED WAS IN PLAY. EVERYTHING WE PUT ON
12	THE TABLE, AND THAT'S THE PLAN THAT WE CAME UP WITH
13	HERE TODAY.
14	SO FIRST THING, WE JUST TALKED ABOUT THIS
15	A SECOND AGO, BUT THE FIRST THING, WE HAD TO ASSESS
16	WHAT ACTUALLY WAS OUR RUNWAY AND HOW MUCH TIME AND
17	MONEY ARE WE TALKING ABOUT. WE WENT OVER THIS
18	PREVIOUSLY, SO I WON'T SPEND TOO MUCH TIME ON IT,
19	BUT WE ARE ESTIMATING THAT WE HAVE ABOUT FIVE YEARS
20	IF WE AIM FOR SOMEWHERE BETWEEN 190 TO \$200 MILLION
21	IN NEW AWARDS EACH YEAR. AGAIN, THAT DOES ASSUME AN
22	AWARD RECAPTURE RATE OF ABOUT 10 PERCENT. IT'S BEEN
23	A LITTLE HIGHER OF LATE. THAT WOULD GIVE US A
24	LITTLE EXTRA MONEY TOWARDS THE END OF THIS PROCESS
25	TO REDISTRIBUTE.

1	SO I WENT THROUGH THIS ON THE FIRST PASS
2	OF PUTTING OUT THE STRATEGIC PLAN, SO I'M GOING TO
3	CONDENSE A WHOLE BUNCH OF SLIDES JUST INTO SORT OF
4	SUMMARIES OF THEM. GETTING A GOOD FEELING FOR WHERE
5	WE WERE TODAY AND HOW THINGS HAVE CHANGED WAS
6	IMPORTANT. SO THE FIRST THING, THE PUNCHLINE THAT
7	JUMPED OFF THE TABLE WAS THAT CIRM INITIALLY EXISTED
8	AS AN INITIATIVE-BASED AGENCY. AND THE REASON FOR
9	THIS CENTERED AROUND, AGAIN, THIS HOW MUCH THE WORLD
10	HAS CHANGED. IN 2004 THERE WASN'T ENOUGH DEMAND IN
11	ALL OF THE DIFFERENT AREAS THAT CIRM FUNDS FOR US TO
12	HAVE REGULARLY SCHEDULED PROGRAMS IN ALL OF THOSE
13	DIFFERENT AREAS. THIS IS MORE ANALOGOUS TO CHARTER
14	FLIGHT. IF YOU WANTED TO START GOING ACROSS THE
15	COUNTRY, WHEN THERE WAS ENOUGH PEOPLE AT THE
16	AIRPORT, WE'D CHARTER A FLIGHT AND WE WOULD TAKE YOU
17	TO SALT LAKE CITY. AND THEN WHEN THERE WAS ENOUGH
18	PEOPLE THERE, WE WOULD SCHEDULE THE NEXT FLIGHT OUT
19	TO OKLAHOMA OR SOMETHING LIKE THAT. THAT WAS AN
20	INITIATIVE-BASED APPROACH.
21	WE NOW HAVE SO MUCH DEMAND THAT WE CAN NOW
22	HAVE REGULARLY SCHEDULED SERVICE BETWEEN THESE
23	VARIOUS POINTS WHERE WE CONTINUOUSLY HAVE FLIGHTS
24	GOING. WHEN YOU SHOW UP TO THE AIRPORT, YOU KNOW
25	THERE'S A PLANE THAT'S GETTING READY TO LEAVE AND
	42

1	IT'S GOING TO TAKE YOU ON TO THE NEXT PLACE.
2	SYSTEMS-BASED VERSUS INITIATIVE IS A BIG CONCEPT
3	THAT CAME OUT OF THIS.
4	THE SECOND WAS WE DID THESE SURVEYS AND
5	POLLS AND WE ASKED A LOT OF PEOPLE. AND I THINK ONE
6	OF THE MOST GRATIFYING THINGS THAT WE FOUND WAS ON
7	ALMOST EVERY ISSUE ALL OF OUR STAKEHOLDERS WERE
8	ALIGNED. WE ARE IN COMPLETE ALIGNMENT AROUND THE
9	MOST IMPORTANT ISSUE, WHICH CENTERS AROUND WHY THE
10	AGENCY EXISTS.
11	WE DID FIND THAT THERE WAS A PARTICULAR
12	STAGE OF RESEARCH THAT NEEDED HELP. SO THIS
13	TRANSLATIONAL STAGE, I'LL BE TALKING MORE ABOUT
14	THIS. THAT IS FROM THE TIME WE DISCOVER A STEM CELL
15	PRODUCT THAT LOOKS PROMISING TO THE TIME WE CAN
16	ACTUALLY GET AN IND APPROVED BY THE FDA TO WHERE WE
17	CAN START DOING CLINICAL TRIALS, THAT RIGHT NOW FOR
18	STEM CELL THERAPIES IS TOO LONG. IT'S SOMEWHERE
19	BETWEEN SIX TO EIGHT YEARS. THE INDUSTRY AVERAGE
20	FOR ANYTHING THAT'S NOT A STEM CELL IS 3.2 YEARS.
21	SO WE LOOK AT THAT AND WE SAY, OKAY, THAT'S A
22	PROBLEM. WE CAN ALSO LOOK AT THAT AND SAY WHAT IF
23	WE GOT IT BACK TO INDUSTRY AVERAGE? WE COULD
24	ACTUALLY CUT IN HALF TRANSLATIONAL TIME. WOULDN'T
25	THAT BE GREAT? SO THAT'S SOMETHING WE DUG IN DEEPER

1	TO.
2	WE ALSO KNOW THAT STEM CELL THERAPIES FROM
3	A COMMERCIAL STANDPOINT CLEARLY ARE DISADVANTAGED.
4	SO BIG PHARMA COMPANIES WILL DISPROPORTIONATELY
5	IN-LICENSE NON-CELL TECHNOLOGIES AT A MUCH GREATER
6	RATE THAN CELL TECHNOLOGIES. ONLY 8 PERCENT OF
7	CIRM'S ACADEMIC PROGRAMS ACTUALLY HAVE PARTNERS.
8	AND WE NEED TO ADDRESS THE BACK END OF THIS PROBLEM
9	IF WE'RE ACTUALLY GOING TO GET THERAPIES ALL THE WAY
10	TO PATIENTS. WE NEED MORE COMMERCIAL INTEREST.
11	AND THEN THE LAST THING, AND THIS WAS
12	FAIRLY SHOCKING, WAS THE REGULATORY ENVIRONMENT IS
13	SEEN AS A MAJOR IMPEDIMENT. IN FACT, 70 PERCENT OF
14	RESPONDENTS TO OUR SURVEYS LISTED FDA AS THE SINGLE
15	BIGGEST IMPEDIMENT TO DEVELOPING STEM CELL
16	THERAPIES. WE'RE GOING TO TALK MORE ABOUT THAT.
17	AS I SAID, THE MOST GRATIFYING THING WE
18	HAD WAS AMONG THE BOARD COMPLETE AGREEMENT WHAT OUR
19	MISSION SHOULD BE, AND WE DIDN'T HAVE TO GO THROUGH
20	THE BURDENSOME EXERCISE OF TRYING TO FIGURE OUT WHAT
21	A NEW MISSION FOR THE ORGANIZATION SHOULD BE. AND
22	SO OUR MISSION TO ACCELERATE STEM CELL TREATMENTS TO
23	PATIENTS WITH UNMET MEDICAL NEEDS WAS 100 PERCENT
24	AGREED UPON BY THE BOARD AND BY ALL OTHER
25	STAKEHOLDERS, ANOTHER 215 PEOPLE OR SO THAT LOOKED

1	AT IT, 219, WAS 95.4 PERCENT. SO THAT'S WHERE WE
2	STARTED. WE VIEW THAT, THEN, AS SORT OF OUR
3	UNMOVABLE, IMMOVABLE POINT OF REFERENCE THAT WE WILL
4	USE FOR EVERYTHING ELSE, AND WE'LL TEST OFF AGAINST
5	IT TO MAKE SURE. IRRESPECTIVE OF WHAT STRATEGIES WE
6	MIGHT USE OR METHODS WE MIGHT USE, WE'RE ALWAYS
7	HEADING OR WORKING TOWARDS THAT DIRECTION.
8	SO THEN WE WORKED ON A VISION, WHICH IS TO
9	EXPONENTIALLY ADVANCE CIRM'S MISSION BY LEADING A
10	COORDINATED CAMPAIGN THAT HOLISTICALLY ATTACKS THE
11	OBSTACLES MEANINGFULLY AFFECTING THE SPEED
12	PROBABILITY AND SUSTAINABILITY OF STEM CELL
13	TREATMENTS TO HELP PATIENTS IN NEED.
14	ONLY A COUPLE OF WORDS HERE THAT I WANT TO
15	POINT OUT. ACTUALLY THE PLAN WE BREAK OUT LITERALLY
16	EVERY WORD AND WHY EVERY WORD WAS CHOSEN. THE WORD
17	"EXPONENTIALLY," THIS IS A THEME THROUGHOUT THIS
18	PLAN. WE ARE NOT LOOKING TO DO A LITTLE BETTER. WE
19	WEREN'T LOOKING TO DO THAT WHICH WAS EASILY
20	ACCOMPLISHABLE. AS JEFF SAID, HIGH RISK, HIGH
21	REWARD. THIS IS DOABLE, BUT IT IS GOING TO BE HARD.
22	AND WHAT WE WANTED THE RESULT TO BE WAS, AS SHERRY
23	LANSING SAID IN HER COMMENTS IN THE PRESS RELEASE,
24	WE WANT IT TO BE WORTH DOING. IF WE PUT ALL THIS
25	HARD WORK AND DEDICATION AND RESOURCE INTO IT, ALL

1	OF THAT EFFORT IS WORTH THIS OUTCOME BECAUSE THE
2	OUTCOME WILL BE TRANSFORMATIONAL. AND THAT'S
3	IMPORTANT.
4	THE SECOND THING THAT'S A LITTLE DIFFERENT
5	HERE IN STRATEGY IS THE WORD "HOLISTICALLY ATTACKS"
6	OR THE PHRASE "HOLISTICALLY ATTACKS." BECAUSE, AS I
7	SAID, WE VIEWED EVERYTHING BETWEEN US AND STEM CELL
8	THERAPIES GETTING TO PATIENTS IN NEED AS FAIR GAME
9	FOR CIRM. THAT'S WHY YOU SEE THINGS LIKE WE HAVE TO
10	FIGURE OUT A WAY TO GET MORE INDUSTRY INVOLVEMENT OR
11	WE HAVE TO FIND A WAY TO FACILITATE CLINICAL
12	OPERATIONS OR WE NEED TO DEAL WITH REGULATORY
13	PROBLEMS THAT ARE SLOWING US DOWN. EVERYTHING WAS
14	FAIR GAME BETWEEN US AND THEM.
15	WITH THAT, WE CAME UP WITH THREE STRATEGIC
16	THEMES. WE TALKED ABOUT THIS BEFORE. THE GIANT
17	STEM CELL BOULDER OF LOVE AND HAPPINESS THAT WE'RE
18	TRYING TO PUSH OVER THE MOUNTAIN TO THE VALLEY OF
19	THE PATIENTS BELOW. BECAUSE WE'RE NOT GOING TO HURT
20	THEM, WE'RE GOING TO CRUSH THEM WITH LOVE AND
21	HAPPINESS AND JOY. THAT'S THE ONE PART OF THE
22	METAPHOR THAT BREAKS DOWN, WHICH IS WHY I KEEP
23	COUCHING IT IN THAT LIGHT. IT IS A STEM CELL
24	BOULDER OF LOVE AND HAPPINESS, JUST TO BE CLEAR.
25	BUT WE ARE GOING TO DO THAT BY WHAT WE'VE
	46

1	DONE HISTORICALLY AS AN AGENCY, WHICH IS PUSH AND
2	PUSH REALLY HARD BECAUSE THAT'S GOOD. AND WE'VE
3	BEEN DOING IT. WE CAN GET BETTER AT IT. WE'RE
4	GOING TO CONTINUE TO GET BETTER AT IT AND COORDINATE
5	OUR EFFORTS SO WE GET MAXIMAL EFFECT THERE. BUT WE
6	ALSO NEED TO ENGAGE DOWNSTREAM DEMAND IN THAT. WE
7	NEED TO HAVE PULL COME INTO THIS. WE NEED TO HAVE
8	OTHER PEOPLE HELPING US PULL THIS BOULDER WHILE
9	WE'RE PUSHING ON IT.
10	AND THEN THE LAST THING IS WE NEED TO TAKE
11	A LONG, HARD LOOK AT THIS MOUNTAIN THAT WE'RE GOING
12	OVER FROM A REGULATORY PERSPECTIVE BECAUSE THIS
13	PARADIGM HAS BEEN IN PLACE FOR 15 YEARS, AND IT
14	STILL HAS A SCORE OF ZERO ON THE SCOREBOARD.
15	NOTHING HAS BEEN APPROVED DESPITE 15 YEARS OF A
16	REGULATORY PARADIGM BEING IN PLACE AND NOTHING'S
17	CLOSE, AND WE NEED TO LOOK AT WHY THAT IS.
18	SO WITH THESE THREE THEMES IN PLACE, PUSH,
19	PULL AND LEVEL, WE THEN ARE ABLE TO GO ON TO
20	SPECIFIC ACTIONS. AND I WILL TALK ABOUT THESE. I
21	HAVE SLIDES ON EACH OF THESE IN DETAIL.
22	SO THE FIRST PUSHING ACTION THAT WE HAVE,
23	IT REALLY GOES TO THIS CONCEPT OF CREATING A
24	SYSTEMS-BASED APPROACH VERSUS AN INITIATIVE-BASED
25	APPROACH. WE'VE SPENT OVER THE FIRST TEN YEARS

1	BUILDING THESE BEAUTIFUL PIECES OF THIS ENGINE, AND
2	NOW WHAT WE'RE DOING IS WE'RE TAKING THOSE BEAUTIFUL
3	PIECES AND WE'RE ASSEMBLING THEM INTO SOMETHING THAT
4	WILL INTEGRATE THEM AND HAVE THEM WORK TOGETHER IN A
5	WAY THAT PRODUCES THE MOST THRUST. IT'S A REALLY,
6	REALLY IMPORTANT PART OF THIS STRATEGIC PLAN, THAT
7	EVERY PROGRAM THAT CIRM HAS WORK TOGETHER AND PUSH
8	IN THE SAME DIRECTION AS EVERY OTHER PROGRAM SO THAT
9	THEY'RE ALL ALIGNED SO WE CAN CREATE THE MOST
10	EFFECT.
11	THE SECOND THING WE'VE DONE IS WORK ON
12	OPERATIONAL EXCELLENCE. NOW, WITH YOUR HELP
13	THROUGHOUT THIS YEAR, WE HAVE NOW FULLY IMPLEMENTED
14	CIRM 2.0 FOR EVERYTHING FROM THE EARLIEST STAGE
15	RESEARCH IN DISCOVERY THROUGH TRANSLATIONAL AND
16	CLINICAL RESEARCH. WE JUST HAD A REVIEW ON HOW THE
17	CLINICAL PART OF IT HAS GONE. IT'S GOING VERY WELL.
18	IT IS DEFINITELY FASTER; IT'S PREDICTABLE. WE HAVE
19	INCENTIVIZED THE RIGHT THINGS, WHICH IS A VERY BOLD
20	THING FOR US TO DO, AND WE HAVE INCREASED OUR LEVEL
21	OF PARTNERSHIP, NOT JUST WITH THE AWARDEES THAT WE
22	WORK WITH, BUT WITH THE PATIENT COMMUNITIES AS WELL.
23	RECALL UNDER THE CIRM 2.0 CLINICAL PROGRAMS, EVERY
24	CLINICAL PROGRAM THAT GETS LAUNCHED HAS A CLINICAL
25	ADVISORY PANEL. EVERY ONE OF THOSE HAS TO HAVE AT

1	LEAST ONE PATIENT REPRESENTATIVE ON IT THAT CAN HELP
2	DRIVE THE PROGRAM.
3	THAT HAS HAPPENED. THAT IS IN PLACE. AND
4	THE INPUT FROM THAT PARTNERSHIP OF CIRM, THE
5	AWARDEES, AND THE PATIENTS HAS MADE VERY, VERY
6	MEANINGFUL DIFFERENCES SO FAR. I THINK DR. DOYLE IS
7	GOING TO TALK MORE ABOUT THAT COMING UP. BUT THIS
8	DRIVE FOR OPERATIONAL EXCELLENCE WE HAVE STARTED.
9	IT IS NOT SOMETHING WE FINISH. WE DON'T EVER GET
10	THERE. WE CAN ALWAYS GET BETTER BASICALLY IN HOW WE
11	PUSH THIS BOULDER UP THE HILL AND HOW WE MAKE THE
12	ENGINE RUN FROM AN EFFICIENCY STANDPOINT. WE DO
13	THAT BY MEASURING, MONITORING, BEING VERY
14	SELF-EFFACED AND OPEN TO IDEAS ON HOW TO MAKE THINGS
15	BETTER.
16	ANOTHER THING WE'RE DOING FROM A PUSH
17	STANDPOINT IS WE CALL THIS THE CIRM PITCHING
18	MACHINE. THIS GOES AROUND TO WE HAVE THIS UNUSUALLY
19	LONG TRANSLATIONAL TIME FOR STEM CELL THERAPY THAT'S
20	DIFFERENT THAN FOR THINGS THAT ARE SMALL MOLECULES.
21	AND WE IDENTIFIED REALLY TWO AREAS WHERE WE COULD
22	HAVE AN IMPACT ON THAT. FIRST IS THROUGH WHAT WE
23	CALL A TRANSLATIONAL CENTER. I DON'T KNOW WHAT
24	WE'RE ACTUALLY GOING TO CALL IT, BUT RIGHT NOW WE
25	CALL IT A TRANSLATIONAL CENTER. AND THE
	40

1	TRANSLATIONAL CENTER IS FOCUSED ON DOING ALL OF
2	THOSE KINDS OF REGULATORY STUDIES THAT TYPICAL
3	ACADEMIC INVESTIGATORS, ONE, HAVE NO EXPERIENCE IN
4	DOING, BUT, MOST IMPORTANTLY, TWO, HAVE NO INTEREST
5	IN DOING. WE WENT AROUND TO, WITHOUT EXCEPTION,
6	EVERY UNIVERSITY, EVERY MAJOR RESEARCH CENTER THAT
7	WE WENT TO AND WE SAT DOWN AND WE TALKED TO SAID
8	THEY WOULD LOVE HELP OR A PLACE WHERE THEY CAN GO
9	DO, IN THEIR TERMS, THE UNINTERESTING, BUT NECESSARY
10	REGULATORY STUDIES THAT ARE REQUIRED IN ORDER TO GET
11	AN IND. THESE ARE THINGS LIKE DOING STABILITY
12	STUDIES OR PRECLINICAL TOX STUDIES THAT ARE SORT OF
13	CHECK-BOX THINGS THAT THE FDA REQUIRES, BUT THE FDA
14	HAS VERY SPECIFIC WAYS IN WHICH THEY WANT THEM DONE.
15	SO THE IDEA OF CREATING THIS TRANSLATING
16	CENTER THAT WOULD THEN WORK IN COORDINATION WITH
17	WHAT WE CALL THE ACCELERATING CENTER. IT'S A FANCY
18	WORD FOR A STEM CELL-SPECIFIC CRO THAT WOULD WORK
19	WITH THE TRANSLATING CENTER, CAPTURE ALL OF THEIR
20	IND-ENABLING STUDIES, AND COMPILE AN IND THAT WOULD
21	HAVE THE BEST CHANCE OF GETTING APPROVAL BY THE FDA
22	IN A TIMELY FASHION.
23	NOW, BECAUSE THESE TWO CENTERS WILL BE
24	FOCUSED ONLY ON THIS, THEY WOULD BE GOOD AT IT, ONE.
25	TWO, WHEN WE TALKED TO FDA ABOUT THIS IDEA, THEY

1	LOVED IT. AND THE REASON THE FDA LOVED IT WAS
2	BECAUSE RIGHT NOW THEY TALK TO A WIDE VARIETY OF
3	RESEARCHERS, MANY OF WHICH ARE FUNDED BY CIRM, THAT
4	HAVE VARYING LEVELS OF UNDERSTANDING OF HOW TO GET
5	AN IND AND THE IND FILING PROCESS AND THE STUDIES
6	THAT WOULD REQUIRE. AND THEY TOLD US IF THERE WAS A
7	SINGLE POINT OF CONTACT THAT WE COULD WORK WITH, WE
8	COULD SET UP WEEKLY CALLS, WE COULD DO A LOT OF
9	DIFFERENT WAYS WHERE WE CAN PARTNER IN ORDER TO
10	BASICALLY BUNDLE UP ALL OF THESE DIFFERENT THINGS
11	AND VERY EFFICIENTLY MOVE THIS RESEARCH FORWARD.
12	AGAIN, THERE'S A LOT OF OPPORTUNITY IN
13	THIS TRANSLATIONAL PHASE FOR US TO GO AFTER. WE CAN
14	LITERALLY CUT THIS PHASE IN HALF. SO THE RESEARCH
15	CENTERS ARE EXCITED ABOUT IT, AND THE FDA WAS
16	EXCITED ABOUT IT, AND I WAS EXCITED ABOUT IT. AND
17	THAT'S A VERY, VERY UNUSUAL TRINITY TO HAVE. AND SO
18	THAT'S SOMETHING DR. MILLAN WILL BE TALKING MORE
19	ABOUT IN THE CONCEPT PLAN.
20	IT ALSO GOES AND FITS, DOVETAILS VERY
21	NICELY WITH THE ALPHA CLINICS. WE NOW HAVE JUST
22	OVER A YEAR OF EXPERIENCE WITH THE ALPHA CLINICS UP
23	AND RUNNING, AND WE ARE SEEING SOME VERY POSITIVE
24	RESULTS. AND SO THE ACCELERATING CENTER, THE
25	TRANSLATIONAL CENTER WOULD FIT, IN A PARTNERSHIP

1	WITH FDA, WOULD FIT VERY NICELY INTO AN EXPANDED
2	ALPHA CLINIC PROGRAM, AGAIN, IF WE ULTIMATELY DEEM
3	IT'S WARRANTED. SO, AGAIN, DR. MILLAN WILL TALK
4	MORE ABOUT THAT IN A SECOND.
5	THE OTHER THING WE HEARD, THIS IS
6	SWITCHING GEARS NOW, TO PULL, BUT EVERY MAJOR
7	RESEARCH CENTER WE TALKED TO TALKED ABOUT THESE
8	LINKAGES THAT WE TALKED ABOUT WHERE RESEARCH A GOES
9	TO B TO C TO D AND SO ON AND SO FORTH. WHAT WE
10	FOUND IS DISCOVERY STAGE RESEARCHERS THAT WANT TO
11	STAY IN THE LAND OF DISCOVERY DON'T KNOW WHO TO TALK
12	TO AND DON'T KNOW HOW TO GET IN TOUCH WITH THE
13	APPROPRIATE TRANSLATIONAL RESEARCHERS THAT ARE
14	INTERESTED IN TAKING THEIR PROGRAMS FORWARD. AND,
15	AGAIN, WITHOUT EXCEPTION EVERY MAJOR UNIVERSITY WE
16	TALKED TO ASKED FOR HELP IN THIS PARTICULAR AREA.
17	SO WE CAME UP WITH THIS CONCEPT OF THE
18	CIRM EXCHANGE WHERE WE CAN ACTUALLY LINK UP
19	INTERESTED DOWNSTREAM RESEARCHERS WITH PEOPLE WHO
20	HAVE PROMISING TECHNOLOGIES IN EARLIER STAGE. THIS
21	WOULD GO ALL THE WAY THROUGH TO LINKING UP COMPANIES
22	WITH RESEARCHERS AS WELL.
23	AND THEN LASTLY, IN THE PULL CATEGORY IS
24	WHAT WE CALL ATP3. WE HAVE A LOT OF THINGS IN THIS
25	PLAN THAT ARE NEW FOR A FUNDING AGENCY TO ATTEMPT,
	5 2

1	BUT THIS ONE STANDS OUT EVEN FOR THIS PLAN. HERE
2	THE CONCEPT IS THAT WE HAVE A WHOLE BUNCH OF
3	TECHNOLOGY. WE HAVE 300 OR SO DIFFERENT PROGRAMS AT
4	CIRM. AS WE SAID, ONLY 8 PERCENT OF OUR ACADEMIC
5	PROGRAMS CURRENTLY HAVE INDUSTRY PARTNERS. SO HOW
6	DO WE FIX THAT? WE THOUGHT ABOUT A NUMBER OF
7	DIFFERENT WAYS, BUT ONE OF THE WAYS IS JUST TO GO
8	DIRECTLY AND DO IT. WE THOUGHT, WELL, WHAT WOULD
9	HAPPEN IF WE PUT OUT A CALL FOR BASICALLY THE
10	CREATION OF A NEW ENTITY OR A NEW COMPANY THAT WOULD
11	BE A CALIFORNIA-BASED COMPANY THAT COULD TAKE THESE
12	TECHNOLOGIES AND AGGREGATE THEM AND FOCUS ON
13	DEVELOPING AND COMMERCIALIZING STEM CELL-SPECIFIC
14	TECHNOLOGIES?
15	SO THE IDEA IS HERE WE WOULD PUT OUT A
16	CALL THAT WOULD REQUIRE A SUCCESSFUL APPLICANT TO
17	PUT TOGETHER A BUSINESS PLAN THAT WOULD DESCRIBE
18	WHAT TYPES OF TECHNOLOGIES FROM OUR PORTFOLIO THAT
19	THEY WOULD LIKE TO AGGREGATE AND THE SYNERGIES
20	ASSOCIATED WITH THOSE, A GREAT MANAGEMENT TEAM THAT
21	COULD ACTUALLY MAKE THAT HAPPEN IN A SUCCESSFUL WAY,
22	AND THEN, VERY IMPORTANTLY, A TREMENDOUS AMOUNT OF
23	UPFRONT CAPITAL THAT THEY'RE GOING TO COMMIT. I
24	THINK WE HAD IN THIS CONCEPT \$75 MILLION IN UPFRONT
25	CAPITAL THAT THEY'RE GOING TO COMMIT INTO TAKING

1	THESE TECHNOLOGIES AND DRIVING THEM FORWARD. THEN
2	WE WOULD THEN PARTNER WITH THEM ON ACTUALLY A VERY
3	EFFICIENT BASIS TO HELP FUND SOME OF THAT RESEARCH
4	GOING FORWARD.
5	AGAIN, THE IDEA BEING WHAT WE WOULD HAVE
6	AT THE END IS AN ENTITY, BASICALLY A POWERHOUSE IN
7	THE STATE OF CALIFORNIA THAT HAVE THESE TECHNOLOGIES
8	THAT THEY'RE ACTIVELY COMMERCIALIZING. IT WOULD BE
9	AN OUTFLOW FOR NEW TECHNOLOGIES THAT ARE COMING OUT
10	OF CIRM THAT NEED AN INDUSTRY HOME AND OBVIOUSLY
11	CREATE JOBS AND EXPAND THE TAX BASE FOR CALIFORNIA.
12	AND THEN, LASTLY, BUT MOST IMPORTANTLY, BE A VEHICLE
13	FOR GETTING THE FINAL SPAN OF THIS BRIDGE WHERE WE
14	GO FROM LATE STAGE RESEARCH ACTUALLY THROUGH
15	COMMERCIALIZATION SO PATIENTS CAN BENEFIT FROM THEM.
16	AGAIN, DR. MILLAN IS GOING TO TALK MORE
17	ABOUT THIS. I DON'T WANT TO COMPLETELY STEAL HER
18	THUNDER, BUT IT'S A COOL PART.
19	THE LAST PIECE OF THIS IS LEVEL. SO WE'RE
20	PUSHING, WE'RE PULLING, AND THEN LEVELING. I WANT
21	TO TALK A LITTLE BIT ABOUT WHY WE THINK THIS IS
22	PARTICULARLY IMPORTANT. SO WHEN YOU LOOK AT THE
23	CURRENT REGULATORY PARADIGM, THE FIRST THING THAT
24	JUMPS OUT IS THERE'S AN EXCESSIVELY LONG
25	TRANSITIONAL PERIOD FOR CELL THERAPIES TO GO FROM WE
	54

1	UNDERSTAND WE HAVE A CELL THERAPY INTO ACTUAL
2	CLINICAL TRIALS. RIGHT NOW IT'S SOMEWHERE BETWEEN
3	SIX TO EIGHT YEARS. IT'S IMPORTANT TO UNDERSTAND
4	THIS IS NOT ALL FDA. WE'RE NOT SAYING THIS IS ALL
5	FDA. THERE'S ACTUALLY A NUMBER OF REASONS FOR THIS,
6	BUT IT IS AN OPPORTUNITY FOR US TO GET BETTER.
7	SECONDLY, THERE IS A CLEAR PERCEIVED BIAS
8	AGAINST RARE DISEASES THAT WORKS OUT STATISTICALLY.
9	IT IS VERY, VERY DIFFICULT TO MEET FDA STANDARDS AND
10	DEVELOP A DRUG IN A FEASIBLE MANNER FOR A DISEASE
11	THAT AFFECTS A SMALL POPULATION OF PEOPLE. AND
12	THEN, LASTLY, THE SYSTEM THAT'S CURRENTLY IN PLACE,
13	AND IT WAS FIRST PROPOSED IN 1997, IT WAS ADOPTED IN
14	2001, CREATES A VERY ARBITRARY AND BINARY SYSTEM.
15	AND SO CELL THERAPIES ARE EITHER ESSENTIALLY
16	UNREGULATED BY FDA, AND I MEAN VERY LITTLE
17	REGULATION. TAKE YOU LESS THAN \$100,000 AND TAKE
18	YOU LESS THAN THREE MONTHS TO COMPLY. OR THEY ARE
19	EXCESSIVELY REGULATED BY FDA WHERE IT COSTS GREATER
20	THAN A BILLION DOLLARS AND TAKES LONGER THAN 12
21	YEARS AND THERE'S NOTHING IN BETWEEN. ALL OF THE
22	OTHER DISCIPLINES OF MEDICINE HAVE SOMETHING IN
23	BETWEEN THOSE TWO PATHWAYS.
24	SO WHY DO WE WANT TO CHANGE? WHY DO WE
25	THINK THIS NEEDS TO BE LOOKED AT? WELL, THE FIRST

1	THING IS IT'S BEEN 15 YEARS SINCE THIS REGULATORY
2	PARADIGM HAS BEEN IN PLACE. STEM CELL RESEARCH HAS
3	BEEN GOING ON LONG BEFORE THAT, AND THERE IS A ZERO
4	ON THE SCOREBOARD. NOTHING HAS BEEN NO STEM CELL
5	THERAPY HAS BEEN APPROVED BY THE FDA SINCE
6	IMPLEMENTING THIS REGULATORY PARADIGM 15 YEARS AGO.
7	SO IT'S IMPORTANT TO UNDERSTAND AS WE TALK
8	ABOUT THIS WHAT CIRM ISN'T. CIRM IS NOT
9	ANTI-REGULATION. WE ARE NOT ANTI-FDA. WE'RE NOT
10	SAYING GET RID OF THE RULES. BUT THE SCOREBOARD
11	ISN'T LYING HERE. AND WHEN PEOPLE LOOK AT THE
12	CURRENT REGULATORY SYSTEM AND SEE THAT IT'S BEEN IN
13	PLACE FOR 15 YEARS AND NOTHING IS APPROVED, A VERY
14	IMPORTANT EFFECT OF THAT IS PEOPLE DON'T ATTEMPT
15	THAT TO WHICH THEY BELIEVE IS IMPOSSIBLE TO BEGIN.
16	WE'RE NOT SAYING THAT IT'S NOT ACTUALLY FEASIBLE TO
17	COMPLY WITH THE ENTIRE 15-YEAR, \$2.6 BILLION, WHICH
18	IS THE CURRENT AVERAGE FOR DRUGS, REGULATORY
19	PARADIGM, BUT IN A LOT OF CASES YOU CAN'T
20	ECONOMICALLY JUSTIFY IT.
21	I WILL TELL YOU IN MY FORMER LIFE AT
22	OSIRUS AS CEO OF OSIRIS, ROUTINELY PEOPLE WOULD COME
23	UP TO US WITH NEW TECHNOLOGIES AND NEW IDEAS,
24	INTERNALLY WE WOULD DEVELOP SOMETHING, AND WE WOULD
25	LOOK AND EVALUATE IT, AND WE WOULD HAVE STRONG
	E.C.

1	CONVICTION AROUND IT WAS POSSIBLE, AND THEN WE WOULD
2	LOOK INTO THE REGULATORY PATHWAY. IF IT GOT PUT
3	DOWN THIS PARTICULAR REGULATORY PATHWAY, WE'D HAVE
4	TO ABANDON IT BECAUSE THERE IS NO WAY WE COULD EVER
5	JUSTIFY THAT KIND OF INVESTMENT AND THAT KIND OF
6	TIME WHEN YOU'RE GOING AFTER, AGAIN, PARTICULARLY
7	SMALLER NUMBERS OF PATIENT POPULATIONS WHICH CELL
8	THERAPIES TEND TO ADDRESS.
9	THE OTHER THING HERE, THERE'S A LOT OF
10	TALK ABOUT THE COST OF HEALTHCARE AND THE COST OF
11	NEW DRUGS. WELL, WHEN THE REGULATORY PARADIGM IS 15
12	YEARS AND \$2.6 BILLION, DRUGS ARE GOING TO BE
13	REALLY, REALLY EXPENSIVE. SO WE NEED TO LOOK AT
14	THIS.
15	LASTLY, IT'S THE PATIENTS. SO WHILE WE
16	SIT HERE AND WE THINK ABOUT THIS, AND THE FDA IS,
17	RIGHTFULLY SO, VERY CONCERNED ABOUT PROTECTING
18	PATIENTS FROM THEORETICAL RISKS OF A NEW THERAPY,
19	OFTENTIMES THOSE PATIENT POPULATIONS ARE DYING OF
20	THEIR VERY REAL DISEASES. AND SO WE ARE NOT
21	PRESCRIPTIVE ON WHAT THE ANSWER FOR THIS IS. I WANT
22	TO MAKE THAT CLEAR. WE'RE NOT SAYING HERE'S WHAT'S
23	GOING TO HAVE TO HAPPEN, BUT WE ARE SAYING THE
24	CURRENT SITUATION IS NOT ACCEPTABLE AND DOING
25	NOTHING ABOUT IT IS NOT OKAY. SO WE'RE PROPOSING TO

1	WORK WITH FDA, AS WE TALKED ABOUT ON A NUMBER OF
2	DIFFERENT FRONTS, AND WORK WITH PATIENTS AND WORK
3	WITH INDUSTRY AND WHOEVER WE HAVE TO WORK WITH IN
4	ORDER TO GET SOMETHING HERE WORKED OUT SO WE CAN
5	START HAVING SUCCESSES GO ON THAT SCOREBOARD.
6	SO THAT'S PUSH, PULL, LEVEL. IF WE'RE
7	SUCCESSFUL WITH ALL OF THAT, WE HAVE SIX MAJOR
8	OUTCOMES THAT WE THINK WE CAN ACHIEVE. FIRST IS WE
9	START AT THE BEGINNING. ONE OF THE THINGS I WANT TO
10	EMPHASIZE BECAUSE I GET ASKED THIS QUESTION
11	SOMETIMES. WELL, IT SEEMS LIKE THIS HAS SHIFTED TO
12	REALLY LATE STAGE RESEARCH. NO. WE ARE TRYING TO
13	CREATE AN ENGINE THAT ACCELERATES EVERYTHING THROUGH
14	IT. AND THAT STARTS WITH THE VERY EARLIEST STAGES
15	OF RESEARCH. SO OUR FIRST EXPECTED RESULT, OUR
16	FIRST METRIC IS 50 NEW DISCOVERY ENTITIES
17	DISCOVERED, CANDIDATES DISCOVERED FOR EITHER
18	THERAPEUTICS OR DEVICES.
19	SECONDLY, THEN WE WANT TO HAVE THE
20	FREQUENCY IN WHICH THESE PROGRAMS MOVE FROM ONE
21	STAGE OF DEVELOPMENT IMPROVE BY 50 PERCENT. WE WANT
22	A NEW REGULATORY PARADIGM WITHIN THE NEXT FIVE
23	YEARS. WE WANT TO REDUCE THE TIME IT TAKES
24	TRANSLATION BY 50 PERCENT. THAT'S THIS SIX TO EIGHT
25	YEARS DOWN TO THE INDUSTRY AVERAGE OF 3.2. WE WANT
	Γ0

1	TO HAVE 50 NEW CLINICAL TRIALS GET INTRODUCED. AND
2	THEN, LASTLY, PARTNER AT LEAST 50 PERCENT OF OUR
3	ACADEMIC PROGRAMS SUCCESSFULLY WITH INDUSTRY
4	PROGRAMS THAT CAN TAKE THOSE THINGS FORWARD AND
5	BRING THEM TO THE PATIENTS WHO NEED THEM.
6	AGAIN, THIS IS NOT AN OPERATIONAL PLAN, SO
7	THIS FINANCIAL SUMMARY IS ONLY HERE TO SHOW THAT
8	THIS IS POSSIBLE. IT DOES ASSUME 890 MILLION IN NEW
9	AWARDS. AS I SAID, THAT WAS PREDICATED ON AN
10	ASSUMPTION OF A 10-PERCENT AWARD RECAPTURE RATE.
11	THAT RIGHT NOW AS OF LAST YEAR WAS 22 PERCENT, SO WE
12	MAY ACTUALLY HAVE MORE MONEY HERE TO DEPLOY.
13	ANOTHER THING I WANT TO SAY ABOUT THIS IS
14	WE ARE NOT HOLDING BACK ON BUILDING AND STARTING
15	THIS ENGINE. WE HAVE NOT TRIED TO SMOOTH THIS OUT
16	LIKE PEANUT BUTTER. WE ARE TRYING TO GET THIS
17	ENGINE UP AND RUNNING AT MAXIMUM POWER AS QUICKLY AS
18	WE ARE CAPABLE OF DOING.
19	LASTLY, I WANT MAKE SURE WE UNDERSTAND, AS
20	WE'VE TALKED ABOUT ALL ALONG, THIS IS A VERY
21	AGGRESSIVE PLAN. IT HAS VERY AMBITIOUS GOALS. IT
22	CLEARLY HAS RISK. IT HAS SOME RISKS THAT ARE VERY
23	UNIQUE TO CIRM. AND SO WE HAVE FIVE OF THEM LISTED
24	HERE. THERE'S MORE IN THE PLAN. THERE'S TWO,
25	THOUGH, THAT I REALLY WANT TO POINT OUT. ONE IS ONE
	50

1	OF THE THINGS WE CAN'T DO IN TRYING TO OBTAIN THESE
2	50 NEW CANDIDATES THAT HAVE BEEN DISCOVERED AND 50
3	NEW PROGRAMS IN CLINICAL TRIALS, ONE OF THE THINGS
4	WE CAN'T DO IS LOWER OUR STANDARDS FOR QUALITY.
5	QUALITY HAS TO STAY HIGH. SO WE NEED TO BE
6	INNOVATIVE. THESE GUYS NEED TO BE REALLY INNOVATIVE
7	ON HOW THEY BRING THINGS INTO CIRM TO BE EVALUATED.
8	BUT IF WE'RE NOT GOING TO LOWER OUR STANDARDS ON
9	QUALITY, THERE IS A RISK THAT THERE MAY BE
10	INSUFFICIENT NUMBERS OF MERITORIOUS PROGRAMS THAT
11	COME TO CIRM THAT ACTUALLY JUSTIFY OUR FUNDING TO
12	REACH OUR GOALS. WE HOPE THAT'S NOT THE CASE.
13	WE'RE GOING TO DO EVERYTHING WE CAN TO MAKE SURE
14	THAT'S NOT THE CASE. BUT THAT IS A UNIQUE RISK, NOT
15	THAT WE DON'T HAVE THE FUNDING FOR IT, BUT THAT
16	THERE SIMPLY AREN'T ENOUGH PROGRAMS OUT THERE THAT
17	MEET OUR STANDARDS WITHOUT US HAVING TO LOWER THEM.
18	THE SECOND RISK THAT I WANT TO POINT OUT,
19	WHICH IS THE THIRD BULLET ON HERE THAT'S, I THINK,
20	UNIQUE TO CIRM IS THAT, AT LEAST RIGHT NOW IN THE
21	CURRENT ENVIRONMENT, CIRM HAS A LIMITED LIFETIME
22	ASSOCIATED WITH IT. THE THING THAT MAKES THIS
23	ENGINE WORK, AND IF THIS IS AT ALL GOING TO BE
24	POSSIBLE, IS THE TEAM THAT WE HAVE RUNNING AND
25	OPERATING THAT ENGINE. WE NEED TO BE VERY COGNIZANT

AS WE GET TOWARDS THE END OF OUR LIFE SPAN THAT IT
WILL BECOME MORE DIFFICULT TO ATTRACT AND RETAIN THE
HIGH QUALITY TALENT THAT WE NEED IN ORDER TO MAKE
THIS WORK. NOW, WE KNOW THAT RISK, AND SO WE'RE
GOING TO DO EVERYTHING WE CAN TO MITIGATE IT, BUT IT
IS A UNIQUE RISK TO CIRM.
SO I WANT TO END ON THIS SLIDE HERE AND
STOP TALKING, OBVIOUSLY TAKE ANY QUESTIONS THAT
ANYONE ABOUT THIS MIGHT HAVE. BUT I PUT UP HERE THE
PHRASE "ALL IN" BECAUSE WHAT I'M SURE OF, THIS IS A
HARD PLAN AND I'M NOT POSITIVE WE WILL ACCOMPLISH
IT, BUT I AM SURE THAT IF WE DON'T HAVE CONVICTION
AROUND IT, WE WILL DEFINITELY NOT ACCOMPLISH IT. SO
WE ARE ALL IN ON THIS. AND I THINK AS A GROUP WE
NEED TO BE ALL IN ON THIS. IF WE ARE, PROBABILITY
OF SUCCESS IS GREAT. AND IF WE ARE SUCCESSFUL, AS
SHERRY SAID, IT WILL HAVE BEEN WORTH DOING. SO I
WILL STOP TALKING AND TAKE QUESTIONS NOW.
DR. DULIEGE: SO A SET OF BRIEF COMMENTS.
I JUST WANT TO SAY, AND I THINK I CAN SPEAK FOR ALL
OF US, CERTAINLY MANY OF US, EXCELLENT. REALLY VERY
INSPIRING. AND THANK YOU TO THE ENTIRE TEAM. I
KNOW YOU ARE PRESENTING IT, BUT I KNOW YOU SPEAK ON
BEHALF OF ALL OF YOU. SO GREAT WORK.
COUPLE OF COMMENTS OF WHAT I'D LIKE TO
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1	HEAR FURTHER DOWN, NOT RIGHT NOW, WHICH IS YOU
2	MENTIONED YOU WILL DO ALL THE EFFORTS TO KEEP THE
3	LIFE OF THIS EFFORT ONGOING AFTER CIRM 2.0, AND THAT
4	WILL BE, I'M SURE, TOPIC OF FUTURE DISCUSSIONS,
5	WHICH WILL BE GREAT.
6	THE SECOND, YOU ALLUDED ABOUT THE COST OF
7	TREATMENT, AND I'D BE CURIOUS TO KNOW, AGAIN DOWN
8	THE ROAD, WHICH IMPACT YOU THINK CIRM MIGHT HAVE ON
9	COST OF TREATMENT, SOMETHING WHICH IS DEAR TO OUR
10	HEARTS.
11	THE THIRD IS THAT I WAS QUITE INTERESTED
12	TO SEE THE COLLABORATION THAT YOU'VE STARTED TO HAVE
13	WITH FDA, AND THIS WILL HAVE IMMENSE IMPACT. DO YOU
14	BELIEVE YOU ALSO WANT TO HAVE AN IMPACT ON HOW FDA
15	REGULATES OR SO FAR THE LACK OF REGULATION OF NEW
16	BOUTIQUE SHOPS ABOUT STEM CELLS THAT ARE DOING A
17	DISSERVICE TO THE FIELD, TO THE RESEARCH, AND TO
18	PATIENTS? A LITTLE BIT MAYBE BEYOND THE SCOPE OF
19	CIRM, BUT WOULD BE INTERESTED TO KNOW THAT.
20	AND FINALLY, I'M SURE THIS EFFORT IS GOING
21	TO BE A GREAT SOURCE OF INSPIRATION BEYOND
22	CALIFORNIA IN THE U.S. AND OUTSIDE THE U.S. AND I
23	WILL BE DOWN THE ROAD VERY INTERESTED TO KNOW WHAT
24	KIND OF INTERNATIONAL COLLABORATIONS MIGHT BE EITHER
25	ONGOING OR PLANNED FOR THE FUTURE. SO MORE FORWARD

1	LOOKING DISCUSSIONS.
2	CHAIRMAN THOMAS: OTHER COMMENTS OR
3	QUESTIONS?
4	DR. PRIETO: YES. THANK YOU FOR THAT
5	PRESENTATION. I HAD SOME QUESTIONS ABOUT THE
6	TRANSLATING CENTER AND THE ATP3 PORTION OF THIS. IN
7	BOTH OF THOSE YOU IMPLY OR MEASURE OR I SHOULD SAY
8	THE PLAN DOES THAT THIS WILL HAVE MINIMAL CIRM
9	FUNDING. WITH ATP3 IT SAYS THAT NO CIRM FUNDS WILL
10	BE USED FOR THE ESTABLISHMENT OR OPERATION OF THE
11	ENTITY. SO I WAS NOT CLEAR EXACTLY WHAT'S OUR ROLE
12	IN SETTING THESE UP, AND THEN ARE THEY
13	SELF-SUPPORTING, AND HOW?
14	DR. MILLS: DR. MILLAN WILL ACTUALLY BE
15	BRINGING THESE CONCEPT PLANS WITH GREATER
16	SPECIFICITY TO THE BOARD. BUT WITH ATP3, THE IDEA
17	IS THEY COME TO US WITH SIGNIFICANT UPFRONT, 75
18	MILLION IN UPFRONT CAPITAL TO CREATE THE ACTUAL
19	ENTITY AND THE BUSINESS. AND WE DO A COUPLE OF
20	THINGS. ONE IS WE HELP THEM AGGREGATE THOSE
21	TECHNOLOGIES, WORK OUT LICENSING ARRANGEMENTS WITH
22	ALL THESE DIFFERENT UNIVERSITIES. THESE ARE ALL
23	CIRM-FUNDED PROGRAMS THAT WE'RE TALKING ABOUT. SO
24	THESE ARE WHAT WE'RE TALKING ABOUT ARE
25	AGGREGATING CURRENT ACTIVE PROGRAMS WITH CIRM SO
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1	THEY WILL HAVE RESIDUAL FUNDING ASSOCIATED WITH THEM
2	WHEN THEY GET AGGREGATED INTO THIS ENTITY.
3	AND THEN ON A PER-PROGRAM BASIS, THE GWG
4	WILL REVIEW THE MERITS OF FUNDING THOSE PROGRAMS
5	FORWARD AND PROVIDE A LINE OF CREDIT TO THE
6	ORGANIZATION ONLY FOR THE CONTINUED FUNDING THAT
7	CIRM WOULD OTHERWISE BE DOING.
8	DR. PRIETO: WON'T THAT INVOLVE SOME CIRM
9	STAFF TIME, CIRM EFFORTS PULLING THESE ELEMENTS
10	TOGETHER?
11	DR. MILLS: AROUND AS MUCH AS A NORMAL GWG
12	REVIEW FOR THOSE PROGRAMS GETTING REUPPED WOULD
13	OTHERWISE.
14	DR. STEWARD: RANDY, THANKS. AND JUST,
15	AGAIN, CONGRATULATIONS TO THE GROUP FOR AN
16	INCREDIBLE AMOUNT OF VERY HARD WORK TO BRING THIS
17	FORWARD.
18	I WANTED TO ACTUALLY EMPHASIZE SOMETHING
19	THAT'S INTRINSIC TO YOUR PRESENTATION, BUT REALLY
20	SAY IT OUT LOUD JUST TO MAKE IT CLEAR. I THINK
21	YOU'VE MADE IT VERY CLEAR THAT YOU'RE RECOGNIZING
22	THE DIFFICULTIES AND ROADBLOCKS THAT HAVE BEEN IN
23	PLACE THAT HAVE MADE IT DIFFICULT TO MOVE STEM CELL
24	THERAPIES FORWARD THROUGH THE REGULATORY PROCESS,
25	BUT I THINK THAT YOU'VE SAID IT AND I JUST WANT TO

1	EMPHASIZE THAT I DON'T THINK YOU OR ANYONE HAS SEEN
2	THE FDA AS AN OPPONENT IN THIS PROCESS, THAT REALLY
3	CIRM IS LOOKING FORWARD TO TRYING TO LOWER SOME OF
4	THESE BARRIERS IN COOPERATION AND COLLABORATION WITH
5	THE FDA. I JUST WANT TO EMPHASIZE THAT.
6	DR. MILLS: ABSOLUTELY. AND JUST TO SORT
7	OF TOUCH ON THAT POINT AND A COMMENT THAT ANNE-MARIE
8	BROUGHT UP ABOUT WHAT CIRM COULD OR COULD NOT DO
9	ABOUT THE BAD ACTORS OF THE UNREGULATED. I THINK
10	THE REASON WE HAVE TO LOOK AT THAT. SO WHY IS
11	THERE UNREGULATED CELL THERAPIES GOING ON IN THE
12	UNITED STATES RIGHT NOW? BECAUSE THEY'RE LOOKING AT
13	THE ONLY PATHWAY IN FRONT OF THEM AND SAYING, WELL,
14	OUR OPTIONS ARE WE EITHER AVOID DETECTION OR WE
15	DON'T DO IT. AND SO IF WE CAN CREATE BETTER
16	REGULATORY PARADIGMS, WE CAN ACTUALLY BRING MORE
17	THINGS INTO REGULATION THAN WOULD OTHERWISE EXIST
18	RIGHT NOW.
19	DR. JUELSGAARD: RANDY, I JUST WANT TO SAY
20	CONGRATULATIONS ON A VERY THOUGHTFUL, VERY THOROUGH,
21	AND VERY BOLD STRATEGIC PLAN. I THINK WE ALL HAVE
22	TO KEEP IN MIND THAT THIS PLAN IS FULL OF
23	EXPERIMENTS, THINGS THAT I'VE NEVER SEEN DONE
24	BEFORE, THE TRANSLATIONAL CENTER, THE ACCELERATING
25	CENTER, ATP3. SO WE'RE GOING TO BE TRYING SOME

1	THINGS OUT, ASSUMING WE GO FORWARD WITH THIS, THAT
2	ARE GOING TO BE FIRST OF KIND, AND WE'LL HAVE TO SEE
3	HOW THEY PLAY OUT.
4	BUT I THINK AT LEAST THIS IS A GREAT
5	OUTLINE OF A DIRECTION TO GO WHERE WE HAVEN'T REALLY
6	BEEN AS DIRECTIONALLY FOCUSED AS WE SHOULD HAVE
7	BEEN. SO THANK YOU VERY MUCH, AND CONGRATULATIONS
8	AT LEAST FROM MY POINT OF VIEW.
9	MS. LANSING: I WANT TO ECHO THIS AGAIN.
10	WE'VE BEEN PART OF THIS AS YOU'VE GONE THROUGH ITS
11	VARIOUS REITERATIONS. I JUST WANT TO REALLY
12	COMPLIMENT YOU, RANDY, AND THE ENTIRE TEAM. AND I
13	STILL REMEMBER WHEN WE FIRST MET YOU AND YOU WERE A
14	CANDIDATE FOR THE JOB, AND GOD KNOWS WE CERTAINLY
15	MADE THE RIGHT DECISION. I JUST WANT TO SAY I
16	REMEMBER I WAS SO STRUCK BY THE SENSE OF URGENCY
17	THAT YOU CONVEYED IN OUR FIRST MEETING AND HOW EVERY
18	DAY THAT WENT BY, UNFORTUNATELY SOMEONE WAS STRUCK
19	BY A DISEASE AND OFTEN DIDN'T SURVIVE IT AND THAT
20	THIS WAS ALL ABOUT SAVING LIVES. I THINK YOU'VE
21	BROUGHT THAT SAME SENSE OF URGENCY THROUGHOUT ALL
22	THE TIME YOU'VE BEEN HERE.
23	I THINK WHAT'S MOST IMPRESSIVE TO ME AMONG
24	THE MANY THINGS ABOUT THIS STRATEGIC PLAN IS THAT IT
25	CONVEYS A SENSE OF URGENCY.

1	LIKE ANY STRATEGIC PLAN, IT'S ALWAYS GOING
2	TO BE A WORK IN PROGRESS, AS JEFF SAID. WE WILL
3	TINKER WITH IT. YOU WILL SEE OPPORTUNITIES THAT NO
4	ONE THOUGHT ABOUT, AND YOU WILL QUICKLY ADAPT TO
5	THEM. SO I'M NOT WORRIED THAT EVERYTHING HASN'T
6	BEEN COVERED BECAUSE IT WILL CHANGE. IT WILL BE
7	FLEXIBLE. I URGE US TO I THINK THE FDA IS OUR
8	FRIEND BECAUSE IT REALLY IS A SITUATION WHERE
9	THEY'RE TRYING TO PROTECT PEOPLE FROM THINGS THAT
10	HAVEN'T BEEN PROVEN. IT'S A FRUSTRATION LEVEL THAT
11	I THINK ALMOST EVERY ADVOCACY GROUP AND EVERY
12	DISEASE GROUP FEELS EVEN THOUGH THEY'RE GETTING
13	THERE HALF THE TIME. I URGE US TO PARTNER BECAUSE I
14	DO BELIEVE THAT THE FDA ALWAYS HAS OUR BEST INTEREST
15	AT HEART. AND IF WE CAN EXPLAIN CERTAIN THINGS AND
16	VIEW THEM THAT WAY WITH OTHER GROUPS, OBVIOUSLY WITH
17	DRUG COMPANIES AS WELL, I THINK THAT'S REALLY GOOD.
18	I ALSO WANT TO SAY THAT THIS IS A
19	FANTASTIC DOCUMENT. AND IT'S, THANKS TO YOU,
20	LAUREN, VERY USER FRIENDLY. WHEN YOU LOOK AT IT,
21	YOU WANT TO READ IT, BUT IT IS VERY LONG. I'M NOT
22	SAYING FOR US AND FOR THE JOURNALISTS, WHATEVER, BUT
23	I URGE US TO GET OUT A ONE- OR TWO-PAGE SHEET THAT
24	JUST HAS THE BULLET POINTS AND REALLY, REALLY
25	DISTRIBUTE THIS BECAUSE THE CITIZENS OF CALIFORNIA
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1	VOTED FOR THIS. THIS IS A SEISMIC SHIFT IN WHAT
2	WE'RE DOING AND A GOOD SEISMIC SHIFT THAT COULD NOT
3	HAVE HAPPENED HAD THE WORK NOT BEEN DONE ALL THOSE
4	YEARS BEFORE AS YOU SO ACKNOWLEDGE, BUT LET'S GET
5	THIS OUT. LET'S REALLY LET PEOPLE KNOW THAT.
6	WITH THAT SAID, I FULLY, FULLY SUPPORT
7	THIS. I THANK THE ENTIRE TEAM WHO IS SITTING THERE
8	AND THERE, AND I WON'T NAME EVERYBODY. AND I WOULD
9	LIKE TO MOVE THE ITEM.
10	MR. TORRES: SECOND.
11	DR. PULIAFITO: YOU STOLE MY MOTION.
12	CHAIRMAN THOMAS: MOVED BY SHERRY,
13	SECONDED BY SENATOR TORRES, THIRDED, THOUGH STOLEN
14	BY SHERRY, BY DR. PULIAFITO.
15	DR. MILLS: TWO THINGS. ONE IS WITH
16	REGARDS TO THE FDA, I WAS RECENTLY IN WASHINGTON.
17	WE WERE TALKING ABOUT THIS. AND SOMEBODY FROM THE
18	FDA WAS SAYING, WELL, THE EXPECTATION IS THAT WE
19	PROTECT. AND, YES, THE EXPECTATION IS TO PROTECT,
20	BUT IT HAS TO BE MORE THAN THAT. WE GET THE FDA.
21	WE ASKED FOR IT. I DON'T MEAN US AS PEOPLE IN
22	CALIFORNIA OR AT CIRM, BUT AS A COUNTRY WE ASKED FOR
23	IT. WE HAVE TO THROW A LITTLE BIT MORE WE NEED
24	STUFF THROUGH IT TOO. BECAUSE SAYING NO TO
25	EVERYTHING WOULD KEEP EVERYONE SAFE, BUT WE NEED TO
	68
	I UA

1	ALSO MAKE PROGRAMS. SO WE'RE GOING TO DO THAT. AND
2	THERE ARE SO MANY OPPORTUNITIES FOR US TO ACTUALLY
3	DIRECTLY, CIRM WITH FDA, PARTNER AND HAVE AND WE
4	ARE.
5	YOUR POINT AND OS AND EVERYONE ELSE IS NOT
6	JUST UNDERSTOOD, IT'S EMBRACED, AND IT IS WHAT WE
7	WILL DO.
8	MS. LANSING: REACH OUT TO OTHER PEOPLE IN
9	OTHER DISEASE PROGRAMS. EVERYONE KNOWS THIS.
10	EVERYONE KNOWS THEY'RE ON THEIR SIDES, BUT WANTS TO
11	HELP GET THE INFORMATION SO THEY CAN MAKE INFORMED
12	DECISIONS.
13	DR. MILLS: WHAT I WANT TO DO IS JUST MAKE
14	SURE WE TALK ABOUT IT, AND WE DON'T VIEW IT AS,
15	WELL, THAT'S JUST WHAT IT IS. IT CAN'T BE JUST WHAT
16	IT IS. WE HAVE TO TALK ABOUT IT.
17	MS. LANSING: I DON'T MEAN TO BELABOR
18	THIS. AND THE BREAKTHROUGH THERAPY THINGS, AND SOME
19	OF THE THINGS THAT THEY HAVE DONE, I THINK, SHOW
20	THAT WE CAN MAKE A DIFFERENCE. AND PATIENT
21	ADVOCATES, OUR BOARD, ALL THE OTHER DISEASE GROUPS,
22	MAYBE THEY'RE SAYING TWO AND A HALF YEARS IS STILL
23	TOO LONG. I THINK WE HAVE TO WORK TOGETHER.
24	DR. MILLS: A COMMENT. WHEN WE GOT THIS
25	THING DONE, WE GOT IT BACK FROM THE PRINTER, IT WAS

1	50 PAGES LONG, WE SAID, AH, CRAP. SO WE MADE THAT.
2	IT'S JUST ONE PAGE.
3	CHAIRMAN THOMAS: OKAY. COMMENTS FROM
4	MEMBERS OF THE BOARD? COMMENTS FROM MEMBERS OF THE
5	PUBLIC? DR. LORING.
6	DR. LORING: I'M JEANNE LORING FROM THE
7	SCRIPPS RESEARCH INSTITUTE. AND I WAS AT THE WORLD
8	STEM CELL SUMMIT AS WERE MANY OF YOU LAST WEEK, AND
9	WE HAD THE OPPORTUNITY TO MEET THE NEW FDA
10	COMMISSIONER, ROBERT CALIFF, JUST BEFORE HE ACTUALLY
11	GETS APPROVED. AND I THINK WE COULD LEVERAGE THE
12	FACT THAT HE WAS AT THE WORLD STEM CELL SUMMIT AND
13	TALKED WITH A LOT OF US ABOUT STEM CELL THERAPIES TO
14	TRY TO CREATE SOME KIND OF CONVERSATION WITH THEM.
15	AS YOU KNOW, THERE ARE A NUMBER OF PATIENT
16	ADVOCATES IN THE AUDIENCE TODAY, AND THE FDA REALLY
17	WANTS TO LISTEN TO THEM. SO I'M REALLY CURIOUS
18	ABOUT THE DETAILS OF THE PROGRAM TO APPROACH THE FDA
19	AND HAVE THEM MORE FULLY UNDERSTAND WHAT THE ISSUES
20	ARE. THAT'S A QUESTION.
21	CHAIRMAN THOMAS: I'M SORRY, DR. LORING.
22	WOULD YOU REPEAT THE QUESTION?
23	DR. LORING: QUESTION WAS I'M ASKING FOR A
24	BIT MORE DETAIL ABOUT HOW, AT LEAST IN BROAD
25	STROKES, HOW CIRM IS PLANNING TO APPROACH THE FDA.
	70

1	CHAIRMAN THOMAS: I THINK, DR. MILLS,
2	WOULD YOU LIKE TO COMMENT ON THAT?
3	DR. MILLS: SO, AGAIN, WITH REGARDS TO THE
4	FDA, IT'S MULTIFACTORIAL. THERE ARE A NUMBER OF
5	THINGS HERE WHERE THERE ARE OPPORTUNITIES TO HAVE
6	ADVANCEMENTS. FIRST IS WORKING THROUGH THIS
7	TRANSLATIONAL ISSUE THAT WE HAVE WHERE IT'S TAKING
8	TOO LONG. THAT, WE'VE ALREADY STARTED DISCUSSIONS
9	WITH FDA ABOUT THEY HAVE AN INTEREST IN PARTNERING.
10	I'M NOT GOING TO GET INTO THE EXTENT OF THEM, THE
11	AMOUNT OF DETAILS OF WHAT WE'RE GOING TO DO
12	SPECIFICALLY OR WHAT WE'RE TALKING ABOUT DOING
13	SPECIFICALLY, BUT THE GOAL OF THAT IS TO FIND A WAY,
14	AND THERE SEEMS TO BE PLENTY OF GROUND, THAT WE CAN
15	HAVE FUNCTIONALLY, AT LEAST CIRM PROGRAMS, GO FROM A
16	PATHWAY THAT IS SOMEWHERE BETWEEN SIX TO EIGHT YEARS
17	DOWN TO MORE THE INDUSTRY STANDARD.
18	THE OTHER CENTERS AROUND ACTUALLY THINKING
19	ABOUT SHOULD THERE IS THERE REALLY JUST A ONE
20	SIZE FITS ALL REGULATORY PARADIGM NO MATTER WHAT THE
21	CELL, NO MATTER WHAT THE DISEASE, NO MATTER WHAT THE
22	DELIVERY SYSTEM? IS IT REALLY ALL THIS MASSIVELY
23	LONG, COMPLICATED, AND EXPENSIVE BLA AND TALK ABOUT
24	THE FACT THAT SINCE THAT STANDARD HAS BEEN IN PLACE,
25	NOTHING HAS GOTTEN THROUGH IT, AND SEE IF WE CAN'T
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1	FIGURE OUT SOMETHING ELSE. SO THAT'S WHAT WE'LL DO.
2	CHAIRMAN THOMAS: THANK YOU. ANOTHER
3	COMMENT FROM A VERY FRIENDLY, FAMILIAR FACE.
4	DR. CHIU: ARLENE CHIU FROM THE CITY OF
5	HOPE. I WANT TO CONGRATULATE CIRM ON SUCH AN
6	AUDACIOUS AND EXCITING PLAN. I THINK WHAT STRUCK
7	ME, LOOKING AT IT FOR THE FIRST TIME, IS THAT YOU'VE
8	PINPOINTED THE AREAS, SELECT AREAS, THAT ARE TRUE
9	BOTTLENECKS. IT'S NOT JUST RESEARCH, BUT THE
10	BOTTLENECKS MOVING FORWARD. I'M A GREAT FRIEND OF
11	THE FDA. AS YOU WERE AT THE WORLD ALLIANCE FORUM,
12	AND I WAS TOO JUST RECENTLY AT GOLDEN GATE PARK,
13	WE'VE NOTICED THAT IN JAPAN THE SCIENTISTS AND THE
14	REGULATORS HAVE BEEN ABLE TO WORK TOGETHER TO CRAFT
15	A PATHWAY FOR CLINICAL STUDIES FOR STEM CELL
16	THERAPIES THAT HAS MADE THE PATHWAY MUCH SHORTER.
17	AND, OF COURSE, INDIVIDUAL SCIENTISTS IN AMERICA
18	HAVE A DIFFICULT TIME DEALING ONE ON ONE WITH THE
19	FDA. BUT THE IDEA OF A TRANSLATIONAL CENTER AND AN
20	ACCELERATING CENTER WHERE THESE ARE GROUPED TO TALK
21	WITH THE FDA, I THINK IT'S A BRILLIANT IDEA. THE
22	FDA NEEDS YOUR HELP TO UNDERSTAND WHAT ARE THE
23	ISSUES, WHAT THE LEVEL OF SAFETY CONCERNS. AND ON
24	THE OTHER HAND, ONCE THEY UNDERSTAND, THEY ARE IN A
25	BETTER POSITION TO CRAFT NOVEL, NEW PATHWAYS TO US.
	70
	72

1	SO I WANT TO THANK EVERYONE FOR SUCH A
2	GREAT IDEA. OF COURSE, THE PROOF OF THE PUDDING IS
3	IN THE TASTING, SO WE'RE ALL EXCITED TO SEE WHAT
4	HAPPENS.
5	CHAIRMAN THOMAS: THANK YOU. ANY OTHER
6	COMMENTS FROM MEMBERS OF THE PUBLIC? HEARING NONE,
7	WE WILL PROCEED TO A ROLL CALL VOTE. BEFORE WE DO
8	THAT, MARIA, I JUST WANT TO SAY, HAVING VIEWED THE
9	DEVELOPMENT OF THE STRATEGIC PLAN FROM THE INSIDE, I
10	WANT TO REITERATE THE ENORMOUS AMOUNT OF WORK THAT
11	WENT INTO THIS BY RANDY, ALL MEMBERS OF THE TEAM,
12	MANY, MANY STAKEHOLDERS OF ALL SORTS OF WALKS ON THE
13	OUTSIDE WHO ADVISED MEMBERS OF THE BOARD, THIS WAS
14	AN EXHAUSTIVE EFFORT THAT TOOK MONTHS AND MONTHS AND
15	IS NOW BEFORE YOU ON A VOTE. SO I WANT TO ADD MY
16	CONGRATULATIONS TO DR. MILLS AND THE TEAM FOR A JOB
17	EXCEPTIONALLY WELL DONE. WITH THAT, MARIA, WILL YOU
18	PLEASE CALL THE ROLL.
19	MS. BONNEVILLE: DAVID BRENNER. LINDA
20	BOXER.
21	DR. BOXER: YES.
22	MS. BONNEVILLE: KEN BURTIS. ANNE-MARIE
23	DULIEGE.
24	DR. DULIEGE: YES.
25	MS. BONNEVILLE: MICHAEL FRIEDMAN.
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1		DR. FRIEDMAN: YES.
2		MS. BONNEVILLE: JUDY GASSON.
3		DR. GASSON: YES.
4		MS. BONNEVILLE: SAM HAWGOOD. DAVID
5	HIGGINS.	
6		DR. HIGGINS: YES.
7		MS. BONNEVILLE: STEPHEN JUELSGAARD.
8		MR. JUELSGAARD: YES.
9		MS. BONNEVILLE: SHERRY LANSING.
10		MS. LANSING: YES.
11		MS. BONNEVILLE: KATHY LAPORTE.
12		DR. LAPORTE: ENTHUSIASTICALLY YES.
13		MS. BONNEVILLE: BERT LUBIN. SHLOMO
14	MELMED.	
15		DR. MELMED: YES.
16		MS. BONNEVILLE: LAUREN MILLER.
17		MS. MILLER: YES.
18		MS. BONNEVILLE: ADRIANA PADILLA.
19		DR. PADILLA: YES.
20		MS. BONNEVILLE: JOE PANETTA.
21		MR. PANETTA: YES.
22		MS. BONNEVILLE: ROBERT PRICE.
23		DR. PRICE: YES.
24		MS. BONNEVILLE: FRANCISCO PRIETO.
25		DR. PRIETO: AYE.
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1	MS. BONNEVILLE: CARMEN PULIAFITO.
2	DR. PULIAFITO: YES.
3	MS. BONNEVILLE: ROBERT QUINT. AL
4	ROWLETT.
5	MR. ROWLETT: YES.
6	MS. BONNEVILLE: JEFF SHEEHY.
7	MR. SHEEHY: YES.
8	MS. BONNEVILLE: OSWALD STEWARD.
9	DR. STEWARD: YES.
10	MS. BONNEVILLE: JONATHAN THOMAS.
11	CHAIRMAN THOMAS: YES.
12	MS. BONNEVILLE: ART TORRES.
13	MR. TORRES: AYE.
14	MS. BONNEVILLE: KRISTINA VUORI.
15	DR. VUORI: YES.
16	MS. BONNEVILLE: DIANE WINOKUR.
17	MS. WINOKUR: I JOIN SHERRY
18	ENTHUSIASTICALLY, YES.
19	CHAIRMAN THOMAS: UNANIMOUSLY AND
20	ENTHUSIASTICALLY SUPPORTED. CONGRATULATIONS, DR.
21	MILLS AND TEAM.
22	(APPLAUSE.)
23	CHAIRMAN THOMAS: WE'RE GOING TAKE A
24	TEN-MINUTE BREAK. BE BACK IN TEN.
25	(A RECESS WAS TAKEN.)
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1	CHAIRMAN THOMAS: WE'RE GOING TO PICK UP
2	NOW WITH THE NEXT AGENDA TOPIC. AS YOU HEARD FROM
3	DR. MILLS, PART AND PARCEL OF THE STRATEGIC PLAN ARE
4	THREE INITIATIVES WE'RE GOING TO CONSIDER IN
5	SEQUENCE HERE, EACH TO BE PRESENTED BY DR. MILLAN.
6	START FIRST WITH THE ACCELERATING CENTER CONCEPT
7	PLAN.
8	DR. MILLAN: THANK YOU, CHAIRMAN THOMAS,
9	MEMBERS OF THE BOARD, THE PUBLIC, AND CIRM
10	COLLEAGUES. I'LL BE PRESENTING THE CONCEPT
11	PROPOSALS FOR TWO STRATEGIC INFRASTRUCTURE PROGRAMS,
12	THE ACCELERATING CENTER AND THE TRANSLATING CENTER.
13	DR. MILLS TOUCHED ON A LOT OF THIS JUST A
14	LITTLE WHILE AGO, BUT I WANTED TO JUST SET THE TONE
15	AND SET THE BACKGROUND FOR THE NEED FOR THESE
16	INITIATIVES.
17	AS HE SUMMARIZED IN HIS REPORT, CIRM'S
18	PIPELINE IS MATURING WITH GROWING NUMBERS OF LATE
19	TRANSLATIONAL, PRECLINICAL, AND CLINICAL STAGE
20	PROJECTS. AND OUR PLAN UNDER THE FIVE-YEAR
21	STRATEGIC PLAN IS TO GROW THESE PROGRAMS TO EVEN
22	MORE ROBUST PROPORTIONS, BUT WE DO IDENTIFY, THROUGH
23	MANAGEMENT OF OUR PORTFOLIO GRANTS AND THROUGH
24	EXTENSIVE DISCUSSIONS WITH STAKEHOLDERS, MULTIPLE
25	CHALLENGES IN BRINGING THESE DEVELOPMENT PROJECTS
	76

1	FORWARD. BUT WE SEE THESE CHALLENGES AS AN
2	OPPORTUNITY. WE SEE IT AS AN OPPORTUNITY TO
3	SIGNIFICANTLY DECREASE THE TIME IT TAKES TO BRING
4	PROJECTS FROM PRECLINICAL TRANSLATIONAL STAGE TO THE
5	CLINICS. AND WE PROPOSE THAT STRATEGICALLY DEPLOYED
6	INFRASTRUCTURE PROGRAMS COULD IDENTIFY AND ATTACK
7	THE UNIQUE CHALLENGES THAT GET IN THE WAY OF
8	BRINGING THESE DEVELOPMENT PROGRAMS FORWARD.
9	SO JUST TO IDENTIFY SOME CONCRETE
10	CHALLENGES THAT WE HAVE NOTED, PROCESS DEVELOPMENT,
11	CREATION OF ROBUST PROCESSES TO PRODUCE THESE CELL
12	THERAPY PRODUCTS THAT WILL GO INTO CLINICAL TESTING
13	AND TO THE PATIENTS IS A HUGE CHALLENGE AND CAN
14	REALLY HOLD THAT PROJECT AND SOMETIMES UNNECESSARILY
15	CAUSE UNDUE DELAY OR MAYBE CAUSE THOSE PROJECTS TO
16	BE STOPPED PREMATURELY.
17	IN ADDITION, THERE'S SOME HUGE HURDLES
18	THAT HAVE TO BE OVERCOME TO MEET THE REGULATORY
19	REQUIREMENTS FOR FILING AND TO ENGAGE WITH THE FDA
20	IN TERMS OF CRAFTING THE BEST PATH FORWARD TO
21	GETTING A PRODUCT INTO CLINICAL TESTING.
22	IN ADDITION, THESE PRODUCTS ARE UNIQUE
23	AND, THEREFORE, THE STEM CELL TRIALS AND DELIVERY OF
24	THESE PRODUCTS REQUIRE SOME SPECIALIZED
25	CONSIDERATIONS. WITH INFRASTRUCTURE PROGRAMS, WE'RE
	77

1	ABLE TO PUT IN PLACE FOCUSED EXPERTISE TO ADDRESS
2	THESE CHALLENGES. AND IN ADDITION, AS THESE
3	CHALLENGES ARE BEING ADDRESSED, THERE'S AN
4	AGGREGATED AND SELF-FEEDING LOOP OF EXPERIENCE AND
5	EXPERTISE THAT'S BUILT WITHIN THESE INFRASTRUCTURES
6	AND CONTINUAL IMPROVEMENT SO THEY CAN BEST SERVE THE
7	SPONSORS AND EVENTUALLY THE PATIENTS.
8	THE CONCEPT OF A ONE-STOP SHOP IS THERE'S
9	A GO-TO PLACE TO HAVE CONVERSATIONS, TO BUILD
10	RELATIONSHIPS SUCH AS CONVERSATIONS WITH THE FDA,
11	FOR INSTANCE. SO THE ENTITIES COULD BRING IN THEIR
12	AGGREGATED EXPERIENCE AND PRESENT ON BEHALF OF THE
13	SPONSORS AND THE STAKEHOLDERS THE PRODUCTS IN THE
14	BEST POSSIBLE WAY.
15	I PUT UP AGAIN THE SCHEMATIC DIAGRAM OF
16	THE JET ENGINE TO REMIND US THAT INFRASTRUCTURE
17	PROGRAMS AT CIRM, EITHER EXISTING OR THOSE THAT ARE
18	BEING PROPOSED, WILL WORK COORDINATELY WITH ALL OF
19	CIRM'S PROGRAMS TO BRING THEM THROUGH THE MACHINERY
20	FROM DISCOVERY TO THE CLINICAL PHASES. AND I'LL
21	FOCUS TODAY MAINLY ON THE CLINICAL AND LATE
22	PRECLINICAL INFRASTRUCTURE PROGRAMS.
23	FIRST, I'D LIKE TO JUST TAKE THIS
24	OPPORTUNITY TO UPDATE THE BOARD ON ONE OF THE
25	INFRASTRUCTURE PROGRAMS ALREADY IN PLACE, THE ALPHA

1	CLINICS NETWORK. ON THIS SLIDE ARE THE PROGRAM
2	DIRECTORS OF THE ALPHA CLINICS NETWORK: DR. JOHN
3	ADAMS, WHO'S ACTUALLY IN THE AUDIENCE, IS HERE FOR
4	THE UCLA/UC IRVINE CONSORTIUM; DR. CATRIONA JAMIESON
5	FROM UC SAN DIEGO; DR. JOHN ZAIA FROM THE CITY OF
6	HOPE.
7	THE GOAL OF THE ALPHA CLINICS NETWORK,
8	WHICH WAS JUST LAUNCHED ACTUALLY IN FEBRUARY OF THIS
9	YEAR, THE FIRST CLINIC WAS OPENED IN FEBRUARY OF
10	THIS YEAR, THE GOAL OF THIS NETWORK IS TO ACCELERATE
11	STEM CELL THERAPY DEVELOPMENT BY PROVIDING
12	EFFICIENT, SCALABLE, AND DURABLE INFRASTRUCTURE THAT
13	DELIVERS HIGH QUALITY CLINICAL TRIALS WITH THE
14	PATIENT EXPERIENCE AND PATIENT SAFETY AT THE
15	FOREFRONT.
16	AS A SUMMARY OF WHAT'S OCCURRED THIS YEAR,
17	THIS BOARD APPROVED FUNDING FOR THE ALPHA CLINICS
18	NETWORK IN OCTOBER OF 2014. IMMEDIATELY AFTER THAT
19	A PRELAUNCH MEETING WAS CONVENED WITH ALL OF THE
20	TEAMS, AND SHORTLY THEREAFTER SITE VISITS WERE
21	CONDUCTED. WE LAUNCHED THE CLINICS BETWEEN FEBRUARY
22	AND APRIL OF THIS YEAR AND ESTABLISHED REGULAR
23	STEERING COMMITTEE MEETINGS QUARTERLY ALONG WITH
24	MONTHLY CALLS IN ADDITION TO THE WEEKLY MEETINGS AND
25	DAILY COMMUNICATIONS THAT OCCUR AT ALL THESE
	70

1	CLINICS.
2	AND CURRENTLY THIS NETWORK SUPPORTS 14
3	ACTIVE CLINICAL TRIALS, INCLUDING DR. KOHN'S TRIAL
4	WHICH YOU HEARD ABOUT THIS MORNING FOR THE SEVERE
5	COMBINED IMMUNODEFICIENCY SYNDROME, AS WELL AS
6	SICKLE CELL ANEMIA, AND OTHER VERY IMPORTANT TRIALS
7	IN HIV/AIDS WITH GENE-MODIFIED CELL THERAPIES, AND
8	OTHER TRIALS THAT WERE JUST RECENTLY APPROVED FOR
9	FUNDING BY THIS BOARD THROUGH THE CIRM 2.0 IN
10	MULTICENTER TRIALS.
11	THESE CLINICS ARE ACTIVELY BUILDING A
12	ROBUST PIPELINE, DEPLOYING WHAT WE CALL ACCELERATING
13	AND VALUE-ADD RESOURCES WE CALL AVARS, AND THESE
14	AVARS ARE MEANT TO BE TOOLS TO ACTUALLY CONTINUE TO
15	BRING ABOUT EFFICIENCIES AND SHARE THESE
16	EFFICIENCIES ACROSS THE NETWORK AND BEYOND.
17	SOME EXAMPLES OF THESE AVARS ARE LISTED
18	HERE, AND THEY ARE TARGETED TOWARD SOME KEY GATING
19	ITEMS AND KEY OBSTACLES IN THE DELIVERY OF CRITICAL
20	STEM CELL TRIALS TO THE CLINICS AND TO PATIENTS.
21	ONE OF THEM LEVERAGES THE IRB RELIANCE RESOURCE FROM
22	THE UC BRAID AND BRINGS IT INTO THE NETWORK FOR THE
23	STEM CELL CLINICAL TRIALS. WHAT THIS ALLOWS THE
24	INVESTIGATORS TO DO IS TO GET SIMULTANEOUS AND
25	RECIPROCAL IRB APPROVALS SO ONE COULD GET A REVIEW

1	BY AN IRB AT ONE OF THE INSTITUTIONS AND IT WOULD
2	TRIGGER AUTOMATIC APPROVAL AT THE OTHER
3	INSTITUTIONS. AND THIS WOULD BRING ABOUT TIME AS
4	WELL AS EFFICIENCY CONSIDERATIONS.
5	IN ADDITION, THE NETWORK IS CONTINUALLY
6	BEING NIMBLE ADDRESSING PROBLEMS AS THEY ARISE,
7	ADDRESSING NEEDS OF THE PATIENTS. AND ONE OF THOSE
8	EXAMPLES IS ACTUALLY ACTIVELY IMPROVING THE PATIENT
9	INFORMED CONSENT PROCESS. AND THIS IS VERY TIMELY
10	GIVEN ALL OF THE NEW POLICIES AND APPROACHES TO
11	INFORMED CONSENT.
12	IN ADDITION, THE NETWORK SHARES
13	EXPERIENCES, KNOW-HOW, RESOURCES, AND ARE DOING THIS
14	IN A SYSTEMATIC WAY AND CAPTURING IT IN SHARED
15	DATABASES. ONE OF THE DATABASES WE'RE UTILIZING IS
16	EXISTING WITHIN THE UC BRAID SYSTEM AS WELL.
17	WITH THE ACCELERATING CENTER, WHICH I'LL
18	PRESENT SHORTLY, THESE AVARS WILL BE OPTIMIZED
19	ACROSS THE NETWORK, BUT ALSO WILL BE MADE AVAILABLE
20	TO ALL STEM CELL CLINICAL TRIALS AROUND CALIFORNIA.
21	SO I'LL MOVE ON, THEN, TO A NEW
22	INFRASTRUCTURE PROGRAM WHICH WE'RE BRINGING TO YOU
23	TODAY FOR CONSIDERATION AND APPROVAL OF THE CONCEPT,
24	WHICH IS THE ACCELERATING CENTER, WHICH IS A
25	STEM-CELL FOCUSED CLINICAL RESEARCH ORGANIZATION.
	0.1

1	THE GOAL OF THE ACCELERATING CENTER IS TO
2	ASSIST SPONSORS IN THE REGULATORY PATH AND THE
3	CLINICAL MANAGEMENT OF CLINICAL TRIALS IN A WAY THAT
4	SPEEDS THE PROGRESSION OF THESE PROJECTS.
5	THE ACTIVITIES OF THE ACCELERATING CENTER
6	ARE TO PROVIDE CUSTOMIZED SUPPORT FOR STEM CELL
7	CLINICAL TRIALS IN THE AREAS OF REGULATORY AFFAIRS,
8	CLINICAL TRIAL MANAGEMENT, AND DATA MANAGEMENT AND
9	ANALYTICS.
10	THEY WILL WORK ACTIVELY IN PARTNERSHIP
11	WITH THE FDA TO EFFICIENTLY MOVE STEM CELL PRODUCTS
12	THROUGH THE REGULATORY PATH. THERE WAS A LOT OF
13	DISCUSSION OF HOW THIS WOULD BE DONE, DATA DRIVES,
14	DATA SHOULD DRIVE, AND THE FDA WANTS DATA. SO THE
15	EXPERIENCES AND THE DATA THAT ARE BEING GATHERED AT
16	A GO-TO PLACE SUCH AS THIS CENTER WOULD VERY MUCH GO
17	A LONG WAY TOWARD THESE DISCUSSIONS AND TOWARD
18	CRAFTING NEW APPROACHES THAT WOULD BRING US TO OUR
19	GOAL.
20	IT ALSO WOULD COORDINATE WITH ALPHA
21	CLINICS NETWORK TO SCALE UP AND SCALE OUT THE ADDED
22	VALUE OF ACCELERATING RESOURCES, SOME OF WHICH WAS
23	JUST DESCRIBED, TO SUPPORT STEM CELL CLINICAL TRIALS
24	THROUGHOUT THE STATE AND BEYOND, AND THEY WOULD
25	CREATE A SUSTAINABLE RESOURCE FOR THE GROWING

1	PIPELINE OF STEM CELL TRIALS IN CALIFORNIA.
2	JUST TO GO INTO A LITTLE BIT MORE DETAIL
3	ABOUT WHAT THE REGULATORY AFFAIRS ACTIVITIES WOULD
4	BE OF THE ACCELERATING CENTER, THEY WOULD ACTUALLY
5	PLAY AN ESSENTIAL ROLE FOR FDA INTERACTIONS. THEY
6	WOULD BE WITH A SPONSOR IN ASSEMBLING THE IND, WOULD
7	BE THERE FOR DISCUSSIONS, WOULD HAVE ACCESS TO THE
8	KNOWLEDGE BASE OF THE AGGREGATED EXPERIENCE OF THE
9	STEM CELL CLINICAL TRIALS IN CIRM'S PORTFOLIO, AND,
10	THEREFORE, WOULD BE WELL POSITIONED TO DRIVE THESE
11	DISCUSSIONS AND TO IMPLEMENT WHAT WE HOPE ARE SOME
12	VERY NEEDED NEW APPROACHES TO BRINGING THESE
13	DEVELOPMENT PROGRAMS THROUGH THE REGULATORY PATH.
14	IN TERMS OF CLINICAL TRIALS SUPPORT, THEY
15	WOULD ASSIST THE SITES IN THEIR CLINICAL OPERATIONS
16	MANAGEMENT AND PROVIDE LOGISTICAL SUPPORT WHERE
17	REQUIRED. AND THIS SUPPORT WOULD BE SCALED
18	ACCORDING TO THE NEEDS OF THE SPONSOR, THE NEEDS OF
19	THE TRIALS.
20	IN ADDITION TO PROVIDING THE DATA
21	MANAGEMENT AND ANALYTICS THAT ARE REQUIRED TO
22	SUPPORT THE CLINICAL TRIALS, THE ACCELERATING CENTER
23	WOULD ALSO PROVIDE A CENTRALIZED REPOSITORY FOR
24	REALLY ESSENTIAL AND CRITICAL DATASETS AND KNOWLEDGE
25	THAT WOULD SERVE IN THE DISCUSSIONS WITH THIRD-PARTY

1	STAKEHOLDERS SUCH AS REGULATORS AND WITH
2	REIMBURSEMENT PARTIES.
3	WE PROPOSE THE TIMELINE THAT'S POSTED
4	HERE. EXPECT THAT WITH APPROVAL WE WOULD BE ABLE TO
5	GET THIS ACCELERATING CENTER RFA OUT IN THE
6	BEGINNING OF 2016, HAVE IT REVIEWED AND BROUGHT BACK
7	TO YOU FOR APPROVAL IN THE SECOND HALF OF 2016,
8	HOPEFULLY HAVING THE CENTER OPERATIONAL BY THE END
9	OF 2016, WITH A PROPOSED BUDGET OF \$15 MILLION, UP
10	TO \$15 MILLION IN FUNDING.
11	CHAIRMAN THOMAS: THANK YOU, DR. MILLAN.
12	ARE THERE QUESTIONS ON THE ACCELERATING CENTER FROM
13	MEMBERS OF THE BOARD?
14	DR. PRICE: WHAT SORT OF ENTITY OR
15	ENTITIES DO YOU ANTICIPATE APPLYING TO BECOME
16	ACCELERATING CENTERS?
17	MS. MILLAN: A CLASSIC CRO, BUT A CRO THAT
18	HAS AN INTEREST IN THE REGENERATIVE MEDICINE STEM
19	CELL SPACE. SO THEY WOULD ACTUALLY COME IN AND SET
20	UP A SPECIALIZED UNIT FOR THAT. WE HAVE HAD
21	DISCUSSIONS WITH A VARIETY OF KIND OF LARGER CRO'S.
22	THERE IS INTEREST IN THIS. THIS WOULD BE AN
23	INCENTIVE FOR THEM TO DO IT. TYPICALLY THEY SERVE
24	VERY LARGE TRIALS. LATER STAGE MULTICENTER TRIALS
25	ARE MORE ACCUSTOMED TO CANCER TRIALS OR SUCH. SO
	0.4

1	THESE SPECIALIZED TRIALS ARE TYPICALLY NOT WITHIN
2	THEY MAY SUPPORT THEM, BUT NOT IN ANY TYPE OF
3	STRATEGIC MANNER.
4	THERE IS INTEREST OUT THERE IN CREATING
5	UNITS FOR THIS BECAUSE THEY SEE THAT THE FIELD IS
6	GOING THAT WAY, AND THEY WOULD LIKE TO BE ABLE TO
7	START PROVIDING THAT TYPE OF SERVICE TO THE
8	COMMUNITY.
9	MR. PANETTA: GREAT PRESENTATION. I WANT
10	TO KIND OF BUILD ON THAT QUESTION AND ASK YOU IF YOU
11	COULD FURTHER CLARIFY THE EXTENT TO WHICH THIS SORT
12	OF CAPABILITY OR AT LEAST THE POTENTIAL TO EXPAND ON
13	EXISTING CAPABILITY IS HERE IN CALIFORNIA. HOW BIG
14	OF AN UNDERTAKING IS THIS GOING TO BE IN ADDITION TO
15	WHAT ALREADY EXISTS OUT THERE IN THE WAY OF
16	CAPABILITY, IF YOU KNOW? MAYBE WE DON'T KNOW TILL
17	WE ASK.
18	MS. MILLAN: WITHIN CALIFORNIA THERE ARE
19	VERY FEW CRO'S WITH THIS SPECIALIZATION. THAT WE
20	KNOW. MANY OF OUR GRANTEES EITHER GO OUT OF STATE
21	OR IN MANY CASES THEY TRY TO JUST WORK WITH WHAT
22	THEY HAVE, AND PUT BITS AND PIECES TOGETHER. SO
23	THERE IS NOTHING THAT WE KNOW OF THAT IS A
24	SPECIALIZED STEM CELL CRO THAT EXISTS, PERIOD, AND
25	DEFINITELY NOT IN CALIFORNIA.
	0.5

1	WHEN WE SPEAK TO OUR GRANTEES, THEY DO
2	BELIEVE THAT THIS WOULD BE USEFUL BECAUSE EVEN JUST
3	THE PROCESS OF IDENTIFYING THE RIGHT CRO WHO WOULD,
4	FIRST OF ALL, TAKE ON THEIR STUDY FOR THE BUDGET
5	THAT THEY HAVE AVAILABLE TO THEM FOR SUCH A SMALL
6	STUDY, AS WELL AS THE SPECIALIZATION THAT ONE WOULD
7	NEED TO KIND OF UNDERSTAND THE TYPES OF STUDIES,
8	THAT DOESN'T CURRENTLY EXIST. SO WE DO BELIEVE
9	WE'RE ADDRESSING SOMETHING THAT IS CRITICAL, BUT NOT
10	CURRENTLY IN EXISTENCE.
11	MR. HIGGINS: THANK YOU. I'M AN
12	ENTHUSIASTIC SUPPORTER OF THIS IDEA, BUT A QUESTION
13	COMES TO MIND THAT YOUR SUCCESS MAY LEAD TO THIS
14	PARTICULAR STRUCTURE SURVIVING BEYOND CIRM. HAVE
15	YOU GIVEN THAT ANY THOUGHT?
16	MS. MILLAN: YES, WE HAVE. AND WHAT WE
17	PLAN TO DO IN TERMS OF THE STRUCTURE OF THIS IS
18	INITIALLY THIS ACCELERATING CENTER WOULD BE REQUIRED
19	TO SUPPORT THE CIRM PROGRAMS FOR THE SETUP BECAUSE
20	WE DO BELIEVE THAT THERE'S SOME JUST NEEDS THAT WE
21	WANT FILLED, BUT ALSO WE BELIEVE THAT THERE ARE
22	STANDARDS THAT WE WANT SET, AND THERE'S A CERTAIN
23	LEVEL OF PRODUCT THAT THEY CAN CREATE DURING THAT
24	INITIAL PERIOD OF TIME. THEN THEREAFTER THAT CENTER
25	WOULD BE ABLE TO GO AFTER ADDITIONAL BUSINESS,

1	LEVERAGING NOW THE PRODUCT THAT THEY'VE CREATED
2	TOWARD A SPECIALIZED SERVICE. CIRM WOULDN'T PAY FOR
3	THAT, BUT THEY WOULD USE THAT TO SUSTAIN THEIR
4	BUSINESS FOR THE FUTURE.
5	CHAIRMAN THOMAS: OTHER COMMENTS,
6	QUESTIONS FROM MEMBERS OF THE BOARD?
7	DR. PRICE: ARE THERE FUNCTIONS ENVISIONED
8	HERE FOR THE ACCELERATING CENTERS THAT ARE REDUNDANT
9	WITH WHAT THE ALPHA CLINICS ARE PROVIDING? IF SO,
10	WOULD IT MAKE SENSE TO THINK ABOUT A FUTURE MERGING
11	OF THIS?
12	MS. MILLAN: THANK YOU FOR THAT QUESTION.
13	IN FACT, THE PROPOSED STRUCTURE OF THE ACCELERATING
14	CENTER IS THAT IT WOULD ACTUALLY WORK
15	SYNERGISTICALLY WITH THE ALPHA CLINIC NETWORKS.
16	AND, IN FACT, THE ALPHA CLINICS NETWORK NEEDS ARE
17	CONSIDERED AS WE HAVE CRAFTED THE CONCEPT, AND THEN
18	WE'LL CRAFT THE RFA AND STRUCTURE.
19	AND IN ADDITION, THE ALPHA CLINICS NETWORK
20	HAVE IDENTIFIED KEY ASSETS AND PRODUCTS THAT WE'RE
21	NAMING AVAR. WE NAME IT AVAR SO THAT WE CAN HAVE
22	CONCRETE THINGS THAT WE CAN POINT TO THAT ACTUALLY
23	ENVISION THAT THE ACCELERATING CENTER WILL WANT TO
24	BRING IN AND DEVELOP THOSE PRODUCTS SO THAT THEY CAN
25	EITHER ASSIST THE ALPHA CLINICS IN OPTIMIZING THOSE
	0.7

1	RESOURCES, EXPAND THOSE ACTIVITIES, AS WELL AS OFFER
2	IT TO ADDITIONAL SITES AND INVESTIGATORS, INITIALLY
3	CIRM AND THEN OTHERWISE.
4	DR. FRIEDMAN: I TOO THINK IT'S A REALLY
5	INTERESTING AND GOOD IDEA. IT SEEMS AS THOUGH IT'S
6	MADE UP OF TWO COMPONENTS ROUGHLY. ONE IS THE
7	SKILLFUL MANAGEMENT OF SOME CONVENTIONAL CRO TASKS,
8	DATA MANAGEMENT, REPORTING, THINGS LIKE THAT. AND I
9	THINK THAT'S REALLY VERY GOOD BECAUSE IT'S A
10	VARIABLE QUALITY THROUGHOUT DIFFERENT GRANTEES AND
11	EVEN INDUSTRIES IN CALIFORNIA. SO THAT MAKES A LOT
12	OF SENSE.
13	THE SECOND COMPONENT IS SOMEWHAT MORE
14	CREATIVE, AND I WONDER IF YOU CAN TALK TO US A
15	LITTLE BIT ABOUT THAT, AGENCY INTERACTIONS, THINGS
16	THAT LIKE THAT. AND THAT'S THE AREA THAT I THINK IS
17	MORE CHALLENGING AND ALSO MORE INTRIGUING.
18	DR. MILLAN: THANK YOU. SO MANY OF OUR
19	SITES, INCLUDING THE ALPHA CLINICS, HAVE REGULATORY
20	TEAMS, FOR INSTANCE. AND THESE REGULATORY TEAMS IN
21	THEIR OWN CENTERS HAVE DEVELOPED SOME RAPPORT AND
22	EXPERTISE SURROUNDING THEIR PARTICULAR PROJECT. SO
23	WE USED THAT AS A NICE EXAMPLE OF HOW IT'S
24	ADVANTAGEOUS TO HAVE THAT ASSET. HOWEVER, THIS IS
25	NOT AVAILABLE TO ALL, AND IT'S NOT AVAILABLE TO THE
	0.0

1	VARIETY OF DIFFERENT TECHNOLOGY PLATFORMS THAT WE
2	HAVE IN OUR PORTFOLIO AND ARE ANTICIPATING COMING
3	IN.
4	SO WHAT WE WILL ASK THE APPLICANT TO BE
5	ABLE TO DEMONSTRATE IS THEY'LL HAVE THE CAPABILITY
6	FOR REGULATORY SUPPORT FOR FILING AN IND FOR
7	REGULATORY STRATEGY. AND TRADITIONAL CRO'S HAVE
8	THAT, BUT THE DIFFERENCE WITH THIS CENTER IS NOW IT
9	WILL BE FOCUSED ON CELL THERAPY, STEM CELL THERAPIES
10	SO THEY WILL NOW REALLY PUT THEIR RESOURCES AND
11	THEIR BRAIN POWER TO IT. AND AS THEY SUPPORT MORE
12	AND MORE OF THESE TYPES OF TRIALS, THEY'LL LEARN
13	MORE AND MORE AND BE ABLE TO BRING THAT AGGREGATED
14	EXPERIENCE EACH TIME THEY REPRESENT THE NEXT
15	SPONSOR.
16	DR. FRIEDMAN: I THINK THAT'S REALLY
17	VALUABLE. IT STRIKES ME, SINCE THE REGULATORY
18	SIMPLIFICATION AND EFFICIENCY IS SUCH A KEY
19	COMPONENT OF OUR STRATEGIC PLAN, THAT IT'S POSSIBLE
20	THAT THE CRO COULD HAVE A DIFFERENT POINT OF VIEW
21	AND TAKE A DIFFERENT TACT THAN THE LEADERSHIP OF
22	CIRM. AND WE JUST NEED I JUST ASK YOU TO THINK
23	ABOUT HOW THAT WILL OCCUR. USUALLY A CRO IS VERY
24	SKILLFUL AT A SET PATHWAY FOR BIOLOGICS, DEVICES, OR
25	DRUGS BECAUSE THEY ALL WALK THE CLIENT THROUGH THAT
	00

1	PATH. HERE YOU ARE DISCOVERING NEW GROUND AND A LOT
2	OF DIFFERENT WAYS OF APPROACHING EVEN DISAGREEMENT.
3	WE SHOULD JUST THINK ABOUT HOW TO ADJUDICATE AND
4	SIMPLIFY THOSE THINGS LATER.
5	DR. MILLAN: ABSOLUTELY. AND PART OF THE
6	APPLICATION AND THE REVIEW WILL BE TO SORT THROUGH
7	WHICH TYPES OF ORGANIZATIONS WOULD BE ABLE TO DO
8	THAT. AND THERE WILL BE A STEERING COMMITTEE THAT'S
9	SET UP AFTER THE AWARD AS PART OF THE TERMS OF THE
10	AWARD WHICH INVOLVES CIRM AND PROBABLY THE
11	TRANSLATING CENTER. AND THERE WILL BE AN
12	ACCELERATING CENTER ALPHA CLINIC STEERING COMMITTEE
13	AS WELL.
14	CHAIRMAN THOMAS: OTHER COMMENTS FROM
15	MEMBERS OF THE BOARD? COMMENTS FROM MEMBERS OF THE
16	PUBLIC?
17	DR. LORING: THIS IS JEANNE LORING FROM
18	THE SCRIPPS RESEARCH INSTITUTE. WHEN I THINK ABOUT
19	STEM CELL CRO'S, ONLY ONE NAME COMES TO MIND, AND
20	THAT'S LONZA BECAUSE I KNOW THAT THEY'RE ALREADY
21	ESTABLISHED. THEY'VE ACTUALLY DONE SOME WORK FOR
22	SOME CIRM INVESTIGATORS. NOW, THEY'RE NOT LOCATED
23	IN CALIFORNIA, BUT ARE YOU CONSIDERING TRYING TO
24	ENCOURAGE OTHER EXISTING CRO'S TO ADOPT THEIR SORT
25	OF BUSINESS PLAN, OR ARE YOU THINKING ABOUT
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1	BRAND-NEW CRO'S BEING INITIATED AS BUSINESSES IN
2	CALIFORNIA?
3	DR. MILLAN: SO THE APPLICANT WE WOULD
4	LIKE TO COME IN WOULD HAVE A BREADTH OF EXPERIENCE
5	AND A TRACK RECORD FOR BEING ABLE TO EXECUTE ON THE
6	TYPES OF ACTIVITIES. CERTAINLY CRO'S THAT HAVE
7	EXPERIENCE IN THE ACTIVITIES WE LAID OUT IN TERMS OF
8	THE REGULATORY FILINGS, CLINICAL OPERATIONS WOULD BE
9	COMPETITIVE FOR THIS TYPE OF AWARD.
10	DR. LORING: THANK YOU.
11	DR. BRATT-LEAL: HI. ANDRES BRATT-LEAL
12	FROM STEM CELL AND SCRIPPS RESEARCH INSTITUTE. I
13	THINK IT'S GREAT. IT WOULD BE A GREAT RESOURCE. MY
14	QUESTION IS CIRM IS GOING TO BE CREATING A HUGE
15	DEMAND FOR CRO WORK WITH THE AMBITIOUS CIRM 2.0
16	PROPOSALS WITH 30 AND 40 AND 45 TRANSLATIONAL AWARDS
17	AND THEN A LOT OF TRANSLATIONAL AWARDS. DOES THIS
18	MATCH THE AMOUNT OF DEMAND THAT YOU EXPECT THAT'S
19	GOING TO BE COMING FOR CRO'S, ESPECIALLY FOR STEM
20	CELL EXPERTISE? I GUESS MY QUESTION WAS DO YOU EVEN
21	NEED MORE THAN THIS TO MATCH THAT DEMAND?
22	DR. MILLAN: WHAT WE HOPE FOR IS THAT
23	WE'LL NEED MORE THAN THIS. THE WAY THIS WOULD BE
24	SET UP HOW IT WILL BE SET UP IS THESE
25	ORGANIZATIONS SHOULD BE SCALABLE AND BE ABLE TO MEET

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1	THE DEMANDS WITHIN CALIFORNIA AND EVENTUALLY FOR
2	MULTICENTER AND INTERNATIONAL TRIALS INVOLVING
3	CALIFORNIA.
4	DR. JUELSGAARD: SO THERE ARE SOME REALLY
5	GREAT CRO'S THAT ARE OUT THERE THAT HAVE BEEN AROUND
6	FOR A LONG TIME AND REALLY KNOW WHAT THEY'RE DOING,
7	AND THEY MAY NOT HAVE A TREMENDOUS AMOUNT OF
8	EXPERIENCE OF STEM CELL THERAPY, BUT THEY KNOW THE
9	NUTS AND BOLTS OF EVERY OTHER PART OF THE PROCESS.
10	SO BECAUSE THIS NEEDS TO BE A CALIFORNIA-BASED
11	ORGANIZATION, MEANING GREATER THAN 50 PERCENT OF
12	THEIR EMPLOYEES RESIDE HERE, ETC., IS IT POSSIBLE
13	THAT A CRO BASED IN CHICAGO OR WHATEVER COULD CREATE
14	A SUBSIDIARY HERE IN CALIFORNIA AND THAT SUBSIDIARY
15	HAS TEN PEOPLE, AND SIX OF THEM LIVE IN CALIFORNIA
16	AND THAT WOULD QUALIFY. DOESN'T HAVE TO BE THE
17	PARENT CRO, RIGHT?
18	DR. MILLAN: ABSOLUTELY. THAT'S WHAT WE
19	HAVE IN MIND AS WE'RE LOOKING AT THIS.
20	CHAIRMAN THOMAS: HEARING NO FURTHER
21	DISCUSSION, DO WE HAVE A MOTION?
22	ANYBODY ON THE PHONE HAVE ANY QUESTIONS?
23	DR. LUBIN: NONE FOR ME.
24	CHAIRMAN THOMAS: HEARING NO QUESTIONS,
25	IT'S BEEN MOVED BY MS. LANSING. IS THERE A SECOND?
	0.2

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1	IS THERE A SECOND? SECOND BY SENATOR TORRES. ANY
2	FURTHER DISCUSSION BY MEMBERS OF THE BOARD? HEARING
3	NONE, MARIA, PLEASE TAKE THE ROLL.
4	MS. BONNEVILLE: DAVID BRENNER. LINDA
5	BOXER.
6	DR. BOXER: YES.
7	MS. BONNEVILLE: KEN BURTIS. ANNE-MARIE
8	DULIEGE.
9	DR. DULIEGE: YES.
10	MS. BONNEVILLE: MICHAEL FRIEDMAN.
11	DR. FRIEDMAN: YES.
12	MS. BONNEVILLE: JUDY GASSON.
13	DR. GASSON: YES.
14	MS. BONNEVILLE: SAM HAWGOOD. DAVID
15	HIGGINS.
16	DR. HIGGINS: YES.
17	MS. BONNEVILLE: STEPHEN JUELSGAARD.
18	MR. JUELSGAARD: YES.
19	MS. BONNEVILLE: SHERRY LANSING.
20	MS. LANSING: YES.
21	MS. BONNEVILLE: KATHY LAPORTE.
22	DR. LAPORTE: YES.
23	MS. BONNEVILLE: BERT LUBIN.
24	DR. LUBIN: YES.
25	MS. BONNEVILLE: SHLOMO MELMED.
	93

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1	DR. MELMED: YES.
2	MS. BONNEVILLE: LAUREN MILLER.
3	MS. MILLER: YES.
4	MS. BONNEVILLE: ADRIANA PADILLA.
5	DR. PADILLA: YES.
6	MS. BONNEVILLE: JOE PANETTA.
7	MR. PANETTA: YES.
8	MS. BONNEVILLE: ROBERT PRICE.
9	DR. PRICE: YES.
10	MS. BONNEVILLE: FRANCISCO PRIETO.
11	DR. PRIETO: AYE.
12	MS. BONNEVILLE: CARMEN PULIAFITO. ROBERT
13	QUINT. AL ROWLETT.
14	MR. ROWLETT: YES.
15	MS. BONNEVILLE: JEFF SHEEHY.
16	MR. SHEEHY: YES.
17	MS. BONNEVILLE: OSWALD STEWARD.
18	DR. STEWARD: YES.
19	MS. BONNEVILLE: JONATHAN THOMAS.
20	CHAIRMAN THOMAS: YES.
21	MS. BONNEVILLE: ART TORRES.
22	MR. TORRES: AYE.
23	MS. BONNEVILLE: KRISTINA VUORI.
24	DR. VUORI: YES.
25	MS. BONNEVILLE: DIANE WINOKUR.
	0.4
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1	MS. WINOKUR: YES.
2	CHAIRMAN THOMAS: MOTION PASSES. THANK
3	YOU. ON TO COMPONENT NO. 2, THE TRANSLATING CENTER
4	CONCEPT PLAN.
5	DR. MILLAN: THANK YOU VERY MUCH. THE
6	TRANSLATING CENTER IS THE PARTNER PROGRAM TO THE
7	ACCELERATING CENTER, AND IT IS A PRECLINICAL
8	RESEARCH ORGANIZATION THAT CIRM HOPES TO PUT IN
9	PLACE AS ANOTHER KEY INFRASTRUCTURE PROGRAM TO
10	ADDRESS THE CHALLENGES THAT I JUST RECENTLY
11	SUMMARIZED.
12	AND THE MAJOR ACTIVITIES OF THIS
13	TRANSLATING CENTER ARE TO EXECUTE ON WHAT WE CALL
14	IND-ENABLING, MEANING THE WORK THAT NEEDS TO BE DONE
15	IN ORDER TO FILE AN IND TO GAIN PERMISSION TO GO
16	INTO CLINICAL TRIALS.
17	THE MAJOR ACTIVITIES OF THE TRANSLATING
18	CENTER WOULD BE PROVIDE CORE SERVICES TO THE
19	SPONSORS, TO THE INVESTIGATOR FOR CREATING PROCESSES
20	THAT THEY CAN REPRODUCIBLY PRODUCE THEIR CELL
21	PRODUCT. AND THIS IS A HUGE CHALLENGE ACTUALLY FROM
22	GETTING DEVELOPMENT CANDIDATES TO THE CLINICS THAT
23	IS A RECURRING CHALLENGE FOR MANY OF OUR
24	INVESTIGATORS AND OTHERS. AND WE KNOW THAT THERE'S
25	DEMAND FOR THIS EVEN WITHIN OUR OWN PORTFOLIO. WE

1	KNOW THERE'S DEMAND FOR THIS FOR FOLKS WHO WANT TO
2	COME INTO CIRM TO DEVELOP THEIR PROMISING
3	CANDIDATES, BUT DON'T HAVE THE SKILL SET.
4	IN ADDITION, THIS TRANSLATING CENTER WOULD
5	OVERSEE, MANAGE, ASSEMBLE THE NECESSARY PRECLINICAL
6	STUDIES THAT WOULD GO INTO THE IND AND THEN WOULD
7	ASSIST THE ACCELERATING CENTER, THE ACCELERATING
8	CENTER BEING THE LEAD ON THE FDA INTERACTIONS, WOULD
9	ASSIST THE ACCELERATING CENTER IN IND PREPARATION
10	AND SUBMISSION.
11	AND, AGAIN, THE SAME IDEA THAT WITH THIS
12	CONTINUAL FOCUSED EXPERIENCE, THAT THE PRODUCT
13	BECOMES BETTER AND BETTER AND THEY CAN BRING THEIR
14	AGGREGATED KNOWLEDGE TO THE TABLE.
15	SO SOME OF THE ACTIVITIES ARE LISTED HERE,
16	AGAIN, RELATED TO WHAT NEEDS TO HAPPEN TO GET A
17	DEVELOPMENT CANDIDATE TO BE A PRODUCT.
18	REPRODUCIBLE, ROBUST PROCESSES, ALL THE THINGS THAT
19	GO INTO THAT, AS WELL AS QUALITY SYSTEMS AND ASSAYS
20	THAT NEED TO BE VALIDATED OR AT LEAST INITIALLY
21	VALIDATED, AS WELL AS LOOKING AT FORWARD PLANNING
22	FOR SUCCESS IN TERMS OF WHAT TYPES OF PROCESSES AND
23	WHAT SYSTEMS AND DOCUMENTATION NEED TO BE IN PLACE
24	IN ORDER TO BE ABLE TO TRANSFER THIS TECHNOLOGY TO A
25	MANUFACTURER, AND WHAT CONSIDERATIONS NEED TO BE

1	INCORPORATED EARLIER ON IN A QUALITY-BY-DESIGN
2	MANNER SO THAT THESE PROCESSES CAN BE SCALED UP AND
3	SCALED OUT WHEN APPROPRIATE SHOULD THE PRODUCT GO
4	THROUGH A SUCCESSFUL PATH IN DEVELOPMENT.
5	THE TRANSLATING CENTER WOULD ALSO ASSEMBLE
6	THE PRECLINICAL DATASETS, THEY WOULD INTERACT WITH
7	THE SPONSOR, INVESTIGATORS WHO HAVE THE PRECLINICAL
8	STUDIES IN TERMS OF THE SCIENCE, THE EFFICACY-TYPE
9	DATASETS, CHARACTERIZATION DATASETS FROM THE SPONSOR
10	AS WELL AS FROM THE INTERNAL WORK DURING PROCESS
11	DEVELOPMENT, AND WITH EXTERNAL CRO'S OR INTERNAL, IF
12	THEY HAVE IT, BUT MOST LIKELY EXTERNAL CRO'S TO DO
13	WHAT DR. MILLS HAD REFERRED TO EARLIER AS WHAT MAY
14	BE PERCEIVED AS UNINTERESTING BUT NECESSARY SAFETY,
15	TOXICITY, DISTRIBUTION STUDIES, THINGS THAT ARE
16	REQUIRED FOR A CELL THERAPY IND. AND THEY WOULD
17	COORDINATE, AGAIN, WITH THE ACCELERATING CENTER TO
18	COMPILE THIS DATASET INTO AN IND.
19	SO FOR THIS CENTER WE ARE PROPOSING AN UP
20	TO \$15 MILLION BUDGET. WE WILL PUT OUT THE RFA FOR
21	THIS IN 2016, AGAIN, HOPE TO BRING, AFTER THE REVIEW
22	PROCESS, RECOMMENDATIONS TO THIS BOARD FOR APPROVAL
23	BY THE END OF 2016.
24	CHAIRMAN THOMAS: THANK YOU, DR. MILLAN.
25	QUESTIONS, COMMENTS FROM MEMBERS OF THE BOARD?
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1	MR. SHEEHY: I MOVE ADOPTION OF THIS FOR
2	APPROVAL.
3	DR. PRICE: SECOND.
4	CHAIRMAN THOMAS: MOVED BY MR. SHEEHY.
5	SECONDED BY DR. PRICE. OKAY. DISCUSSION? COMMENTS
6	FROM MEMBERS OF THE PUBLIC? ANY COMMENTS FROM
7	MEMBERS ON THE PHONE? HEARING NONE, MARIA, PLEASE
8	CALL THE ROLL.
9	MS. BONNEVILLE: DAVID BRENNER. LINDA
10	BOXER. KEN BURTIS. ANNE-MARIE DULIEGE.
11	DR. DULIEGE: YES.
12	MS. BONNEVILLE: MICHAEL FRIEDMAN.
13	DR. FRIEDMAN: YES.
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25	DR. MELMED: YES.
	98
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21	MR. TORRES: AYE.
22	MS. BONNEVILLE: KRISTINA VUORI.
23	DR. VUORI: YES.
24	MS. BONNEVILLE: DIANE WINOKUR.
25	WE'RE GOING TO CHECK THE PHONE LINES. I
	99
	33

1	DON'T THINK WE CAN HEAR THEM.
2	CHAIRMAN THOMAS: THOSE OF YOU ON THE
3	PHONE, I THINK WE LOST YOU FOR A SECOND. MARIA WAS
4	IN THE PROCESS OF CALLING THE ROLL ON THIS ITEM.
5	LET'S TRY ONE MORE TIME.
6	MS. BONNEVILLE: LINDA BOXER.
7	DR. BOXER: YES.
8	CHAIRMAN THOMAS: ARE THOSE ON THE LINE
9	MS. BONNEVILLE: WE'RE GOING TO HOLD THE
10	ROLL OPEN AND MOVE ON TO THE NEXT ITEM UNTIL WE GET
11	THE AUDIO TAKEN CARE OF.
12	CHAIRMAN THOMAS: THANK YOU. WE'RE GOING
13	TO GO ON TO THE NEXT ITEM. I MIGHT GIVE, BY WAY OF
14	PRELUDE, THE COMMENT THAT OVER A YEAR AGO DR. MILLS
15	AND I WERE SITTING AROUND TALKING ABOUT A VARIETY OF
16	THINGS. AND WE HAD SORT OF INDEPENDENTLY COME TO
17	THE NOTION THAT IT WOULD BE AN IDEA TO CONSIDER AN
18	ARM'S LENGTH, ADJUNCT ENTITY THAT COULD DRAW
19	INDUSTRY IN TO HELP WITH THE ACCELERATION AND
20	COMMERCIALIZATION OF SOME OF OUR MOST PROMISING
21	TECHNOLOGIES. AND WE TALKED ABOUT IT AT THAT POINT,
22	AND FROM THAT CONVERSATION CAME WHAT TURNED OUT TO
23	BE ABOUT 15 MONTHS OF VERY HARD WORK BY MANY PEOPLE
24	THAT HAS ENDED WITH THE NOTION OF ACCELERATING
25	THERAPIES PUBLIC PRIVATE PARTNERSHIP OR ATP3 THAT'S
	100
	100

1	GOING TO BE PRESENTED TO YOU TODAY.
2	SO LET'S TURN NOW TO THAT THIRD COMPONENT
3	OF THE STRATEGIC PLAN, ATP3. DR. MILLAN.
4	DR. MILLAN: CAN EVERYBODY SEE THE SCREEN.
5	THE PRESENTATION IS ALSO ON THE AGENDA AND MAY BE
6	PRINTED OUT. I'LL NOW PRESENT THE ACCELERATING
7	THERAPIES THROUGH A PUBLIC PRIVATE PARTNERSHIP WHICH
8	I'LL REFER TO AS ATP3.
9	BY WAY OF BACKGROUND, AGAIN, THIS IS A
10	LITTLE BIT REPETITIVE FROM WHAT DR. MILLS HAD
11	ALREADY LAID OUT. WE HAVE IDENTIFIED AND WE KNOW AS
12	A FIELD THAT THERE IS A LACK OF INDUSTRY PULL FOR
13	STEM CELL THERAPEUTICS. THOUGH CIRM HAS INVESTED
14	APPROXIMATELY \$2 BILLION SO FAR IN DEVELOPING A
15	PORTFOLIO OF APPROXIMATELY 300 TECHNOLOGIES, WE KNOW
16	THAT ONLY 6 PERCENT OF CIRM'S ACADEMIC PROJECTS HAVE
17	BEEN LICENSED BY INDUSTRY. AND IN DISCUSSIONS WITH
18	THE UNIVERSITY OF CALIFORNIA SYSTEM, WE KNOW THAT OF
19	THE 3400 TECHNOLOGIES BEING MARKETED, WE'RE NOT EVEN
20	TALKING ABOUT ALL TECHNOLOGIES, BUT JUST THOSE THAT
21	ARE BEING ACTIVELY MARKETED, LESS THAN 2 PERCENT OF
22	THOSE ARE STEM CELL PROGRAMS.
23	SO WE ARE PROPOSING TO THE BOARD TODAY AN
24	INITIATIVE, THE ATP3 INITIATIVE, AS A MEANS OF
25	ENGAGING INDUSTRY BY CREATING AN OPPORTUNITY FOR TOP

1	TIERED LEADERSHIP AND MANAGEMENT TEAMS TO COME IN,
2	AND COMPETITIVELY BE EVALUATED IN THEIR ABILITY TO
3	FORM AN ENTITY WHICH WOULD AGGREGATE CIRM'S MOST
4	PROMISING TECHNOLOGIES. BY AGGREGATION, IT WOULD
5	OFFER MULTIPLE SHOTS ON GOAL ON THESE PRODUCT
6	DEVELOPMENT CANDIDATES WHICH INCREASES THE
7	PROBABILITY OF SUCCESS, SO CALLED DERISKING THE
8	PROPOSITION. AND WHAT WE ANTICIPATE IS THIS WOULD
9	MAKE IT MORE SIGNIFICANTLY PALATABLE AND ACTUALLY
10	INCENTIVIZE INDUSTRY TO COME IN IN PARTNERSHIP.
11	IN ADDITION, WHAT'S BAKED INTO THIS
12	INITIATIVE IS THAT CIRM WOULD LEVERAGE ITS
13	CAPACITIES IN TERMS OF ADMINISTRATIVE REVIEW
14	STRUCTURE AND ADVISORS TO HELP THIS ENTITY COME UP
15	WITH THE BEST POSSIBLE PORTFOLIO. AND CIRM WOULD
16	CONTINUE TO BE INVOLVED BY FUNDING THE DEVELOPMENT
17	OF THESE IN-LICENSED TECHNOLOGIES.
18	SO AS A GENERAL STRUCTURE, THE
19	ACCELERATING THERAPIES TO PUBLIC PRIVATE
20	PARTNERSHIP, ATP3, THE MAJOR GOAL OF THIS IS TO GET
21	THE CIRM-FUNDED STEM CELL TECHNOLOGY CANDIDATES TO
22	THE PATIENTS, GET THE TECHNOLOGIES TO THE PATIENTS.
23	AND HOW DO WE DO THAT? WE PULL INDUSTRY IN, WE GET
24	A PRIVATE PARTNER THROUGH THIS COMPETITIVE PROCESS
25	WHO WILL IN-LICENSE, DEVELOP, AND DRIVE TOWARD
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1	COMMERCIALIZATION THE AGGREGATED PORTFOLIO. AND AS
2	I JUST STATED, CIRM WILL BE ACTIVELY INVOLVED IN
3	THIS AND CHOOSING AND ENABLING THE LICENSING AND IN
4	HELPING TO FUND THESE PROGRAM'S DEVELOPMENTS.
5	IN ADDITION, THE RESEARCHERS WOULD HAVE
6	CONTINUED FUNDING FOR THE ADVANCEMENTS OF THEIR
7	PROJECT. JUST TO BACK UP A BIT, WHEN THESE
8	IN-LICENSED PROGRAMS COME IN, THEY COME IN WITH
9	CURRENT FUNDING FOR THESE PROGRAMS TO GO TO A
10	CERTAIN VALUE INFLECTION POINT. IF THEY'RE CHOSEN
11	BY CIRM AND BY THE ATP3 AWARDEE TO COME INTO THEIR
12	PORTFOLIO, THEN THE PROJECT WOULD GET ADDITIONAL
13	FUNDING. FOR UNIVERSITIES, BY DESIGN OF THIS
14	INITIATIVE, THERE WOULD BE A DEMAND CREATION FOR
15	OUT-LICENSING CIRM-FUNDED TECHNOLOGIES AND,
16	THEREFORE, A GREATER OPPORTUNITY FOR FINANCIAL
17	RETURN WHICH THEN COULD GO ON TO FUND FUTURE
18	PROJECTS AND EFFORTS.
19	AND FOR CITIZENS OF CALIFORNIA, AS DR.
20	MILLS STATED, THIS IS AN OPPORTUNITY TO CREATE A
21	THERAPEUTIC POWERHOUSE THAT INCREASES THE LIKELIHOOD
22	OF GETTING STEM CELL THERAPEUTICS TO THE PATIENTS.
23	THE PRIVATE PARTNER OR THE AWARDEE, THE
24	APPLICANT, COULD BE AN ESTABLISHED COMPANY, A
25	SPIN-OFF, OR A NEW COMPANY ALTOGETHER THAT'S FORMED
	103

1	BY A TEAM OF PROFESSIONALS THAT HAVE COME OUT OF
2	EITHER PHARMA, BIOTECH, OR COULD BE INVESTORS. THEY
3	WILL BE JUDGED ON AND WILL PROPOSE AN EXCEPTIONAL
4	BUSINESS PLAN TO AGGREGATE THESE TECHNOLOGIES, GIVE
5	THE RATIONALE FOR THIS, PROPOSE HOW THIS WILL CREATE
6	VALUE AND BRING RETURN TO THE STAKEHOLDERS. AND
7	THEY WOULD COME IN WITH A LEADERSHIP TEAM THAT WOULD
8	BE JUDGED ON THEIR TRACK RECORD AND THEIR STRENGTH
9	THAT THEY BRING TO THE INITIATIVE AND THE LIKELIHOOD
10	THEY'LL BE ABLE TO EXECUTE ON THE BUSINESS PLAN AND
11	BRING ABOUT THE GOALS OF THIS INITIATIVE.
12	THE ENTITY WILL BE REQUIRED TO COME IN
13	WITH SIGNIFICANT INVESTMENT UPFRONT AND UTILIZE THIS
14	TO EXECUTE ON THE BUSINESS PLAN WHILE CIRM WILL FUND
15	THE SUPPORT OF THE DEVELOPMENT OF THE IN-LICENSED
16	PROJECTS.
17	THE CIRM AWARD IS ANTICIPATED TO BE
18	APPROXIMATELY \$75 MILLION OF FUNDING OVER A
19	FIVE-YEAR PERIOD. IT COULD BE IN THE FORM OF A
20	LOAN, BUT THE APPLICANT, THE AWARDEE, WOULD BE
21	REQUIRED TO MATCH THE TOTAL AWARD AMOUNT, REGARDLESS
22	OF HOW MUCH OF THE LOAN THEY TAKE ON, DOLLAR FOR
23	DOLLAR UPFRONT. THE AWARDEE WOULD ALSO BE REQUIRED
24	TO COMPLY WITH THE PRICING ACCESS AND MARCH-IN
25	PROVISIONS OF CIRM'S IP REGULATIONS AND TO PROVIDE
	104

1	THE LICENSOR OF THE CIRM PROJECTS WITH THE RIGHT OF
2	FIRST REFUSAL SHOULD THEY DECIDE TO SHELF OR CEASE
3	DEVELOPMENT OF THAT PARTICULAR TECHNOLOGY.
4	CHAIRMAN THOMAS: THANK YOU, DR. MILLAN.
5	ARE THERE QUESTIONS, COMMENTS FROM MEMBERS OF THE
6	BOARD?
7	MR. JUELSGAARD: SO THE PRESUMPTION
8	UNDERNEATH ALL OF THIS IS THAT THE ENTITY THAT GETS
9	FORMED WILL HAVE ACCESS TO CERTAIN PROJECTS AND THAT
10	THEY WILL BE AVAILABLE AND THEY WILL BE PROVIDED BY
11	THE ORGANIZATIONS THAT ARE CURRENTLY CONDUCTING
12	CLINICAL TRIALS. SO BEFORE ONE GOES TO THE EFFORT
13	OF PUTTING TOGETHER RAISING \$75 MILLION AND A
14	BUSINESS PLAN, ETC., ETC., BECAUSE THAT'S A
15	FAIR AMOUNT WORK, FAIR AMOUNT OF TIME INVOLVED, IT
16	WOULD SEEM TO ME THAT THAT ORGANIZATION WOULD WANT
17	TO KNOW WHAT'S ON THE TABLE FROM CIRM'S POINT OF
18	VIEW. WHAT PROJECTS ARE GOING TO BE THE ONES THAT
19	WOULD BE AVAILABLE TO THEM TO IN-LICENSE IN TERMS OF
20	THEIR BUSINESS PLAN.
21	SO WHAT IS THE GAME PLAN IS THERE A
22	GAME PLAN FOR DOING THAT? AND IF SO, WHAT IS THAT
23	GAME PLAN, AND HOW DOES IT GET EFFECTED?
24	DR. MILLAN: SO WE CURRENTLY HAVE A
25	PROGRAM NEAL LITTMAN IS HERE TODAY FOR

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1	INFORMING POTENTIAL PARTNERS, INDUSTRY, OF WHAT'S IN
2	OUR PORTFOLIO. AND WE WOULD JUST MAKE SURE THAT WE
3	DEPLOY THAT PROGRAM IN A MORE EXTENSIVE WAY SO THAT
4	THAT INFORMATION COULD BE PROVIDED TO THE POTENTIAL
5	APPLICANTS. AND THE APPLICANTS THEMSELVES CAN
6	CONTACT THE INSTITUTIONS OR THE PI'S DIRECTLY EVEN
7	DURING THE APPLICATION PROCESS TO DO THEIR OWN DUE
8	DILIGENCE OF WHETHER THEY'D COME IN FOR THIS.
9	CURRENTLY THE INFORMATION IS WE DO HAVE
10	PUBLICLY AVAILABLE INFORMATION ON THE PROJECTS, AND
11	WHAT WE'LL PLAN TO DO IS TAKE THAT AND CREATE, I
12	WOULD SAY, AN APPENDIX-TYPE THING SO IT'S ACTUALLY
13	PROVIDED TO THE APPLICANTS IN TERMS OF WHAT OUR
14	PORTFOLIO LOOKS LIKE. WE ALSO EXPECT TO PROVIDE AT
15	LEAST LINKS TO OUR INFORMATION ABOUT WHAT STANDARD
16	LICENSING TERMS CURRENTLY EXIST WITHIN OUR FUNDED
17	INSTITUTIONS SO THAT THEY KNOW THEIR BASE WHEN THEY
18	NEGOTIATE THESE LICENSES.
19	DR. JUELSGAARD: IF I CAN JUST ASK A
20	FOLLOW-UP QUESTION, AND I'LL USE MY GOOD FRIEND
21	MICHAEL FRIEDMAN AS AN EXAMPLE. SO THE CITY OF HOPE
22	HAS CERTAIN PROJECTS IN DEVELOPMENT. SO WHAT IS IT
23	THAT YOU WOULD EXPECT THE CITY OF HOPE TO AGREE TO
24	UPFRONT SO THAT SOME PERSON WHO'S GOING TO FILE AN
25	APPLICATION TO BE THIS ATP3 WILL KNOW WHICH, IF ANY,
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1	OF THE CITY OF HOPE'S PROJECTS ARE GOING TO BE IN
2	THE BAG, SO TO SPEAK, THAT THEY'LL HAVE ACCESS TO IF
3	THEY WERE SUCCESSFULLY SELECTED?
4	DR. MILLAN: AT THIS TIME WE DON'T EXPECT
5	TO HAVE THOSE TYPE OF COMMITMENTS FROM THE
6	INSTITUTIONS THEMSELVES. WHAT WE DO PLAN TO DO IS
7	CONTINUE TO HAVE CONVERSATIONS BECAUSE WE HAVE HAD
8	CONVERSATIONS WITH THE TECHNOLOGY AND LICENSING
9	OFFICES ABOUT WHAT THE GENERAL TERMS COULD LOOK LIKE
10	IN TERMS OF THESE LICENSE AGREEMENTS. THE
11	MANAGEMENT TEAM OF THIS ENTITY ATP3 WOULD NEGOTIATE
12	DIRECTLY WITH THOSE TECHNOLOGY AND LICENSING
13	OFFICES.
14	MR. PANETTA: THANK YOU. FIRST OF ALL,
15	ALL OF THESE INITIATIVES ARE EXCITING AND
	ALL OF THESE INITIATIVES ARE EXCITING AND AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE
15	
15 16	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE
15 16 17	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT
15 16 17 18	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT YOU'RE LOOKING TO IMPLEMENT HERE.
15 16 17 18 19	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT YOU'RE LOOKING TO IMPLEMENT HERE. FIRST OF ALL, TWO QUESTIONS. ON THIS
15 16 17 18 19 20	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT YOU'RE LOOKING TO IMPLEMENT HERE. FIRST OF ALL, TWO QUESTIONS. ON THIS AGGREGATION STRATEGY, I'M A LITTLE BIT UNCLEAR AS TO
15 16 17 18 19 20 21	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT YOU'RE LOOKING TO IMPLEMENT HERE. FIRST OF ALL, TWO QUESTIONS. ON THIS AGGREGATION STRATEGY, I'M A LITTLE BIT UNCLEAR AS TO WHETHER THIS MEANS THAT YOU WOULD AGGREGATE SIMILAR
15 16 17 18 19 20 21	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT YOU'RE LOOKING TO IMPLEMENT HERE. FIRST OF ALL, TWO QUESTIONS. ON THIS AGGREGATION STRATEGY, I'M A LITTLE BIT UNCLEAR AS TO WHETHER THIS MEANS THAT YOU WOULD AGGREGATE SIMILAR PROJECTS AND THEN OFFER THEM INDIVIDUALLY TO
15 16 17 18 19 20 21 22 23	AGGRESSIVE. AND TO REITERATE WHAT SHERRY SAID, THE SENSE OF URGENCY IS OBVIOUS IN ALL THE PROGRAMS THAT YOU'RE LOOKING TO IMPLEMENT HERE. FIRST OF ALL, TWO QUESTIONS. ON THIS AGGREGATION STRATEGY, I'M A LITTLE BIT UNCLEAR AS TO WHETHER THIS MEANS THAT YOU WOULD AGGREGATE SIMILAR PROJECTS AND THEN OFFER THEM INDIVIDUALLY TO INDUSTRY PARTNERS OR IF THE CONCEPT IS THAT THEY ARE

1	AND THEN THE SECOND QUESTION THAT I HAVE
2	GOES TO THE WHOLE TECHNOLOGY TRANSFER INTELLECTUAL
3	PROPERTY ISSUE. AND THAT IS THE EXTENT TO WHICH
4	THERE WOULD BE IN-HOUSE SERVICES WITHIN CIRM TO BE
5	ABLE TO ASSIST IN THE TRANSFER OF THESE
6	TECHNOLOGIES, OR IF THIS IS GOING TO BE THE
7	RESPONSIBILITY OF THE LICENSOR AND THE LICENSEE.
8	DR. MILLAN: THE LICENSING WILL BE THE
9	RESPONSIBILITY OF THE ENTITY THAT'S FUNDED BY CIRM.
10	CIRM WILL ASSIST IN THE DUE DILIGENCE-TYPE
11	ACTIVITIES BY WAY OF OUR GWG. SO IF THERE ARE
12	POTENTIAL MPT'S THAT ARE OF INTEREST TO THE ENTITY,
13	WE WOULD PUT IT THROUGH OUR REVIEW PROCESS TO GIVE A
14	SENSE OF IS THIS READY FOR THEM TO TAKE UP. AND SO
15	THOSE WHO WE'VE SPOKEN TO FELT THAT THIS KIND OF
16	PREREVIEW OR INSIDE LOOK IS OF GREAT VALUE. SO
17	THERE ARE ENTITIES OUT THERE WHO WANT TO MAKE A
18	MEANINGFUL INVESTMENT INTO THIS SPACE AND DO VALUE
19	WHAT CIRM COULD BRING IN TERMS OF ITS EXPERTISE AND
20	SYSTEMS FOR EVALUATING THESE PROJECTS.
21	WE PROPOSE THROUGH THIS FUNDING TO FUND
22	ONLY THE DEVELOPMENT COST FOR THE PROJECTS THAT ARE
23	IN-LICENSED. THE OPERATIONAL COSTS FOR THIS ENTITY
24	WILL BE PAID FOR BY THE ENTITY ITSELF AND, THUS, THE
25	UPFRONT CAPITAL INVESTMENT.
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	100

1	DR. MILLS: YOU HAD A FIRST PART TO THAT
2	QUESTION AROUND AGGREGATION.
3	MR. PANETTA: ACTUALLY I HAVE ANOTHER
4	QUESTION TO GO WITH THAT.
5	DR. MILLS: IN THE AGGREGATION, WE DON'T
6	DO THE AGGREGATING. AND IT'S ONE OF THE, I THINK,
7	REALLY IMPORTANT PARTS OF THE RFA. WHAT WE SAY IS
8	COME LOOK AT THE PORTFOLIO OF PRODUCTS THAT CIRM
9	HAS, AND YOU TELL US WHAT YOU WANT TO DRAW A LINE
10	AROUND TO COME UP WITH YOUR OWN BASKET THAT'S
11	SYNERGISTIC. THAT MIGHT BE WE WANT TO DO EVERYTHING
12	CARDIOVASCULAR, BUT IT MIGHT BE WE WANT TO DO
13	EVERYTHING IPS AND WE'RE DISEASE AGNOSTIC. IT'S FOR
14	THEM TO PROPOSE A PLAN THAT MAKES THE MOST SENSE TO
15	THE GWG ON HOW THEY ENVISION SYNERGIES IN SOMETHING
16	BEING SUCCESSFUL.
17	MR. PANETTA: CAN I ASK A FOLLOW-UP?
18	CHAIRMAN THOMAS: CERTAINLY. ALTHOUGH
19	BEFORE YOU DO, I'D JUST LIKE TO ADD THAT THE GWG
20	WILL BE EVALUATING THESE PROPOSALS. THIS WILL NOT
21	BE YOUR TYPICAL GWG. WE WON'T HAVE THE USUAL STEM
22	CELL RESEARCH SCIENTISTS EVALUATING. THIS WILL BE
23	MORE OF A BUSINESS ORIENTED GWG THAT WILL BE DRAWN
24	TOGETHER SPECIFICALLY BECAUSE THEY ARE EQUIPPED TO
25	EVALUATE THE BUSINESS ASPECTS OF THE PROPOSAL.
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1	MR. PANETTA: SO THAT'S A PERFECT LEAD-IN			
2	TO THIS BECAUSE ONE OF THE THINGS THAT I'VE HEARD			
3	FOR YEARS FROM THE VENTURE CAPITAL COMMUNITY IS THAT			
4	THEY'RE NOT READY TO INVEST YET IN STEM CELL			
5	TECHNOLOGIES. THIS MIGHT PROVIDE AN OPPORTUNITY AS			
6	WELL FOR VENTURE CAPITAL TO COME IN AND MAKE			
7	INVESTMENTS IN SOME OF THESE TECHNOLOGIES. COULD A			
8	VENTURE CAPITAL GROUP, FOR EXAMPLE, COME IN AND			
9	APPLY FOR THE OPPORTUNITY TO LICENSE SOME OF THESE			
10	TECHNOLOGIES AT LEAST OR LICENSE THEM TO THEIR			
11	COMPANIES?			
12	DR. MILLS: I THINK FROM OUR STANDPOINT,			
13	THE MOST SUCCESSFUL APPLICANT, THE ONE THAT PUTS			
14	TOGETHER THE ENTIRETY OF THE PACKAGE THAT SAYS			
15	THEY'RE GOING TO BE ABLE TO TAKE THESE THEY HAVE			
16	AN AGGREGATION STRATEGY THAT MAKES SENSE, THEY HAVE			
17	A BUSINESS PLAN THAT MAKES SENSE, THEY HAVE A			
18	MANAGEMENT TEAM THAT'S TOPNOTCH, AND THEY HAVE THE			
19	CAPITAL TO PUT INTO IT, AND THEY CAN PULL IT OFF.			
20	WHOEVER THAT IS IS FINE.			
21	CHAIRMAN THOMAS: I WOULD ALSO ADD TO			
22	THAT, MR. PANETTA, THAT IN THE COURSE OF MANY, MANY			
23	DISCUSSIONS ON THE ATP3 TOPIC, WE'VE VISITED WITH A			
24	NUMBER OF POTENTIAL APPLICANT TYPES TO SORT OF GAUGE			
25	THEIR LEVEL OF INTEREST TO SEE IF WE'RE ONTO			
	110			

1	SOMETHING HERE OR NOT AND GOTTEN VERY POSITIVE
2	RESPONSES, INCLUDING FROM MEMBERS OF THE VENTURE
3	CAPITAL COMMUNITY.
4	DR. FRIEDMAN: I SUSPECT THAT IT WILL BE
5	WELL RECEIVED. COUPLE OF QUESTIONS PLEASE. ONE IS
6	IF THIS IS AS SUCCESSFUL AS YOU WOULD LIKE IT TO BE,
7	THERE ARE GOING TO BE MULTIPLE COMPETITORS FOR THE
8	SAME PROJECT. THAT'S A DIFFERENT SITUATION THAN
9	WHAT WE USUALLY ENCOUNTER. AND I WONDER HOW YOU
10	THOUGHT ABOUT HOW YOU WILL COMPARE AND CONTRAST
11	DIFFERENT OFFERS THAT ARE MADE. THAT'S PART A. I'M
12	LEARNING ABOUT HOW TO ASK TWO-PART QUESTIONS.
13	PART B IS, WHEREAS MOST OF THE TIME WHEN
14	SOMEONE SUBMITS SOMETHING, WE DECIDE YES OR NO AND
15	THEN MOVE ON, AND THEY GET A CHANCE TO REVISE IT.
16	HERE'S A SITUATION WHERE YOU MIGHT WANT TO DO IT
17	RATHER DIFFERENTLY, WHICH IS INSTEAD OF SAYING YES
18	OR NO, YOU WANT TO HAVE A COMPETITIVE, ONGOING
19	DYNAMIC PROCESS, AND YOU WILL THEN SOLICIT THE BEST
20	APPLICATION FROM VC OR WHOEVER BECAUSE THEY GET A
21	CHANCE TO REITERATE AND TO COMPARE AND, IF YOU WILL,
22	HAVE A BIDDING PROCESS, BUT IT'S GOT TO BE FAIR AND
23	IT'S GOT TO BE TRANSPARENT. AND THAT'S A DIFFERENT
24	WAY OF ACTING THAN WE'VE DONE IN THE PAST. PLEASE
25	SHARE WITH US HOW YOU THOUGHT ABOUT THAT.
	111

1	DR. MILLAN: SO FOR THIS ROUND, FOR THE
2	CONCEPT WE'RE BRINGING FORWARD TO YOU TODAY, IT WILL
3	BE DIFFERENT IN TERMS OF THE COMPOSITION PERHAPS OF
4	THE REVIEWERS, BUT WHAT WE ARE CURRENTLY PLANNING IS
5	THAT IT WILL BE A COMPETITIVE PROCESS WHERE THEY
6	BRING THEIR BEST BUSINESS PLAN FORWARD TO LOOK AT.
7	WE UNDERSTAND THAT WHEN THEY ACTUALLY START, WHEN
8	THEY ACTUALLY START OPERATIONS AND THEY START TRYING
9	TO LICENSE, THERE ARE GOING TO BE THINGS THAT
10	HAPPEN, THAT IT MAY NOT BE EXACTLY AS THEY PLANNED.
11	BUT THE IDEA IS THAT THE REVIEWERS WHO WILL HAVE
12	EXPERTISE IN TERMS OF THIS WORLD WILL BE ABLE LOOK
13	AT THIS BUSINESS PLAN, LOOK AT WHAT THEY PROPOSE TO
14	DO, LOOK AT THEIR MITIGATION STRATEGY, AND DETERMINE
15	IS THIS A RATIONAL APPROACH GIVEN THAT NOT
16	EVERYTHING IS KNOWN AT THAT POINT.
17	DR. FRIEDMAN: HERE'S THE THING. WHAT YOU
18	ASKED FOR IS A BEST AND FINAL OFFER THAT THEN WILL
19	BE REVIEWED AND UNDERSTOOD THAT IT MIGHT CHANGE AND
20	SO ON AND SO FORTH. I JUST ASK YOU TO THINK ABOUT
21	WHETHER YOU'D WANT TO, SINCE WE'RE DOING SO MANY
22	EXPERIMENTS, WHETHER WE WANT TO EXPERIMENT AND SEE
23	ABOUT HAVING A COMPETITIVE BIDDING PROCESS WHICH
24	WOULD BE TO THE BENEFIT OF THE GRANTEE BECAUSE HE OR
25	SHE WOULD BE GETTING MORE RESOURCES OR BETTER
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1	EXPERTISE OR WHATEVER AND TO THE BENEFIT OF THE			
2	CITIZENS OF CALIFORNIA. IT'S JUST A LITTLE BIT			
3	DIFFERENT WAY OF DOING IT, AND MUCH MORE			
4	COMPLICATED, MAYBE NOT WORTH THE TROUBLE, BUT I ASK			
5	YOU JUST TO CONSIDER IT.			
6	DR. MILLAN: OKAY. WE WILL. THANK YOU.			
7	CHAIRMAN THOMAS: OTHER COMMENTS,			
8	QUESTIONS FROM MEMBERS OF THE BOARD?			
9	MR. SHEEHY: I'D LIKE TO MOVE APPROVAL OF			
10	THE CONCEPT.			
11	MS. LANSING: I'LL SECOND IT.			
12	CHAIRMAN THOMAS: MOVED BY MR. SHEEHY.			
13	SECONDED BY MS. LANSING. ARE THERE ANY COMMENTS			
14	FROM MEMBERS ON THE PHONE?			
15	MS. LAPORTE: COULD YOU JUST CLARIFY. IS			
16	THIS A ONE-TIME COMPETITIVE PROCESS THEN, OR IS			
17	THERE AN OPPORTUNITY TO DO MULTIPLE DEALS HERE? NOT			
18	EVERY DEAL IS GOING TO MAKE \$75 MILLION OF SENSE TO			
19	INVESTORS. I'M JUST WONDERING HOW YOU KIND OF			
20	OPTIMIZE.			
21	DR. MILLS: SO THE ORIGINAL VERSION IS			
22	INTENDED TO BE ONE TIME. AND DEPENDING ON THE			
23	TECHNOLOGIES THAT ARE PUT INTO THE BUNDLE AND HOW			
24	MANY AND OF WHAT SCOPE AND WHAT'S REMAINING, WE HAVE			
25	CONTEMPLATED DOING IT AGAIN. THAT WOULD BE			
	113			
	-			

1	SOMETHING WE WOULD COME BACK TO THE BOARD FOR
2	OBVIOUSLY AFTER WE LEARNED LESSONS LEARNED FROM THE
3	FIRST GO OF IT IF WE THOUGHT IT WOULD BE A GOOD
4	THING TO TRY AGAIN. OUT OF THE GATE, IT'S INTENDED
5	TO BE FOR THE FIRST TIME JUST ONE AWARD.
6	CHAIRMAN THOMAS: OTHER QUESTIONS FROM
7	MEMBERS ON THE PHONE? HEARING NONE, COMMENTS FROM
8	MEMBERS OF THE PUBLIC?
9	MS. MC CLAREN: SO I HAVE A STATEMENT
10	PREPARED THAT I'D LIKE TO READ TO YOU GUYS.
11	CHAIRMAN THOMAS: PLEASE STATE YOUR NAME.
12	MS. MC CLAREN: MY NAME IS MALLORY
13	MCCLAREN. SO I'M HERE BECAUSE I MUST EXPRESS MY
14	CONCERN THAT THE CURRENT ATP3 PROPOSAL FOR A CIRM
15	PUBLIC PRIVATE INVESTMENT PARTNERSHIP IS LIKELY TO
16	STIFLE NEW BANKING AND FINANCE INNOVATIONS WHICH
17	CARRY POTENTIAL TO EXPONENTIALLY ACCELERATE THE
18	GROWTH OF THE REGENERATIVE MEDICINE FIELD.
19	THE PROPOSAL UNDER DISCUSSION TODAY WILL
20	FUND UP TO \$75 MILLION, ONE, A CALIFORNIA COMPANY;
21	TWO, WHICH HAS A DEMONSTRATED TRACK RECORD OF
22	INVESTING IN BIOTECHNOLOGY; AND, THREE, THAT ALREADY
23	HAS THE CAPITAL OR CAN ACCESS THE CAPITAL TO MATCH
24	THE CIRM GRANT. THIS GRANT WILL NOT FUND EITHER
25	FORMATION, OPERATION, OR MANAGEMENT COSTS OF THE
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1	APPLICANT ENTITY AS HAS BEEN DISCUSSED.
2	THESE CRITERIA EFFECTIVELY EXCLUDE NEARLY
3	ALL OTHERS BUT ESTABLISHED VENTURE CAPITAL FIRMS IN
4	CALIFORNIA. THE HIGH RISK, HIGH RETURN STRUCTURE OF
5	VC, WHICH GENERALLY REQUIRES THE LIKELIHOOD OF A 30
6	TO 100 X RETURN, AND WHICH FAILS TO PROCURE THE
7	CAPITAL TO MAKE A SUFFICIENT NUMBER OF INVESTMENTS
8	TO DERISK A BIOTECHNOLOGY PORTFOLIO, HAS A DUBIOUS
9	TRACK RECORD WITH BRINGING ENOUGH BIOTECH IDEAS TO
10	FRUITION TO MAKE A DIFFERENCE.
11	REGENERATIVE MEDICINE'S GOALS ARE SIMPLY
12	TOO IMPORTANT FOR OUR STATE TO DEPEND UPON ENTITIES
13	THAT ARE TOO FIRMLY ENTRENCHED IN STRATEGIES THAT
14	ARE NOT APPROPRIATE FOR BIOTECHNOLOGY TO ACCOMPLISH
15	THE RELEVANT GOALS. BEFORE CIRM INVESTS AROUND 8
16	PERCENT OF ITS REMAINING FUNDS INTO JUST ONE
17	COMPANY, BE IT VC OR OTHERWISE, THE COMMITTEE SHOULD
18	HOLD OFF ON APPROVING THE ATP3 PROPOSAL TODAY AND
19	COMMIT TO UNDERTAKING FURTHER INQUIRY ON HOW CIRM
20	CAN SPONSOR INVESTMENT IN A MANNER THAT WILL BOTH
21	CULTIVATE THE DEVELOPMENT OF FRESH BANKING AND
22	FINANCE INNOVATIONS FOR REGENERATIVE MEDICINE AND
23	SUPPORT IDEAS WHICH DO NOT NECESSARILY EMERGE FROM
24	ESTABLISHED INVESTMENT OUTFITS.
25	ETERNA BIOCAPITAL, OF WHICH I AM THE

1	FOUNDER, IS ONE SUCH COMPANY. WE ARE BASED IN
2	CALIFORNIA AND ARE PROPOSING TO BUILD A SECURITIZED
3	LOWER RISK AND LOWER RETURN MEGA FUND DESIGNED TO
4	ATTRACT INSTITUTIONAL CAPITAL AND TO SPUR
5	REGENERATIVE MEDICINE RESEARCH GLOBALLY. OUR
6	COMPANY IS BUILT UPON IDEAS ADAPTED FROM AN MIT
7	WHITEPAPER ON HOW TO BUILD SUCH AN INVESTMENT
8	VEHICLE. ETERNA BIOCAPITAL'S PLAN CAN RESPONSIBLY
9	AND SUSTAINABLY SUPPORT TRANSLATIONAL R&D OVER THE
10	LONG TERM. OURS IS AN IDEA AMONG OTHER IDEAS, SUCH
11	AS CROWD FUNDING, THAT ARE AT LEAST AS WORTHY AS
12	VENTURE CAPITAL OF CIRM'S CONSIDERATION AND SUPPORT.
13	I RESPECTFULLY IMPLORE THE COMMITTEE TO
14	FURTHER CONSIDER HOW THESE PROPOSALS PUT FORTH TODAY
15	HAVE THE POWER TO EITHER MAKE OR BREAK FLEDGLING
16	CALIFORNIA COMPANIES LIKE ETERNA BIOCAPITAL WHICH
17	AIM TO IMPLEMENT NEW AND USEFUL INVESTMENT PLATFORMS
18	FOR REGENERATIVE MEDICINE. I APPRECIATE YOUR
19	CONSIDERATION.
20	CHAIRMAN THOMAS: THANK YOU. OTHER
21	COMMENTS BY MEMBERS OF THE PUBLIC?
22	MR. BONDY: GOOD MORNING. MY NAME IS KEN
23	BONDY. I'VE BEEN A RESIDENT OF CALIFORNIA FOR MORE
24	THAN 70 YEARS NOW. I GREW UP IN NORTH HOLLYWOOD.
25	I'M A RETIRED STRUCTURAL ENGINEER. I HAD A LONG 50

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1	PLUS YEAR CAREER SPECIALIZING IN THE DESIGN OF			
2	CONCRETE BUILDING STRUCTURES, ONE OF WHICH,			
3	COINCIDENTALLY, WAS THE VERY FIRST PARKING STRUCTURE			
4	THAT WAS EVER BUILT NEXT DOOR AT LAX. IT'S AT 501			
5	WORLD WAY. IT WAS COMPLETED IN 1965, AND IT'S IN			
6	DAILY USE TODAY.			
7	I ALSO HAVE A LONG HISTORY WITH UCLA. I			
8	WENT TO SCHOOL THERE IN THE LATE '50S. AND FOR A			
9	LITTLE MORE THAN 20 YEARS NOW, I'VE TAUGHT SENIOR			
10	UNDERGRADUATE STRUCTURAL ENGINEERING DESIGN CLASSES.			
11	I'M OBVIOUSLY NOT A BIOTECHNOLOGY PROFESSIONAL, BUT			
12	I AM INTENSELY INTERESTED IN THE ACTIVITIES OF YOUR			
13	AGENCY.			
14	CANCER HAS TOUCHED MY LIFE WAY TOO OFTEN.			
15	I'VE LOST CLOSE RELATIVES AND DEAR FRIENDS, EACH FAR			
16	TOO SOON. EVEN NOW MY WIFE IS BATTLING THYROID			
17	CANCER. IT HAS BECOME CLEAR TO ME THAT STEM CELL			
18	RESEARCH OFFERS OUR BEST CHANCE TO FINALLY DISCOVER			
19	CURES FOR THESE HORRIBLE DISEASES.			
20	I'M HERE TO SPEAK TO THE ATP3 PROPOSAL. I			
21	STRONGLY AGREE WITH WHAT MS. MCCLAREN JUST SAID. AS			
22	A CALIFORNIA TAXPAYER, I STRONGLY SUPPORT CIRM'S			
23	DESIRE TO PROMOTE A WORKABLE AND SUSTAINABLE			
24	ENVIRONMENT FOR REGENERATIVE BIOMEDICAL TECHNOLOGY,			
25	BUT WHAT I CAN'T SUPPORT IS WHAT APPEARS TO BE A			

1	PLAN TO GIVE \$75 MILLION IN PUBLIC FUNDS TO JUST ONE
2	VENTURE CAPITAL COMPANY. AND EVEN IF IT REQUIRES A
3	MATCHING GRANT, THEN JUST HOPE THAT EVERYTHING TURNS
4	OUT OKAY. IN MY OPINION, THAT WOULD RETARD OR
5	PREVENT THE DEVELOPMENT OF OTHER NEW AND INNOVATIVE
6	IDEAS FOR REGENERATIVE MEDICINE.
7	IF ATP3 IS APPROVED IN ITS PRESENT FORM,
8	CIRM, I BELIEVE, WILL BE MARRYING ITSELF TO ONE
9	INVESTMENT SCHEME AND REJECTING EVERY OTHER VALID
10	INVESTMENT IDEA. TO SUMMARIZE, I THINK VENTURE
11	CAPITAL IS SIMPLY NOT RIGHT FOR BIOTECHNOLOGY. I
12	URGE CIRM TO REJECT THE ATP3 PROPOSAL TODAY AND
13	REVISE IT TO MAKE IT FAIR TO ALL CALIFORNIA BANKING
14	AND FINANCE INNOVATORS. THANK YOU VERY MUCH.
15	CHAIRMAN THOMAS: OTHER COMMENTS BY
16	MEMBERS OF THE PUBLIC? BEFORE WE VOTE ON THIS, I DO
17	WANT TO REMIND THE COMMITTEE FROM DR. MILLAN'S
18	PRESENTATION, WE HAVE SPOKEN TO A NUMBER OF
19	DIFFERENT STAKEHOLDERS REPRESENTING DIFFERENT
20	POTENTIAL TYPES OF ENTITIES THAT CAN APPLY FOR THIS.
21	WE ARE BY NO MEANS CONFINED TO VENTURE CAPITAL. SO
22	WITH THAT, MARIA, WILL YOU PLEASE CALL THE ROLL.
23	DR. JUELSGAARD: I WOULD JUST LIKE TO
24	EXPOUND ON THAT A LITTLE BIT. THAT GOES BACK TO THE
25	QUESTIONS I WAS ASKING. YOU KNOW, IT'S NOT EASY
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1	WHEN YOU REALLY DON'T KNOW WHAT IT IS THAT YOU ARE			
2	GOING TO GET WITH ANY CERTAINTY TO PUT TOGETHER A			
3	FUNDING PLAN OR \$75 MILLION, THAT'S A FAIR AMOUNT OF			
4	MONEY, PUT TOGETHER A BUSINESS TEAM, AND GET ALL THE			
5	STUFF SET UP IN HOPES THAT YOU'LL GET AWARDED AN			
6	APPLICATION. SO THAT'S A TREMENDOUS HILL TO CLIMB			
7	OVER IF YOU'RE IN THE VENTURE CAPITAL INDUSTRY.			
8	HOWEVER, IF YOU'RE AN ESTABLISHED INDUSTRY, IT'S A			
9	LOT EASIER. YOU ALREADY HAVE ALL OF THOSE RESOURCES			
10	AVAILABLE. AND IF IT'S INTERESTING ENOUGH, YOU CAN			
11	ESTABLISH A SUBSIDIARY BASICALLY TO SERVE AS THE			
12	APPLICANT.			
13	SO I THINK THERE'S AS MUCH LIKELIHOOD, IF			
14	THIS GOES FORWARD, THAT THE APPLICANTS MAY ACTUALLY			
15	COME FROM ESTABLISHED INDUSTRY AS THEY NECESSARILY			
16	WOULD FROM THE VENTURE CAPITAL COMMUNITY.			
17	CHAIRMAN THOMAS: OKAY.			
18	MS. BONNEVILLE: DAVID BRENNER. LINDA			
19	BOXER.			
20	DR. BOXER: YES.			
21	MS. BONNEVILLE: KEN BURTIS. ANNE-MARIE			
22	DULIEGE.			
23	DR. DULIEGE: YES.			
24	MS. BONNEVILLE: MICHAEL FRIEDMAN.			
25	DR. FRIEDMAN: YES.			
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1		MS. BONNEVILLE: JUDY GASSON.
2		DR. GASSON: YES.
3		MS. BONNEVILLE: SAM HAWGOOD. DAVID
4	HIGGINS.	
5		DR. HIGGINS: YES.
6		MS. BONNEVILLE: STEPHEN JUELSGAARD.
7		MR. JUELSGAARD: YES.
8		MS. BONNEVILLE: SHERRY LANSING.
9		MS. LANSING: YES.
10		MS. BONNEVILLE: KATHY LAPORTE.
11		DR. LAPORTE: YES.
12		MS. BONNEVILLE: BERT LUBIN.
13		DR. LUBIN: YES.
14		MS. BONNEVILLE: SHLOMO MELMED.
15		DR. MELMED: YES.
16		MS. BONNEVILLE: LAUREN MILLER.
17		MS. MILLER: YES.
18		MS. BONNEVILLE: ADRIANA PADILLA.
19		DR. PADILLA: YES.
20		MS. BONNEVILLE: JOE PANETTA.
21		MR. PANETTA: YES.
22		MS. BONNEVILLE: ROBERT PRICE.
23		DR. PRICE: YES.
24		MS. BONNEVILLE: FRANCISCO PRIETO.
25		DR. PRIETO: AYE.
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		120

1	MS. BONNEVILLE: CARMEN PULIAFITO. ROBERT
2	QUINT. AL ROWLETT.
3	MR. ROWLETT: YES.
4	MS. BONNEVILLE: JEFF SHEEHY.
5	MR. SHEEHY: YES.
6	MS. BONNEVILLE: OSWALD STEWARD.
7	DR. STEWARD: YES.
8	MS. BONNEVILLE: JONATHAN THOMAS.
9	CHAIRMAN THOMAS: YES.
10	MS. BONNEVILLE: ART TORRES.
11	MR. TORRES: AYE.
12	MS. BONNEVILLE: KRISTINA VUORI.
13	DR. VUORI: YES.
14	MS. BONNEVILLE: DIANE WINOKUR.
15	CHAIRMAN THOMAS: MOTION PASSES. THAT
16	CONCLUDES THE SEGMENT.
17	MS. BONNEVILLE: I'D LIKE TO GO BACK TO
18	THE TRANSLATING CENTER. WE HAVEN'T FINISHED THAT.
19	AND MEMBERS ON THE PHONE, IF I COULD JUST CAPTURE
20	THEIR VOTE.
21	MS. BONNEVILLE: LINDA BOXER.
22	DR. BOXER: YES.
23	MS. BONNEVILLE: DIANE WINOKUR. BERT
24	LUBIN.
25	DR. LUBIN: YES.
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1	MS. BONNEVILLE: KATHY LAPORTE.
2	MS. LAPORTE: YES.
3	CHAIRMAN THOMAS: THANK YOU, MARIA.
4	MR. HARRISON, I PRESUME BOTH MOTIONS
5	PASSED. SO THAT CONCLUDES THE THREE MAJOR
6	COMPONENTS TO BE VOTED ON FROM THE STRATEGIC PLAN.
7	I WANT TO THANK DR. MILLS, DR. MILLAN, MR. LITTMAN,
8	EVERYBODY WHO HAD A MAJOR HAND IN THIS. THIS HAS
9	BEEN AN UNDERTAKING WITH A GREAT DEAL OF THOUGHT,
10	AND I'M PERSONALLY VERY EXCITED TO MOVE FORWARD WITH
11	THESE THREE COMPONENTS AS WELL AS THE REST OF THE
12	STRATEGIC PLAN.
13	SO HAVING SAID THAT, LET'S NOW GO OUT AND
14	BREAK TO GET OUR LUNCH. PLEASE BRING IT IT'S
15	RIGHT OUTSIDE THE DOOR THERE. PLEASE BRING IT IN,
16	AND WE WILL CONTINUE AS A WORKING LUNCH TO GO
17	THROUGH THE BALANCE OF OUR AGENDA. THANK YOU.
18	(A RECESS WAS TAKEN.)
19	CHAIRMAN THOMAS: WE'RE GOING TO RESUME.
20	THOSE OF YOU ON THE PHONE, WE'VE HAD A FINE
21	FIVE-STAR CUISINE BREAK HERE. WE HOPE YOU'VE HAD
22	THE SAME.
23	WE'RE GOING TO MOVE ON NOW TO ITEM NO. 12
24	ON THE ACTION ITEM AGENDA, WHICH IS AMENDMENTS TO
25	THE GOVERNING BOARD BYLAWS AND INTERNAL GOVERNANCE

1	POLICY. MR. HARRISON.
2	MR. HARRISON: GOOD AFTERNOON. THIS IS
3	WHEN THE MEETING REALLY STARTS TO GET EXCITING, SO
4	HOLD ON TO YOUR SEATS. IT'S GOING TO BE A WILD
5	RIDE.
6	SO AS YOU KNOW, WE SPENT A LOT OF TIME
7	THINKING THROUGH AND REVISING THE PROCESSES BY WHICH
8	CIRM AWARDS AND MANAGES GRANTS AND LOANS. AND THE
9	RESULT OF THAT IS CIRM 2.0, BUT PRESIDENT MILLS WAS
10	NOT SATISFIED WITH THAT. HE CHALLENGED US TO APPLY
11	THE SAME LEVEL OF SCRUTINY TO OUR INTERNAL POLICIES
12	AND PRACTICES, OUR ADMINISTRATIVE POLICIES AND
13	PROCEDURES. SO AS PART OF THAT PROCESS, WE HAVE
14	BEGUN A PROCESS THAT WE CALL CORE 2.0 IN WHICH WE
15	ARE EVALUATING ALL OF THE INTERNAL OPERATING
16	PRACTICES AND PROCEDURES, SOME OF WHICH HAVE BEEN IN
17	EFFECT FOR A LONG TIME AND HAVE NOT BEEN REVISITED.
18	TODAY WE ARE BRINGING SIX POLICY CHANGES
19	FOR YOUR CONSIDERATION. THEY INCLUDE THE BOARD
20	BYLAWS, THE INTERNAL GOVERNANCE POLICY, THE EMPLOYEE
21	CONFLICT OF INTEREST POLICY, THE COMPENSATION
22	POLICY, THE RELOCATION POLICY, AND LAST, BUT NOT
23	LEAST, THE GRANTS WORKING GROUP BYLAWS. WE'LL TRY
24	TO MAKE THESE PRESENTATIONS BRIEF AND SUCCINCT
25	BECAUSE THE WHOLE POINT OF OUR REVIEW OF CORE 2.0
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IS TO LOOK FOR OPPORTUNITIES TO MAKE OUR POLICIES
AND PROCEDURES MORE EFFECTIVE AND MORE EFFICIENT.
SO FIRST I'D LIKE TO START WITH THE BOARD
BYLAWS AND WITH AN ITEM THAT I WOULD LIKE THE BOARD
TO TAKE UP SEPARATE FROM THE REMAINDER OF THE
AMENDMENTS TO THE BYLAWS AND TO THE INTERNAL
GOVERNANCE POLICY. THIS HAS TO DO WITH THE PER DIEM
PAID TO PATIENT ADVOCATE MEMBERS OF THE BOARD WHO
SERVE ON THE GRANTS WORKING GROUP.
WHEN THE LEGISLATURE ENACTED SB 1064, IT
GAVE THE BOARD THE AUTHORITY TO SET A PER DIEM RATE
FOR MEMBERS OF THE BOARD WHO SERVE AS PATIENT
ADVOCATES ON THE GRANTS WORKING GROUP. AND THAT WAS
IN 2009. AT THE TIME THE BOARD APPROVED A PER DIEM
RATE OF 75 PERCENT OF WHAT IS PAID TO THE SCIENTIFIC
MEMBERS OF THE GRANTS WORKING GROUP, BUT CAPPED IT
AT \$15,000 PER YEAR.
NOW, FOR THOSE OF YOU WHO WERE MEMBERS OF
THE BOARD BACK IN 2009 AND 2010, YOU WILL RECALL
THAT ON AVERAGE THE GWG MET PERHAPS THREE OR FOUR
TIMES A YEAR, AND THE PATIENT ADVOCATES WHO SERVED
ON THE GRANTS WORKING GROUP WERE, I GUESS WHAT I
WOULD CALL FROM TODAY'S PERSPECTIVE, PLAYING A MORE
PASSIVE ROLE.
FAST FORWARD TO 2015 WITH THE
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1	IMPLEMENTATION OF CIRM 2.0, AND THE GRANTS WORKING
2	GROUP HAS MET MORE THAN A DOZEN TIMES ALREADY THIS
3	YEAR. FURTHERMORE, THE PATIENT ADVOCATE MEMBERS ARE
4	NOW ACTIVELY ENGAGED IN THE GRANTS WORKING GROUP
5	REVIEW. EACH APPLICATION IS ASSIGNED TO A PATIENT
6	ADVOCATE REVIEWER WHO PROVIDES THE GWG WITH HIS OR
7	HER INPUT DURING THE GWG'S EVALUATION OF THAT
8	APPLICATION.
9	TO SAY THAT THE OBLIGATIONS AND
10	RESPONSIBILITIES IMPOSED ON THE PATIENT ADVOCATE
11	MEMBERS OF THE GWG HAS INCREASED WOULD BE AN
12	UNDERSTATEMENT. TO USE A WORD DR. MILLS LIKES, IT'S
13	INCREASED EXPONENTIALLY, AND YET THE PER DIEM CAP
14	REMAINS AT \$15,000.
15	SO THE FIRST THING WE'D LIKE TO PROPOSE TO
16	THE BOARD, IN RECOGNITION OF THE INCREASED
17	RESPONSIBILITY AND TIME COMMITMENTS IMPOSED ON THE
18	PATIENT ADVOCATE MEMBERS OF THE BOARD, IS TO
19	INCREASE THE CAP TO \$30,000 PER YEAR AND TO APPLY
20	THAT RETROACTIVELY TO JANUARY 1, 2015.
21	I'D BE HAPPY TO ANSWER ANY QUESTIONS ABOUT
22	THAT ITEM.
23	CHAIRMAN THOMAS: ANY QUESTIONS FROM
24	MEMBERS OF THE BOARD?
25	DR. PRICE: HOW MANY?
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1	MR. HARRISON: THERE ARE SEVEN PATIENT
2	ADVOCATE MEMBERS OF THE GWG. BY THE WAY, I SHOULD
3	POINT OUT THAT THE REASON WE ARE CONSIDERING THIS
4	ITEM SEPARATELY FROM THE REMAINDER OF THE PROPOSED
5	AMENDMENTS IS BECAUSE THE PATIENT ADVOCATES WHO ARE
6	ELIGIBLE TO SERVE, WHO INCLUDE ALL THE PATIENT
7	ADVOCATES ON THE BOARD WITH THE EXCEPTION OF THE
8	CHAIR AND THE VICE CHAIR WHO DO NOT RECEIVE A PER
9	DIEM, WILL RECUSE THEMSELVES FROM VOTING ON THIS
10	PARTICULAR MATTER.
11	CHAIRMAN THOMAS: QUESTIONS FROM MEMBERS
12	OF THE BOARD? THIS ONE WE NEED A MOTION AND A ROLL
13	CALL VOTE. SO CAN I GET A MOTION FIRST PLEASE?
14	DR. VUORI: SO MOVED.
15	CHAIRMAN THOMAS: SO MOVED BY DR. VUORI,
16	SECONDED BY
17	DR. DULIEGE: I CAN SECOND IT, BUT I ALSO
18	HAVE A QUESTION.
19	CHAIRMAN THOMAS: SECONDED BY DR. DULIEGE
20	WITH A QUESTION.
21	DR. DULIEGE: MAYBE I MISSED THAT, BUT I'M
22	CERTAINLY TOTALLY IN AGREEMENT WITH THIS CAP TO THE
23	PER DIEM. WHAT I'M UNCLEAR, WHAT'S THE PER DIEM
24	ITSELF?
25	MR. HARRISON: THE PER DIEM IS SET AT 75
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1	PERCENT OF THE RATE THAT IS PAID TO THE SCIENTIFIC
2	MEMBERS OF THE GRANTS WORKING GROUP, WHICH VARIES
3	DEPENDING UPON THE EXPECTATIONS OF THE NUMBER OF
4	APPLICATIONS FOR ANY PARTICULAR REVIEW.
5	CHAIRMAN THOMAS: I'D JUST LIKE TO NOTE,
6	JUST TO GIVE YOU MY PERSPECTIVE, TO ECHO WHAT JAMES
7	IS SAYING, THE ROLE OF THE PATIENT ADVOCATE HAS
8	TRULY INCREASED DRAMATICALLY THIS YEAR. AND I THINK
9	THIS MOTION IS REFLECTIVE OF THAT AND ABSOLUTELY
10	WARRANTED GIVEN THE GREAT TIME AND ENERGY PUT INTO
11	IT. DR. MILLS.
12	DR. MILLS: I WOULD LIKE TO STRONGLY ECHO
13	CHAIRMAN THOMAS' COMMENTS. WE CHANGED VERY
14	SIGNIFICANTLY THE ROLE OF THE PATIENT ADVOCATES AT
15	THE GRANTS WORKING GROUP, AND THE RESULTS OF THAT
16	AREN'T DECORATIVE. THEY'RE VERY, VERY SUBSTANTIAL.
17	IT'S WORKING THE WAY THAT WE HAD HOPED AND PLANNED
18	IT WOULD WORK. THEY'RE BRINGING A VERY UNIQUE
19	PERSPECTIVE TO THE GWG AND A VERY IMPORTANT
20	PERSPECTIVE THAT NEEDS TO BE HEARD AND CONSIDERED.
21	AND THEIR WORKLOAD HAS, AS JAMES SAID, IT HAS GONE
22	UP EXPONENTIALLY BECAUSE WHEN WE SAID WE'RE GOING TO
23	HOLD REVIEWS EVERY 30 DAYS SO WE CAN HAVE A
24	TURNAROUND TIME THE WAY WE HAVE THE TURNAROUND TIME,
25	THAT PLACED JUST TREMENDOUS DEMAND ON THE PATIENT
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1	ADVOCATES. THAT'S ALL.
2	CHAIRMAN THOMAS: ANY OTHER COMMENTS BY
3	MEMBERS OF THE BOARD? COMMENTS BY MEMBERS ON THE
4	PHONE? COMMENTS FROM MEMBERS OF THE PUBLIC?
5	HEARING NONE, MARIA, PLEASE CALL THE ROLL.
6	MS. BONNEVILLE: DAVID BRENNER. LINDA
7	BOXER.
8	DR. BOXER: YES.
9	MS. BONNEVILLE: KEN BURTIS. ANNE-MARIE
10	DULIEGE.
11	DR. DULIEGE: YES.
12	MS. BONNEVILLE: MICHAEL FRIEDMAN.
13	DR. FRIEDMAN: YES.
14	MS. BONNEVILLE: JUDY GASSON.
15	DR. GASSON: YES.
16	MS. BONNEVILLE: SAM HAWGOOD. STEPHEN
17	JUELSGAARD.
18	MR. JUELSGAARD: YES.
19	MS. BONNEVILLE: KATHY LAPORTE.
20	DR. LAPORTE: YES.
21	MS. BONNEVILLE: BERT LUBIN.
22	DR. LUBIN: YES.
23	MS. BONNEVILLE: SHLOMO MELMED.
24	DR. MELMED: YES.
25	MS. BONNEVILLE: JOE PANETTA.
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	BARRISTERS REPORTING SERVICE
1	MR. PANETTA: YES.
2	MS. BONNEVILLE: ROBERT PRICE.
3	DR. PRICE: YES.
4	MS. BONNEVILLE: CARMEN PULIAFITO.
5	JONATHAN THOMAS.
6	CHAIRMAN THOMAS: YES.
7	MS. BONNEVILLE: ART TORRES.
8	MR. TORRES: AYE.
9	MS. BONNEVILLE: KRISTINA VUORI.
10	DR. VUORI: YES.
11	CHAIRMAN THOMAS: THANK YOU. THE MOTION
12	PASSES. ON TO THE NEXT ITEM, MR. HARRISON.
13	MR. HARRISON: THANK YOU. SO NOW I'D LIKE
14	TO TURN TO A SERIES OF OTHER AMENDMENTS THAT WE
15	WOULD PROPOSE TO MAKE BOTH TO THE BYLAWS AND THE
16	INTERNAL GOVERNANCE POLICY WHICH AT TIMES OVERLAP.
17	FIRST, AS SOME OF YOU MAY RECALL, IN 2009 THE BOARD
18	CREATED THE POSITION OF BYLAWS VICE CHAIR. THE
19	BOARD CREATED THIS POSITION IN LIGHT OF THE RATHER
20	UNIQUE SET OF CIRCUMSTANCES WHEN THE BOARD WAS
21	BLESSED WITH TWO VERY FINE AND DISTINGUISHED
22	CANDIDATES FOR VICE CHAIR, SENATOR TORRES AND THE
23	LATE DUANE ROTH, WHO WAS ALREADY SERVING AS A MEMBER
24	OF THE BOARD.
25	THE POSITION WAS CREATED REALLY IN
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1	RECOGNITION OF DUANE'S UNIQUE ROLE AND CONTRIBUTIONS
2	TO THE BOARD. SINCE HIS UNTIMELY DEATH IN 2013, THE
3	BOARD HAS NOT REPLACED DUANE IN THE POSITION OF
4	BYLAWS VICE CHAIR. AS A RESULT, WE HAVE PROVISIONS
5	IN BOTH THE INTERNAL GOVERNANCE POLICY AND THE
6	BYLAWS THAT MAKE REFERENCE TO THIS POSITION. IN
7	LIGHT OF THE UNIQUE SET OF CIRCUMSTANCES PURSUANT TO
8	WHICH THE BOARD CREATED IT AND DUANE'S DEATH, WE
9	WOULD REQUEST THAT THE BOARD ELIMINATE THAT
10	POSITION.
11	THE NEXT ITEM RELATES TO TELEPHONIC
12	MEETINGS. THOSE OF YOU WHO WERE HERE AT THE
13	BEGINNING WILL REMEMBER THAT BOARD MEETINGS OFTEN
14	LASTED TWO DAYS AND WERE HELD NEARLY ON A MONTHLY
15	BASIS. AS A RESULT OF THE DEMANDS THAT WERE PLACED
16	ON BOARD MEMBERS AT THAT TIME, THE BOARD CREATED A
17	TELEPHONIC PARTICIPATION POLICY, WHICH CONTAINED A
18	NUMBER OF RESTRICTIONS, A CAP ON THE NUMBER OF
19	MEMBERS WHO COULD PARTICIPATE TELEPHONICALLY IN ANY
20	ONE MEETING. AT TIMES THIS POSED CHALLENGES FOR US
21	IN OBTAINING AND MAINTAINING A QUORUM.
22	IN LIGHT OF THE FACT THAT THE BOARD IS NOW
23	MEETING LESS FREQUENTLY IN PERSON, FOUR TIMES PER
24	YEAR, AND MORE FREQUENTLY VIA THE TELEPHONE, WE'D
25	PROPOSE TO ENCOURAGE RATHER THAN REQUIRE MEMBERS TO

1	ATTEND IN-PERSON MEETINGS IN PERSON AND LIFT THE
2	RESTRICTIONS ON TELEPHONIC PARTICIPATION SO THAT WE
3	CAN ENSURE THAT WE HAVE THE QUORUM NECESSARY TO TAKE
4	ACTIONS.
5	NEXT, THERE IS A PROVISION RELATING TO
6	SUBCOMMITTEES WHICH IS SUPERFLUOUS IN THAT IT
7	REQUIRES THE APPROVAL OF THE CHAIR TO EXPAND THE
8	SIZE OF A SUBCOMMITTEE, BUT THE POLICY ALREADY
9	REQUIRES THE CHAIR TO CONCUR WITH THE APPOINTMENT OF
10	THE MEMBERS TO A SUBCOMMITTEE. SO WE PROPOSE TO
11	DELETE THAT AS SUPERFLUOUS.
12	LIKEWISE, THERE IS A PROVISION THAT
13	SPECIFIES THE NUMBER OF MEETINGS AND NUMBER OF
14	MEMBERS FOR THE GOVERNANCE AND LEGISLATIVE
15	SUBCOMMITTEES WHICH WE BELIEVE IS UNNECESSARY.
16	NEXT ITEM RELATES TO THE ORGANIZATIONAL
17	CHART. IN THE PAST WHEN CIRM WAS UNDERGOING, I'D
18	SAY, MORE FREQUENT EVOLUTIONS IN TERMS OF ITS
19	ORGANIZATIONAL STRUCTURE, THE INTERNAL GOVERNANCE
20	POLICY AND THE BYLAWS REQUIRED THAT PROPOSED CHANGES
21	TO THE ORGANIZATIONAL CHART BE PRESENTED FIRST TO
22	THE GOVERNANCE SUBCOMMITTEE AND THEN TO THE BOARD.
23	WE HAVE OBVIOUSLY COMPLETED A REORGANIZATION WITH
24	THE APPOINTMENT OF DR. MILLS. WE HAVE ADOPTED A
25	STRATEGIC PLAN, AND WE DON'T ANTICIPATE MAKING
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1	SIGNIFICANT CHANGES TO THE ORGANIZATIONAL CHART
2	GOING FORWARD. BUT IF WE DO, WE PROPOSE TO TAKE
3	THOSE DIRECTLY TO THE BOARD SO THAT WE CAN HANDLE
4	ANY SUCH CHANGES MORE EFFICIENTLY.
5	FINALLY, THE BYLAWS SET FORTH THE
6	FUNCTIONS OF THE WORKING GROUPS. THIS JUST REPEATS
7	WHAT'S IN STATUTE, SO WE PROPOSE TO DELETE IT. THE
8	INTERNAL GOVERNANCE POLICY SPECIFIES THE NUMBER OF
9	EMPLOYEES IN THE OFFICE OF THE CHAIR. THIS
10	PROVISION WAS ADOPTED BY THE BOARD AT A TIME WHEN
11	THERE WAS A 50-EMPLOYEE CAP ON THE AGENCY, WHICH HAS
12	SINCE BEEN LIFTED THROUGH THE LEGISLATURE'S ADOPTION
13	OF SB 1064. SO WE BELIEVE IT'S NO LONGER NECESSARY
14	TO SPECIFY THE NUMBER OF EMPLOYEES IN THE OFFICE OF
15	THE CHAIR. WE ARE NOW FUNCTIONING VERY WELL AS A
16	SINGLE TEAM. WE THINK THAT PROVISION IS NO LONGER
17	NECESSARY.
18	FINALLY, WITH RESPECT TO THE COMPENSATION
19	OF EMPLOYEES, THE INTERNAL GOVERNANCE POLICY
20	CURRENTLY REQUIRES THAT CIRM LEADERSHIP TO SEEK THE
21	APPROVAL OF THE GOVERNANCE SUBCOMMITTEE BEFORE
22	APPOINTING AN EMPLOYEE AT A SALARY THAT EXCEEDS 80
23	PERCENT OF THE RANGE.
24	IN LIGHT OF CIRM'S CURRENT SITUATION, THE
25	LIMITED FUNDING THAT WE HAVE LEFT, RECRUITMENT, AS
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1	DR. MILLS MENTIONED EARLIER, IS ONE OF OUR
2	CHALLENGES. AND WE NEED TO BE NIMBLE AND MOVE
3	QUICKLY SOMETIMES IN ORDER TO CAPTURE THE BEST
4	CANDIDATE. SO WE'D REQUEST THAT THE BOARD GIVE CIRM
5	LEADERSHIP THE AUTHORITY TO SET SALARIES UP TO 100
6	PERCENT OF THE RANGE APPROVED BY THE BOARD AND, OF
7	COURSE, THE BOARD WOULD RETAIN AUTHORITY TO APPROVE
8	ANY SALARIES THAT EXCEED THE RANGE.
9	WITH THAT, WE'D ASK FOR A MOTION APPROVING
10	THESE PROPOSED AMENDMENTS. AND I'D BE HAPPY TO
11	ANSWER ANY QUESTIONS.
12	DR. JUELSGAARD: SO MOVED.
13	DR. PRIETO: SECOND.
14	MR. PANETTA: SECOND.
15	CHAIRMAN THOMAS: MOVED BY MR. JUELSGAARD,
16	SECONDED BY MR. PANETTA. ANY FURTHER DISCUSSION,
17	QUESTIONS, COMMENTS? DR. PRICE, YOU LOOK
18	PARTICULARLY PENSIVE.
19	DR. PRICE: THIS IS AN OMNIBUS.
20	CHAIRMAN THOMAS: THAT'S CORRECT. THIS IS
21	AN OMNIBUS MOTION.
22	DR. PRICE: SO I'M ALLOWED TO COMMENT ON
23	JUST A PART OF IT, I PRESUME.
24	CHAIRMAN THOMAS: ABSOLUTELY.
25	DR. PRICE: SO THE POINT I WANT TO TALK
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1	ABOUT IS THE TELEPHONIC, THE CHANGE IN THE
2	TELEPHONIC, WHICH I GATHER LIFTS THE RESTRICTIONS.
3	SO IN PRINCIPLE IF WE ADOPT THESE BYLAWS IN ANY OF
4	THESE FOUR MEETINGS OR ALL OF THEM, WE COULD END UP
5	WITH A MEETING WITH TWO PEOPLE SITTING HERE AND
6	EVERYBODY ELSE ON THE PHONE. OKAY. SO WE'VE
7	ALREADY ELIMINATED, AS FAR AS I CAN TELL, ALL OF THE
8	MEETINGS OF THE BOARD EXCEPT FOR THESE FOUR. THOSE
9	MEETINGS WILL BE TELEPHONIC.
10	MS. BONNEVILLE: THERE WILL BE MONTHLY
11	MEETINGS.
12	DR. PRICE: THEY'LL BE TELEPHONIC.
13	EXACTLY. WHAT THIS AMENDMENT SEEMS TO ME TO DO IS
14	TO TURN THEM ALL OR POTENTIALLY TURN THEM ALL INTO
15	TELEPHONIC MEETINGS BY LIFTING THAT RESTRICTION. I
16	KIND OF I WON'T COMMENT ON THE COMPARATIVE VALUE
17	OF TELEPHONIC VERSUS IN-PERSON MEETINGS. I'LL JUST
18	LEAVE THAT ASIDE FOR THE MOMENT. BUT I JUST HAVE TO
19	SAY, FRANKLY, THAT IF MEMBERS, IF PEOPLE ARE EITHER
20	UNWILLING OR UNABLE TO MEET IN PERSON FOUR TIMES A
21	YEAR, I QUESTION WHY ACCEPT MEMBERSHIP ON THE BOARD.
22	SO THAT'S MY I THINK IN-PERSON MEETINGS ARE MUCH
23	MORE VALUABLE THAN TELEPHONIC ONES. AND, THEREFORE,
24	I'M OPPOSED TO THE IDEA OF INTRODUCING THIS NEW
25	AMENDMENT WHICH WILL ALLOW ALL, IN PRINCIPLE OR

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1	HYPOTHETICALLY, ALL MEETINGS TO BECOME LARGELY, IF
2	NOT ENTIRELY, TELEPHONIC.
3	CHAIRMAN THOMAS: I THINK THE IDEA IS
4	ABSOLUTELY NOT TO END UP WITH THAT RESULT. THIS IS
5	JUST A WAY OF TRYING TO MAKE IT WORK BECAUSE WE HAVE
6	MORE BOARD MEETINGS NOW UNDER CIRM 2.0 THAN EVER.
7	BOARD LITERALLY MEETS MONTHLY IF THERE ARE
8	APPLICATIONS TO CONSIDER, AND SO WE DIDN'T WANT TO
9	HAVE MONTHLY MEETINGS. SO WHAT WE'VE DONE IS WE'VE
10	ACTUALLY INCREASED THE NUMBER OF TOTAL BOARD
11	MEETINGS, BUT REDUCED THE NUMBER OF IN-PERSON, BUT
12	WE BY NO MEANS ARE ANGLING TOWARDS NO IN-PERSON
13	MEETINGS.
14	MR. HARRISON: I'LL JUST ADD THE INTENT,
15	OUR HOPE AND EXPECTATION, IS THAT THE MEMBERS WILL
16	ATTEND THE FOUR REGULAR MEETINGS IN PERSON. WHAT WE
17	WERE LOOKING FOR WAS SOME ADDITIONAL FLEXIBILITY IN
18	THE EVENT THAT AT THE LAST MINUTE A MEMBER CALLED
19	MARIA BONNEVILLE AND SAID HE OR SHE WAS SICK OR
20	DETAINED IN SOME OTHER WAY, AND WE THEN PERHAPS LOSE
21	A QUORUM AND CAN'T MOVE FORWARD.
22	SO I DEFER TO YOU, BUT IT WAS IN THAT
23	SPIRIT NOT TO DISCOURAGE PEOPLE FROM PARTICIPATING
24	IN PERSON.
25	DR. PRICE: I UNDERSTAND THE SPIRIT. IT'S
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1	THE CONSEQUENCES THAT I'M CONCERNED ABOUT. YOU
2	COULD EXPAND THE CAP WITHOUT LIFTING IT ENTIRELY.
3	CHAIRMAN THOMAS: APPRECIATE THAT. THANK
4	YOU, DR. PRICE.
5	DR. DULIEGE: HAVING EXPERIENCED THAT
6	SYSTEM FOR ABOUT SIX MONTHS, I THINK BY NOW, I LIKE
7	IT A LOT BECAUSE IT ACTUALLY FORCES PEOPLE TO COME
8	MORE TO THE IN-PERSON MEETING. IT WAS SORT OF EASY
9	TO SKIP BEFORE BECAUSE YOU HAD SO MANY IN-PERSON
10	MEETINGS. NOW THERE'S ONLY FOUR. YOU REALLY WANT
11	TO BE THERE, NO. 1.
12	NO. 2, FOR THE PHONE MEETINGS, IT'S MOSTLY
13	ABOUT GIVING OUR POSITION ON SPECIFIC GRANTS, WHICH
14	IF PEOPLE PREPARE THEIR HOMEWORK, WE CAN DO IT VERY
15	WELL OVER THE PHONE. SPECIFICALLY THE CIRM STAFF
16	HAS MADE AN EFFORT TO MAKE ALL THE PREPARATION FOR
17	THESE DECISIONS TO BE MADE MUCH MORE
18	STRAIGHTFORWARD. SO IF YOU DO YOUR HOMEWORK, YOU
19	CAN BE VERY ACTIVE MEMBER OVER THE PHONE WHILE
20	WHAT'S DISCUSSED DURING THE IN-PERSON ARE THESE KIND
21	OF THINGS, WHICH ARE THE OVERALL STRATEGIC
22	DIRECTIONS OF CIRM TOGETHER.
23	SO OVERALL STRATEGIC DIRECTION, IN-PERSON
24	MEETING. SPECIFIC APPLICATIONS, PHONE. AND I THINK
25	THAT'S FINE. FOR ME THAT HAS WORKED EXTREMELY WELL.
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1	CHAIRMAN THOMAS: WHEN ARE THE FOUR BOARD
2	MEETINGS IN PERSON ARE GOING TO BE HELD?
3	MS. BONNEVILLE: THEY'LL BE HELD
4	QUARTERLY, MARCH, JUNE I'VE SENT THOSE OUT. I'LL
5	SEND THEM OUT AGAIN, AND WE CAN HOPEFULLY GET THEM
6	ON EVERYONE'S CALENDAR.
7	DR. DULIEGE: I JUST WANTED TO FINISH AND
8	SAY I THINK THE ONLY DOWNSIDE OF THIS PLAN, ONLY
9	ONE, IS I BELIEVE THAT IT MAY BE A LITTLE LESS EASY
10	FOR THE MEMBERS OF THE PUBLIC TO INTERVENE OVER THE
11	PHONE. AND SO IF THEY WANTED TO INTERVENE ABOUT A
12	PARTICULAR APPLICATION, THAT'S A LITTLE BIT MORE
13	DIFFICULT.
14	MS. LANSING: THE WAY I'M UNDERSTANDING
15	THIS, AND I HOPE I'M UNDERSTANDING IT CORRECTLY, IS
16	WE STILL HAVE TO ABIDE BY BAGLEY-KEENE. SO YOU
17	STILL HAVE TO POST WHERE YOU ARE. AND SO I'VE BEEN
18	ON THE PHONE, UNFORTUNATELY OR FORTUNATELY,
19	WHATEVER, BUT IT HAS ALLOWED ME TO CONTINUE TO
20	PARTICIPATE. BUT I ALWAYS HAVE TO POST IT, AND I
21	ALWAYS HAVE TO OPEN IT TO THE PUBLIC AND HAVE THE
22	PUBLIC MEMBERS THERE AND THEY COMMENT, AND YOU
23	ACTUALLY END UP HAVING MORE TIME WITH THEM.
24	CHAIRMAN THOMAS: OTHER COMMENTS? OKAY.
25	WE DID A HAVE A MOTION AND SECOND.

1	MR. SHEEHY: JUST ONE COMMENT ON THE
2	SUBCOMMITTEES. MAYBE YOU COULD JUST CLARIFY WHAT
3	THE PROCESS IS FOR GETTING ON A SUBCOMMITTEE BECAUSE
4	I KNOW WE HAVE SOME NEW MEMBERS. AND I THINK JUST
5	TO MAKE SURE THAT WE'RE REALLY POROUS TO
6	PARTICIPATION BY ANY MEMBER, HOW DO PEOPLE BECOME A
7	MEMBER OF A SUBCOMMITTEE IF SOMEONE HAS AN INTEREST
8	BECAUSE I PERSONALLY THINK THAT THEY SHOULD BE OPEN
9	TO ANYONE WHO WANTS TO PARTICIPATE. AND THEN ALSO
10	TO ENCOURAGE FOLKS IF THEY WANT TO BE ON A
11	SUBCOMMITTEE, ONCE WE SAY WHAT THE PROCESS IS,
12	PLEASE, IF YOU'D LIKE TO, FOLLOW THAT PROCESS.
13	MR. HARRISON: THE CHAIR OF EACH
14	SUBCOMMITTEE HAS THE AUTHORITY TO APPOINT THE
15	MEMBERS OF THE SUBCOMMITTEE WITH THE CONCURRENCE OF
16	THE CHAIR OF THE BOARD. SO IT IS OPEN TO MEMBERS
17	WHO'D LIKE TO JOIN. ONE OF THE THINGS WE'VE DONE IS
18	TO LIFT THE SPECIFICITY WITH RESPECT TO THE NUMBER
19	OF MEMBERS ON THE LEGISLATIVE AND GOVERNANCE
20	SUBCOMMITTEES. SO IT'S UP TO THE DISCRETION OF THE
21	CHAIR OF THE SUBCOMMITTEE AND THE BOARD, BUT I'M
22	SURE THEY WOULD WELCOME PARTICIPATION.
23	CHAIRMAN THOMAS: I THINK YOU RAISE A
24	POINT, MR. SHEEHY. I THINK WHAT WE WOULD LIKE TO
25	DO, HEADING INTO A NEW YEAR, IS TO CIRCULATE TO
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1	EVERYBODY JUST SO YOU KNOW WHAT THE RANGE IS OF THE
2	SUBCOMMITTEES THAT ARE AVAILABLE; AND IF YOU ARE
3	INTERESTED IN ANY OF THEM, BY ALL MEANS PLEASE LET
4	US KNOW BECAUSE THE MORE HELP WE CAN GET ON ALL
5	TOPICS, THE BETTER. SO I WOULD ECHO WHAT MR. SHEEHY
6	SAID, ACTIVELY ENCOURAGE PARTICIPATION, BECAUSE A
7	LOT OF THINGS HAPPEN AT THESE SUBCOMMITTEES BEFORE
8	THEY MAKE IT TO THE BOARD.
9	ANY OTHER COMMENTS? THIS IS A VOICE VOTE,
10	MR. HARRISON, EXCEPT FOR THOSE ON THE PHONE. ALL
11	THOSE IN FAVOR OF THE OMNIBUS MOTION SET FORTH,
12	PLEASE SAY AYE. OPPOSED? ABSTENTIONS?
13	MARIA, WILL YOU PLEASE POLL THOSE ON THE
14	PHONE?
15	MS. BONNEVILLE: LINDA BOXER.
16	DR. BOXER: YES.
17	MS. BONNEVILLE: KATHY LAPORTE.
18	MS. LAPORTE: YES.
19	MS. BONNEVILLE: BERT LUBIN.
20	DR. LUBIN: YES.
21	MS. BONNEVILLE: DIANE WINOKUR.
22	CHAIRMAN THOMAS: OKAY. THAT PASSES.
23	THANK YOU VERY MUCH.
24	SCOTT, ARE YOU TAKING THIS NEXT ITEM?
25	MR. TOCHER: THAT'S CORRECT. THE CIRM
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1	CONFLICT OF INTEREST POLICY FOR CIRM EMPLOYEES.
2	CHAIRMAN THOMAS: OKAY. WITHOUT FURTHER
3	ADO, MR. TOCHER.
4	MR. TOCHER: THANK YOU. AS MENTIONED,
5	THIS NEXT ITEM ON OUR CORE 2.0 REVIEW CONCERNS
6	CIRM'S CONFLICT OF INTEREST POLICY FOR CIRM
7	EMPLOYEES. UNDER EXISTING STATE LAW, ALL CIRM
8	EMPLOYEES ARE PUBLIC OFFICIALS, JUST AS MEMBERS OF
9	THIS BOARD, AND ARE GOVERNED BY SEPARATE STATE LAW
10	BY THE POLITICAL REFORM ACT WHICH GOVERNS CONFLICTS
11	OF INTEREST AND FINANCIAL DISCLOSURES. HOWEVER,
12	BECAUSE OF CIRM'S UNIQUE MISSION AND PROFILE, EARLY
13	IN THE AGENCY'S EXISTENCE, WE ADOPTED ADDITIONAL
14	RULES TO AUGMENT THE STATE LAW TO IDENTIFY
15	ADDITIONAL CIRCUMSTANCES WHERE EMPLOYEES MUST
16	REFRAIN FROM PARTICIPATING. AND THERE ARE SIX
17	CIRCUMSTANCES.
18	WE ARE PROPOSING IMPROVEMENTS TO ALL OF
19	THOSE SIX. MEMBERS OF THE GOVERNANCE SUBCOMMITTEE
20	MET EARLIER THIS MONTH AND RECOMMENDED UNANIMOUSLY
21	THESE AMENDMENTS BEFORE YOU, AND I'LL ITEMIZE THEM
22	NOW. THESE ARE ATTACHED IN YOUR BINDERS.
23	THE FIRST AND SECOND CIRCUMSTANCES GOVERN
24	PARTICIPATION BY AN EMPLOYEE WHERE A FAMILY MEMBER
25	MAY RECEIVE A FINANCIAL BENEFIT FROM A GRANT OR THE
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1	MEMBER IS AN EMPLOYEE OF THE APPLICANT INSTITUTION.
2	AND THE AMENDMENTS THAT WE PROPOSE SIMPLY CONFORM
3	WITH STATE LAW THE DEFINITION OF IMMEDIATE FAMILY
4	MEMBER TO INCLUDE SPOUSE AND DEPENDENT CHILDREN.
5	IN THE THIRD CIRCUMSTANCE, IT CONSIDERS
6	WHERE THE PI ON AN APPLICATION OR GRANT IS OR HAS
7	BEEN A RECENT COLLABORATOR OF THE EMPLOYEE. AND
8	HERE AGAIN WE ARE CONFORMING OUR DEFINITION OF
9	RESEARCH COLLABORATOR WITH THE BOARD'S RECENT
10	ADOPTION OF THE DEFINITION OF THAT TERM IN THE
11	CONTEXT OF OUR GRANTS WORKING GROUP CONFLICT OF
12	INTEREST POLICY. SO THIS SHOULD BE A MORE EFFICIENT
13	AND EASIER TO FOLLOW DEFINITION ACROSS THE AGENCY'S
14	COI POLICIES.
15	THE NEXT CIRCUMSTANCE PREVENTS CIRM
16	EMPLOYEES FROM OWNING STOCK EXCEEDING A VALUE OF
17	\$10,000 IN STEM CELL COMPANIES THAT HAVE A
18	SUBSTANTIAL INTEREST IN CELL THERAPIES. THE
19	EXISTING DEFINITION IN OUR POLICY DEFINES
20	SUBSTANTIAL INTEREST AS ONE WHERE THE COMPANY
21	DEVOTES 5 PERCENT, AT LEAST 5 PERCENT OF ITS
22	RESEARCH BUDGET TO STEM CELL RESEARCH. IN PRACTICE,
23	THIS HAS PROVEN SOMEWHAT DIFFICULT FOR EMPLOYEES TO
24	DETERMINE BASED ON PUBLICLY AVAILABLE INFORMATION.
25	AS A RESULT, WE HAVE PROPOSED TO INCREASE
	1.41

1	THE THRESHOLD TO 20 PERCENT OF THE COMPANY'S
2	RESEARCH BUDGET, WHICH THROUGH PUBLICLY AVAILABLE
3	INFORMATION, WE BELIEVE, WILL BE EASIER TO ASCERTAIN
4	AND, THUS, MAKE COMPLIANCE AND ADHERENCE MORE
5	CONSISTENT.
6	AND FINALLY, THE LAST CIRCUMSTANCE IS ONE
7	IN WHICH IT ADDRESSES EMPLOYEES WHO ARE PERFORMING
8	CONSULTING, TEACHING, OR ADVISORY BOARD SERVICES FOR
9	AWARDEES. AND HERE OUR AMENDMENTS PROPOSE TO
10	CONFORM TO A RECENT INTERPRETATION OF STATE LAW BY
11	THE FAIR POLITICAL PRACTICES COMMISSION IN THE
12	CONTEXT WHERE SUCH SERVICE IS VOLUNTARY AND THE
13	EMPLOYEE RECEIVES NO COMPENSATION OR FINANCIAL
14	BENEFIT.
15	SO AS I MENTIONED, THESE AMENDMENTS WERE
16	APPROVED AND RECOMMENDED TO YOUR APPROVAL BY THE
17	GOVERNANCE SUBCOMMITTEE. AND I'M HAPPY TO TAKE ANY
18	QUESTIONS ABOUT THE SPECIFICS.
19	CHAIRMAN THOMAS: QUESTIONS? DO I HEAR A
20	MOTION?
21	MR. ROWLETT: SO MOVED.
22	CHAIRMAN THOMAS: SO MOVED BY MR. ROWLETT.
23	DR. GASSON: SECOND.
24	CHAIRMAN THOMAS: SECONDED BY DR. GASSON.
25	ANY COMMENTS? ANY COMMENTS BY MEMBERS ON THE PHONE?
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1	ANY COMMENTS BY MEMBERS OF THE PUBLIC? HEARING
2	NONE, THIS IS ANOTHER VOICE VOTE ITEM. ALL THOSE IN
3	FAVOR PLEASE SAY AYE. OPPOSED? ABSTENTIONS?
4	MARIA, POLL THOSE ON THE PHONE PLEASE.
5	MS. BONNEVILLE: LINDA BOXER.
6	DR. BOXER: YES.
7	MS. BONNEVILLE: KATHY LAPORTE.
8	MS. LAPORTE: YES.
9	MS. BONNEVILLE: BERT LUBIN.
10	DR. LUBIN: YES.
11	MS. BONNEVILLE: DIANE WINOKUR.
12	CHAIRMAN THOMAS: MOTION IS APPROVED. MR.
13	HARRISON, YOU'RE BACK UP FOR THE NEXT ITEM, WHICH I
14	BELIEVE IS ON EMPLOYEE COMPENSATION.
15	MR. HARRISON: AS MR. TOCHER SAID, THE
16	GOVERNANCE SUBCOMMITTEE REVIEWED ALL OF THESE
17	POLICIES, INCLUDING WHAT I'D LIKE TO DISCUSS WITH
18	YOU NOW, WHICH ARE PROPOSED AMENDMENTS TO THE
19	EMPLOYEE COMPENSATION POLICY AND THE RELOCATION
20	POLICY.
21	WITH RESPECT TO THE EMPLOYEE COMPENSATION
22	POLICY, WHICH IS SET FORTH UNDER TAB 14, WE PROPOSE
23	TO MAKE THREE CHANGES OF SIGNIFICANCE. ONE, WE
24	PROPOSE TO DELETE THE PROVISION ALLOWING CIRM TO
25	MAKE SPOT AWARDS. SPOT AWARDS WERE AWARDS OF \$75 OR
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1	LESS. THEY RAISE TAX ISSUES AND DID NOT PROVE TO BE
2	PARTICULARLY EFFECTIVE IN TERMS OF EMPLOYEE MORALE,
3	RETENTION, AND RECRUITMENT.
4	RATHER THAN USING SPOT AWARDS TO RECOGNIZE
5	EXCEPTIONAL PERFORMANCE, WE WOULD PROPOSE TO EXPAND
6	THE PROVISION PROVIDING FOR PERFORMANCE AWARDS.
7	CURRENTLY PERFORMANCE AWARDS ARE LIMITED TO SALARY
8	RANGES 1 THROUGH 6. WE WOULD PROPOSE TO ALLOW CIRM
9	LEADERSHIP TO USE PERFORMANCE AWARDS FOR ALL
10	EMPLOYEES UP TO SALARY RANGE LEVEL 9. SO THAT WOULD
11	EXCLUDE EMPLOYEES IN SALARY RANGE 10, WHICH MEANS
12	THE PRESIDENT, THE CHAIR, THE VICE CHAIR.
13	WE THINK THIS WILL HELP US ADDRESS ONE OF
14	THE CHALLENGES DR. MILLS IDENTIFIED EARLIER, WHICH
15	IS EMPLOYEE RETENTION. AND OBVIOUSLY, AS HAS BEEN
16	NOTED EARLIER, WE'VE SEEN SOME EXCEPTIONAL
17	PERFORMANCE AT CIRM OVER THE COURSE OF ITS EXISTENCE
18	AND IN PARTICULAR OVER THE COURSE OF THE LAST YEAR.
19	AND CIRM'S LEADERSHIP WOULD LIKE THE ABILITY TO
20	RECOGNIZE THAT.
21	NEXT WE PROPOSE TO ELIMINATE THE REFERENCE
22	TO PROFESSIONAL DEVELOPMENT IN THE COMPENSATION
23	POLICY. DR. MILLS AND THE CIRM LEADERSHIP ARE VERY
24	INTERESTED IN PROFESSIONAL DEVELOPMENT, BUT IT JUST
25	DOESN'T APPEAR TO BELONG IN THE COMPENSATION POLICY.
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1	SO WE PROPOSE TO HANDLE PROFESSIONAL DEVELOPMENT
2	SEPARATELY. IT IS A LINE ITEM IN THE BUDGET, AND WE
3	PLAN TO ADDRESS IT IN THAT CONTEXT.
4	AND THEN, FINALLY, IN ALIGNMENT WITH THE
5	CHANGES THE BOARD APPROVED WITH RESPECT TO
6	GOVERNANCE SUBCOMMITTEE APPROVAL AT APPOINTMENTS
7	OVER 80 PERCENT OF THE SALARY RANGE, WE PROPOSE TO
8	ELIMINATE THAT PROVISION IN THE COMPENSATION POLICY
9	AS WELL.
10	THERE'S ONLY ONE REAL SIGNIFICANT CHANGE
11	IN THE RELOCATION POLICY WHICH ALLOWS CIRM TO
12	PROVIDE A RELOCATION ALLOWANCE TO EMPLOYEES WHO MOVE
13	FROM OUT OF STATE TO JOIN THE AGENCY, WHICH IS
14	LIMITED TO THE LESSER OF 25 PERCENT OF THE
15	EMPLOYEE'S SALARY OR \$75,000. CURRENTLY THAT AMOUNT
16	WHICH IS PAID OUT OVER FOUR YEARS IS ADDED IN THE
17	FIRST YEAR'S SALARY FOR PURPOSES OF DETERMINING
18	WHETHER OR NOT BOARD APPROVAL IS REQUIRED. IN OTHER
19	WORDS, IT'S ADDED TO THE BASE SALARY TO DETERMINE
20	WHETHER OR NOT THE SALARY EXCEEDS 100 PERCENT OF THE
21	RANGE, WHICH DOESN'T REALLY MAKE A LOT OF SENSE TO
22	US. IT IS ESSENTIALLY A ONE-TIME PAYMENT REFLECTING
23	THE COSTS ASSOCIATED WITH RELOCATING TO CALIFORNIA.
24	SO WE PROPOSE TO ELIMINATE THAT FROM THE
25	CALCULATION OF THE EMPLOYEE'S BASE SALARY FOR

1	PURPOSES OF DETERMINING WHEN WE HAVE TO BRING SALARY
2	TO THE BOARD FOR ITS APPROVAL. SO THAT'S A QUICK
3	SUMMARY OF THE SUBSTANTIVE CHANGES TO THE
4	COMPENSATION AND RELOCATION POLICY. I'D BE HAPPY TO
5	ANSWER ANY QUESTIONS.
6	CHAIRMAN THOMAS: QUESTIONS FROM MEMBERS
7	OF THE BOARD? QUESTIONS, COMMENTS
8	MR. SHEEHY: MOVE APPROVAL.
9	DR. STEWARD: SECOND.
10	CHAIRMAN THOMAS: MOVED BY MR. SHEEHY,
11	SECONDED BY DR. STEWARD. ANY FURTHER COMMENTS FROM
12	MEMBERS OF THE BOARD, INCLUDING THOSE ON THE PHONE?
13	DR. LUBIN: YOU DON'T HAVE ANY RELOCATION
14	UPGRADING FROM SAN FRANCISCO TO OAKLAND, DO YOU?
15	MR. TORRES: YOU SHOULD HAVE GIVEN US
16	SOME.
17	CHAIRMAN THOMAS: THAT'S AN EXCELLENT
18	QUESTION, DR. LUBIN.
19	OKAY. THIS IS ANOTHER VOICE VOTE. ANY
20	COMMENTS FROM MEMBERS OF THE PUBLIC? HEARING NONE,
21	ALL THOSE IN FAVOR PLEASE SAY AYE. OPPOSED?
22	ABSTENTIONS? GLAD TO HEAR DR. LUBIN APPROVES.
23	MARIA, DO YOU WANT TO POLL THOSE, EVEN THOUGH WE
24	KNOW HIS RESPONSE?
25	MS. BONNEVILLE: LINDA BOXER.
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1	DR. BOXER: YES.
2	MS. BONNEVILLE: KATHY LAPORTE.
3	MS. LAPORTE: YES.
4	CHAIRMAN THOMAS: MOTION PASSES. OKAY.
5	JAMES, DO YOU HAVE ONE MORE UP HERE? IT'S ON THE
6	GWG BYLAWS. NOW WE'RE REALLY GETTING INTO THE
7	RIVETING SUBJECT MATTER.
8	MR. HARRISON: THANK YOU. THIS IS THE
9	LAST ITEM IN THIS LINE OF VERY EXCITING ITEMS FOR
10	YOUR CONSIDERATION. AS PART OF OUR REVIEW OF
11	INTERNAL POLICIES, WE'VE ALSO TAKEN ANOTHER LOOK AT
12	THE GWG BYLAWS TO DETERMINE WHETHER THERE ARE
13	OPPORTUNITIES FOR IMPROVEMENT, CLARIFICATION, ETC.
14	AND WE WANT TO BRING A COUPLE OF ITEMS TO YOUR
15	ATTENTION TODAY.
16	FIRST, PROP 71 IMPOSES A TWO CONSECUTIVE
17	TERM LIMIT ON MEMBERS OF THE GRANTS WORKING GROUP,
18	AND IT ALSO SPECIFIES THAT AFTER THE INITIAL TERMS
19	OF THE SCIENTIFIC MEMBERS OF THE GRANTS WORKING
20	GROUP, THAT SCIENTIFIC MEMBERS ARE THEREAFTER
21	APPOINTED ONE-THIRD TO TWO-YEAR TERMS, ONE-THIRD TO
22	FOUR-YEAR TERMS, AND ONE-THIRD TO SIX-YEAR TERMS.
23	PROP 71 DOES NOT SPECIFY HOW LONG A PERSON
24	WHO HAS SERVED TWO CONSECUTIVE TERMS MUST BE A
25	NONMEMBER BEFORE HE OR SHE IS ELIGIBLE FOR

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1	REAPPOINTMENT. AS SOME OF YOU KNOW, WE'VE HAD THE
2	BENEFIT OF THE SERVICE OF SOME REALLY WONDERFUL
3	GRANTS WORKING GROUP MEMBERS WHO WE WOULD LIKE TO
4	PERHAPS HAVE THE OPPORTUNITY TO APPOINT IN THE
5	FUTURE. SO WE'D PROPOSE TO DEFINE THE GAP TERM, IF
6	YOU WILL, TO THE SHORTEST TERM POSSIBLE, WHICH IN
7	THIS CASE WOULD BE TWO YEARS. IN OTHER WORDS, A
8	MEMBER OF THE GRANTS WORKING GROUP WHO SERVED TWO
9	CONSECUTIVE TERMS WOULD BE ELIGIBLE FOR
10	REAPPOINTMENT TWO YEARS AFTER HIS OR HER LAST TERM
11	ENDS.
12	THE NEXT ITEM RELATES TO THE REVIEW OF THE
13	APPLICATIONS FOR NONCLINICAL PROGRAMS. AS YOU WILL
14	RECALL, WE DEVELOPED A NEW SCORING SYSTEM FOR THE
15	CLINICAL PROGRAMS WHERE A SCORE OF ONE MEANS THE
16	APPLICATION IS RECOMMENDED FOR FUNDING, A SCORE OF
17	TWO INDICATES THAT THE APPLICATION HAS SOME MERIT,
18	BUT HAS SOME FLAWS THAT COULD BE ADDRESSED, AND A
19	SCORE OF THREE INDICATES THAT THE APPLICATION IS
20	SUFFICIENTLY FLAWED THAT THE SAME APPLICATION OR
21	SAME PROJECT, RATHER, SHOULD NOT BE SUBMITTED.
22	WE THINK THIS SCORING SYSTEM WORKS VERY
23	WELL FOR THE PURPOSES OF THE CLINICAL PROGRAM
24	BECAUSE WE HAVE A MONTHLY REVIEW CYCLE AND THE
25	VOLUME OF APPLICATIONS IS RELATIVELY SMALL. WE
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DON'T THINK THAT THE SAME SCORING SYSTEM WOULD BE
EFFECTIVE FOR PURPOSES OF THE NONCLINICAL PROGRAM,
FOR EXAMPLE, THE TRANSLATION AND DISCOVERY PROGRAMS,
AND THE PROGRAMS THAT DR. MILLAN PRESENTED TO YOU
EARLIER TODAY.
SO AS TO THOSE PROGRAMS, WE PROPOSE TO
REVERT TO A HYBRID OF OUR OLD SYSTEM. SO
APPLICATIONS WOULD BE SCORED ON A SCALE OF 1 TO 100
AS THEY HAVE TRADITIONALLY. A SCORE OF 85 OR ABOVE
WOULD MEAN THE APPLICATION IS IN TIER I, WHICH
SIGNIFIES THAT IT'S RECOMMENDED FOR FUNDING IF FUNDS
ARE AVAILABLE. IF THE APPLICATION HAD AN AVERAGE
SCIENTIFIC SCORE OF 84 OR BELOW, IT WOULD BE DEFINED
TO BE IN TIER II, WHICH SIGNIFIES THAT IT'S NOT
RECOMMENDED FOR FUNDING. IN PARTICULAR, FOR
APPLICATIONS FOR WHICH WE ANTICIPATE ONLY ONE AWARD,
NAMELY, THE THREE CONCEPT PLANS YOU APPROVED EARLIER
TODAY, WE'D PROPOSE TO SPECIFY THAT THE APPLICATION
THAT RECEIVES THE HIGHEST AVERAGE SCORE FROM THE GWG
IS DEEMED TO BE THE GWG'S RECOMMENDATION FOR
FUNDING.
THE LAST PROPOSAL WE'D LIKE YOU TO
CONSIDER RELATES TO THE FINAL MOTION. AS THOSE OF
YOU WHO HAVE OBSERVED GWG REVIEW MEETINGS KNOW, AT
THE END OF THE REVIEW, THE GWG, INCLUDING THE
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1	PATIENT ADVOCATE MEMBERS, TAKES A VOTE TO FORWARD
2	THE GROUP'S RECOMMENDATIONS ON TO THE BOARD.
3	CURRENTLY THIS MOTION REALLY IS NOTHING MORE THAN AN
4	INDICATION THAT THE SCORES THAT THEY ARE FORWARDING
5	TO YOU ARE AN ACCURATE REFLECTION OF WHAT HAPPENED
6	AT THE MEETING.
7	WE HAVE NOTICED OVER THE COURSE OF TIME
8	THAT FREQUENTLY MEMBERS OF THE BOARD WILL ASK THEIR
9	COLLEAGUES WHO SERVE ON THE WORKING GROUP ABOUT THE
10	TENOR OF THE REVIEW, HOW RIGOROUS IT WAS, ETC.
11	BECAUSE ONLY THE PATIENT ADVOCATE MEMBERS OF THE
12	BOARD ARE ELIGIBLE TO SERVE ON THE GRANTS WORKING
13	GROUP, THEY REALLY PLAY A SPECIAL ROLE IN THE SENSE
14	THAT THEY ARE A BRIDGE BETWEEN THE GRANTS WORKING
15	GROUP AND THE BOARD. AND TO REFLECT THAT, WE'D LIKE
16	TO ENHANCE THAT FINAL MOTION THAT THE GWG CONSIDERS.
17	FIRST, WHAT WE PROPOSE IS THAT THE FULL
18	GWG CONSIDER A MOTION THAT WOULD FORWARD THE
19	RECOMMENDATIONS ALONG TO THE BOARD ALONG WITH THE
20	DETERMINATION OF WHETHER THE REVIEW WAS
21	SCIENTIFICALLY RIGOROUS, WHETHER THERE WAS
22	SUFFICIENT TIME FOR ALL VIEWPOINTS TO BE HEARD, AND
23	THAT THE SCORES ACCURATELY REFLECT WHAT HAPPENED.
24	WE'D PROPOSE A SECOND PART TO THAT MOTION,
25	WHICH WOULD BE LIMITED TO THE PATIENT ADVOCATE
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1	MEMBERS OF THE GWG WHO REALLY IN SOME SENSE FUNCTION
2	NOT ONLY AS A BRIDGE, BUT AS OBSERVERS OF THE BOARD
3	TO THE CONDUCT OF THE MEETING, FOR THE PATIENT
4	ADVOCATES TO CONSIDER WHETHER THE REVIEW WAS CARRIED
5	OUT IN A FAIR MANNER THAT WAS FREE FROM UNDUE BIAS.
6	SO WE PROPOSE TO FORMALIZE THIS IN THE BYLAWS AND
7	MAKE THIS PART OF A STANDING MOTION THAT THE GWG
8	WOULD CONSIDER AT THE CLOSE OF EACH REVIEW.
9	SO WE REQUEST THAT THE BOARD ADOPT THESE
10	PROPOSED AMENDMENTS TO THE GRANTS WORKING GROUP
11	BYLAWS, AND I'D BE HAPPY TO ANSWER ANY QUESTIONS.
12	DR. JUELSGAARD: LAST OF THESE, THE GWG
13	MOTIONS, SO ARE THEY TO BE APPROVED UNANIMOUSLY OR
14	SIMPLY BY A MAJORITY? SO IF THERE'S A STRONG
15	DIVISION IN SCORES AND, SAY, IT'S, PICK A NUMBER, 8
16	TO 7, SOMETHING LIKE THAT, IT WILL STILL REFLECT
17	THAT THE REVIEW WAS SCIENTIFICALLY RIGOROUS, THERE
18	WAS SUFFICIENT TIME FOR ALL VIEWPOINTS, THE SCORES
19	REFLECT THE RECOMMENDATIONS, ETC.?
20	MR. HARRISON: YES. SO TO BE CLEAR,
21	WHATEVER THE VOTE IS FIRST OF ALL, WE WOULD HOPE
22	THAT IT'S UNANIMOUS; BUT IF IT'S NOT, WHATEVER THE
23	VOTE IS WOULD BE BROUGHT TO THE BOARD FOR THE BOARD
24	TO MAKE ITS JUDGMENT ON WHETHER OR NOT IT FELT THE
25	REVIEW WAS SUFFICIENT OR WHETHER OR NOT A REREVIEW
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1	IS REQUIRED.
2	CHAIRMAN THOMAS: OTHER QUESTIONS?
3	DR. PRIETO: I GUESS I'M WONDERING ABOUT
4	THE PREVIOUS SECTION THAT YOU MENTIONED. ARE WE
5	ANTICIPATING THAT DOING IT THE WAY, THE ONE, TWO,
6	THREE SCORING SYSTEM, WOULD REQUIRE US TO GO BACK
7	AND REREVIEW AND THAT THAT WOULD BE AN UNDUE BURDEN?
8	MR. HARRISON: NO. SO FIRST OF ALL, LET
9	ME JUST TAKE THE TRANSLATIONAL PROGRAM AS AN
10	EXAMPLE. THE TRANSLATIONAL PROGRAM ANNOUNCEMENTS
11	WILL NOW BE OFFERED EVERY SIX MONTHS. SO IF AN
12	APPLICANT RECEIVES A SCORE OF 2, NOT RECOMMENDED FOR
13	FUNDING, THAT APPLICANT WOULD BE FREE TO TRY TO
14	IMPROVE THE APPLICATION AND RESUBMIT IT SIX MONTHS
15	LATER.
16	WHAT WE WERE CONCERNED ABOUT WITH RESPECT
17	TO THE IDEA OF APPLYING THE CLINICAL SYSTEM TO THE
18	TRANSLATIONAL AND DISCOVERY PROGRAMS, GIVEN THE
19	VOLUME OF APPLICATIONS, IS THAT IN ORDER TO
20	DETERMINE WHETHER AN APPLICATION WAS A THREE RATHER
21	THAN A TWO, MEANING IT'S SO FLAWED THAT IT SHOULDN'T
22	COME BACK, WOULD REQUIRE THE GWG TO SPEND AN AWFUL
23	LOT OF TIME ON APPLICATIONS THAT THE CONSENSUS IS
24	SHOULD NOT BE FUNDED AT THAT TIME IN THAT STATE
25	BECAUSE THE THREE, THE SAME APPLICATION OF THE SAME
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1	PROJECT SHOULD NOT BE RESUBMITTED.
2	SO WE WERE COGNIZANT OF THE ADDITIONAL
3	TIME AND BURDEN THAT THAT SYSTEM WOULD IMPOSE ON THE
4	GWG, AND THAT'S WHY WE PROPOSED TO HAVE ONLY TWO
5	TIERS, A TIER I AND A TIER II.
6	CHAIRMAN THOMAS: NO OTHER COMMENTS.
7	MEMBERS ON THE PHONE? MEMBERS OF THE PUBLIC?
8	HEARING NONE, ALL THOSE IN FAVOR PLEASE SAY AYE. WE
9	DON'T HAVE A MOTION.
10	MS. LANSING: MOVE IT.
11	MR. SHEEHY: SECOND.
12	CHAIRMAN THOMAS: MOVED BY MS. LANSING.
13	SECONDED BY MR. SHEEHY. ALL THOSE IN FAVOR PLEASE
14	SAY AYE. OPPOSED? ABSTENTIONS? MARIA, PLEASE
15	POLL.
16	MS. BONNEVILLE: LINDA BOXER.
17	DR. BOXER: YES.
18	MS. BONNEVILLE: KATHY LAPORTE.
19	MS. LAPORTE: YES.
20	MS. BONNEVILLE: BERT LUBIN.
21	DR. LUBIN: YES.
22	CHAIRMAN THOMAS: OKAY. WE HAVE APPROVAL.
23	THAT CONCLUDES THE JAMES AND SCOTT SHOW. WE'VE NOW
24	GOTTEN THROUGH ALL OF OUR AMENDMENTS. WE'RE GOING
25	TO MOVE ON TO DR. SAMBRANO TO TALK ABOUT A NUMBER OF
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1	UPDATES TO THE 2.0 CONCEPT PLANS.
2	DR. SAMBRANO: THANK YOU, MR. CHAIRMAN.
3	GOOD AFTERNOON, EVERYONE. I HOPE MY PRESENTATION IS
4	AS EXCITING AS THE PREVIOUS ONES. SO LET ME JUST
5	TELL YOU THAT ABOUT A YEAR AGO WE LAUNCHED CIRM 2.0
6	WITH THREE NEW PROGRAM ANNOUNCEMENTS TO OFFER
7	FUNDING OPPORTUNITIES FOR OUR CLINICAL STAGE
8	PROGRAMS. AND OVER THE LAST SEVERAL MONTHS, WE HAVE
9	LAUNCHED SEVERAL NEW PROGRAM ANNOUNCEMENTS INCLUDING
10	OUR TRANSLATIONAL PROGRAM AND TWO OF OUR DISCOVERY
11	PROGRAM ANNOUNCEMENTS.
12	NOW, UNLIKE OUR PREVIOUS RFA'S, THESE CIRM
13	2.0 PROGRAM ANNOUNCEMENTS ARE ONGOING SOLICITATIONS
14	THAT REMAIN OPEN. AND OUR EXPERIENCE WITH EACH
15	ITERATION OF THESE WILL HIGHLIGHT IMPROVEMENTS AND
16	ADJUSTMENTS THAT CAN BE MADE TO IMPROVE THEM AND TO
17	ACHIEVE OUR INTENDED RESULTS. SO WE WILL BE
18	BRINGING TO YOU ON OCCASION SUGGESTIONS FOR
19	ADJUSTMENTS TO THESE CONCEPTS.
20	AND SO TODAY WE'RE GOING TO BRING FOR YOUR
21	CONSIDERATION FOUR PROPOSED CHANGES THAT WOULD BE
22	APPLIED TO ALL CIRM 2.0 CONCEPT PLANS, MEANING THOSE
23	THAT WE'VE ALREADY ISSUED, THE CLINICAL PROGRAM, AND
24	THOSE THAT ARE YET TO COME EXCEPT WHERE NOTED
25	OTHERWISE. SO THERE'S ONE THAT APPLIES ONLY TO THE
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1	CLINICAL PROGRAM, AND I'LL HIGHLIGHT THAT.
2	SO LET ME GO THROUGH EACH ONE OF THEM SO
3	THAT WE CAN PROVIDE A LITTLE CONTEXT AND
4	UNDERSTANDING OF THEM. THE FIRST ONE, THE FIRST
5	PROPOSAL IS TO MAKE AVAILABLE TO THE GRANTS WORKING
6	GROUP OBJECTIVE INFORMATION FROM PREVIOUS CIRM
7	AWARDS FOR CONSIDERATION IN THE EVALUATION OF A NEW
8	AND RELATED APPLICATION.
9	SO OVER THE COURSE OF SUPPORTING AND
10	MANAGING AWARDS OR FUNDED PROJECTS, CIRM HAS
11	ACQUIRED EXPERIENCE AND INFORMATION ABOUT EACH OF
12	THOSE PROJECTS AND ABOUT THE APPLICANTS WHO MAY NOW
13	BE INTERESTED IN APPLYING FOR ADDITIONAL FUNDING IN
14	MANY CASES TO CONTINUE A PROJECT AND MOVE IT
15	FORWARD. AND QUITE OFTEN INFORMATION FROM THE
16	PREVIOUS AWARDS IS KEY TO PROPERLY ASSESSING THE
17	MERIT OF ANY NEW PROPOSAL THAT COMES OUR WAY.
18	SO, FOR EXAMPLE, DATA THAT WAS OBTAINED
19	FROM A PREVIOUS AWARD THAT HELPS ESTABLISH A PROOF
20	OF CONCEPT, THE ACHIEVEMENT OF A SPECIFIC MILESTONE,
21	OR OTHER OUTCOMES FROM THAT AWARD MAY HELP REVIEWERS
22	JUDGE THE RATIONALE AND FEASIBILITY THAT WE'RE
23	ASKING THEM TO DO FOR US.
24	SIMILARLY, INFORMATION ON THE USE OR THE
25	SUCCESS OR FAILURE OF THE SAME PRODUCT FROM THE

1	PREVIOUS AWARD MAY ALSO HELP ADDRESS QUESTIONS OR
2	CONCERNS OR HELP ADDRESS OR SATISFY A REVIEWER
3	QUESTION OF A NEW PROPOSAL USING THAT SAME PRODUCT.
4	OVERALL THE GOAL HERE OF THIS PARTICULAR
5	AMENDMENT OR CHANGE TO THE CONCEPT PROPOSALS IS TO
6	PROVIDE REVIEWERS ALL OF THE INFORMATION THAT'S
7	AVAILABLE TO CIRM TO EFFECTIVELY EVALUATE A
8	PROPOSAL. IN DOING THIS, WE WANT TO DO IT IN A
9	CONSISTENT WAY, AND WE WANT TO PROVIDE INFORMATION
10	THAT IS OBJECTIVE. SO WE PROPOSE TO INCLUDE, IN
11	QUOTES, PAST CIRM AWARD INFORMATION, IF APPLICABLE,
12	AS A FORMAL COMPONENT THAT IS UTILIZED BY THE GRANTS
13	WORKING GROUP TO EVALUATE A NEW PROPOSAL. THAT'S
14	THE FIRST ONE.
15	THE SECOND ONE IS TO INCLUDE ACCURACY AND
16	COMPLETENESS OF AN APPLICATION WITHIN THE
17	ELIGIBILITY CRITERIA. SO WHEN SUBMITTING AN
18	APPLICATION, WE NORMALLY ASK AN APPLICANT TO ATTEST
19	TO THE ACCURACY AND COMPLETENESS OF THE INFORMATION
20	THAT THEY SUBMITTED TO US. HOWEVER, THERE ARE CASES
21	WHERE WE RECEIVE APPLICATIONS THAT MAY BE LESS THAN
22	ACCURATE AND LESS THAN COMPLETE. SO WE WANT TO
23	ESTABLISH A MECHANISM BY WHICH TO ADDRESS THIS.
24	SO, FOR EXAMPLE, PROPOSALS THAT WE GET MAY
25	HAVE SECTIONS THAT ARE NOT APPROPRIATELY COMPLETED
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1	OR MAY BE MISSING REQUESTED DOCUMENTS THAT ARE NOT
2	INCLUDED, SUCH AS LETTERS OF SUPPORT OR FDA
3	CORRESPONDENCE AND SUCH. AND SO THESE, AT LEAST ON
4	CIRM'S PART, ARE VERY OBJECTIVE DETERMINATIONS THAT
5	WE CAN MAKE. AND THE INTENT IS TO INFORM THE
6	APPLICANT THAT THEIR APPLICATION CANNOT BE REVIEWED
7	AND IS, THEREFORE, INELIGIBLE DUE TO THIS
8	DEFICIENCY.
9	WE WANT TO PROVIDE THE APPLICANT AN
10	OPPORTUNITY TO REMEDY THE ISSUE. SO FOR A SIMPLE,
11	MINOR OMISSION, WE MAY ALLOW THE APPLICANT A VERY
12	SHORT TIME WINDOW TO PROVIDE THE MISSING INFORMATION
13	IF IT'S READILY AVAILABLE. SO THEY FORGOT TO PUT IN
14	THEIR FDA CORRESPONDENCE. WE CAN ASK THEM TO DO
15	THAT WITHIN A 24-HOUR PERIOD AND WE MOVE FORWARD
16	WITH THEIR APPLICATION. IN OTHER CASES THE
17	APPLICANT WILL BE ASKED TO RESUBMIT AT THE NEXT
18	DEADLINE FOR THE CLINICAL PROGRAM THAT OCCURS EVERY
19	MONTH. THAT'S USUALLY NOT AN ISSUE, BUT CERTAINLY
20	IF IT'S SUBSTANTIAL INFORMATION, WE WOULD JUST DEEM
21	IT INELIGIBLE AND ASK THEM TO RESUBMIT AT THE NEXT
22	OPPORTUNITY.
23	THE SAME WOULD APPLY IF WE IDENTIFY
24	INFORMATION OR STATEMENTS THAT ARE NOT ACCURATE.
25	SO, FOR EXAMPLE, AN APPLICANT PROVIDES AN ERRONEOUS

1	BUDGET, MAKES A CLAIM THAT WE KNOW TO BE INCORRECT,
2	OR WE FIND A MATERIAL OMISSION OF DATA. CIRM'S
3	DETERMINATION ON SOME OF THESE MAY BE MORE
4	SUBJECTIVE IN SOME OF THESE CASES.
5	SO IF THE APPLICANT'S ATTEMPT TO REMEDY
6	THE PROBLEM IS NOT SATISFACTORY TO CIRM, THE
7	APPLICATION WILL REMAIN INELIGIBLE, BUT THE
8	APPLICANT MAY APPEAL THAT ELIGIBILITY DETERMINATION
9	TO THE GRANTS WORKING GROUP. THIS IS CONSISTENT
10	WITH OTHER SUBJECTIVE CRITERIA THAT WE'VE
11	IMPLEMENTED IN THE CLINICAL PROGRAM WHERE THERE IS
12	AN OPPORTUNITY FOR THEM TO APPEAL THAT
13	DETERMINATION. WE ANTICIPATE THIS WOULD BE A VERY
14	RARE OCCURRENCE.
15	THE THIRD ONE IS RELATED TO THE IND
16	SPONSOR. WE WERE SILENT ON THIS WHEN WE IMPLEMENTED
17	THE CLINICAL PROGRAM. AND THIS APPLIES ONLY TO THE
18	CLINICAL PROGRAM; THAT IS, THE PROGRAM ANNOUNCEMENTS
19	THAT WE NOW REFER TO AS CLIN 1, CLIN 2, AND CLIN 3.
20	SO THESE ARE LATE STAGE PRECLINICAL THROUGH PHASE
21	III CLINICAL PROJECTS. AND THE REQUIREMENT HERE IS
22	THAT THE INTENDED OR CURRENT IND SPONSOR BE THE
23	APPLICANT ORGANIZATION IF IT IS AN
24	ORGANIZATION-SPONSORED IND OR THE PI IF IT'S AN
25	INVESTIGATOR-INITIATED IND FOR APPLICATIONS IN THE
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1	CLINICAL PROGRAM.
2	SO THE REASON FOR THIS IS THAT IN MANY
3	CASES WE HAVE THE ACCOUNTABILITY TO THE
4	ACCOUNTABILITY THEY HAVE TO CIRM AND HAVING A DIRECT
5	RELATIONSHIP WITH THE IND HOLDER HELPS US HAVE
6	ACCESS TO FDA CORRESPONDENCE, INFORMATION THAT IS
7	OFTEN NEEDED IN ORDER TO ASSESS MILESTONES, ENSURE
8	THAT OUTCOMES ARE ACHIEVED, AND HAVE A PROPER
9	OVERSIGHT OF THE WORK.
10	OVER THE COURSE OF THE YEAR THAT THE
11	PROGRAM HAS LAUNCHED, THERE HAS NOT REALLY BEEN A
12	CIRCUMSTANCE THAT WE'VE SEEN WHERE THE APPLICANT
13	THAT IS MOST APPROPRIATE TO COME IN BEING THE IND
14	HOLDER. SO IT IS SOMETHING THAT WE ENCOURAGE, AND
15	SO WE FEEL THAT IT'S SOMETHING THAT WE WOULD LIKE TO
16	HAVE IN ALL CASES TO HELP US WITH OVERSIGHT.
17	THE FOURTH PROPOSED CHANGE IS TO ALLOW
18	INDIVIDUALS UNDER CONTRACT TO ACT ON BEHALF OF THE
19	APPLICANT ORGANIZATION TO QUALIFY AS A PI. SO
20	CURRENTLY TO BE ELIGIBLE TO SERVE AS A PI ON A CIRM
21	AWARD, THE INDIVIDUAL MUST ACCOMPLISH THREE THINGS.
22	ONE, BE AN EMPLOYEE OF THE APPLICANT ORGANIZATION,
23	THEY MUST COMMIT THE MINIMUM REQUIRED EFFORT FOR
24	THAT PROGRAM ANNOUNCEMENT, AND THEN BE AUTHORIZED BY
25	THE APPLICANT ORGANIZATION TO CONDUCT THE RESEARCH

1	AND ASSUME THE RESPONSIBILITIES OF THE PI.
2	THE REQUIREMENT FOR THE PI TO BE AN
3	EMPLOYEE, THE UNDERLYING REASON FOR THAT, IT WAS
4	ESTABLISHED TO ENSURE ACCOUNTABILITY OF THE PI TO
5	THE APPLICANT ORGANIZATION WHICH, IN TURN, IS
6	ACCOUNTABLE TO CIRM TO CONDUCT THE WORK. AND SO
7	THAT WAS A SIMPLE WAY OF ENSURING THAT WE HAD THAT
8	DIRECT RELATIONSHIP. HOWEVER, WE'VE ENCOUNTERED
9	SITUATIONS AND CIRCUMSTANCES WHERE THE MOST
10	APPROPRIATE PERSON TO LEAD A PROJECT IS NOT
11	NECESSARILY AN EMPLOYEE OF THE APPLICANT
12	ORGANIZATION, PARTICULARLY IF THE MAJORITY OF THE
13	WORK IS TO BE CONDUCTED OFFSITE AT A COLLABORATING
14	INSTITUTION.
15	SO AN EXAMPLE OF THIS MIGHT BE IF WE HAVE
16	AN IND HOLDER APPLICANT ORGANIZATION THAT'S OUTSIDE
17	OF CALIFORNIA, THEY INTEND TO DO WORK THAT IS IN
18	CALIFORNIA, AND WE CAN ONLY COVER COSTS THAT OCCUR
19	IN CALIFORNIA. THE BEST PERSON TO LEAD THAT STAGE
20	OF WORK MIGHT BE SOMEONE WHO IS LOCATED IN
21	CALIFORNIA, BUT MAY NOT NECESSARILY BE THE EMPLOYEE
22	OF THAT ORGANIZATION.
23	SO WHAT WE'RE PROPOSING HERE IS TO EXTEND
24	THE ELIGIBILITY AS A PI TO INDIVIDUALS WHO ARE
25	ACCOUNTABLE TO THE APPLICANT ORGANIZATION IN ONE OF
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1	TWO WAYS. EITHER THEY ARE EMPLOYED BY THE APPLICANT
2	ORGANIZATION OR THEY HAVE A CONTRACTUAL AGREEMENT TO
3	ACT AS AN AGENT ON BEHALF OF THE ORGANIZATION IN
4	THAT CAPACITY.
5	SO THOSE ARE THE FOUR PROPOSED CHANGES,
6	AND SO WE'RE REQUESTING APPROVAL OF THOSE
7	MODIFICATIONS. AND I AM HAPPY TO TAKE ANY
8	QUESTIONS.
9	CHAIRMAN THOMAS: QUESTIONS FOR DR.
10	SAMBRANO?
11	DR. MILLS: J.T., I WOULD JUST LIKE TO
12	I APPRECIATE THAT HIGH LEVEL REVIEW, GIL. I'D LIKE
13	TO PROVIDE THE BOARD WITH ACTUALLY A LOT MORE DETAIL
14	ON EACH OF THESE POINTS.
15	MR. TORRES: SECOND MR. MILLS' MOTION.
16	CHAIRMAN THOMAS: ANY COMMENTS BY MEMBERS
17	ON THE PHONE? ANYBODY WHO FAINTED FROM DR. MILLS'
18	COMMENT?
19	MR. SHEEHY: I'D LIKE TO MOVE ADOPTION.
20	DR. JUELSGAARD: SECOND.
21	CHAIRMAN THOMAS: MOVED BY MR. SHEEHY,
22	SECONDED BY MR. JUELSGAARD. ANY FURTHER DISCUSSION,
23	MEMBERS OF THE BOARD? MEMBERS OF THE PUBLIC?
24	ANYBODY ELSE? OKAY.
25	ALL THOSE IN FAVOR PLEASE SAY AYE.
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	BARRISTERS REPORTERS SERVICE
1	OPPOSED? ABSTENTIONS?
2	MS. BONNEVILLE: LINDA BOXER.
3	DR. BOXER: YES.
4	MS. BONNEVILLE: KATHY LAPORTE.
5	MS. LAPORTE: YES.
6	MS. BONNEVILLE: BERT LUBIN.
7	DR. LUBIN: YES.
8	CHAIRMAN THOMAS: MOTION APPROVED. IN ALL
9	SERIOUSNESS, THE REVIEWS UNDERTAKEN BY
10	MESSRS. HARRISON, TOCHER, AND SAMBRANO WERE NOT
11	EASY. THEY WERE CHARGED WITH A CRITICAL REVIEW OF
12	THEIR RESPECTIVE TOPICS AND UNDERSTAND IT'S NOT A
13	SIMPLE TASK AND CAME OUT WITH GOOD RECOMMENDATIONS
14	TO BRING THEIR END OF CIRM 2.0 UP TO THE REST OF
15	EVERYTHING. SO THANK YOU, GENTLEMEN, VERY MUCH.
16	OKAY. WE'RE GOING TO GO ON TO THE NEXT
17	ITEM, WHICH I GUESS IS ME. THE ITEM IS LISTED ON
18	THE AGENDA AS CONSIDERATION OF ACCEPTANCE OF DONOR
19	FUNDS.
20	I'M GOING TO TAKE A BIT OF LIBERTY HERE TO
21	EXPAND THAT TOPIC TO SAY SOME COMMENTS ABOUT
22	SUSTAINABILITY. WE, AS YOU KNOW, FOR A WHILE
23	BELIEVED, BASED ON OUR EXPENDITURE RATE, THAT WE
24	WOULD RUN OUT OF FUNDS IN 2017. WE'VE NOW KNOWN FOR
25	QUITE SOME TIME, BASED ON HISTORICAL DATA THE LAST
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1	TWO TO THREE YEARS, THAT WE BELIEVE OUR FUNDING WILL
2	LAST UNTIL 2020. AND THEN THE ANSWER TO THAT IS
3	WHAT THEN.
4	SO WE HAVE A FEW SLIDES HERE. YOU'VE SEEN
5	THE FIRST FEW OF THESE THINGS BEFORE. THE 2.75
6	BILLION FOR AWARDS, SECOND BUCKET, OUR
7	ADMINISTRATIVE BUCKET, WHICH IS A 180 MILLION. WE
8	ANTICIPATE WE HAVE AWARDS FOR FIVE YEARS THROUGH
9	2020, FUNDING UP TO EIGHT YEARS. WHAT THAT MEANS,
10	OF COURSE, IS WE HAVE MULTIYEAR AWARDS AND WE
11	ANTICIPATE MAKING AWARDS ALL THE WAY UP THROUGH
12	2020, SO THAT CARRIES US A BIT BEYOND. AT THIS
13	POINT WE HAVE 760-ISH MILLION UNCOMMITTED. YOU CAN
14	SEE UP THERE, SORT OF FOLLOW THAT ALONG, ALL OF
15	WHICH ENDS UP, AFTER YOU'VE TAKEN BACK MONEY FROM
16	PROJECTS THAT WE ANTICIPATE BEING TERMINATED ON AN
17	ANNUAL BASIS, SPEND ABOUT 170 MILLION PER YEAR,
18	WHICH IS HOW WE GET WITH WHAT WE HAVE LEFT THROUGH
19	то 2020.
20	THEN WE HAVE THE ADMINISTRATIVE BUCKET.
21	AT OUR CURRENT STAFFING LEVELS, WE FIGURE WE HAVE
22	FIVE MORE YEARS, ALSO THROUGH 2020, FOR OUR TEAM.
23	AND THAT OBVIOUSLY IS A CRITICAL COMPONENT OF THE
24	WHOLE PROGRAM. OUR CURRENT SPEND RATE IS 15 TO 16
25	MILLION PER YEAR, AS WE ARE EXPERTLY ADVISED OF BY
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1	CHILA AT VARIOUS MEETINGS.
2	SO WE GET TO 2020, AND THE QUESTION IS
3	WHAT ARE WE THINKING ABOUT? NOW, THIS IS FIVE YEARS
4	IN ADVANCE OF THINGS. SO THIS IS KIND OF A LITTLE
5	EARLY TO BE HAVING THIS, BUT IT'S NEVER TOO EARLY, I
6	THINK. IF YOU LOOK AT 2020, ONE OF THE BIG GOALS
7	AND A LOT OF THE THINGS WE'VE BEEN TALKING ABOUT,
8	WHETHER IT'S ATP3 OR PARTNERING OR WHATEVER, IS
9	TRYING TO GET OUR PROJECTS FAR ENOUGH ALONG THAT WE
10	WILL HAVE PROOF OF CONCEPT IN AS MANY OF THOSE
11	PROJECTS AS WE CAN WHICH IS WHAT'S GOING TO DRIVE
12	THE INTEREST OF THE SO-CALLED BIG MONEY AT THE END
13	OF THE TUNNEL, BE IT BIG PHARMA, BIG BIOTECH,
14	VENTURE, OR WHATEVER.
15	WE GET TO 2020, THE FACTS ARE THAT, THOUGH
16	WE'LL HAVE A VERY LARGE TRANSLATIONAL PORTFOLIO AT
17	THAT STAGE, THE VAST MAJORITY OF PROJECTS WILL NOT
18	HAVE HIT THE END OF PHASE II, AT WHICH TIME THEY
19	HAVE ESTABLISHED PROOF OF CONCEPT. SO IF WE DON'T
20	HAVE ADDITIONAL FUNDING AT THAT POINT, WE WILL HAVE
21	ONLY PARTIALLY MET OUR OBLIGATION TO DEVELOP
22	THERAPIES AND CURES FOR THE BENEFIT OF PATIENTS. SO
23	WE AS A GROUP REALLY NEED TO FIGURE OUT A WAY TO
24	SUSTAIN US BEYOND THAT.
25	SO LET'S TALK ABOUT THIS SLIDE. IT'S GOT
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1	THREE BULLET POINTS, BUT THERE ARE A NUMBER OF
2	SUBSET TOPICS HERE. SO AS WE ALL KNOW, CIRM WAS
3	FUNDED BY PROP 71 WHICH WAS AN INITIATIVE PUT ON THE
4	BALLOT THROUGH SIGNATURE GATHERING. THERE ARE TWO
5	WAYS TO GET ON THE BALLOT. THAT'S ONE WAY. THE
6	OTHER IS FOR THE LEGISLATURE TO PUT A BALLOT MEASURE
7	ON THE BALLOT ITSELF. AND THERE HAS CERTAINLY BEEN
8	IN THE PRESS DISCUSSION ABOUT THE POTENTIAL FOR A
9	BOND MEASURE TO PRODUCE INCREASED FUNDING IN 2018.
10	WE'VE ALL SEEN THAT IN THE PRESS.
11	I THINK THAT CERTAINLY GOING FORWARD ANY
12	SERIES OF OPTIONS THAT WE'RE LOOKING AT INVOLVING
13	ONGOING FUNDING HAS TO CONTEMPLATE THE INITIATIVE
14	PROCESS, WHETHER IT IS THROUGH SIGNATURE GATHERING
15	OR THROUGH SOMETHING DEVELOPED BY THE LEGISLATURE.
16	SO BOTH OF THOSE WILL REQUIRE CONSIDERATION AS WE GO
17	FORWARD HERE.
18	I HAVE TO POINT OUT, WE, AS A PUBLIC
19	AGENCY, CANNOT ENGAGE IN A POLITICAL CAMPAIGN.
20	SO
21	MR. SHEEHY: I GUESS WE'VE NOT REALLY
22	THOUGHT SERIOUSLY ABOUT GOING BACK TO THE
23	LEGISLATURE TO GET BACK ON THE BALLOT, BUT IT DOES
24	SEEM TO ME THAT THAT'S SOMETHING THAT WE SHOULD
25	ACTUALLY HAVE A WIDER DISCUSSION ABOUT AND A PLANNED

1	DISCUSSION ABOUT IT. IT SEEMS VERY RATIONAL TO ME
2	TO ENTERTAIN THAT. CERTAINLY I WOULDN'T BE LOOKING
3	AT THE NEXT ELECTION, BUT MAYBE IN 2018. WE'RE AN
4	EXISTING STATE AGENCY. I THINK TO HAVE SOMEBODY
5	ELSE CONTROL OUR FATE BY GOING OUT AND COLLECTING
6	SIGNATURES TO EXTEND US SEEMS UNUSUAL. AND I THINK
7	BEING ACCOUNTABLE TO THE LEGISLATURE BY FORMALLY
8	I DON'T KNOW IF IT'S APPROPRIATE FOR US TO
9	FORMALLY MAYBE MR. HARRISON HAS AN OPINION ON
10	THAT TO GO TO THE LEGISLATURE AND SAY IN 2018
11	THIS IS WHERE WE ARE, THESE ARE OUR SUCCESSES, THIS
12	IS WHAT WE'RE DOING, AND FORMALLY ASK IT ONLY
13	TAKES 50 PLUS ONE OF THE LEGISLATURE TO GET ON THE
14	BALLOT AND ACTUALLY BE A NORMAL STATE AGENCY IN THAT
15	WAY.
16	OTHERWISE, I THINK WE'RE LEAVING THE FATE
17	OF THE AGENCY UP TO OTHER ACTORS. AND I JUST THINK,
18	HAVING HEARD ALL WE'VE HEARD TODAY AND THE MOMENTUM
19	AND PROGRESS OF THIS AGENCY, I THINK IT'S VERY
20	REASONABLE TO TALK TO THE LEGISLATURE. I MEAN WHEN
21	I TALK TO MY LEGISLATORS IN SAN FRANCISCO, THEY'RE
22	ENTHUSIASTIC ABOUT THE WORK WE'RE DOING. THEY DON'T
23	ALL KNOW ABOUT IT. SO WHEN I TALK TO THEM ABOUT IT,
24	THEY'RE VERY POSITIVE. I DON'T KNOW WHAT OTHER
25	FOLKS' EXPERIENCE IS, BUT THAT SEEMS TO ME TO BE A
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1	VERY CREDIBLE PATH FOR US TO ENGAGE IN. AND IF IT'S
2	LEGAL, IT CERTAINLY IS DIFFERENT FROM HAVING SOME
3	OTHER ENTITY NOT CONNECTED WITH THE AGENCY GO OUT
4	AND COLLECT SIGNATURES TO EITHER KEEP US GOING OR TO
5	HAVE SOME OTHER THING HAPPEN WITH WHAT WE'VE
6	ACCOMPLISHED. THAT SEEMS WEIRD TO ME. SO
7	CHAIRMAN THOMAS: THANK YOU, MR. SHEEHY.
8	ANY OTHER COMMENTS?
9	MR. SHEEHY: THAT WASN'T A COMMENT. THAT
10	WAS A QUESTION. I HOPE THAT THE QUESTION WAS
11	SHOULD I BE MORE EXPLICIT? WHAT ARE THE PLANS OF
12	THE LEADERSHIP OF THE BOARD TO HAVE THIS DISCUSSION
13	ABOUT THE POSSIBILITY OF GOING TO THE LEGISLATURE?
14	AND IF IT IS INDEED MR. HARRISON, IS THAT
15	SOMETHING THAT WE CAN DO? CAN WE SPEAK WITH
16	LEGISLATORS AS A BOARD TO ASK THEM TO REVIEW OUR
17	AGENCY AND CONSIDER PUTTING A MEASURE ON TO GET
18	ADDITIONAL BOND FUNDS TO CONTINUE OUR EXISTENCE?
19	MR. HARRISON: YES. AS A STATE AGENCY,
20	JUST TO CLARIFY ONE POINT CHAIRMAN THOMAS MADE, THE
21	AGENCY DOES HAVE THE ABILITY TO INTERFACE WITH THE
22	LEGISLATURE. THE AGENCY ALSO HAS THE ABILITY, NOT
23	THAT I'M SUGGESTING IT, BUT JUST SO THAT THE LEGAL
24	LINES ARE CLEAR, UNDER CALIFORNIA LAW PUBLIC
25	AGENCIES CAN GO SO FAR AS TO USE PUBLIC FUNDS TO
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1	DRAFT A BALLOT MEASURE THAT CAN THEN BE CIRCULATED
2	BY THE VOTERS FOR SIGNATURE. PUBLIC FUNDS CAN'T BE
3	USED TO CIRCULATE IT, BUT THE DRAFTING IS A
4	PERMISSIBLE PUBLIC EXPENDITURE.
5	MR. SHEEHY: AND THE COST OF CIRCULATING
6	THE PETITION AND GETTING IT APPROVED IS ABOUT TWO
7	AND A HALF MILLION THESE DAYS; AM I CORRECT?
8	MR. HARRISON: THAT'S RIGHT.
9	MR. SHEEHY: AND THEN THERE WILL HAVE TO
10	BE A COMMITTEE ESTABLISHED INDEPENDENT OF THIS
11	AGENCY IN ORDER TO COLLECT FUNDS AND DO THAT PLUS IN
12	ORDER TO PAY FOR AN EVENTUAL CAMPAIGN.
13	MR. HARRISON: THAT'S TRUE. I DON'T KNOW
14	WHAT YOU WERE REFERRING TO IN TERMS OF ADDITIONAL
15	FUNDING BY THE LEGISLATURE. BUT IF THE LEGISLATURE
16	WERE TO DO IT BY BOND FUNDING, IT WOULD BE A BOND
17	MEASURE PLACED BY THE LEGISLATURE ON THE BALLOT.
18	MR. SHEEHY: I THINK THERE MIGHT BE A
19	DIFFERENT FLAVOR, AT LEAST IN PUBLIC PERCEPTION, IF
20	THIS WAS SOMETHING THAT WAS PUT ON THE BALLOT BY THE
21	LEGISLATURE TO EXTEND OUR AGENCY AFTER A FULL AND
22	PUBLIC DISCUSSION AT THE LEGISLATURE.
23	DR. FRIEDMAN: I THINK JEFF RAISES SOME
24	GOOD QUESTIONS, BUT MAY I JUST ASK YOU. GIVE US A
25	SENSE WHERE YOUR DISCUSSION IS GOING. IF YOU'RE
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1	RAISING IT TO SAY THIS IS SOMETHING WE SHOULD PAY
2	ATTENTION TO, WE CAN THEN TALK ABOUT THE VARIOUS
3	COMPONENTS THAT WE'D LIKE TO SEE EVALUATED IN
4	GREATER DETAIL, ON WHAT TIME FRAME. THAT'S A
5	PERFECTLY VALID DISCUSSION TO HAVE. IF YOU HAVE A
6	PROPOSAL FOR US, PARTICULARLY SOMETHING THAT YOU
7	WANT US TO VOTE ON TODAY, THAT'S A DIFFERENT SORT OF
8	THING. I'M JUST TRYING TO GET A SENSE OF WHERE THIS
9	IS GOING BECAUSE THESE ARE THREE VERY RICH,
10	COMPLICATED TOPICS, EACH OF WHICH COULD TAKE,
11	WITHOUT EXAGGERATION, A FULL BOARD MEETING. SO GIVE
12	US YOUR GUIDANCE ABOUT HOW YOU'D LIKE TO SEE THIS
13	DISCUSSION PROGRESS.
14	CHAIRMAN THOMAS: SO WHAT I'M TRYING TO DO
15	HERE IS TO LAY OUT SORT OF A SERIES OF OPTIONS FOR
16	US TO CONSIDER. I DON'T WANT ANYBODY TO VOTE ON
17	ANYTHING BECAUSE EACH OF THESE HAVE THEIR
18	COMPLEXITIES, SOME OF WHICH ARE SUITABLE FOR VOTING
19	ON, SOME OF WHICH AREN'T. I JUST WANT THE BOARD TO
20	UNDERSTAND KIND OF MY PERSPECTIVE OF THE PLAYING
21	FIELD OF OPTIONS THAT WE SHOULD BE LOOKING AT TO
22	DEVELOP AS WE GO ALONG.
23	DR. FRIEDMAN: I THINK THAT'S ENORMOUSLY
24	HELPFUL. THANK YOU FOR DOING THAT. MY OWN
25	SUGGESTION IS I GUESS WHAT I WOULD SUGGEST IS
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1	YOUR INPUT AND YOUR THOUGHTS ARE GOING TO BE VERY
2	IMPORTANT, VERY FORMATIVE IN THIS. BUT THAT MY OWN
3	PREFERENCE WOULD BE TO LAY OUT A PLAN FOR HOW WE
4	WANT TO LOOK AT THESE. IF YOU WANT TO SET UP
5	SUBCOMMITTEES, THAT'S FINE. IF YOU WANT TO DRAW UP
6	DRAFT PROS AND CONS FOR EACH ONE THAT WE CAN
7	CONSIDER AHEAD OF TIME. OTHERWISE, I'M AFRAID THAT
8	THE DISCUSSION MAY BE A LITTLE MEANDERING IS MY ONLY
9	CONCERN.
10	CHAIRMAN THOMAS: OKAY. SO, MR. SHEEHY,
11	GETTING BACK TO YOUR QUESTION, I THINK THE ISSUE OF
12	CONTEMPLATING THIS AS A PARTICULAR OPTION IS
13	SOMETHING THAT WOULD REQUIRE FURTHER DISCUSSION. WE
14	HAVE A LEGISLATIVE SUBCOMMITTEE THAT'S SET UP, IF WE
15	SO CHOOSE TO TAKE IT TO THAT TO HAVE A DISCUSSION
16	THERE, WHICH WOULD SEEM TO BE THE APPROPRIATE FORUM
17	IN TERMS OF TRYING TO LAY OUT WHAT THE STEPS WOULD
18	BE IF WE WERE TO DO THIS, THE ADVISABILITY OF IT,
19	ALL THAT SORT OF THING. SO I THINK THAT THAT'S
20	SOMETHING WE COULD ABSOLUTELY DO IF THAT WERE THE
21	INTEREST OF THE BOARD HERE.
22	MR. SHEEHY: I KIND OF AGREE WITH DR.
23	FRIEDMAN'S SUGGESTION. WE HAVE THREE OPTIONS THAT
24	ARE LISTED HERE. MAYBE AN EXHAUSTIVE DISCUSSION
25	STARTING AT A SUBCOMMITTEE LEVEL AND THEN BRINGING

1	THAT INPUT FOR AN AGENDA ITEM AT THE BOARD SOMETIME
2	IN THE NEAR FUTURE WOULD SEEM A VERY, VERY
3	REASONABLE OPTION FOR US TO TAKE TO MOVE FORWARD. I
4	ACTUALLY LIKE THAT IDEA BECAUSE I DO THINK THIS IS
5	SOMETHING WE SHOULD DISCUSS AND WE SHOULD TAKE UP
6	AND NOT KIND OF LEAVE TO THE FUTURE. A NICE,
7	DETAILED DISCUSSION WITH THESE THREE OPTIONS WOULD
8	BE GREAT.
9	CHAIRMAN THOMAS: BY THE WAY, I DON'T
10	THINK THE THREE OPTIONS NECESSARILY ARE MUTUALLY
11	EXCLUSIVE, AT LEAST SOME COMPONENT OF IT.
12	MR. TORRES: THE ISSUE IS COMPLEX. IT'S
13	POLITICALLY COMPLEX, IT'S SUBSTANTIVELY COMPLEX, AND
14	FROM A PUBLIC POLICY PERCEPTION, IT'S COMPLEX. THE
15	REASON MOST LEGISLATORS SEEKING REELECTION CHOOSE
16	NOT TO PAY THE FILING FEE TO COLLECT SIGNATURES IS
17	TO CREATE A CAMPAIGN THEMSELVES AND TO CREATE A
18	NETWORK. SO THE COLLECTION OF SIGNATURES HAS ALWAYS
19	BEEN VIEWED AS A WAY TO ESTABLISH AND DEVELOP A
20	NETWORK WHICH CAN THEN BE UTILIZED FOR A CAMPAIGN
21	MORE EFFECTIVELY THAN JUST BY PAYING YOUR FILING
22	FEE.
23	THE LEGISLATIVE PROCESS IS VERY COMPLEX
24	SIMPLY BECAUSE OF WHAT WE'VE EXPERIENCED IN DEALING
25	WITH THIS WHOLE ISSUE OF AN AUDIT OF US ON THE FETAL

1	TISSUE ISSUE. THAT WAS NOT A PLEASANT EXPERIENCE
2	FOR ME. AND IT WAS A VERY DIFFICULT ISSUE BECAUSE
3	YOU ALSO HAVE A LOT OF PEOPLE THAT DON'T BELIEVE IN
4	WHAT WE DO WHO ARE MEMBERS OF THE LEGISLATURE AND
5	COULD INVARIABLY IMPACT ANY PROPOSAL THAT WOULD COME
6	BEFORE THE LEGISLATURE.
7	I THINK WE NEED TO TAKE TIME TO EXAMINE
8	ALL THESE VARIABLES. WE'RE LOOKING AT 2018
9	BASICALLY AND A FILING DEADLINE, BUT ALSO KEEP IN
10	MIND IT'S NOT REALLY JAMES, CORRECT ME IF I'M
11	WRONG IT MAY NOT COST TWO AND A HALF OR THREE
12	MILLION SIMPLY BECAUSE THE NUMBER OF SIGNATURES
13	REQUIRED HAVE BEEN REDUCED BECAUSE OF THE VOTER
14	TURNOUT IN THE LAST GUBERNATORIAL ELECTION, WHICH IS
15	THE CRITERIA BY WHICH YOU DETERMINE THE NUMBER OF
16	SIGNATURES THAT YOU NEED.
17	SO I GUESS I'M AGREEING WITH JEFF AND WITH
18	MICHAEL, THAT WE NEED TO MOVE FORWARD ON AN
19	EXHAUSTIVE REVIEW, AND LOT OF IT MAY NOT BE AS EASY
20	AS WE WOULD THINK INITIALLY.
21	MS. LANSING: SO AS I'M LOOKING AT THIS, I
22	THINK I AGREE WITH, I GUESS IT WAS, MICHAEL AND JEFF
23	AND NOW ART, THAT WE NEED TO HAVE A REVIEW OF
24	THINGS. I THINK THE PRIVATE FUNDING AND THE
25	PARTICIPATE LEVERAGING CIRM'S FUNDS, THOSE ARE NOT
	4=0

1	MUTUALLY EXCLUSIVE. BUT I THINK IF YOU GO OUT TO
2	GET ADDITIONAL FUNDING AS ANOTHER BOND ISSUE, YOU
3	WILL FIND THAT PRIVATE FUNDERS OR PARTNERSHIPS WILL
4	STAY AWAY BECAUSE THEY'LL SAY, WELL, YOU'RE GOING
5	OUT FOR A PUBLIC THING. WHY SHOULD I GET INVOLVED?
6	OR THEY'LL GIVE YOU MONEY TO HELP YOU WITH YOUR
7	CAMPAIGN.
8	SO, J.T., IF YOU ESTABLISH A SUBCOMMITTEE
9	REALLY TO LOOK AT THE FEASIBILITY, WHAT ARE THE
10	CONSTITUENT'S FEELINGS, WHAT'S THE STATE'S FEELINGS
11	TO SEE HOW REALISTIC IT IS TO DO ANOTHER BALLOT
12	INITIATIVE. I DON'T KNOW THE ANSWER TO THAT. I DO
13	BELIEVE THAT THE LAST TWO, THAT THERE ARE A LOT OF
14	PEOPLE NOT A LOT, BUT SEVERAL OPPORTUNITIES WITH
15	INDIVIDUALS AND PARTNERSHIPS WHO REALLY BELIEVE IN
16	STEM CELL RESEARCH. AND I THINK THAT WE HAVE AN
17	OPPORTUNITY TO RAISE SOME MONEY FROM THAT.
18	CHAIRMAN THOMAS: OKAY.
19	MR. SHEEHY: I JUST WANT TO MAKE A COUPLE
20	OF COMMENTS ABOUT THE SIGNATURE ROUTE. ONE, THOSE
21	NAMES WE COLLECT, SIGNATURES ARE COLLECTED BY
22	PRIVATE ENTITIES BY PAID GATHERERS. SO THOSE REALLY
23	IN A WAY DON'T NECESSARILY CREATE A CAMPAIGN. AND
24	THE NAMES OF THOSE FOLKS ARE NOT AVAILABLE TO A
25	CAMPAIGN FOR FURTHER CONTACT AFTER THEY'RE

1	COLLECTED.
2	SECOND, IT IS BECOMING INCREASINGLY
3	CHALLENGING TO COLLECT SIGNATURES BECAUSE THE
4	LOCATIONS OF BIG BOX STORES, WHERE HISTORICALLY
5	PEOPLE HAVE COLLECTED A LOT OF SIGNATURES, ARE NO
6	LONGER ALLOWING SIGNATURE GATHERERS ON THE GROUND
7	AND COURTS HAVE SUPPORTED THAT. SO I DO THINK THE
8	WHOLE I HATE TO SAY I HAVE A BIT OF EXPERIENCE
9	WITH SIGNATURE GATHERING AT MULTIPLE LEVELS, AND I
10	DO THINK RELYING ON SIGNATURES MAY NOT BE A GOOD
11	OPTION.
12	CHAIRMAN THOMAS: OKAY. LET ME, IF I
13	MIGHT, KEEP GOING HERE. LET'S GO ON TO THE I
14	UNDERSTAND THE SUBCOMMITTEE IDEA HAS BEEN PROPOSED
15	AND SOUNDS LIKE IT'S SOMETHING OF INTEREST TO THE
16	BOARD.
17	LET'S KEEP GOING ON THE PRIVATE FUNDING.
18	THERE'S A LOT OF STUDIES THESE DAYS THAT SHOWS THAT
19	WITH THE PROBLEMS IN FUNDING AT NIH, WHETHER IT WAS
20	THROUGH SEQUESTRATION OR JUST GENERAL LACK OF WILL
21	BY CONGRESS TO PUT AS MUCH MONEY AS NEEDED INTO
22	DISEASE RESEARCH AND PREVENTION, THAT THE PRIVATE
23	SECTOR HAS INCREASINGLY STEPPED INTO THE VOID TO
24	FUND MEDICAL RESEARCH. YOU SEE STORIES ABOUT IT ALL
25	THE TIME. AND YOU HAVE MORE AND MORE
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1	PHILANTHROPISTS WHO ARE INTERESTED IN THAT, EITHER
2	ON A SPECIFIC BASIS WHERE THEY PUT MONEY INTO THE
3	FUNDING FOR A TARGETED DISEASE OR CONDITION, OR ON
4	AN UNRESTRICTED BASIS WHERE THEY ARE MORE INTERESTED
5	IN A TECHNOLOGY REGARDLESS OF THE PARTICULAR
6	APPLICATION OF THE DISEASE.
7	SO THERE ARE A NUMBER OF VERY
8	MISSION-DRIVEN PHILANTHROPISTS OUT THERE THAT ARE
9	TARGETING MEDICAL RESEARCH AS AN AREA THEY WANT TO
10	PUT THEIR MONEY INTO. AND INCREASINGLY,
11	INTERESTINGLY ENOUGH, WE'RE SEEING AN INTEREST IN
12	REGENERATIVE MEDICINE SPECIFICALLY. SO AN OPTION
13	THAT WE HAVE, AND IT'S SOMETHING THAT I AND AMY
14	LEWIS AND MARIA HAVE TALKED ABOUT QUITE A BIT AND
15	HAVE BEEN PURSUING, AN OPTION IS TO HAVE A
16	SIGNIFICANT PRIVATE FUNDING ELEMENT TO THIS, WHETHER
17	IT'S THROUGH LARGE GRANTS, WHETHER IT'S THROUGH
18	CHALLENGE GRANTS, WHATEVER FORM THAT MAY TAKE,
19	TOWARDS GENERATING ADDITIONAL RESEARCH DOLLARS
20	EITHER GOING STRAIGHT TO CIRM OR TO A FOUNDATION
21	THAT COULD BE ESTABLISHED AS AN ADJUNCT ENTITY THAT
22	WOULD PUT MONEY IN PARI-PASSU WITH CIRM OR WHATEVER.
23	SO THAT IS A STRATEGY THAT WE'VE BEEN
24	PURSUING NOW FOR A NUMBER OF MONTHS. AND AS YOU CAN
25	APPRECIATE, I CAN'T GET INTO DETAILS ON THAT BECAUSE
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1	IT INVOLVES A LOT OF DISCUSSIONS WITH PEOPLE THAT
2	AREN'T RIPE FOR PUBLIC DISCUSSION AT THIS POINT.
3	AT THE SAME TIME THERE ARE A NUMBER OF
4	PHILANTHROPISTS WHO ARE QUITE INTERESTED IN SPECIFIC
5	SUBJECT MATTER, AND WE'VE HAD A NUMBER OF
6	DISCUSSIONS WITH DIFFERENT SUCH FOLK ON THE TOPIC OF
7	TRYING TO GENERATE RESEARCH DOLLARS FOR SPECIFIC
8	EITHER PROJECTS THAT WE HAVE OR FAMILIES OF
9	PROJECTS. SO, FOR EXAMPLE, SOMEBODY WHO MIGHT BE
10	INTERESTED IN OUR OCULAR PORTFOLIO OR OUR HEART
11	PORTFOLIO OR WHATEVER. SO THAT AS AN ONGOING
12	STRATEGY, I THINK, IS A VERY GOOD ONE. WE'RE
13	INCREASING OUR VISIBILITY OUT THERE, NOT JUST IN
14	CALIFORNIA, BUT NATIONALLY. INDEED, WE'VE HAD
15	DISCUSSIONS WITH PHILANTHROPISTS AROUND THE COUNTRY.
16	AND I DO THINK THERE IS THE OPPORTUNITY HERE FOR A
17	BIG TICKET, SIGNIFICANT PRIVATE FUNDING COMPONENT OF
18	WHATEVER WE DO GOING FORWARD AS PART OF OUR PLAN.
19	ON THE ISSUE OF PARTNERSHIP AND LEVERAGING
20	CIRM FUNDS, HERE, WE ARE IN DISCUSSIONS WITH PHARMA,
21	WITH DISEASE FOUNDATIONS. THERE ARE VARIOUS MODELS
22	FLOATING AROUND OUT THERE ON THE DISEASE FOUNDATION
23	SIDE. EVERYBODY KNOWS THE CYSTIC FIBROSIS MODEL
24	WHERE FOUNDATIONS HAVE PUT MONEY INTO RESEARCH IN A
25	VERY ENTREPRENEURIAL FASHION AND HAVE GENERATED
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1	SIGNIFICANT RETURNS FOR THEIR PARTICULAR FOUNDATION.
2	THIS WOULD BE ON A SORT OF PROJECT OR
3	CATEGORY-SPECIFIC BASIS, BUT IT'S SOMETHING THAT WE
4	NEED TO DO BECAUSE IF WE HAPPEN TO GET TO THE END OF
5	THE ROAD AND WE HAVEN'T SUCCEEDED IN GENERATING
6	ADDITIONAL BOND FUNDS, WE HAVEN'T SUCCEEDED IN
7	RAISING PRIVATE FUNDS, AND WE AT THAT POINT WILL
8	WANT TO HAVE AS MANY OF OUR PROJECTS PARTNERED UP
9	WITH PHARMA, WITH FOUNDATIONS, OR WHATEVER GOING
10	BACK TO OUR STRATEGIC PLAN, WHICH HAS THE GOAL OF 50
11	PERCENT OF OUR CURRENTLY NONPARTNERED PROJECTS,
12	PARTNERED UP BY 2020. IT WOULD ALL BE PART AND
13	PARCEL OF THAT.
14	UNDER THIS CATEGORY ALSO WOULD BE ATP3,
15	WHICH, WITH CIRM PUTTING FUNDING INTO THAT
16	PROJECT AND THE DETAILS OF THE STRUCTURE OF THAT,
17	BY THE WAY, ARE GOING TO BE DISCUSSED IN JANUARY.
18	WE ARE DOING A GREAT DEAL OF ANALYSIS ON THAT
19	PARTICULAR TOPIC THERE IS THE POTENTIAL FOR A
20	RETURN TO CIRM OF FUNDS INVESTED AT SOME MULTIPLE
21	THAT WOULD GIVE US THE OPPORTUNITY TO HAVE FURTHER
22	FUNDING FOR RESEARCH.
23	SO I THINK WHEN YOU SORT OF LUMP ALL THIS
24	TOGETHER, WE HAVE A VARIETY OF OPTIONS THAT NEED TO
25	BE CONSIDERED HERE AND A SORT OF COHESIVE PLAN
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1	DEVELOPED, A LOT OF WHICH IS ALREADY IN PROGRESS,
2	AND SO WITH THAT, I'LL OPEN IT UP TO ADDITIONAL
3	COMMENTS.
4	NO ADDITIONAL COMMENTS? PEOPLE ARE TIRED.
5	OKAY. SO, AGAIN, THERE'S NO NEED FOR A MOTION HERE.
6	I THINK ON THE ISSUE OF A SUBCOMMITTEE, WE CAN
7	FIGURE OUT HOW WE WANT TO HANDLE THAT AND WHO MIGHT
8	BE ON, ETC. WELL, LET ME ASK. JEFF WAS
9	SPECIFICALLY TALKING ABOUT THE LEGISLATIVE ISSUES.
10	MICHAEL WAS TALKING ABOUT SOMETHING, I THINK, BEYOND
11	THAT. WHEN YOU WERE SAYING SUBCOMMITTEE, WE HAVE
12	THE LEGISLATIVE SUBCOMMITTEE FOR THE BOND MATTER.
13	MICHAEL, WERE YOU CONTEMPLATING ANOTHER SUBCOMMITTEE
14	HERE?
15	DR. FRIEDMAN: NOT NECESSARILY. I THINK
16	IT SHOULD BE A COMMITTEE OR IT SHOULD BE A GROUP
17	THAT LOOKS AT ALL THE OPTIONS. THE POINT YOU MADE
18	ABOUT THESE BEING NOT MUTUALLY EXCLUSIVE IS TRUE.
19	THERE MAY EVEN BE A FOURTH OPTION THAT WE HAVEN'T
20	CONSIDERED. AND SO PICK ANY COMMITTEE YOU WANT.
21	JUST HAVING IT THOUGHTFULLY LAID OUT AND THEN HAVING
22	ENOUGH TIME HERE FOR A GOOD, INTENSE DISCUSSION WAS
23	ALL I WAS SUGGESTING.
24	CHAIRMAN THOMAS: WELL, THE PURPOSE OF
25	THIS WAS JUST TO SORT OF PAINT A BROAD BRUSH OF
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1	OPTIONS JUST TO LET THE BOARD AND OTHERS KNOW THAT,
2	EVEN THOUGH WE'RE FIVE YEARS OUT, IS A TOP PRIORITY,
3	AND WE ARE ALREADY BUSILY AT WORK. MR. HARRISON, DO
4	YOU HAVE A COMMENT?
5	MR. HARRISON: JUST ONE. THERE IS ONE
6	ACTION WE WOULD LIKE THE BOARD TO TAKE, WHICH IS TO
7	ACCEPT
8	CHAIRMAN THOMAS: WAIT. WAIT. I
9	HAVEN'T GOTTEN TO THAT YET. I'M TRYING TO GET
10	COMMENTS ON THIS. WE HAVEN'T HIT THAT YET.
11	MS. LANSING: I THINK WE GOT THE BOARD TO
12	TAKE THE NEXT STEP AND OBVIOUSLY THE SOONER THE
13	BETTER. SO WHATEVER KIND OF COMMITTEE, HAVE THE
14	LEGISLATURE COMMITTEE, YOU WANT TO ADD PEOPLE TO IT,
15	YOU WANT TO NOT ADD PEOPLE TO IT, I THINK THAT
16	SHOULD BE UP TO YOU TO DECIDE.
17	CHAIRMAN THOMAS: EVERYBODY GOOD WITH
18	THAT? OKAY. THAT WILL BE THE ORDER OF THE DAY.
19	THANK YOU, SHERRY.
20	SO THE LAST ITEM HERE, SO WE GET TO THE
21	QUESTION THAT WE HOPEFULLY DON'T HAVE TO DEAL WITH,
22	WHICH IS WHAT HAPPENS IF WE DON'T GET ADDITIONAL
23	BOND FUNDS OR RAISE MONEY IN ANY OF THE OTHER
24	VARIOUS WAYS WE'VE DISCUSSED? WE HIT 2020, WE HAVE
25	NO ADDITIONAL FUNDING, WE STILL HAVE ABOUT FOUR

1	YEARS WORTH OF ADMINISTRATION TO GO TO ESSENTIALLY
2	WIND DOWN CIRM AND ADMINISTER THE GRANTS THAT HAVE
3	BEEN AWARDED ALL THE WAY UP UNTIL 2020 ITSELF.
4	SO WE'VE TAKEN A LOOK AT THIS, AND WE'VE
5	DETERMINED THAT WE NEED TO HAVE AMY, CORRECT ME
6	IF I'M WRONG ON THIS 30 MILLION. SO WE NEED TO
7	HAVE \$30 MILLION OF FUNDS IN HAND TO MEET THE
8	ADMINISTRATIVE GOAL. SO EVEN THOUGH WE'RE FIVE
9	YEARS OUT ON THIS AS WELL, WE HAVE BEEN BUSILY AT
10	WORK AND HAVE GONE OUT AND SORT OF DEVELOPED A LIST
11	OF FOLKS WE FEEL WE CAN TALK TO WHO WE MIGHT BE ABLE
12	TO GET A COMMITMENT FROM FOR ADMINISTRATIVE FUNDING.
13	AND THE COMMENT TO THEM WAS THAT THAT FUNDING WOULD
14	BE CONTINGENT ONLY IF AND WHEN WE DON'T HAVE THE
15	FUNDING AS OF 2020, WOULD THEIR PLEDGE KICK IN? IT
16	WOULD BE FOUR YEARS, ETC.
17	SO HAPPY TO REPORT TO THE BOARD THAT WE'VE
18	HAD TWO DISCUSSIONS, ONE WITH BILL BOWES, ONE WITH
19	PITCH JOHNSON. THOSE ARE TWO NAMES MOST OF YOU ARE
20	VERY FAMILIAR WITH, BOTH LONGTIME SUPPORTERS OF
21	CIRM, LONGTIME SUPPORTERS OF MEDICAL RESEARCH AND
22	REGENERATIVE MEDICINE. MR. BOWES HAS GENEROUSLY
23	AGREED TO A CONTINGENT GIFT OF \$5 MILLION.
24	MR. JOHNSON, ACTUALLY PITCH AND KATHY JOHNSON HAVE
25	COMMITTED TO A \$2 MILLION CONTINGENT GIFT. SO THE
	180

1	30 MILLION, WE ALREADY HAVE SEVEN TO KICK-START OUR
2	ADMINISTRATIVE CAMPAIGN.
3	WE ARE REQUIRED, ACCORDING TO MR.
4	HARRISON, TO VOTE TO ACCEPT DONOR FUNDING OF 3
5	MILLION OR GREATER. SO WE HAVE BEFORE US A MOTION
6	FOR APPROVAL TO ACCEPT THE BOWES GIFT OF 5 MILLION.
7	DO I HAVE ANY QUESTIONS ON THIS, OR WOULD SOMEBODY
8	LIKE TO MOVE THIS ITEM?
9	MR. SHEEHY: I'D LIKE TO MOVE. I WOULD
10	JUST LIKE TO SAY CONGRATULATIONS, CHAIRMAN THOMAS.
11	THIS IS TREMENDOUS. THANK YOU. IT'S WELL DESERVED.
12	MS. LANSING: SECOND.
13	(APPLAUSE.)
14	CHAIRMAN THOMAS: THANK YOU, MR. SHEEHY.
15	SECONDED BY MS. LANSING.
16	DR. JUELSGAARD: SO JUST A QUESTION. SO I
17	THOUGHT I UNDERSTOOD YOU TO SAY THAT, IN ESSENCE,
18	THESE ARE PLEDGES. THEY'RE NOT REALLY AT THIS POINT
19	DONATIONS AND THEY'RE DEPENDENT UPON DOWNSTREAM
20	THINGS HAPPENING. SO WE'RE ACCEPTING A GIFT WHICH
21	REALLY ISN'T BEING MADE AT THIS POINT. IS THAT
22	OKAY, MR. HARRISON?
23	MR. HARRISON: YES, THAT'S OKAY.
24	ESSENTIALLY THE BOARD IS APPROVING THE TERMS OF THE
25	PLEDGE WHICH ARE SET FORTH IN A GIFT COMMITMENT
	181
	

1	SIGNED BY MR. BOWES AND DR. MILLS.
2	DR. JUELSGAARD: I HAVEN'T LOOKED AT THE
3	FORM OF THE GIFT COMMITMENT, BUT MANY TIMES THE
4	PLEDGES ARE NOT BINDING. IN OTHER WORDS, YOU COULD
5	DECIDE TO WALK AWAY FROM IT AS THE DONOR DOWNSTREAM.
6	IS THAT THE CASE IN THESE?
7	MR. HARRISON: NO. THESE ARE BINDING.
8	THE TWO CONTINGENCIES, ONE IS THAT, IN THE CASE OF
9	COMMITMENT FROM THE BOWES FOUNDATION, THAT CIRM
10	RAISES AN ADDITIONAL 25 MILLION. AND SECOND, THAT
11	IT DOES NOT HAVE ACCESS TO ADDITIONAL STATE FUNDING
12	AS OF JUNE 30, 2019.
13	DR. JUELSGAARD: THANK YOU.
14	CHAIRMAN THOMAS: I WILL SAY THAT BOTH
15	MR. BOWES AND PITCH AND KATHY JOHNSON ARE VERY
16	ENTHUSIASTIC ABOUT THIS. THEY'RE HUGE BELIEVERS IN
17	WHAT WE'RE DOING, AND WE'RE VERY HAPPY TO HAVE THEM
18	CONTINUE THEIR INTEREST IN THIS FASHION. IT WAS
19	VERY, VERY GENEROUS OF THEM. REALLY APPRECIATE IT.
20	DR. MILLS: IF WE'RE IN THE COMMENT
21	PERIOD, MAY I?
22	CHAIRMAN THOMAS: YES.
23	DR. MILLS: THANK YOU. HOPEFULLY THEY'LL
24	BE A LITTLE MIXED IN, BUT IT WON'T ALL BE HUMOR. IT
25	NEVER IS.
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1	I'M A FIRST THINGS FIRST KIND OF GUY. AND
2	WE LAID OUT A STRATEGIC PLAN THAT IS PRETTY COOL.
3	YOU CAN READ ABOUT IT AT CIRM.CA.GOV IF YOU'RE
4	INTERESTED IN IT. AS PART OF THAT STRATEGIC PLAN
5	WAS THE SPENDING OF FUNDS, AND WE TALKED TODAY ABOUT
6	WE'RE A 150 WE HAVE 150 MILLION MORE TO SPEND
7	THAN WE ANTICIPATED, BUT THE LITTLE BUCKET, THE
8	ADMINISTRATIVE FUNDS, DIDN'T GET ANY BIGGER JUST
9	BECAUSE WE HAVE 150 MILLION MORE TO SPEND. AND WE
10	LISTED SPECIFICALLY IN THIS PLAN AS A RISK FACTOR,
11	AS WE GET CLOSER AND CLOSER TO THE END, WE MAY NOT
12	BE ABLE TO ATTRACT THE TEAM.
13	WE TALKED TO J.T. ABOUT THAT AND SAID,
14	J.T., YOU KNOW, \$30 MILLION WOULD GO A LONG WAY IN
15	MAKING SURE WE COULD PROPERLY ADMINISTER, NOT JUST
16	THE FUNDS WE HAVE, BUT POTENTIALLY THIS OTHER 150
17	MILLION AND KEEPING THE TEAM TOGETHER. AND J.T.
18	WENT, "GOT IT. I'LL GO OUT. I'LL DO IT." AND IN A
19	VERY SHORT PERIOD OF TIME HAS ALREADY BROUGHT BACK
20	SEVEN OF THOSE \$30 MILLION TO KEEP IT GOING.
21	SO YOU DO KNOW I LOVE THE GAME BALLS, AND
22	SO, J.T., FOR THE \$7 MILLION, I'D LIKE TO GIVE YOU A
23	GAME BALL, BUDDY.
24	(APPLAUSE.)
25	CHAIRMAN THOMAS: THANK YOU, DR. MILLS.
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1	IS THIS A DODGER OR A GIANT BALL? THAT'S WHAT I
2	WANT TO KNOW. SO THANK YOU. I APPRECIATE THAT.
3	AND THANK YOU, EVERYBODY.
4	SO DO WE HAVE A MOTION? WE DO, YES.
5	WE'VE HAD SO MANY MOTIONS TODAY, I CAN'T REMEMBER
6	WHAT'S MOTIONED AND NOT MOTIONED. SO THAT
7	REQUIRES DO WE HAVE COMMENTS FROM MEMBERS OF THE
8	PUBLIC? HEARING NONE, THIS IS A VOICE VOTE. ALL
9	THOSE IN FAVOR PLEASE SAY AYE. OPPOSED?
10	ABSTENTIONS? MARIA.
11	MS. BONNEVILLE: KATHY LAPORTE. LINDA
12	BOXER.
13	DR. BOXER: YES.
14	MS. BONNEVILLE: DIANE, ARE YOU BACK?
15	DIANE WINOKUR.
16	MS. WINOKUR: YES. THANK YOU.
17	CHAIRMAN THOMAS: OKAY. THANK YOU. THAT
18	CONCLUDES THE ACTION ITEMS. WE'RE NOW
19	MS. LANSING: I JUST HAVE A QUESTION
20	BECAUSE I'M THRILLED THAT YOU RAISED THIS MONEY.
21	BECAUSE OF THE NATURE OF US, I'M NOT ASKING LIKE FOR
22	A DEBATE ABOUT THIS SO MUCH AS FOR THE SUBCOMMITTEE,
23	WHO'S GOING TO DO THIS. IF WE CHOOSE TO CONTINUE
24	RAISING MONEY FROM DONORS, ARE WE ALLOWED TO WHAT I
25	WOULD REFER TO AS DONOR-DIRECTED FUNDS? IN OTHER
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1	WORDS, A DONOR CAN GIVE TO CIRM, THAT I KNOW, BUT
2	CAN A DONOR SPECIFICALLY SAY, I'M GOING TO GIVE \$100
3	MILLION AND I TRUST I'M MAKING THIS UP. I'M
4	REALLY LIKE DOING IT FOR A REASON. BECAUSE WHAT
5	WE'RE KNOWN FOR IS THE BEST SCIENCE IN THIS FIELD.
6	BUT I WANT I'M NOT GOING TO INTERFERE WITH THE
7	SCIENCE, I'M NOT GOING TO ASK YOU ANYTHING, BUT I
8	WANT MY GRANT TO BE USED OR MY GIFT TO BE USED FOR
9	ALZHEIMER'S. ARE WE ALLOWED TO DO THAT?
10	MR. HARRISON: I'M NOT SURE I UNDERSTAND
11	ENTIRELY.
12	MS. LANSING: TO HAVE A DONOR-DIRECTED
13	GRANT THAT ISN'T JUST TO CIRM, BUT IS TO A SPECIFIC
14	DISEASE.
15	MR. HARRISON: SURE.
16	MS. LANSING: OKAY. WHEN YOU HAVE YOUR
17	SUBCOMMITTEE, I'VE OFTEN FOUND THAT'S THE BEST
18	WAY JUST TO END WITH THIS, DONORS ARE VERY
19	SOPHISTICATED TODAY. AND GRATEFUL PATIENTS OR
20	WHATEVER YOU WANT TO SAY IS USUALLY THE WAY YOU
21	CAN RAISE MONEY. AND I BELIEVE THAT THE FOUNDATIONS
22	OUT THERE ARE LOOKING AT THE WORK THAT'S BEING DONE
23	BY THIS GROUP OF SCIENTISTS, AND WE CAN PROBABLY GET
24	A LOT OF FOUNDATION SUPPORT, BUT IT MIGHT BE DISEASE
25	SPECIFIC.
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1	CHAIRMAN THOMAS: ABSOLUTELY.
2	MS. MILLER: I WILL ADD THAT IT SHOULD BE.
3	AS SOMEONE WHO RUNS A FOUNDATION, WHO DOES GIVE TO
4	VERY SPECIFIC STUDIES, THAT IS THE WAY TO INSPIRE
5	PEOPLE TO DO IT, MAKING IT SPECIFIC.
6	CHAIRMAN THOMAS: THAT'S ABSOLUTELY TRUE.
7	THANK YOU. OTHER COMMENTS? OKAY.
8	MR. SHEEHY: IF WE GOT DONOR FUNDS, WOULD
9	WE STILL HAVE THAT SAME LET'S SAY WE'VE GOT I
10	THINK THIS IS A GREAT IDEA, DONOR-DIRECTED FUNDING.
11	WOULD WE STILL BE UNDER THE LIMITATION TO HAVE THE
12	SAME STATE-BASED LIMITATION TO DO THE WORK WITHIN
13	THE STATE? MAYBE THAT'S PART OF THE ITEM AS WE
14	START TO DISCUSS IT FOR THIS LARGER DISCUSSION.
15	WOULD THAT KIND OF GIVE US A LITTLE MORE FREEDOM ON
16	HOW AT LEAST THE DONOR-DIRECTED FUNDS ARE SPENT? NO
17	ANSWER NECESSARY NOW.
18	MR. HARRISON: WE CAN EXPLORE THAT IN
19	GREATER DETAIL.
20	CHAIRMAN THOMAS: OTHER COMMENTS? OKAY.
21	THANK YOU, EVERYBODY.
22	NOW, WE'LL MOVE ON TO THE CLINICAL
23	ADVISORY PANEL UPDATE, DR. DOYLE.
24	DR. DOYLE: THANK YOU. THANK YOU, MR.
25	CHAIRMAN. THANK ALL OF YOU WHO HAVE REMAINED. I'LL
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	1

1	MAKE THIS VERY BRIEF.
2	I JUST WANT TO, FIRST OF ALL, REVIEW THE
3	RATIONALE FOR THE CAP PROGRAM WHICH WAS APPROVED BY
4	THE BOARD IN THE BEGINNING OF 2015 BEFORE I ARRIVED.
5	THE PURPOSE IS TO HELP ACCELERATE THE SUCCESSFUL
6	DEVELOPMENT OF THE PROGRAMS THAT WE'VE ALREADY
7	FUNDED. AND THE IDEA IS TO BRING ADDED EXPERTISE
8	AND RESOURCES TO THE FUNDED CLINICAL PROJECTS.
9	HOPEFULLY THE BENEFITS ARE THAT WE IMPROVE
10	THE LIKELIHOOD OF SUCCESS. WE ALSO KEEP REALLY
11	CLOSE TABS ON THE PROJECTS AND HOW THEY'RE DOING AND
12	HOW THEY'RE PROGRESSING OR WHAT CHALLENGES THEY'RE
13	FACING. IT ALSO GIVES US ANOTHER AVENUE TO ENGAGE
14	EXPERTS AS WELL AS PATIENT ADVOCATES IN THE AREAS
15	WHERE WE ARE FUNDING.
16	BASICALLY THE PHILOSOPHY OF THE CAPS IS A
17	STRATEGIC PARTNERSHIP. AND WHAT WE SAY WHEN WE MEET
18	WITH OUR GRANTEES IS WE'RE IN, WE'RE INVESTED,
19	CLEARLY WE'RE HERE TO HELP. AND THE IDEA, THE
20	IDEAL, I WOULD SAY, CAP MEETING IS ONE WHICH IS AN
21	OPEN, HONEST DISCUSSION. WE TRY TO IDENTIFY
22	OPPORTUNITIES TO IMPROVE PROJECTS, HELP THEM BE MORE
23	SUCCESSFUL, OR GO MORE QUICKLY. IMPORTANTLY, AGREE
24	ON WHAT THE OPPORTUNITIES AND CHALLENGES ARE AND
25	ACTUALLY WORK TOGETHER TO ADDRESS THEM. AND I THINK
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1	FUNDAMENTALLY WHAT THAT COMES DOWN TO IS AGREEING
2	WHAT SUCCESS LOOKS LIKE IN THE TRIAL OR IN THE
3	EXPERIMENT, RATHER, THAT WE'RE FUNDING.
4	THE WAY WE MEASURE SUCCESS IS GOOD
5	SCIENCE, A WELL-CONDUCTED TRIAL, GOOD
6	DECISION-MAKING WHICH IS DRIVEN BY SCIENCE AND BY
7	THE DATA. WHAT WE WANT FROM THE THINGS THAT WE
8	FUND, OF COURSE, WE WANT THINGS TO BE YES AND WE GET
9	THERAPIES TO PATIENTS, BUT A LOT OF TIMES A NEGATIVE
10	RESULT FROM A TRIAL CAN HELP DRIVE THE FIELD FORWARD
11	BY AVOIDING TIME SPENT DOWN AN AVENUE WHERE THINGS
12	AREN'T LIKELY TO RESULT IN A THERAPY. SO A CLEAR
13	ANSWER IS REALLY HOW WE MEASURE SUCCESS IN THE
14	CONDUCT OF THE TRIAL.
15	THE CAPS, EACH CAP HAS A STANDING GROUP OF
16	REPRESENTATIVES, INCLUDING SOME FOLKS FROM CIRM. WE
17	ALWAYS HAVE AN EXTERNAL SCIENTIFIC ADVISOR OR TWO,
18	DEPENDING ON THE NEEDS OF THE PROGRAM, AND WE ALWAYS
19	HAVE A PATIENT REPRESENTATIVE. I WOULD JUST SAY
20	THAT THE PATIENTS AND PATIENT REPRESENTATIVES BRING
21	AN IMPORTANT, UNIQUE, AND REALLY VALUABLE
22	PERSPECTIVE, AND I'M NOT JUST SAYING THAT. IN FACT,
23	THE PATIENT'S POINT OF VIEW CAN HELP TRIALS RUN MORE
24	EFFICIENTLY, CAN HELP ENROLLMENT, CAN HELP RAISE
25	CONCERNS OR OPPORTUNITIES THAT OTHER PEOPLE MIGHT
	100

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1	NOT HAVE THOUGHT OF.
2	WE ALSO CAN BRING IN AD HOC SPECIALISTS.
3	SOME OF OUR GROUPS EXPERIENCE PROBLEMS WITH CMC OR
4	MAY NEED SOME REGULATORY ADVICE OR PERHAPS NEED A
5	SECOND STATISTICAL OPINION ON POWERING THEIR TRIALS.
6	WE HAVE THE ABILITY TO BRING IN AD HOC SPECIALISTS
7	AS WELL, WHICH HAS BEEN VERY HELPFUL WITH SEVERAL OF
8	THE PROJECTS.
9	SO THE EXTERNAL MEMBERS ARE TYPICALLY, AS
10	I MENTIONED, DISEASE AREA EXPERTS, PEOPLE WITH
11	SPECIFIC EXPERTISE THAT WE AND THE GRANTEES HAVE
12	IDENTIFIED AS AN AREA OF NEED FOR THAT PARTICULAR
13	TRIAL, AND ALSO SOMETIMES A CLINICAL TRIAL
14	SPECIALIST. AS YOU KNOW, WE FUND COMPANIES, BUT WE
15	FUND A LOT OF ACADEMIC CENTERS. AND THIS ISN'T
16	NECESSARILY THEIR AREA OF EXPERTISE. THEY DON'T DO
17	HIGH VOLUME CLINICAL TRIALS A LOT OF THE TIME, AND
18	THEY NEED JUST BASICALLY THE NUTS AND BOLTS OF HOW
19	TO DO IT.
20	AND I MENTIONED ALREADY STATISTICIANS,
21	REGULATORY SPECIALISTS, TOXICOLOGISTS, WHATEVER WE
22	NEED, WHATEVER THEY NEED, AND WE AGREE THEY NEED TO
23	ACTUALLY MOVE THINGS FORWARD.
24	BASICALLY THE LOGISTICS RIGHT NOW, WE MEET
25	QUARTERLY. THE FIRST MEETING IS FACE TO FACE.
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1	AFTER THAT THEY OCCUR JUST BY TC, BUT WE CAN DO FACE
2	TO FACE AGAIN IF WE DEEM THAT NECESSARY. WE TRY TO,
3	IN THE SPIRIT OF TRYING TO BE EFFICIENT IN CIRM 2.0,
4	THE MEETINGS OCCUR THREE TO FOUR WEEKS AFTER A
5	PROGRESS REPORT IS DUE. AND SO WE USE THAT AS A
6	BRIEFING DOCUMENT TO KIND OF INFORM THE DISCUSSION
7	THAT WE'RE GOING TO HAVE.
8	JUST SO YOU KNOW WHERE WE ARE, THANKS TO
9	YOU GUYS, WE'VE HAD SIX FACE-TO-FACE CAP KICKOFF
10	MEETINGS WITH TRIALS RANGING FROM CALADRIUS
11	BIOSCIENCE, WHICH IS A TRIAL FOR METASTATIC
12	MELANOMA, TO DR. WANG, WHICH IS FOR A BLINDING EYE
13	DISEASE, RETINITIS PIGMENTOSA. DR. KOHN, WHO, OF
14	COURSE, WAS FEATURED IN THE VIDEO EARLIER ON, IS
15	DOING TWO REALLY EXCITING PROJECTS, ONE IN SICKLE
16	CELL DISEASE USING ZINC FINGER NUCLEASES TO ACTUALLY
17	PRODUCE BASICALLY TO GENE EDIT BETA GLOBIN AND TO
18	ALLOW PATIENTS WITH SICKLE CELL TO HAVE PRODUCTION
19	OF SOME NORMAL BLOOD CELLS, AT LEAST TO THE POINT
20	THAT THEY DON'T HAVE TO HAVE TRANSFUSIONS. THE
21	OTHER TRIAL OF DR. KOHN THAT WE'RE SPONSORING IS FOR
22	CHRONIC GRANULOMATOUS DISEASE, AN EXTREMELY RARE
23	DISEASE, IN THAT INSTANCE IS USING LENTIVIRAL VECTOR
24	GENE INSERTION.
25	I CAN SAY HONESTLY I DON'T THINK ANYBODY
	100

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1	ELSE WOULD DO THAT TRIAL. IT'S AN EXTREMELY RARE
2	DISEASE. NO COMPANY WOULD INVEST IN A DISEASE WHERE
3	I THINK THERE MAY BE 500 PEOPLE PER YEAR IN THE
4	UNITED STATES THAT HAVE IT. SO I'M REALLY PROUD OF
5	THAT EFFORT. AND, OF COURSE, I THINK THE PROMISE
6	THAT YOU SAW IN THE VIDEO, I THINK, HOPEFULLY CAN BE
7	REALIZED IN THESE TWO OTHER DISEASES WITH GENE
8	EDITING.
9	DR. ZAIA AT CITY OF HOPE IS DOING A TRIAL,
10	AGAIN, WITH SOME ZINC FINGER NUCLEASE INSERTIONS IN
11	CCR 5. THAT'S GOING RIGHT NOW WE DON'T HAVE ANY
12	ENROLLMENT, BUT WE'RE PUSHING AND TRYING TO DO WHAT
13	WE CAN TO HELP THEM OPTIMIZE ENROLLMENT.
14	AND FINALLY, VIACYTE, WHICH IS A COMPANY
15	THAT'S DOING WORK IN DIABETES, MUCH OF WHICH YOU'RE
16	FAMILIAR WITH. AND WE HAVE OTHER PROJECTS COMING UP
17	PROBABLY IN THE FIRST QUARTER. IMMUNOCELLULAR
18	THERAPEUTICS IS DOING A VACCINE TRIAL FOR
19	GLIOBLASTOMA. I THINK THE PROMISE OF CHECKPOINT
20	INHIBITORS PLUS CANCER VACCINES IS SOMETHING THAT
21	WE'RE PRETTY EXCITED ABOUT.
22	ASTERIAS, OF COURSE, IS DOING THE STEM
23	CELL TREATMENT FOR SPINAL CORD INJURY. DR. ABEDI IS
24	DOING AN HIV LYMPHOMA TRIAL, AND DR. HUMAYAN IS
25	ANOTHER PROJECT IN VISION WHERE THEY ACTUALLY ARE

1	USING A SCAFFOLD TO DELIVER RETINAL PIGMENT
2	EPITHELIAL CELLS.
3	SO THAT'S WHERE WE ARE. WE'VE HAD SIX
4	MEETINGS, AND I THINK THEY'VE GONE WELL. I WANTED
5	TO JUST SHARE A COUPLE OF COMMENTS FROM THE
6	GRANTEES. THEY FOUND IT VERY PRODUCTIVE. WE WANT
7	TO BE ALL IN WITH THEM AND HELPING THEM MOVE
8	FORWARD. I THINK WE LIKE I SAID, WE CAN BRING
9	ELEMENTS TO BEAR THAT THEY MIGHT NOT HAVE IN A SMALL
10	START-UP OR EVEN IN A SMALL ACADEMIC COMMUNITY TO
11	HELP MOVE THINGS FORWARD.
12	ACTUALLY SEVERAL OF THE PATIENT
13	REPRESENTATIVES HAD SOME REALLY GOOD FEEDBACK, WHICH
14	I WANTED TO SHARE WITH YOU. ONE WAS THE MOTHER OF A
15	PATIENT WHO WAS HEARING ABOUT THE CONSENTING PROCESS
16	AND SAID, "I'M AFRAID THAT THE WAY YOU'RE PRESENTING
17	THIS IS CREATING FALSE HOPE FOR PATIENTS. WHEN THIS
18	IS JUST A PROOF OF CONCEPT TRIAL, THEY'RE NOT
19	SUPPOSED TO BE BENEFITING." THAT WAS ACTUALLY
20	REALLY POWERFUL FOR THE GRANTEES TO HEAR THAT AND
21	MADE AN ADJUSTMENT TO THE CONSENTING PROCESS.
22	ANOTHER TIME WE HAD A PATIENT ADVOCATE
23	SAY, "THERE'S NO WAY I COULD DO ALL OF THIS IN A
24	DAY. YOUR SCHEDULE OF ASSESSMENTS IS NOT REALISTIC,
25	AND IT'S GOING TO WEAR ANYBODY OUT AND PEOPLE WON'T
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	± <i>36</i>

1	SIGN UP." THAT'S INCREDIBLY USEFUL INFORMATION FOR
2	SOMEBODY DOING A CLINICAL TRIAL.
3	AND ALSO ANOTHER REALLY GOOD POINT WAS THE
4	PATIENTS WHO FAIL SCREENING GETTING INTO A CLINICAL
5	TRIAL WILL GET CURIOUS. WHY DID PEOPLE DECIDE NOT
6	TO ENROLL IN THE TRIAL? WHAT WERE THEY WORRIED
7	ABOUT? WHAT WERE THEY FEARFUL OF? HOW COULD WE
8	HAVE MADE IT EASIER FOR THEM TO ENROLL? SO, AGAIN,
9	THE VALUE OF THE PATIENT AND PATIENT REPRESENTATIVE
10	FEEDBACK COMING TO BEAR RIGHT AWAY.
11	I THINK THE OPPORTUNITIES AND A LITTLE BIT
12	OF MY ASK FROM THE BOARD IS THAT WE WOULD LIKE TO
13	EXPAND OUR POOL OF EXPERT ADVISORS AND PATIENTS AND
14	ENGAGE MORE WITH PATIENT COMMUNITIES. I KNOW MANY
15	OF YOU HAVE CONNECTIONS TO THOSE ADVISORS. MANY OF
16	YOU ARE THOSE ADVISORS, BUT YOU MAY HAVE FACULTY OR
17	COLLEAGUES IN THE INSTITUTIONS THAT YOU REPRESENT.
18	PLEASE LET ME KNOW IF THESE ARE PEOPLE THAT YOU
19	THINK COULD HELP OUR PROCESS. WE'RE ALWAYS WORKING
20	TO IMPROVE THE VALUE OF THE INTERACTION WITH THE
21	GRANTEES. SO WE'RE OPEN FOR SUGGESTIONS.
22	I THINK THE OTHER IMPORTANT PIECE IS WE
23	ARE TRYING TO EDUCATE FOLKS AS WE GO ALONG HOW TO
24	ACTUALLY CONDUCT A CLINICAL TRIAL. THERE'S A RANGE
25	OF EXPERIENCE AROUND THIS IN OUR GRANTEES, AND I

1	THINK WE HAVE A TEAM THAT CAN OFFER A LOT OF
2	EXPERTISE.
3	THE OTHER THING I'LL SAY IN THE SPIRIT OF
4	CIRM 2.0, WE ARE DOING QUALTRICS QUESTIONNAIRES AND
5	METRICS AFTER EACH MEETING. I'LL HAVE THOSE FOR
6	YOU. WHEN WE HAVE A FEW MORE MEETINGS, WE'LL
7	PRESENT THAT AT OUR NEXT OPPORTUNITY TO TALK. SO
8	I'LL STOP THERE AND SEE IF YOU HAVE ANY QUESTIONS OR
9	COMMENTS.
10	CHAIRMAN THOMAS: QUESTIONS, COMMENTS FROM
11	MEMBERS OF THE BOARD? I THINK THE CAP STRUCTURE IS
12	OBVIOUSLY A CRITICAL COMPONENT OF OUR ONGOING
13	WORKING WITH ALL OF OUR GRANTEES IN THE CLINICAL
14	TRIAL PROCESS, AND IT'S FURTHER EVIDENCE OF THE VERY
15	PROACTIVE STANCE THAT WE'RE TAKING UNDER CIRM 2.0 TO
16	HELP IMPROVE AND TWEAK AND DO WHATEVER WE CAN TO
17	MAKE THE PROJECTS MORE SUCCESSFUL. SO I THINK WE
18	SHOULD ALL BE VERY HAPPY WITH THIS AS A CENTRAL
19	COMPONENT, AND MOST PARTICULARLY THE INVOLVEMENT OF
20	THE PATIENT IN THE CAP ITSELF TO GIVE REAL-TIME AND
21	REAL FEEDBACK ON WHATEVER IT IS THAT THE TRIAL IS
22	ABOUT AND HOW IT CAN BE IMPROVED TO HELP THEM, WHICH
23	IS, OF COURSE, WHAT IT'S ALL ABOUT. THANK YOU,
24	DR. DOYLE.
25	WE HAVE AN ITEM 20 ON HERE, WHICH IS
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1	ANOTHER ONE OF THE REALLY RIVETING ITEMS, SUMMARY OF
2	CONTRACTS. IT'S ACTUALLY NOT AN AGENDA ITEM. WE
3	HAVE IT ONLINE AND INVITE ANYBODY WHO HAS ANY
4	QUESTIONS TO GO LOOK AND TALK TO JAMES OR TO SCOTT,
5	BUT IS NOT SOMETHING WE NEED TO DISCUSS HERE.
6	SO WE'VE NOW GOTTEN TO THE END OF THE
7	AGENDA. WE'RE IN THE PUBLIC COMMENT, GENERAL PUBLIC
8	COMMENT PHASE. IS THERE ANY MEMBER OF THE PUBLIC
9	WHO WOULD LIKE TO SAY SOMETHING? MR. SENATOR.
10	MR. TORRES: I JUST WANT TO EXTEND MY
11	THANKS TO PRESIDENT MILLS AND THE STAFF. WHAT WE
12	ACHIEVED TODAY WAS ABSOLUTELY PHENOMENAL GIVEN WHAT
13	WE HAD ON THE AGENDA AND HANDLED THROUGH OUR CHAIR
14	IN SUCH AN EXPEDITIOUS AND EFFICIENT WAY. WE MADE
15	SOME SUBSTANTIAL DECISIONS TODAY WHICH I THINK BODE
16	WELL FOR THE FUTURE OF THIS AGENCY. I JUST WANT TO
17	SAY THANK YOU.
18	CHAIRMAN THOMAS: VERY WELL SAID, MR.
19	SENATOR. ON THAT VERY HIGH NOTE AND HAVING GOTTEN
20	THROUGH A MOST SUBSTANTIVE AGENDA, I WANT TO WISH
21	EVERYBODY HAPPY HOLIDAYS, AND WE WILL SEE YOU IN
22	PERSON NEXT MARCH. EVERYBODY HAVE A GREAT HOLIDAY
23	SEASON, AND THANK YOU VERY MUCH FOR COMING.
24	(APPLAUSE.)
25	(THE MEETING WAS THEN CONCLUDED AT 2:14 P.M.)
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2	
3	REPORTER'S CERTIFICATE
4	
5	
6	T RETUC DRAIN A CERTIFIED CHORTHAND REPORTED IN
7	I, BETH C. DRAIN, A CERTIFIED SHORTHAND REPORTER IN AND FOR THE STATE OF CALIFORNIA, HEREBY CERTIFY THAT THE FOREGOING TRANSCRIPT OF THE PROCEEDINGS BEFORE
8	THE FOREGOING TRANSCRIPT OF THE PROCEEDINGS BEFORE THE INDEPENDENT CITIZEN'S OVERSIGHT COMMITTEE OF THE CALIFORNIA INSTITUTE FOR REGENERATIVE MEDICINE IN
9	THE MATTER OF ITS REGULAR MEETING HELD AT THE LOCATION INDICATED BELOW
10	SHERATON GATEWAY LOS ANGELES HOTEL
11	6101 WEST CENTURY BOULEVARD LOS ANGELES, CALIFORNIA
12	ON DECEMBER 17, 2015
13	WAS HELD AS HEREIN APPEARS AND THAT THIS IS THE
14	ORIGINAL TRANSCRIPT THEREOF AND THAT THIS IS THE THAT APPEAR IN THIS TRANSCRIPT WERE REPORTED
15	STENOGRAPHICALLY BY ME AND TRANSCRIBED BY ME. I ALSO CERTIFY THAT THIS TRANSCRIPT IS A TRUE AND
16	ACCURATE RECORD OF THE PROCEEDING.
17	
18	
19	BETH C. DRAIN, CSR 7152 BARRISTERS' REPORTING SERVICE
20	160 S. OLD SPRINGS ROAD SUITE 270
21	ANAHEIM, CALIFORNIA (714) 444-4100
22	(714) 444-4100
23	
24	
25	
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