#### BEFORE THE

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

REGULAR MEETING

LOCATION: CLAREMONT HOTEL

41 TUNNEL ROAD

BERKELEY, CALIFORNIA

DATE: MARCH 26, 2015

9 A.M.

REPORTER: BETH C. DRAIN, CSR

CSR. NO. 7152

BRS FILE NO.: 97347

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1	BERKELEY, CALIFORNIA; THURSDAY, MARCH 26, 2015
2	9 A.M.
3	
4	CHAIRMAN THOMAS: IF EVERYBODY COULD TAKE
5	YOUR SEATS, PLEASE. LIVE FROM THE SONOMA ROOM AT
6	THE BEAUTIFUL CLAREMONT HOTEL. THE MARCH 26TH,
7	2015, MEETING OF THE ICOC IS CALLED TO ORDER.
8	MARIA, WILL YOU PLEASE LEAD US IN THE PLEDGE OF
9	ALLEGIANCE.
10	(THE PLEDGE OF ALLEGIANCE.)
11	CHAIRMAN THOMAS: MARIA, WILL YOU PLEASE
12	CALL THE ROLL.
13	MS. BONNEVILLE: LINDA BOXER.
14	DR. BOXER: HERE.
15	MS. BONNEVILLE: DAVID BRENNER. KEN
16	BURTIS.
17	DR. BURTIS: HERE.
18	MS. BONNEVILLE: ANNE-MARIE DULIEGE.
19	DR. DULIEGE: HERE.
20	MS. BONNEVILLE: ELIZABETH FINI.
21	DR. FINI: HERE.
22	MS. BONNEVILLE: MICHAEL FRIEDMAN. JUDY
23	GASSON.
24	DR. GASSON: HERE.
25	MS. BONNEVILLE: DAVID HIGGINS.
	$oldsymbol{a}$

1	DR. HIGGINS: HERE.
2	MS. BONNEVILLE: STEPHEN JUELSGAARD.
3	MR. JUELSGAARD: HERE.
4	MS. BONNEVILLE: SHERRY LANSING. KATHY
5	LAPORTE.
6	DR. LAPORTE: HERE.
7	MS. BONNEVILLE: JACOB LEVIN.
8	DR. LEVIN: HERE.
9	MS. BONNEVILLE: BERT LUBIN. SHLOMO
10	MELMED. LAUREN MILLER.
11	MS. MILLER: HERE.
12	MS. BONNEVILLE: JOE PANETTA. FRANCISCO
13	PRIETO.
14	DR. PRIETO: HERE.
15	MS. BONNEVILLE: ROBERT QUINT. AL
16	ROWLETT.
17	MR. ROWLETT: HERE.
18	MS. BONNEVILLE: JEFF SHEEHY.
19	MR. SHEEHY: HERE.
20	MS. BONNEVILLE: OSWALD STEWARD. JONATHAN
21	THOMAS.
22	CHAIRMAN THOMAS: HERE.
23	MS. BONNEVILLE: ART TORRES.
24	MR. TORRES: HERE.
25	MS. BONNEVILLE: KRISTINA VUORI.
	5
	<u> </u>

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

	BARRISTERS REFORTING SERVICE
1	DR. VUORI: HERE.
2	MS. BONNEVILLE: DONNA WESTON.
3	DR. WESTON: HERE.
4	MS. BONNEVILLE: DIANE WINOKUR.
5	MS. WINOKUR: HERE.
6	MS. BONNEVILLE: BRUCE WINTROUB.
7	DR. WINTROUB: HERE.
8	CHAIRMAN THOMAS: THANK YOU, MARIA. WE'LL
9	GO TO THE CHAIR'S REPORT. FIRST, AT THE TOP HERE
10	NEED TO MAKE A COUPLE OF ANNOUNCEMENTS. NO. 1 IS WE
11	HAVE A TIGHT SITUATION ON QUORUM TODAY. SO WE
12	REALLY NEED ALL MEMBERS TO BE PRESENT BOTH IN PERSON
13	AND THOSE ON THE PHONE UNTIL THE END OF THE MEETING,
14	WHICH I DON'T THINK IS GOING TO BE VERY LATE, BUT
15	NONETHELESS WE REALLY NEED EVERYBODY TO STICK
16	AROUND.
17	THE SECOND ANNOUNCEMENT IS DON REED IS
18	DOING A BOOK ABOUT CIRM AND PROP 71 AND STEM CELL
19	RESEARCH AND WOULD LOVE TO HAVE A PICTURE OF THE
20	BOARD AND OUR TEAM FOR THE BOOK. AND WE WOULD LIKE
21	TO HAVE THAT PICTURE TAKEN AT OUR BREAK, WHICH WILL
22	BE IN A BIT, BUT PLEASE EVERYBODY LISTEN UP FOR THAT
23	AND JOIN IN. IT'S A GREAT CHANCE TO GET EVERYBODY
24	DOWN IN HISTORY IN DON'S EXCELLENT WORK THAT HE'S
25	PUTTING TOGETHER.

6

1	NOW ON SPECIFICALLY TO THE CHAIRMAN'S
2	REPORT. THERE WAS A LOT OF ACTIVITY OVER THE LAST
3	COUPLE OF MONTHS. ONE THING THAT'S ALWAYS VERY
4	INTERESTING IS A NUMBER OF OUR INSTITUTIONS THAT WE
5	FUND HOST ANNUAL STEM CELL SYMPOSIA WHICH BRING
6	TOGETHER EXTRAORDINARILY TALENTED SPEAKERS BOTH FROM
7	THOSE INSTITUTIONS AND FROM OUTSIDE. AND I HAD THE
8	PRIVILEGE OF REPRESENTING CIRM AT THREE OF THOSE.
9	UCSD HAD THEIR FIRST ANNUAL WHICH WAS HELD
10	IN THE DUANE ROTH AUDITORIUM DOWN AT THE CONSORTIUM
11	IN LA JOLLA. UCLA HAD, I BELIEVE, ITS ELEVENTH
12	AND THAT IS RIGHT, JUDY?
13	DR. GASSON: YES.
14	CHAIRMAN THOMAS: AND CHILDREN'S
15	HOSPITAL LOS ANGELES HAD ITS ANNUAL AS WELL. HAD
16	THE OPPORTUNITY TO SAY A FEW WORDS AT THE FIRST
17	COUPLE AND TO GIVE THE KEYNOTE SPEECH AT LUNCH AT
18	THE CHILDREN'S HOSPITAL EVENT LAST WEEK.
19	IT'S ALWAYS VERY INTERESTING TO HEAR THE
20	PRESENTATIONS THERE. IT'S ALL ABOUT THE
21	CUTTING-EDGE SCIENCE THAT THE SPEAKERS ARE ENGAGED
22	IN, AND IT'S FURTHER EVIDENCE OF NOT ONLY THE GREAT
23	WORK THAT CIRM IS FUNDING, BUT THE GREAT WORK GOING
24	ON IN THE FIELD BOTH IN CALIFORNIA AND NATIONALLY.
25	ANOTHER EVENT THAT WAS VERY NICE, IN THE
	<del>-</del>

1	LAST TOOLS AND TECHNOLOGIES ROUND, THREE AWARDS WENT
2	TO USC. AND THERE WAS A VERY NICE EVENT HOSTED BY
3	DEAN PULIAFITO AND ANDY MCMAHON WHO RUNS THE STEM
4	CELL PROGRAM THERE WHERE THE THREE AWARDEES SPOKE
5	ABOUT THE PROJECTS THAT THEY RECEIVED FUNDING FOR TO
6	AN AUDIENCE OF MED STUDENTS AND DOCTORAL STUDENTS.
7	AND THAT WAS, AGAIN, A CHANCE TO HEAR WHAT GREAT
8	SCIENCE WE ARE FUNDING AND ENJOYED THAT.
9	WE HAD A GOOD DAY IN SACRAMENTO. SENATOR
10	TORRES TEED UP A FULL DAY. WE HAD OUR ANNUAL
11	APPEARANCE BEFORE THE STEM CELL FINANCE COMMITTEE
12	OVERSEEN BY THE STATE TREASURER WHERE THE BONDS THAT
13	ARE TO BE ISSUED SEMIANNUALLY ON BEHALF OF CIRM ARE
14	PROPERLY AUTHORIZED. SO SENATOR TORRES, MARIA, AND
15	I, REPRESENTING CIRM, WENT UP FOR THAT
16	AUTHORIZATION. THAT WAS FOLLOWED BY A NUMBER OF
17	MEETINGS WITH THE GOVERNOR'S OFFICE, WITH SOME KEY
18	MEMBERS OF THE LEGISLATURE, AND TOPPED OFF WITH
19	STATE TREASURER JOHN CHIANG AND SENIOR MEMBERS OF
20	HIS TEAM.
21	WE HAD SOME ACTIVITY IN THE LAST MONTH
22	WITH OUR COLLABORATIVE FUNDING PROGRAM. WE HAD A
23	DELEGATION FROM GERMANY. GEOFF LOMAX SET UP A
24	MEETING LAST WEEK WHERE WE MET TO TALK ABOUT CIRM
25	2.0 AND HOW THAT WILL FACTOR INTO THE COLLABORATIVE

8

FUNDING PROGRAM. HAD FIVE REPRESENTATIVES FROM
GERMANY, A VERY LIVELY DISCUSSION, AND I THINK THEY
CAME AWAY FROM THAT VERY IMPRESSED WITH 2.0 AND
LOOKING FORWARD TO CONTINUED RELATIONS WITH CIRM
GOING FORWARD.
WE'VE HAD A LOT OF WORK, WHICH WE'LL HEAR
A BIT ABOUT FROM SENATOR TORRES, HAVING TO DO WITH
OUR OFFICE SPACE SITUATION, TOURING VARIOUS PLACES,
NUMEROUS MEETINGS. THAT'S STILL A WORK IN PROGRESS.
IN THE CONTINUING EFFORT TO EXPOSE CIRM TO
MAJOR STAKEHOLDERS IN THE FIELD, I SET UP A LUNCH
WITH THE CEO OF AMGEN, BOB BRADWAY, AND SENIOR
MEMBERS OF HIS TEAM. AND RANDY AND MARIA, AMY
LEWIS, AND I WENT DOWN TO THEIR THOUSAND OAKS CAMPUS
AND HAD A VERY GOOD EXCHANGE WITH THEM ALL TOWARDS
LOOKING TO DEVELOP RELATIONSHIPS, IN THIS CASE,
BETWEEN AMGEN AND CERTAIN OF OUR GRANTEES AND
PROJECTS THAT THEY MAY BE WORKING ON THAT COULD BE
OF MUTUAL INTEREST.
I THINK, AS ALWAYS, HAD A NUMBER OF
MEETINGS WITH MEMBERS OF THE BOARD, VARIOUS LUNCHES
OVER THIS PERIOD, AND THEN A LOT OF WORK WHICH RANDY
HAS SPENT HUGE AMOUNTS OF TIME ON IN ADDING TO OUR
SENIOR TEAM HERE AT CIRM.
SO WITH THAT, THAT CONCLUDES THE CHAIR'S
9

1	REPORT. I WANT TO TURN IT OVER NOW TO DR. MILLS FOR
2	THE PRESIDENT'S REPORT.
3	DR. MILLS: THANK YOU, MEMBERS OF THE
4	BOARD, CHAIRMAN THOMAS. I WILL HOPEFULLY KEEP MY
5	REMARKS SOMEWHAT PITHY TODAY, BUT THERE ARE A COUPLE
6	OF IMPORTANT THINGS TO GO OVER.
7	JUST AN OVERVIEW OF THE FIVE THINGS THAT I
8	WANT TO MENTION. FIRST IS, AS ALWAYS, GO OVER OUR
9	MISSION THERE AND THE FIVE-PART TEST. I WANT TO
10	GIVE AN UPDATE NEXT ON THE ACTUAL WORKINGS OF CIRM
11	2.0 FOR THE CLINICAL STAGE AND ACTUALLY JUST WALK
12	THROUGH WHAT WE'VE EXPERIENCED SO FAR. I FIND IT
13	INTERESTING AND NEAT AND I HOPE YOU GUYS DO, AND
14	TALK A LITTLE BIT ABOUT SOME OF THE LESSONS WE'RE
15	LEARNING ON THAT. WE DEVELOPED WHAT WE CALL A
16	PUNCHLIST. SO AS WE GO THROUGH THIS PROCESS, WHAT
17	ARE THE THINGS THAT WE NEED TO BASICALLY IMPROVE
18	UPON FOR MAYBE WHAT WE WOULD CALL CIRM 2.1?
19	I'M GOING TO TELL YOU A LITTLE BIT ABOUT
20	WHERE WE'RE GOING WITH REGARDS TO OUR DISCOVERY AND
21	TRANSLATIONAL PORTIONS OF CIRM 2.0. I WANT TO MAKE
22	SOME BRIEF COMMENTS ON CONSIDERATION AND DISCUSSIONS
23	THAT WE'RE HAVING REGARDING MODIFYING THE GRANTS AND
24	LOANS PROGRAMS, AND THEN END WITH JUST A RUNDOWN ON
25	WHAT'S TO COME.
	10

1	SO, AS ALWAYS, OUR MISSION, ACCELERATING
2	STEM CELL TREATMENTS TO PATIENTS WITH UNMET MEDICAL
3	NEEDS. THE VALIDATION TESTING THAT WE'RE USING FOR
4	CIRM 2.0, WHEN WE CONSIDER THESE CENTERS, AROUND
5	ASKING OURSELVES FOUR QUESTIONS. ONE, IS THE ACTION
6	WE'RE TAKING GOING TO SPEED UP THE DEVELOPMENT OF A
7	STEM CELL TREATMENT? TWO, IS IT GOING TO INCREASE
8	THE LIKELIHOOD OF SUCCESS? THREE, IS IT ADDRESSING
9	AN ACTUAL UNMET MEDICAL NEED? AND THEN LASTLY, IS
10	IT EFFICIENT? AND IF WE CAN ANSWER YES TO THESE
11	QUESTIONS, THEN WE KNOW WE'RE GENERALLY DOING
12	SOMETHING THAT'S GOING TO BE POSITIVE. AND THAT'S
13	BEEN BASICALLY A STANDARD THAT WE'VE BEEN USING NOW
14	AS WE MOVE THROUGH CIRM AND LOOK NOW IN DISCOVERY
15	AND TRANSLATIONAL PROGRAMS AND CERTAINLY IN THE
16	CLINICAL PROGRAM ON HOW WE COULD MAKE CIRM BETTER
17	AND MORE RESPONSIVE TO ITS MISSION.
18	SO WITH THAT, LET ME JUMP INTO HOW CIRM
19	2.0 FOR THE CLINICAL STAGES IS GOING. SO RECALL
20	THAT WE LAUNCHED JANUARY 1ST, JANUARY 1ST OF THIS
21	YEAR, THE FIRST RFA'S UNDER CIRM 2.0 WHICH WERE FOR
22	THE CLINICAL SECTION. IT'S ALSO IMPORTANT TO
23	REMEMBER, AS WE'VE SAID ALL ALONG, WHILE THAT IS
24	WHERE WE STARTED, THAT IS NOT WHERE WE ARE ENDING.
25	WE ARE ALSO CURRENTLY AND ACTIVELY DEVELOPING THE
	11

1	2.0 COUNTERPARTS TO TRANSLATIONAL AND DISCOVERY
2	PHASE WORK AS WELL. BUT WE NEEDED TO START
3	SOMEWHERE, SO WE STARTED WITH CLINICAL.
4	WE DID SO BY ISSUING THESE THREE RFA'S,
5	AND WE USE, AS ANY GOOD GOVERNMENT AGENCY DOES, WE
6	USE CODE NAMES, AND SO 15-01, 15-02, AND 15-03, AS
7	THEY ARE CALLED. I WILL DECODE THEM FOR YOU, BUT
8	YOU CAN'T TELL ANYBODY. 15-01 STANDS FOR A PROGRAM
9	THAT IS ABOUT TO GET AN IND, SO APPROXIMATELY 24
	MONTHS AWAY FROM GETTING AN IND, BUT IT'S NOT YET IN
10	, '
11	A CLINICAL TRIAL. 15-02 IS FOR ANY PROGRAM'S
12	CLINICAL TRIAL, PHASE I, PHASE II, OR PHASE III.
13	AND THEN 15-03 BASICALLY TAKES THE CONCEPT THAT WE
14	HAD WITH THE ACCELERATED DEVELOPMENT PATHWAY AND
15	SAYS IF ANYONE HAS A 15-01 OR A 15-02 GRANT, WE CAN
16	BOLT ON 15-03 TO MAKE IT FASTER OR BETTER OR
17	STRONGER, KIND OF THE \$6-MILLION-MAN APPROACH.
18	SO THESE ARE THE THREE PROGRAMS THAT ARE
19	UP AND ACTIVE. AND THESE, AS I SAID, WE OPENED
20	THESE UP JANUARY 1ST. WE REVIEW THESE IN BATCHES ON
21	A MONTHLY BASIS, AND SO THE FIRST BATCH CLOSED
22	JANUARY 30TH AND WHATEVER THE LAST BUSINESS DAY OF
23	THE MONTH IN FEBRUARY IS WHEN THAT ONE CLOSED. I
24	CAN'T REMEMBER. I'LL TAKE YOU THROUGH HOW THEY'RE
25	ACTUALLY GOING.
	17

1	SO IN JANUARY WE HAD TWO SUBMISSIONS.
2	BOTH OF THEM WERE 15-02, WHICH IS THE CLINICAL TRIAL
3	STAGE. THE FIRST THING WE DO WITHIN THE FIRST 7
4	DAYS OF AN APPLICATION COMING IN IS WE DO A
5	SCREENING FOR IT. AND THAT SCREENING IS IS IT
6	BASICALLY COMPLIANT IN SCOPE WITH THE RFA THAT THEY
7	ARE APPLYING FOR? WITH REGARDS TO THE TWO THAT WE
8	RECEIVED IN JANUARY, ONE WAS COMPLIANT AND ONE WAS
9	NOT COMPLIANT. AND IT WASN'T ACTUALLY COMPLIANT FOR
10	A NUMBER OF REASONS. IT WILL BE INTERESTING. I'LL
11	TALK A LITTLE BIT ABOUT IT ON THE NEXT SLIDE. IT
12	WAS AN APPLICATION I THINK WE'VE ACTUALLY HAD THE
13	ABILITY TO HELP OUT AND WE WILL SEE AGAIN IN A
14	BETTER FORM.
15	AFTER THE SEVEN-DAY SCREENING, WE SEND IT
16	OFF TO BUDGET REVIEW. THESE ARE ALL TARGET DATES
17	YOU CAN SEE UNDERNEATH THE DIFFERENT STEPS. SO WE
18	SET A TARGET DATE OF 21 DAYS AFTER THE SUBMISSION
19	FOR THE BUDGET REVIEW TO BE COMPLETE. SO THE ONE
20	APPLICATION THAT WAS SENT TO BUDGET REVIEW PASSED
21	THAT BUDGET REVIEW WITHIN 21 DAYS AND THEN WITHIN 30
22	DAYS WAS SENT TO THE GRANTS WORKING GROUP FOR
23	REVIEW.
24	THAT'S WHERE WE ARE RIGHT NOW ON THAT.
25	THE GRANTS WORKING GROUP WILL REVIEW THAT FIRST
	12

1	APPLICATION ON MARCH 30TH, SO NEXT MONDAY. WE WILL
2	THEN HAVE A CIRM RECOMMENDATION BY APRIL 10TH, AND
3	IT WILL GO TO THE ICOC ON APRIL 23D, WITH A 120-DAY
4	CONTRACT SIGN DATE OBVIOUSLY STILL PENDING THE
5	OUTCOMES OF ALL OF THAT.
6	IN FEBRUARY WE HAD TWO MORE APPLICATIONS
7	COME ON. ONE OF THESE WAS FOR AN EARLIER, THE
8	PRECLINICAL STAGE AWARD. ONE OF THEM WAS FOR A
9	CLINICAL TRIAL STAGED AWARD. THE PRECLINICAL AWARD
10	PASSED THE SCREENING REVIEW. THE CLINICAL TRIAL
11	AWARD DID NOT PASS. THEY DIDN'T PASS THAT REVIEW
12	BECAUSE THERE WERE, AGAIN, A NUMBER OF PROBLEMS WITH
13	IT, BUT THE MOST OBVIOUS WAS IT HAD NOTHING TO DO
14	WITH STEM CELLS. SO THAT WAS AN ISSUE. IT THEN
15	WENT THE ONE THAT DID PASS WENT ON TO BUDGET
16	REVIEW. IT PASSED ITS BUDGET REVIEW AND WILL BE
17	SENT TO THE GRANTS WORKING GROUP BY APRIL 1ST.
18	AND SO SO FAR IT'S YOU CAN SEE THE
19	REMAINDER OF THE TARGET DATES FROM THERE ON OUT. SO
20	FAR MECHANISTICALLY IT'S WORKING THE WAY WE HOPED.
21	WE WERE ABLE TO, SOME OF IT BY BRUTE FORCE, BUT WE
22	WERE ABLE TO KEEP EVERYTHING ON SCHEDULE AND WITHIN
23	THE TIMELINES. BUT THERE ARE A COUPLE OF THINGS
24	THAT WE HAVE SEEN SO FAR THAT WE WANT TO CHANGE AND
25	IMPROVE, AND WE CALL THIS THE PUNCHLIST.
	14

1	THIS IS LIKE IF YOU BUILD A HOUSE, YOU GO
2	THROUGH THE HOUSE RIGHT AFTER YOU'VE DONE IT AND YOU
3	MAKE A LIST OF ALL THE THINGS THAT ARE NOT QUITE
4	RIGHT THAT YOU WANT TO MAKE BETTER. AND SO THIS IS
5	OUR 2.0 CLINICAL PUNCHLIST SO FAR. IT'S A LIST THAT
6	WILL CONTINUE TO GROW ALL FOR THE PURPOSE OF TRYING
7	TO MAKE THE PROGRAM BETTER.
8	THE FIRST ONE THAT WE'VE SEEN, JUST AS AN
9	OVERSIGHT, WE NEED TO CLARIFY THE TYPES OF THERAPIES
10	AND PARTICULARLY CELL THERAPIES THAT ARE ELIGIBLE.
11	WE SAID SMALL MOLECULES COULD BE USED TO ATTACK
12	CANCER STEM CELLS, BUT WE DIDN'T SAY THAT CELLS
13	COULD BE USED TO ATTACK CANCER STEM CELLS. THAT WAS
14	JUST AN OVERSIGHT ON OUR PART.
15	WE NEED TO CLARIFY SOME OF THE CO-FUNDING
16	DOCUMENTATION THAT'S REQUIRED. SO ONE OF THE GRANTS
17	THAT WAS RECEIVED IN JANUARY THAT ENDED UP FAILING
18	HAD AN ISSUE ASSOCIATED WITH THE CO-FUNDING
19	DOCUMENTATION JUST SIMPLY WASN'T REQUIRED, AND WE
20	NEED TO BE MORE CLEAR THAT THAT DOCUMENTATION IS
21	NEEDED ON THE FRONT END.
22	THERE WAS A LITTLE BIT OF CONFUSION ON
23	MATCHING, THAT WE'RE MATCHING TO THE TOTAL PROJECT
24	COST, NOT JUST THE DIRECT. AGAIN, I THINK THAT'S
25	SOMETHING WE CAN BE MORE CLEAR ABOUT.

WE ALSO NEED TO REQUEST MORE DETAILED
ENROLLMENT WHEN WE'RE TALKING ABOUT A CLINICAL
TRIAL. SO WHAT WE GET RIGHT NOW IS WE PROJECT THE
TRIAL WILL BE FULLY ENROLLED BY BLANK DATE. WHAT WE
REALLY NEED IS WHAT WE EXPECT THE ENROLLMENT OVER
THE TRIAL TO BE IN SOME MORE DETAILED FASHION THAN
WE CURRENTLY GET.
AND THEN LASTLY, I THINK THIS IS ONE OF
THE MORE INTERESTING THINGS, IS WE'RE IN THE PROCESS
OF ADDING AN IMMEDIATE WHAT WE CALL GRANT COUNSELING
STEP. AND THIS ONE WE DID NOT BY ANY PRESCRIBED
PROCEDURE, BUT ACTUALLY JUST IN TERMS OF TRYING TO
PROVIDE GOOD CUSTOMER SERVICE TO OUR APPLICANTS ON
THE FIRST APPLICATION THAT CAME IN WHERE WE HAD A
GRANT THAT ACTUALLY HAD A REASON FOR IT NOT BEING
WITHIN SCOPE. BESIDES THAT, WE HAD A GRANT THAT WAS
WRITTEN AND CONSTRUCTED IN A WAY TO WHERE IF IT
WOULD HAVE GONE TO THE GRANTS WORKING GROUP, IT
WOULD HAVE GOTTEN A VERY BAD SCORE. AND THAT HAD TO
DO WITH PARTS OF THE APPLICATION WERE JUST IN THE
WRONG SECTION. THEY DIDN'T PUT ANY CLINICAL DATA IN
THE CLINICAL DATA SECTION. AND IN THE CLINICAL
OPERATIONS PLAN THEY DIDN'T QUITE UNDERSTAND WHAT
THAT MEANS, AND SO THEY PUT THE WRONG DOCUMENTATION
INTO THERE.
16

1	AND BECKY AND GIL FIGURED THIS OUT IS WHAT
2	WE SHOULD PROBABLY DO IS JUST CALL THEM INFORMALLY
3	AND SAY, LOOK, IF THIS GOES TO THE GWG, THIS IS NOT
4	GOING TO GO WELL FOR YOU. YOU MAY WANT TO CONSIDER
5	PULLING YOUR GRANT, REVISING IT, GETTING THE STUFF
6	IN THE RIGHT PLACES. WE'LL TALK YOU THROUGH THAT.
7	CLEARLY YOU MISUNDERSTOOD SOME THINGS. WE'LL HELP
8	YOU WITH THAT. SO WE DON'T PUT THE GWG THROUGH AN
9	ENTIRE REVIEW PROCESS FOR AN APPLICATION THAT WE
10	COULD HAVE HELPED CLEAN UP ON THE FRONT END.
11	SO IT'S JUST ANOTHER THING WE'RE DOING IN
12	ORDER, AS I SAID, TO PROVIDE BETTER CUSTOMER SERVICE
13	TO OUR APPLICANTS BECAUSE IF OUR APPLICANTS AREN'T
14	SUCCESSFUL, OBVIOUSLY OUR PATIENTS AREN'T GOING TO
15	BENEFIT FROM THAT KIND OF WORK. SO THAT'S OUR
16	PUNCHLIST RIGHT NOW.
17	WE ARE KEEPING A VERY CRITICAL EYE ON WHAT
18	WE'RE DOING. I EXPECT THE LIST TO GROW, AND I WANT
19	THE LIST TO GROW BECAUSE I WANT THE PRODUCT TO GET
20	BETTER OBVIOUSLY.
21	NOW TURNING TO WHAT WE'RE DOING ON
22	DISCOVERY AND TRANSLATIONAL, SO THE EARLIER TWO
23	PHASES OF PROGRAMS THAT WE OFFER. HERE, I THINK,
24	WE'VE DONE A NICE JOB AND I SAY WE, I MEAN THE
25	DISCOVERY AND TRANSLATIONAL TEAM LED BY PAT OLSON
	17

1	IN REACHING OUT WITHIN THE ENTIRETY OF CIRM AND
2	SOLICITING IDEAS AND INFORMATION. WE'RE ALSO DOING
3	THE SAME THING EXTERNALLY. SO IF WE HAVEN'T GOTTEN
4	TO YOUR CITY YET, WE WILL BE GETTING TO YOUR CITY
5	AND SETTING UP MEETINGS WITH STAKEHOLDERS IN ORDER
6	TO REALLY, REALLY TRY TO FIND OUT WHAT IS DRIVING
7	AND WHAT MOTIVATES OUR STAKEHOLDERS TO PERFORM THE
8	BEST AND HOW WE CAN FIGURE OUT INNOVATIVE WAYS OF
9	ALIGNING THAT WITH WHAT CIRM NEEDS, WHICH I THINK
10	SOMETIMES ARE NOT ALWAYS ALIGNED. WE NEED TO JUST
11	TAKE AN HONEST LOOK AT THAT AND SEE IF WE CAN FIGURE
12	OUT A WAY WHERE WE BOTH GET WHAT WE NEED.
13	SOME OF THE ATTRIBUTES SO FAR THAT WE HAVE
14	IDENTIFIED AGAIN, THIS PROCESS IS STILL VERY
15	EARLY AND VERY FORMATIVE. BUT SOME OF THE
16	ATTRIBUTES THAT WE ALREADY DISCOVERED THAT WE LIKE,
17	FIRST AND FOREMOST, PREDICTABLE REQUESTS FOR
18	APPLICATIONS. WE THINK THIS IDEA OF MAKING SURE OUR
19	RESEARCHERS KNOW THAT THE PROGRAM IS OPEN AND WILL
20	REMAIN OPEN OR WILL OPEN AGAIN WITH SOME PREDICTABLE
21	FASHION LET'S THEM SUBMIT AN APPLICATION WHEN
22	THEY'RE READY TO SUBMIT THE APPLICATION AND NOT
23	SUBMIT AN APPLICATION THAT'S NOT READY FOR PRIME
24	TIME. SO WE WANT TO HAVE PREDICTABILITY THERE.
25	WE ALSO WANT OUR APPLICATION AND OUR CALLS

1	TO BE RESPONSIVE. THAT'S BOTH RESPONSIVE TO NEW
2	DEVELOPMENTS THAT MIGHT BE TAKING PLACE IN THE
3	FIELD, BUT ALSO RESPONSIVE TO NEEDS THAT CIRM MIGHT
4	HAVE OR CIRM AWARD HOLDERS MIGHT HAVE. AND SO, FOR
5	EXAMPLE, IF SOMETHING IN TRANSLATIONAL OR CLINICAL
6	RUNS INTO A PROBLEM THAT A DISCOVERY-TYPE PROGRAM
7	COULD ANSWER, WE WANT TO HAVE SOME FEEDBACK
8	MECHANISM TO WHERE WE CAN USE DISCOVERY IN ORDER TO
9	ADDRESS THAT.
10	ONE OF THE MORE EYE-OPENING THINGS THAT
11	WE'VE COME ACROSS IS THIS IDEA OF MULTIPLE OUTFLOW
12	PATHWAYS FROM DISCOVERY. SO ACTUALLY THE WAY WE
13	TALK ABOUT IT AND I THINK AS WE'VE TALKED ABOUT IT
14	MIGHT JUST BE WRONG IN THIS LINEAR DISCOVERY,
15	TRANSLATION, CLINICAL PARADIGM THAT WE HAD BECAUSE
16	THAT'S TRUE, BUT THAT'S ONLY TRUE IF THE ONLY THING
17	THAT CAN COME OUT OF DISCOVERY IS A DRUG THAT GOES
18	THROUGH TRANSLATION AND INTO THE CLINIC. BUT WE
19	HAVE OPPORTUNITIES TO HAVE THINGS COME OUT OF
20	DISCOVERY THAT ARE MEDICAL DEVICES WHICH ARE A
21	DIFFERENT PATHWAY. WE HAVE THINGS THAT COME OUT
22	THAT ARE MODELS WHICH IS A DIFFERENT PATHWAY, HIGH
23	THROUGHPUT DRUG SCREENING TECHNOLOGIES. WE HAVE
24	PROGRAMS IN PLACE TO HELP DISCOVER THOSE THINGS, BUT
25	WE DON'T HAVE PROGRAMS IN PLACE TO TAKE THOSE THINGS

1	TO THEIR ULTIMATE SUCCESSFUL OUTCOME. AND SO WE'RE
2	LOOKING AT ADDING MULTIPLE OUTFLOW PATHWAYS FROM
3	DISCOVERY, NOT JUST THOSE THAT DIRECTLY GO TO AND
4	ASSUME WHAT YOU ARE DOING IS TAKING A STEM CELL AND
5	PUTTING IT INTO A PATIENT.
6	ANOTHER CONCEPT IS LINKING GRANTS
7	SEAMLESSLY. SO IF WE HAVE A RESEARCHER THAT'S DOING
8	REALLY, REALLY GREAT WORK, WE NEED TO MAKE SURE WE
9	ARE NOT THE THING THAT SLOWS THEM DOWN. THEY NEED
10	TO BE ABLE TO CONTINUE DOING THAT GREAT WORK AS
11	SEAMLESSLY AS POSSIBLE.
12	THIS IS LINKED TO ANOTHER CONCEPT THAT
13	WE'VE DISCOVERED WHICH IS A LOT OF TIMES AND
14	OFTENTIMES THERE COMES A POINT IN RESEARCH WHERE THE
15	EARLY STAGE RESEARCHER DOES NOT HAVE THE INTEREST,
16	ABILITY, INCLINATION, TALENT, LAB, WHATEVER IT MIGHT
17	BE, JUST ABILITY TO DO THE NEXT STAGE OF RESEARCH.
18	AND SO WE NEED TO FIND WAYS TO WHERE WE CAN MOTIVATE
19	PEOPLE THAT HAVE SOMETHING IN AN EARLIER STAGE WHERE
20	THEY'VE TAKEN IT AS FAR AS THEY WANT TO TAKE IT OR
21	CAN TAKE IT TO BE ABLE TO PASS THE BATON TO THE NEXT
22	PERSON SO THEY CAN TAKE IT FORWARD QUICKLY AND WE
23	CAN GET FURTHER DOWN THE ROAD TOWARDS TREATING
24	PATIENTS.
25	WE TALKED ABOUT THIS AT THE LAST MEETING.

1	I WILL TELL YOU IT IS A CENTERPIECE OF WHAT'S GOING
2	TO COME OUT OF OUR DISCOVERY AND TRANSLATION SEE,
3	WE HAVE A CODE FOR THAT. IT'S CALLED D & T IS
4	SIGNIFICANT OPPORTUNITIES FOR TRAINING WHERE WE'RE
5	GOING TO BE VERY, VERY EXPLICIT IN THE DISCOVERY AND
6	TRANSLATIONAL CALLS THAT TRAINING IS GOING TO BE A
7	STATED OBJECTIVE OF THAT. AND SO WE'RE GOING TO DO
8	WHAT WE CAN, WHEN WE HAVE PROMISING PROGRAMS, TO
9	ALSO MAKE SURE THAT WE HAVE PROMISING SCIENTISTS
10	BEING TRAINED AND THAT CIRM IS SUPPORTING THOSE
11	PROMISING SCIENTISTS BEING TRAINED IN THOSE
12	PROGRAMS.
13	THE LAST THING IS BASICALLY ALSO WHAT I
14	STARTED WITH, WHICH CENTERS AROUND HONESTLY AND
15	OBJECTIVELY TAKING A LOOK AT THOSE FACTORS THAT
16	INTEREST AND MOTIVATE AND INCENTIVIZE THE
17	RESEARCHERS AND FIGURE OUT HOW WE CAN ALIGN THOSE
18	WITH WHAT CIRM ULTIMATELY NEEDS FOR THAT PARTICULAR
19	STAGE.
20	SO THAT'S WHERE WE ARE WITH DISCOVERY AND
21	TRANSLATION. AS I SAID, IT IS STILL IN ITS EARLY
22	FORMATIVE STAGES. WE ARE NOT NEARLY DONE COLLECTING
23	INFORMATION FROM STAKEHOLDERS. AS I SAID, IF WE
24	HAVEN'T GOTTEN THERE, WE'LL BE COMING TO A CITY NEAR
25	YOU SHORTLY. ULTIMATELY THIS IS SCHEDULED TO COME

1	TO THE BOARD AS A CONCEPT PLAN IN JULY.
2	OKAY. THEN REALLY THE LAST DISCRETE TOPIC
3	THAT I WANT TO TALK ABOUT CENTERS AROUND OUR LOAN
4	AND GRANTS PROGRAM AND MODIFICATIONS THAT WERE
5	CONSIDERING HERE. SO RECALL THAT CURRENTLY IF YOU
6	GET AN AWARD FROM CIRM, YOU HAVE TWO OPTIONS IN
7	ORDER TO TAKE THAT AWARD. YOU CAN TAKE IT AS A LOAN
8	AND PAY IT BACK AND NOT HAVE ROYALTY OBLIGATIONS OR
9	YOU CAN TAKE IT AS A GRANT, NOT HAVE TO PAY IT BACK,
10	BUT HAVE ROYALTY OBLIGATIONS. AND SO WHAT WE'RE
11	LOOKING TO DO IS WE'RE LOOKING TO SEE IF THERE'S A
12	WAY OF COMBINING THOSE TWO. AND THE PURPOSE OF THIS
13	IS TO ADDRESS WHAT I BELIEVE IS OUR PRIMARY
14	OBJECTIVE, WHICH IS TO PROVIDE A FAIR FUNDING
15	MECHANISM THAT DOES NOT INADVERTENTLY DISSUADE
16	PEOPLE FROM PARTNERING WITH CIRM.
17	SO CLEARLY IT HAS TO BE FAIR TO THE STATE
18	OF CALIFORNIA AND THE TAXPAYERS, BUT WE ALSO DON'T
19	WANT SOMETHING THAT INADVERTENTLY CREATES TOO HIGH
20	OF A SPEED BUMP THAT PEOPLE WITH PROMISING
21	TECHNOLOGIES ELECT NOT TO PARTICIPATE WITH CIRM
22	BECAUSE OF SOME INADVERTENT FEATURE TO EITHER ONE OF
23	THOSE TWO MECHANISMS.
24	AND SO WHAT WE'RE CONSIDERING AGAIN,
25	THIS IS STILL ON THE DRAWING TABLE IS IF WE
	22

1	COMBINED THE TWO WHERE WE ALLOWED THE RECIPIENT TO
2	ELECT WHETHER OR NOT THEY WOULD CONVERT IT WHERE
3	THE GRANT WOULD GET CONVERTED INTO A LOAN, WE COULD
4	ADDRESS THE TWO MAJOR COMPLAINTS THAT WE HEAR. ONE
5	OF THEM, WHICH IS SOMETIMES, AND THIS IS NOT ALWAYS,
6	BUT SOMETIMES LARGER COMPANIES DO NOT LIKE TO TAKE
7	ON PROJECTS THAT ARE ENCUMBERED WITH ROYALTY
8	OBLIGATIONS FROM SMALLER COMPANIES. THE OTHER
9	PROBLEM WE HAVE IS THAT SMALLER COMPANIES OFTENTIMES
10	DON'T LIKE HAVING A HUGE AMOUNT OF DEBT ON THEIR
11	BALANCE SHEET, EVEN AS IN CIRM'S CASE, IF THAT DEBT
12	IS FORGIVABLE IF THE PROGRAM DOESN'T WORK. SO WE'RE
13	HOPING THAT IF WE CAN GET THIS ELECTION PROCESS TO
14	WORK, THAT WE COULD BASICALLY HAVE THE BEST OF BOTH
15	WORLDS. THESE COMPANIES WOULDN'T HAVE TO CARRY IT
16	ON THEIR BALANCE SHEETS AS DEBT; BUT IF THEY WANTED
17	TO, THEY COULD CONVERT IT INTO DEBT AND PAY IT OFF.
18	IT'S MORE OF A JAMES ISSUE, BUT THIS
19	MECHANISM OF COMBINING THESE TWO AND BASICALLY
20	GETTING RID OF THE LOAN PROGRAM AS WE HAVE NOW WOULD
21	SIGNIFICANTLY STREAMLINE THE PROCESS THAT WE HAVE
22	AND WOULD MAKE IT MORE SIMPLE FROM AN ADMINISTRATIVE
23	BURDEN STANDPOINT. I THINK IT'S PROBABLY GOOD TO
24	LOOK INTO THIS BECAUSE WE'VE HAD THIS PROGRAM FOR A
25	WHILE NOW. WE'VE ONLY HAD FIVE PEOPLE EVER TAKE
	22

1	LOANS. WE ONLY HAVE TWO OUTSTANDING NOW, AND THAT'S
2	OUT OF 600 AWARDS THAT WE'VE GIVEN. SO I THINK WE
3	NEED TO SEE IF WE CAN FIND A WAY TO MAKE THIS
4	PROGRAM BETTER. IF NOT, MAYBE THE PROGRAM DOESN'T
5	NEED TO EXIST. I THINK THIS IS AN INTERESTING WAY
6	OF APPROACHING IT. WE DON'T HAVE ALL THE ANSWERS
7	HERE. WE HAVE A LOT MORE WORK TO DO ON IT IN TERMS
8	OF ECONOMIC MODELING AND THERE'S SOME LEGAL ISSUES
9	THAT JAMES AND HIS GROUP ARE WORKING WITH. BUT I
10	WANTED TO GIVE YOU THE IDEA THAT WE'RE CONSIDERING
11	HERE.
12	AND THEN LASTLY, I'LL JUST END WITH THINGS
13	TO COME. SO WE WILL HAVE THE FIRST CONSIDERATION OF
14	AWARDS UNDER CIRM 2.0 APRIL 23D. WE'RE VERY EXCITED
15	ABOUT THAT. THE REVIEW, AS I SAID, IS GOING TO BE
16	NEXT MONDAY. IF THAT GOES WELL, IT WILL BE COMING
17	TO THE BOARD ON APRIL 23D.
18	WE ALSO HAVE OUR FIRST IN-PERSON GWG
19	MEETING. SO THE ONE ON THE 30TH OF THIS MONTH IS
20	GOING TO BE TELEPHONIC, BUT OUR FIRST IN-PERSON GWG
21	MEETING WILL ALSO TAKE PLACE IN APRIL.
22	COMING UP IN MAY WE HAVE TO GET OUR
23	BUDGETS DONE, GO TO THE FINANCE SUBCOMMITTEE. IN
24	MAY WE'RE ALSO GOING TO HAVE OUR FIRST INTERACTIONS
25	WITH THE BOARD AROUND STRATEGIC PLANNING AND HOW

1	THAT'S GOING, AND HOPEFULLY ADOPT THE 2016 BUDGET IN
2	THAT MEETING.
3	LASTLY, IN JULY, VERY BUSY MEETING, IS
4	WHEN WE WILL BE BRINGING THE CONCEPT PLANS FORWARD
5	FOR THE DISCOVERY AND TRANSLATIONAL STAGE PROGRAMS.
6	WE WILL ALSO BE BRINGING FORWARD OUR CREATIVITY AND
7	BRIDGES CONCEPT PLAN, AGAIN, WHICH IS EDUCATION FOR
8	HIGH SCHOOL AND UNDERGRADUATE STUDENTS IN STEM
9	CELLS. AND THEN, LASTLY, AND HOPEFULLY FINALIZE OUR
10	STRATEGIC PLAN. SO WE HAVE A REALLY, REALLY BUSY
11	THREE MONTHS COMING UP, BUT THAT'S THE BUSINESS
12	WE'RE IN. WE'RE IN THE TIME BUSINESS, AND SO WE
13	HAVE TO ACT WITH THE APPROPRIATE SENSE OF URGENCY.
14	WITH THAT, I WILL STOP TALKING AND ANSWER
15	ANY AND ALL QUESTIONS THAT YOU MIGHT HAVE.
16	CHAIRMAN THOMAS: SO ELOQUENT AND
17	PERSUASIVE
18	DR. MILLS: COVERED ALL OF IT. AND I KNOW
19	SOME OF YOU ARE THINKING YOU SAID YOU WERE GOING TO
20	BE PITHY. AND THAT WASN'T. THAT WAS VERBOSE.
21	CHAIRMAN THOMAS: JUST WANT TO NOTE, DR.
22	MILLS, ANECDOTALLY AND FOR THE BOARD, WHEN MR. LOMAX
23	AND I MET WITH, AND PAT AND A UTA MET WITH THE
24	GERMAN DELEGATION, ONE OF THEM, UPON HEARING ABOUT
25	CIRM 2.0, EXPRESSED ASTONISHMENT AND GREAT
	25

1	ADMIRATION FOR THE IDEA THAT WE COULD EVER HAVE A
2	PROCESS THAT COULD STREAMLINE DOWN TO 121 DAYS. SO
3	THE CONCEPT HAS NOW GONE VIRAL INTERNATIONALLY AND
4	IS PLAYING TO VERY GOOD REVIEWS.
5	DR. MILLS: THAT WAS MY PLAN. I WANT TO
6	BE A ROCK STAR FOR MY DAUGHTER. SHE'S LOOKING AT
7	ALL THE GERMAN WEBSITES RIGHT NOW. YES, SIR.
8	DR. LEVIN: YOU DISCUSSED THE INTEREST TO
9	HAVE A CONSULTATION PERIOD AFTER THE GRANTS ARE
10	SUBMITTED IN THESE ROLLING DEADLINES. DO YOU DO
11	THAT ALSO BEFOREHAND?
12	DR. MILLS: ABSOLUTELY. AND THAT'S WAY
13	WHERE WE PREFER IT. GIL AND BECKY TALK TO LOTS AND
14	LOTS AND LOTS OF PEOPLE THAT HAVEN'T APPLIED YET
15	ABOUT CONSTRUCTING THEIR GRANTS. I'LL BET HIS NAME
16	IS EVEN HERE. YEAH. SO PRETTY MUCH ON THE BACK OF
17	EVERY PRESENTATION WE GIVE IS GIL'S CONTACT
18	INFORMATION SO THAT WHEN YOU HAVE A PROGRAM AND
19	YOU'RE CONTEMPLATING APPLYING, THE FIRST THING WE
20	RECOMMEND EVERYONE DOING IS CALLING US AND JUST
21	HAVING A PHONE CONVERSATION ABOUT IT SO WE CAN DO
22	THINGS LIKE IS IT IN SCOPE? DOES IT INVOLVE A STEM
23	CELL? IS IT RESPONSIVE? AND THEN, SECONDLY, IF
24	THERE'S ANY CLARIFICATION THAT CAN BE DONE, WE WANT
25	TO DO IT ON THE FRONT END BECAUSE WE DON'T WANT THEM

1	WASTING IT'S A SUBSTANTIAL APPLICATION TO HAVE TO
2	PUT TOGETHER. WE'D RATHER HAVE IT COME IN WELL
3	CONSTRUCTED AS OPPOSED TO AMEND IT. YES IS A VERY
4	LONG ANSWER TO A SIMPLE QUESTION.
5	I AM NOT DOING WELL WITH THIS PITHY THING
6	TODAY.
7	CHAIRMAN THOMAS: ANY OTHER QUESTIONS,
8	COMMENTS FROM MEMBERS OF THE BOARD? THANK YOU, DR.
9	MILLS.
10	WE'LL NOW TURN IT OVER FOR THE FINANCE
11	REPORT TO MS. SILVA-MARTIN.
12	MS. SILVA-MARTIN: GOOD MORNING, MR.
13	CHAIR, MEMBERS OF THE BOARD, MEMBERS OF THE PUBLIC,
14	AND CIRM TEAM. THIS MORNING I'LL PROVIDE A BRIEF
15	REPORT ON CIRM FINANCES. I WILL BRIEFLY COVER OUR
16	GRANT DISBURSEMENTS AND AVAILABLE CASH, AS WELL AS
17	WHERE WE ARE ON OUR ACTUAL EXPENDITURES FOR THIS
18	FISCAL YEAR, AND ALSO ON OUR BUDGET DEVELOPMENT FOR
19	THE '15-'16 FISCAL YEAR.
20	BEFORE I ACTUALLY GO INTO THE DETAILS, I
21	DO WANT TO POINT OUT THAT THERE HAS BEEN NO MAJOR OR
22	SIGNIFICANT CHANGES SINCE THE LAST TIME THAT I
23	REPORTED. OUR FINANCES ARE ON TRACK. OUR
24	EXPENDITURES ARE WITHIN BUDGET, AND WE CONTINUE TO
25	MAINTAIN A VERY HEALTHY CASH RESERVE.
	27

1	LOOKING AT SOME OF THE DETAILS, AS YOU CAN
2	SEE, OUR GRANT DISBURSEMENTS FOR THE FIRST EIGHT
3	MONTHS OF THE FISCAL YEAR WERE \$133.6 MILLION. AND
4	IN COMPARISON TO LAST YEAR, THEY WERE VIRTUALLY THE
5	SAME DURING THAT SAME PERIOD. WE DISBURSED \$133.1
6	MILLION. OUR CASH RESERVES AS OF FEBRUARY ARE \$72.2
7	MILLION, SO WE HAVE SUFFICIENT FUNDS TO MEET OUR
8	FINANCIAL OBLIGATIONS FOR THE NEXT SEVERAL MONTHS.
9	WE ARE ALSO SCHEDULED TO RECEIVE
10	ADDITIONAL FUNDS FROM THE SPRING BOND SALE. THE
11	STATE TREASURER AND DEPARTMENT OF FINANCE HAVE
12	AGREED TO PROVIDE US WITH UP TO \$115 MILLION
13	ADDITIONAL FUNDS EITHER FROM THE BOND SALE OR FROM
14	COMMERCIAL PAPER. SO, AGAIN, WE'RE IN A VERY GOOD
15	FINANCIAL POSITION WITH OUR CASH RESERVES.
16	NOW LOOKING AT OUR OPERATIONAL
17	EXPENDITURES FOR THE FIRST EIGHT MONTHS OF THE
18	FISCAL YEAR, SO, AS YOU CAN SEE FROM THE CHART
19	ABOVE, WE WERE ALLOCATED FOR THE FIRST EIGHT MONTHS
20	OF THE FISCAL YEAR ABOUT \$11 MILLION, AND WE'VE
21	EXPENDED JUST UNDER \$10 MILLION. SO WE ARE
22	UNDERRUNNING THE BUDGET BY A LITTLE BIT OVER A
23	MILLION DOLLARS. THE MAJORITY OF THAT UNDERRUN IS
24	IN OUR EMPLOYEE EXPENSES, SO SALARIES AND WAGES AND
25	BENEFITS, SUCH AS RETIREMENT AND HEALTHCARE, FOR
	28

1	POSITIONS. THE REASON FOR THE EMPLOYEE UNDERRUN IS,
2	AS YOU MAY RECALL, AT THE BEGINNING OF THE FISCAL
3	YEAR, WE HAD SEVERAL POSITIONS THAT BECAME VACANT.
4	A DETERMINATION WAS MADE THAT THOSE POSITIONS WERE
5	NO LONGER NEEDED AND SO THEY WERE NOT FILLED, AND SO
6	WE'RE SEEING A FAIRLY SIGNIFICANT SAVINGS IN THAT
7	AREA.
8	SIMILARLY, WE'RE SEEING SAVINGS IN ALL OF
9	OUR OTHER CATEGORIES, SO NONE OF OUR CATEGORICAL
10	BUDGETS ARE BEING OVERRUN FOR EXTERNAL SERVICES. AS
11	YOU KNOW, WE IMPLEMENTED CIRM 2.0 AND WE SPENT A LOT
12	OF EFFORT DURING THE FIRST HALF OF THE YEAR
13	IMPLEMENTING THAT. AND SO SOME OF OUR MEETINGS,
14	SUCH AS CDAP MEETINGS, THAT WE HAD BUDGETED FOR DID
15	NOT MATERIALIZE, AND SO OUR COSTS ARE DOWN FOR BOTH
16	EXTERNAL SERVICES AND REVIEW MEETINGS IN SUPPORT OF
17	THOSE TYPE OF EXPENDITURES.
18	THE OTHER CATEGORY I WANTED TO POINT OUT
19	AGAIN IS OUR TRAVEL. OUR TRAVEL CONTINUES TO BE
20	UNDER BUDGET. AND BASICALLY THE TWO COST CENTERS
21	THAT ARE REALLY CONTRIBUTING TO THAT UNDERRUN ARE
22	THE OFFICE OF THE PRESIDENT AND OUR LEGAL TEAM.
23	AND THEN FINALLY, AS DR. MILLS POINTED
24	OUT, WE ARE RIGHT IN THE MIDDLE OF OUR DEVELOPMENT
25	OF OUR '15-'16 BUDGET. WE HAVE RECEIVED INFORMATION

1	FROM OUR COST CENTERS. WE'VE ACCUMULATED THAT DATA,
2	AND IT'S BEING REVIEWED NOW BY THE PRESIDENT AND THE
3	CHAIR. AND AS DR. MILLS INDICATED, WE WILL SHARE IT
4	WITH THE FINANCE SUBCOMMITTEE CHAIRS SOMETIME IN
5	APRIL AND THEN BRING IT BEFORE A FINANCE
6	SUBCOMMITTEE IN MAY AS WELL TO THIS BOARD IN MAY.
7	AND THAT REALLY CONCLUDES MY PRESENTATION.
8	ARE THERE ANY QUESTIONS? THANK YOU.
9	CHAIRMAN THOMAS: THANK YOU, CHILA. I
10	JUST WANTED TO HIGHLIGHT ONE POINT SHE MADE, WHICH
11	IS THE SPRING BOND SALE THAT WILL BE COMING UP IN
12	WHICH WE WILL BE A PART. EVERYBODY SORT OF ASSUMES
13	THAT THE MONEY FROM THE STATE FLOWS SEAMLESSLY TO US
14	AND DOESN'T REALLY INVOLVE A LOT OF WORK ON OUR
15	PART. CHILA DOES A WONDERFUL JOB ON THE FINANCIAL
16	SIDE. AND I'D ALSO LIKE TO MENTION AND SINGLE OUT
17	AMY LEWIS WHO IS THE PRINCIPAL PERSON HERE WHO DEALS
18	WITH THE DEPARTMENT OF FINANCE AND THE GOVERNOR'S
19	OFFICE AND TALKS TO THEM ABOUT WHAT OUR SEMIANNUAL
20	NEEDS ARE GOING TO BE, WALKS THROUGH THEM, AND HELPS
21	ARRIVE AT THE AMOUNT THAT WE NEED TO HAVE FUNDED,
22	WHICH MESSAGE THEN GETS TRANSLATED TO THE STATE
23	TREASURER WHO INCLUDES OUR PRO RATA SHARE IN THEIR
24	SEMIANNUAL BOND OFFERING. AND IT'S THROUGH THAT
25	MECHANISM THAT WE ARE ABLE TO OPERATE. AND SO JUST

1	WANTED TO REPORT THAT THAT IS A VERY IMPORTANT
2	PROCESS, AND WE SPEND A LOT OF TIME MAKING SURE THAT
3	WE DO EVERYTHING WE NEED TO TO MAKE THAT HAPPEN
4	FLAWLESSLY. SO THANK YOU TO AMY FOR THAT.
5	GOING TO MOVE NOW ON TO THE ACTION ITEMS
6	ON THE AGENDA. ITEM NO. 7, CONSIDERATION OF
7	APPLICATIONS FOR RFA 14-02, CIRM PRECLINICAL
8	DEVELOPMENT AWARDS. DR. KADYK WILL PRESENT.
9	DR. KADYK: THANK YOU. GOOD MORNING TO
10	MR. CHAIRMAN, MEMBERS OF THE BOARD, MEMBERS OF THE
11	PUBLIC, AND MY CIRM COLLEAGUES. I'M HERE TO PRESENT
12	THE RECOMMENDATIONS FROM CIRM AND THE GRANTS WORKING
13	GROUP FOR THE RECENTLY REVIEWED PRECLINICAL
14	DEVELOPMENT AWARDS FOR WHICH THE CONCEPT WAS
15	APPROVED ONE YEAR AGO IN MARCH 2014.
16	AND SO THE INTENT OF THIS RFA IS TO FUND
17	DEVELOPMENT OF THERAPEUTIC CANDIDATES THAT HAVE BEEN
18	SUCCESSFULLY IDENTIFIED AS SO-CALLED DEVELOPMENT
19	CANDIDATES, WHICH YOU CAN SEE ON THE DIAGRAM IS
20	ABBREVIATED AS A DC. AND THIS IS AN IMPORTANT STAGE
21	OF PRECLINICAL DEVELOPMENT IN THAT IT MARKS THE
22	TRANSITION, THE FIRST TRANSITION, BETWEEN AN
23	EXPLORATORY STAGE PRECLINICAL RESEARCH AND THE MUCH
24	MORE HIGHLY REGULATED AND COSTLY PRECLINICAL
25	DEVELOPMENT.

1	SO AS YOU CAN SEE, THE IDENTIFICATION OF A
2	DEVELOPMENT CANDIDATE HAS BEEN THE END GOAL OF MANY
3	OF OUR EARLIER STAGE EARLY TRANSLATION RESEARCH
4	AWARDS FOR WHICH WE'VE HAD FOUR ROUNDS. AND THESE
5	AWARDS WOULD THEN ALLOW DEVELOPMENT, THE EARLY
6	DEVELOPMENT, OF THESE CANDIDATES UP TO THE STAGE OF
7	HOLDING A PRE-IND MEETING, AT WHICH POINT THEY WOULD
8	BE READY TO APPLY FOR OUR NEWLY ANNOUNCED PROGRAM
9	ANNOUNCEMENT 15-01, WHICH WOULD FUND PIVOTAL
10	IND-ENABLING STUDIES UP TO THE IND STAGE.
11	AND I ALSO WANT TO POINT OUT THAT THIS THE
12	FINAL RFA THAT WAS ISSUED UNDER THE CIRM 1.0
13	PROCESS.
14	SO SINCE ENTRY INTO PRECLINICAL
15	DEVELOPMENT IS A PRETTY IMPORTANT MILESTONE, IT IS
16	IMPORTANT THAT THE CANDIDATES HERE MEET CERTAIN KEY
17	SCIENTIFIC ELIGIBILITY CRITERIA. SO FIRST OF ALL,
18	THE DEVELOPMENT CANDIDATE SHOULD BE VERY
19	SPECIFICALLY CHARACTERIZED AS A SINGLE AND
20	WELL-DEFINED CANDIDATE THAT DERIVES FROM OR TARGETS
21	STEM CELLS, AND THERE SHOULD BE SUFFICIENT EVIDENCE
22	THAT IT IS READY TO ENTER PRECLINICAL DEVELOPMENT,
23	SUCH AS HAVING A FAIR AMOUNT OF DATA SHOWING
24	CONVINCING, REPRODUCIBLE DISEASE MODIFYING ACTIVITY
25	IN RELEVANT PRECLINICAL MODELS, PRELIMINARY

1	ASSESSMENTS OF SAFETY AND MECHANISM OF ACTION, AS
2	WELL AS REPRODUCIBLE AND SCALABLE RESEARCH GRADE
3	PRODUCTION OF THE CANDIDATE.
4	SO THE OBJECTIVE OF THESE AWARDS IS TO
5	CARRY OUT THE ACTIVITIES NEEDED TO CONDUCT A
6	WELL-PREPARED PRE-IND MEETING WITH THE FDA AT THE
7	END OF THE AWARD. SO THERE'S A NUMBER OF IN-SCOPE
8	ACTIVITIES THAT COULD BE NEEDED TO BE DONE TO GET TO
9	THAT PRE-IND MEETING, INCLUDING, VERY IMPORTANTLY,
10	DEVELOPING A GMP MANUFACTURING PROCESS INCLUDING
11	DEVELOPMENT AND QUALIFICATION OF RELEASE AND
12	IN-PROCESS ASSAYS, IDENTIFYING AN APPROPRIATE DOSE
13	REGIMEN OF DOSING AND ROUTE OF ADMINISTRATION FOR
14	THE THERAPEUTIC, PHARMACOKINETIC STUDIES, BY WHICH I
15	MEAN STUDYING THE EFFECTS OF THE BODY ON THE
16	THERAPEUTIC AFTER THE THERAPEUTIC IS ADMINISTERED,
17	FOR EXAMPLE, BIODISTRIBUTION IN THE CASE OF A CELL
18	THERAPY, HOW LONG THE CELLS SURVIVE IN THE BODY,
19	PILOT SAFETY AND MECHANISM OF ACTION STUDIES,
20	SELECTION OF THE TARGET INDICATION, AND PREPARATION
21	OF A CLINICAL DEVELOPMENT PLAN AND DRAFT PROTOCOL,
22	AND FINALLY, CULMINATING IN THE CONDUCT OF A PRE-IND
23	MEETING WITH THE FDA.
24	SO LAST MARCH THE BOARD APPROVED UP TO \$40
25	MILLION TO FUND THESE AWARDS FOR WHICH WE

1	ANTICIPATED ABOUT FIVE TO EIGHT AWARDS WOULD BE
2	FUNDED. THIS IS A RELATIVELY SHORT-TERM AWARD OF 30
3	MONTHS, AND WE ESTIMATED A BUDGET THAT SHOULD BE IN
4	THE FIVE TO \$8 MILLION RANGE FOR JUSTIFIABLE TOTAL
5	PROJECT COSTS.
6	AND I JUST WANTED TO MENTION THAT WE ALSO
7	HAD A STIPULATION THAT IF THE DEVELOPMENT CANDIDATE
8	WAS NOT IDENTIFIED WITH PRIOR CIRM FUNDING, THEN THE
9	APPLICANT WOULD INDICATE NEED TO PROVIDE TO
10	ONE-TO-ONE MATCHING FUNDING.
11	SO THESE APPLICATIONS WERE REVIEWED BY THE
12	GRANTS WORKING GROUP WITH A NUMBER OF KEY REVIEW
13	CRITERIA THAT CAN BE PUT INTO THREE MAJOR
14	CATEGORIES. FIRST OF ALL, SHOULD THE PROPOSED
15	THERAPEUTIC BE DEVELOPED? SECOND, CAN THE PROPOSED
16	PLAN ACHIEVE THE RFA OBJECTIVE? AND THIRD, IS THIS
17	THE RIGHT TEAM TO EXECUTE THE PLAN?
18	SO SHOULD IT BE DEVELOPED, THE REVIEWERS
19	LOOKED AT ARE THE SIGNIFICANCE OF THE PROPOSAL. IS
20	IT COMPETITIVE WITH STANDARD OF CARE? AND ALSO THE
21	SCIENTIFIC RATIONALE FOR THE PROPOSAL. DO WE REALLY
22	BELIEVE THAT THERE'S A POTENTIAL FOR CLINICAL
23	BENEFIT IN THE TARGETED INDICATION?
24	CAN THE PROPOSED PLAN ACHIEVE THE RFA
25	OBJECTIVE? SO, AGAIN, LOOKING AT THE READINESS THAT

1	I MENTIONED EARLIER OF THE CANDIDATE TO ENTER
2	PRECLINICAL DEVELOPMENT, LOOKING AT THE DESIGN AND
3	FEASIBILITY OF THE PROJECT PLAN AND THE TIMELINE,
4	AND LOOKING AT ASSETS, COLLABORATIONS, AND
5	ENVIRONMENT, ALL THE THINGS THAT ARE NEEDED, SUCH AS
6	MTA'S, PATENTS, CONTRACTS, APPROPRIATE EQUIPMENT,
7	AND FACILITIES TO CARRY OUT THE PLAN.
8	AND THEN FINALLY, IS THIS THE RIGHT TEAM
9	TO EXECUTE THE PLAN? THE REVIEWERS LOOKED AT THE PI
10	AND THE TEAM AND LOOKED AT THEIR EXPERIENCE IN
11	LEADING A TEAM SUCH AS THIS AS WELL AS PRECLINICAL
12	DEVELOPMENT EXPERTISE.
13	SO THE SCORING BY THE GRANTS WORKING GROUP
14	WAS A DETERMINATIVE SCORING MECHANISM WHERE
15	REVIEWERS WERE INSTRUCTED THAT IF SCORES WERE
16	GREATER THAN OR EQUAL TO 75, THAT WOULD INDICATE
17	THAT THE REVIEWER INTENDED TO RECOMMEND THAT THE
18	APPLICATION BE FUNDED. FOR SCORES 65 TO 74 IS
19	CONSIDERED TIER II, MEANING THAT THE APPLICATION WAS
20	PERHAPS OF MODERATE QUALITY AND THE OVERALL AVERAGE
21	SCORE MAY IN A TIER II CATEGORY INDICATES THERE MAY
22	NOT HAVE BEEN CONSENSUS ON THE MERIT OF THE
23	PROPOSAL. AND TIER III, SCORES LESS THAN OR EQUAL
24	64 WERE NOT RECOMMENDED FOR FUNDING AT THIS TIME.
25	I WANT TO POINT OUT THAT THIS RFA DID LIST

1	CERTAIN PRIORITIES THAT COULD BE TAKEN INTO ACCOUNT
2	DURING PROGRAMMATIC CONSIDERATION FOR FUNDING.
3	SO WE CALLED OUT SEVERAL PRIORITIES IN
4	THIS RFA. FIRST OF ALL, THERAPEUTIC CANDIDATES THAT
5	ARE CELL THERAPIES, ESPECIALLY IF THEY'RE DERIVED
6	FROM PLURIPOTENT STEM CELLS OR DIRECTLY REPROGRAMMED
7	CELLS. SECOND, PROJECTS THAT SEEM TO BE POTENTIALLY
8	TRANSFORMATIVE TO MEETING UNMET MEDICAL NEEDS. FOR
9	PROJECTS WERE PROPOSING DEVELOPING A DEVELOPMENT
10	CANDIDATE THAT WAS IDENTIFIED USING CIRM FUNDING, IF
11	THE APPLICANT COULD BRING IN 25-PERCENT CO-FUNDING,
12	THAT WOULD BE CONSIDERED A PRIORITY. AND FOR THOSE
13	DEVELOPMENT CANDIDATES NOT IDENTIFIED WITH PRIOR
14	CIRM FUNDING, THEY'RE ALL REQUIRED TO HAVE MATCHING
15	FUNDING, BUT IF THAT MATCHING FUNDING COMES FROM
16	INDUSTRY, THAT WOULD BE IN THE PRIORITY CATEGORY.
17	SO THIS SLIDE IS A HIGH LEVEL VIEW OF THE
18	GRANTS WORKING GROUP AND CIRM RECOMMENDATIONS AND
19	THE RATIONALE. AFTER THIS GIL SAMBRANO WILL HAVE A
20	MORE DETAILED SPREADSHEET LISTING THE DETAILS OF THE
21	APPLICATIONS.
22	BUT IN SUMMARY, FOR TIER I APPLICATIONS
23	WE'RE RECOMMENDING AND CONCUR WITH THE GRANTS
24	WORKING GROUP ASSESSMENT THAT THESE APPLICATIONS BE
25	FUNDED. THERE WERE FIVE AWARDS IN THE TIER I
	36
	36

1	CATEGORY WITH BUDGETS TOTALING ABOUT 16 MILLION, AND
2	THOSE FIVE APPLICATIONS ALSO BRING IN \$3.25 MILLION
3	OF CO-FUNDING.
4	FOR THE TIER II AND TIER III APPLICATIONS,
5	THAT RECOMMENDATION CURRENTLY IS DO NOT FUND. WE DO
6	ENCOURAGE APPLICANTS TO CONSIDER RESUBMITTING
7	PROPOSALS TAKING INTO ACCOUNT THE REVIEWERS'
8	COMMENTS. AS RANDY HAS OUTLINED AND AS YOU KNOW
9	WITH CIRM 2.0, WE ANTICIPATE HAVING MORE
10	OPPORTUNITIES AND MORE REGULAR OPPORTUNITIES FOR
11	APPLICANTS TO REAPPLY. CURRENTLY THERE IS THE
12	PROGRAM ANNOUNCEMENT 15-01, WHICH MAY BE A LITTLE
13	BIT TOO LATE, TOO ADVANCED FOR SOME OF THESE
14	PROJECTS, BUT, IN FACT, SOME OF THE APPLICATIONS ARE
15	CLOSE TO A PRE-IND MEETING AND MAY BE ABLE TO APPLY
16	FOR THAT, AND THEY SHOULD EVALUATE THAT PROGRAM
17	ANNOUNCEMENT. AND THEN WE DO ALSO ANTICIPATE LATER
18	THIS SUMMER TO HAVE ANOTHER OPPORTUNITY FOR A
19	PROGRAM ANNOUNCEMENT FOR EARLY PRECLINICAL
20	DEVELOPMENT-TYPE AWARDS.
21	SO WITH THAT, I WOULD ASK IF THERE ARE ANY
22	QUESTIONS.
23	CHAIRMAN THOMAS: NO QUESTIONS. THANK YOU
24	VERY MUCH, DR. KADYK. NOW TURN THIS OVER FOR
25	PROGRAMMATIC REVIEW TO MR. SHEEHY.

37

1	MR. SHEEHY: THANK YOU, CHAIRMAN THOMAS.
2	SO I THINK THE FIRST THING IS THAT WE HAVE
3	A NEW WRINKLE IN HOW WE'RE GOING TO PROCESS THESE.
4	IN RESPONSE TO A SENSE THAT APPLICATIONS THAT WERE
5	NOT CONSIDERED BY THE COMMITTEE, WE'RE NOT GIVING
6	APPLICANTS FROM APPLICATIONS THAT WERE NOT
7	CONSIDERED BY THE COMMITTEE WEREN'T GIVEN AN
8	OPPORTUNITY TO EXPRESS AN OPINION ON THOSE GRANTS
9	DIRECTLY. THE WAY WE DID IT BEFORE IS WE'D CONSIDER
10	THE GRANTS. IF THE COMMITTEE DIDN'T TAKE UP A GRANT
11	TO EITHER FUND OR NOT FUND A SPECIFIC MOTION, THEN
12	THOSE GRANTS THEN THERE WAS NOT AN OPPORTUNITY
13	BEFORE THE FINAL MOTION FOR APPROVAL, THE BLANKET
14	APPROVAL, FOR APPLICANTS OR INTERESTED PARTIES TO BE
15	ABLE TO COMMENT TO TRY TO ATTRACT THE ATTENTION OF
16	MEMBERS TO POTENTIALLY MAKE A MOTION IN FAVOR OR TO
17	GET SOME ACTION ON THOSE GRANTS.
18	SO WE'VE INTRODUCED A NEW PROCESS, A
19	SLIGHT VARIATION, AND MAYBE COUNSEL COULD KIND OF
20	WALK US THROUGH IT.
21	MR. HARRISON: THANKS, JEFF. SO AS JEFF
22	INDICATED, WE WILL BE OFFERING AN OPPORTUNITY FOR
23	PUBLIC COMMENT AT THIS STAGE OF THE PROCEEDING
24	RATHER THAN, AS JEFF INDICATED, BEFORE THE FINAL
25	OMNIBUS MOTION. WE HAVE SET GUIDELINES TO ENSURE
	20

1	THAT IT RUNS SMOOTHLY, AND LET ME JUST WALK YOU
2	THROUGH THOSE.
3	FIRST, WE WILL STRICTLY LIMIT PUBLIC
4	COMMENTS TO THREE MINUTES PER SPEAKER. WE WOULD
5	DISCOURAGE ANY DIALOGUE BETWEEN MEMBERS OF THE BOARD
6	AND APPLICANTS OR MEMBERS OF THE PUBLIC ABOUT
7	SCIENTIFIC POINTS. TO THE EXTENT MEMBERS OF THE
8	BOARD HAVE QUESTIONS, THOSE QUESTIONS CAN BE
9	ADDRESSED ONCE THE BOARD BEGINS TO UNDERTAKE
10	CONSIDERATION OF SPECIFIC APPLICATIONS. AND THEN
11	FINALLY, WE WILL NOT ACCEPT MOTIONS DURING THE
12	PUBLIC COMMENT PERIOD. WE'LL RESERVE MOTIONS FOR
13	THE STAGE AT WHICH THE BOARD BEGINS TO CONSIDER
14	INDIVIDUAL APPLICATIONS.
15	AND, OF COURSE, MEMBERS OF THE PUBLIC WILL
16	CONTINUE TO HAVE THE OPPORTUNITY TO OFFER PUBLIC
17	COMMENTS BEFORE ANY INDIVIDUAL VOTE IS TAKEN.
18	MR. SHEEHY: THANK YOU, JAMES.
19	SO IS THAT WELL UNDERSTOOD? ARE THERE ANY
20	MEMBERS OF THE PUBLIC WHO WOULD LIKE TO ADDRESS ANY
21	OF THE GRANTS THAT WE HAVE COMING UP? AND MAYBE
22	KIND QUEUE UP. PLEASE INTRODUCE YOURSELVES.
23	MR. REED: DON REED, MEMBER OF THE PUBLIC.
24	FIRST OF ALL, THANK YOU VERY MUCH FOR THAT
25	INCLUSION OF THE PUBLIC WHICH HAS ALWAYS BEEN A PART
	20

1	OF THIS THING, AND THIS IS TREMENDOUS. THANK YOU
2	VERY MUCH.
3	MY INTEREST IS IN PC1-08128, THE EMBRYONIC
4	AND OSTEOCHONDRAL. I LIVE IN A TWO-STORY HOUSE WITH
5	17 STEPS BETWEEN THE FIRST AND SECOND FLOORS. THERE
6	ARE ALWAYS OBJECTS ON THE STAIRS WAITING FOR ME TO
7	CARRY THEM UP. MY WIFE GLORIA LEAVES THEM THERE FOR
8	ME. SHE TRIES TO PLAN HER DAY SHE'LL ONLY HAVE TO
9	GO DOWN THE STAIRS ONCE IN THE MORNING AND GO UP
10	ONCE AT NIGHT BECAUSE SHE HAS ARTHRITIS IN BOTH
11	KNEES. THE CARTILAGE CUSHION IN THE JOINT OUT, SO
12	BENDING THE KNEE GRINDS BONE AGAINST BONE.
13	FINANCIALLY ARTHRITIS COSTS AMERICA MORE
14	THAN 200 BILLION A YEAR AND MAY AFFECT AS MANY AS 18
15	PERCENT OF ALL WOMEN OVER SIXTY. GRANTED THIS
16	PROJECT IS AIMED AT YOUNGER SUFFERERS OF ARTHRITIS
17	UNLIKE THE MORE COMMON AGING RELATED. HOWEVER, JUST
18	AS THE FIRST ATTEMPTS TO CURE PARALYSIS ARE BEING
19	DONE WITH NEW INJURIES AND ONLY LATER TO BE TRIED IN
20	THE OLDER CHRONIC INJURIES, EVEN SO, ADVANCING A
21	CURE FOR ANY FORM OF ARTHRITIS OPENS THE DOOR FOR
22	UNIVERSAL BELIEF?
23	A CENTRAL TENET OF PROP 71 WAS TO
24	ENCOURAGE EMBRYONIC STEM CELL RESEARCH IN WHICH I
25	HAVE GREAT FAITH. TO THE BEST OF MY KNOWLEDGE, WE
	40

1	DO NOT HAVE AN EMBRYONIC STEM CELL ARTHRITIS THERAPY
2	BEING DEVELOPED, AND THIS SEEMS LIKE A GOOD ONE.
3	THERE IS REASON TO BELIEVE THIS IS A
4	WINNER. STUDYING THE PROPOSAL REMINDS ME OF THE
5	JUST EMERGING, BUT HIGHLY SUCCESSFUL PROJECT BY DR.
6	SUE KIMBER AT THE UNIVERSITY OF MANCHESTER IN THE
7	UK. PRECURSOR CARTILAGE CELLS, EMBRYONIC STEM
8	CELLS, THESE WERE IMPLANTED IN DAMAGED CARTILAGE OF
9	RATS. THE RESULTS, QUOTE, AFTER FOUR WEEKS THE
10	CARTILAGE WAS PARTIALLY REPAIRED. AFTER TWELVE
11	WEEKS, THE CARTILAGE SURFACE WAS SMOOTH AND SIMILAR
12	IN APPEARANCE TO NORMAL CARTILAGE. ANY SUBSTANTIVE
13	PROGRESS TOWARD A CURE OF ARTHRITIS WOULD BE A HOME
14	RUN FOR THE CALIFORNIA STEM CELL PROGRAM AND MIGHT
15	ENCOURAGE OTHER STATES AND NATIONS TO INVEST
16	SIMILARLY. I THEREBY REQUEST THAT PC1-08128 BE
17	MOVED INTO TIER I AND APPROVED FOR FUNDING. THANK
18	YOU VERY MUCH.
19	DR. D'LIMA: THANKS FOR THIS OPPORTUNITY
20	TO SPEAK. MY NAME IS DARRYL D'LIMA. I'M THE PI ON
21	THE SAME GRANT THAT MR. REED SPOKE ABOUT. IN THE
22	INTEREST OF FULL DISCLOSURE, I DON'T KNOW MR. REED
23	AND I HAVEN'T SOLICITED HIS OPINION.
24	IN 2009 CIRM GRANTED US AN AWARD TO
25	TRANSLATE OUR BASIC SCIENCE AND DISCOVERIES INTO
	41

1	STEM CELL TREATMENT. THREE YEARS LATER, AFTER CLOSE
2	COLLABORATIONS WITH CIRM STAFF, WE FOUND A
3	COMPELLING CANDIDATE FOR TREATING OSTEOCHONDRAL
4	DEFECTS WHICH ARE THE MAJOR CAUSE OF OSTEOARTHRITIS.
5	SINCE 2012 SCRIPPS HEALTH HAS BEEN GIVING
6	US ANNUAL BRIDGE FUNDING TO CONTINUE OUR WORK. AND
7	WE'VE CONTINUED TO COLLABORATE WITH CIRM AND
8	FOLLOWED THE RECOMMENDATIONS TO MAKE US CLINICALLY
9	COMPETITIVE FOR THIS PRECLINICAL AWARD.
10	WE'VE CONTACTED THE FDA AND PRESENTED OUR
11	CASE. IN FACT, IN THE GRANT APPLICATION ALL THE
12	TESTING WE PROPOSE IS EITHER RECOMMENDED BY THE FDA
13	OR HAS BEEN APPROVED BY THE FDA.
14	WE'VE ALSO RECEIVED DONATIONS FROM
15	PATIENTS THAT BELIEVE THAT OUR TREATMENT CAN
16	TRANSFORM THE TREATMENT OF OSTEOARTHRITIS.
17	IT IS TRUE THAT YOU CAN REPLACE THE JOINTS
18	OF OLDER PATIENTS WITH ARTHRITIS. JOINT REPLACEMENT
19	IS A 50-YEAR-OLD TECHNOLOGY. THIS IS ONE OF THE
20	IMPLANTS THAT GOES INTO THE KNEE JOINTS. THIS IS
21	THE BOX THAT THE IMPLANT COMES IN. THERE ARE SIX
22	SUCH BOXES THAT ARE IMPLANTED IN A TYPICAL KNEE
23	REPLACEMENT, WHICH HAS METAL, PLASTIC, AND CEMENT.
24	THESE ARE THE INSTRUMENTS THAT WE USE. WE HAVE
25	THREE TRAYS OF THESE INSTRUMENTS IN THE OPERATING
	42

1	ROOM. WE STILL USE HAMMERS, SAWS, AND DRILLS.
2	MR. TORRES: YOU HEAR THAT?
3	DR. D'LIMA: THIS IS OUR TECHNOLOGY, AND
4	THIS IS WHAT WE NEED TO IMPLANT OUR TECHNOLOGY
5	TODAY.
6	NOW, I UNDERSTAND THAT CIRM 2.0 GIVES US
7	BETTER OPPORTUNITIES, BUT I CANNOT GO BACK TO
8	SCRIPPS FOR A FOURTH YEAR OF FUNDING. I HAVE
9	CANNIBALIZED FUNDS FROM OTHER PROJECTS TO PAY FOR
10	THIS PROJECT. I'VE PERSONALLY TAKEN A 20-PERCENT
11	PAY CUT OVER THE LAST TWO YEARS. I'M ON THE VERGE
12	OF LOSING MY TEAM, AND I WILL LOSE \$4 MILLION OF
13	MATCHING FUNDS THAT SCRIPPS HAS COMMITTED TO THIS
14	APPLICATION.
15	TEN OUT OF THE 15 REVIEWERS GAVE US A
16	SCORE THAT RECOMMENDED FUNDING IN TIER I. BY USING
17	THE AVERAGE AS A SCORE FOR RANKING APPLICATIONS, YOU
18	ARE GIVING THE MINORITY A GREATER VOTING POWER THAN
19	THE MAJORITY. SO IF YOU WANT TO GIVE US A FIGHTING
20	CHANCE AT REPLACING 50-YEAR-OLD TECHNOLOGY WITH
21	TODAY'S TECHNOLOGY, PLEASE CONSIDER OUR APPLICATION.
22	THANK YOU.
23	MR. TORRES: I WOULD LIKE TO FIRST OF
24	ALL, THANK YOU FOR BEING HERE TODAY. HAVING JUST
25	UNDERGONE THAT OPERATION 30 DAYS AGO, I WAS HAPPY TO
	43
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1	SEE THAT MY NEW KNEE CAME FROM LEIPZIG, GERMANY,
2	SINCE WE HAVE A BILATERAL AGREEMENT WITH GERMANY,
3	AND I THINK THAT HAS WORKED OUT WELL FOR US AND FOR
4	GERMANY.
5	I DON'T KNOW WHETHER THE GOOD PEOPLE AT
6	THE IOM WOULD OBJECT TO MY MAKING A MOTION TO MOVE
7	THIS FOR FUNDING, OR WOULD I BE CONFLICTED OUT
8	BECAUSE I HAVE ENDURED THE SURGERY?
9	MR. SHEEHY: SENATOR TORRES, AT THIS TIME
10	THE PROCESS WE'VE ENVISIONED DOESN'T ALLOW FOR
11	MOTIONS. THIS IS AN OPPORTUNITY FOR PUBLIC COMMENT;
12	BUT AS WE GET INTO THE PROCESS OF CONSIDERING THE
13	GRANTS, WE'LL TAKE MOTIONS.
14	MR. TORRES: I'M GETTING USED TO THE NEW
15	PROCESS, SO PLEASE INCORPORATE MY REMARKS INTO THE
16	THREE MINUTES THAT I HAD.
17	DR. GROGAN: GOOD MORNING. I'D LIKE TO
18	FOLLOW WITH FURTHER COMMENTS ON THE PROPOSAL THAT
19	DR. D'LIMA HAS JUST DISCUSSED. MY NAME IS SEAN
20	GROGAN, AND I AM A SENIOR RESEARCH SCIENTIST AT
21	SCRIPPS HEALTH, AND I'D LIKE TO THE CDC INDICATES
22	THAT ONE IN FIVE PEOPLE, ONE IN FIVE ADULTS SUFFER
23	FROM THE DEBILITATING AND PAINFUL DISEASE CALLED
24	OSTEOARTHRITIS. THIS HAS A \$128 BILLION PRICE TAG
25	ON THE NATIONAL ECONOMY.
	4.4

1	THESE PATIENTS NOT ONLY HAVE A REDUCED
2	QUALITY OF LIFE, BUT ALSO HAVE AN INCREASED
3	INCIDENCE OF HEART DISEASE, DIABETES, AND OBESITY
4	BASED UPON IMMOBILITY. AT PRESENT THERE ARE NO
5	THERAPIES FOR THE TREATMENT OF OSTEOARTHRITIS OR
6	EVEN TO SLOW IT DOWN.
7	I'VE BEEN WORKING IN THIS FIELD FOR THE
8	PAST 16 YEARS TO DEVELOP THERAPIES FOR CARTILAGE
9	REPAIR OF DAMAGED AND DISEASED TISSUE, AND THIS IS
10	THE FIRST TIME THAT WE HAVE GREAT HOPE TO TRANSLATE
11	THIS FROM THE LAB INTO THE CLINIC TO HELP MILLIONS
12	OF PEOPLE.
13	WE ARE GRATEFUL TO THE PREVIOUS FUNDING
14	THAT WE RECEIVED IN 2009 FROM CIRM WHERE WE
15	DEVELOPED CHONDROPROGENITOR CELLS FROM AN EMBRYONIC
16	STEM CELL SOURCE, AND WE COMPARED THESE TO OTHER
17	HUMAN CELLS IN THE CLINIC AND ALSO COMPARED TO THOSE
18	IN DEVELOPMENT. AND OUR CHONDROPROGENITOR CELLS
19	OUTPERFORMED ANY OF THESE TREATMENT OPTIONS.
20	WE HAVE BEEN IN CONTACT WITH THE FDA TO GO
21	THROUGH THEIR REGULATORY PROCESS, AND WE BELIEVE
22	THAT WE ARE VERY CLOSE TO PRE-IND. AS WELL, WE HAVE
23	PEOPLE ON OUR TEAM THAT ARE MEMBERS OF THE ADVISORS
24	TO THE FDA FOR CELLULAR AND DRUG TREATMENT.
25	WE NEED TO TREAT DAMAGED CARTILAGE BEFORE
	4.5

1	IT PROGRESSES TO OSTEOARTHRITIS. WE HAVE CONVINCING
2	DATA. WE HAVE A TEAM THAT HAS CLINICAL EXPERIENCE
3	TO TRANSLATE BASIC TECHNOLOGIES. WE HAVE MATCHING
4	FUNDS, AND WE HAVE PATIENTS ASKING US TO DEVELOP
5	THIS TREATMENT. WE NEED TO KEEP THE MOMENTUM WHILE
6	WE HAVE OUR TEAM IN PLACE AND WHILE WE HAVE MATCHING
7	FUNDS.
8	SO I APPEAL TO THE COMMITTEE TO CONSIDER
9	GRANTING THIS APPLICATION. THANK YOU.
10	DR. HELMS: MY NAME IS JILL HELMS. I'M A
11	PROFESSOR AT STANFORD AND THE PI OF THE GRANT
12	PC1-08105. WE'VE NEVER MET, BUT YOU AND I HAVE
13	WORKED TOGETHER FOR THE LAST MANY YEARS BUILDING A
14	PROGRAM. WE BEGAN WITH AN EARLY TRANSLATIONAL
15	AWARD. WE GOT THE FIRST EVER BRIDGE FUNDING FROM
16	THIS BOARD, AND WE'VE ALSO GOTTEN EXTRAORDINARY
17	SUPPLEMENTAL FUNDING FROM CIRM.
18	WE'RE NOW ARRIVING AT THE PRECIPICE OF A
19	THERAPY. TOGETHER WITH SOME REALLY EXCEPTIONAL CIRM
20	STAFF, WE BUILT A PROGRAM THAT ADDRESSES A LARGE AND
21	UNMET NEED OF SKELETAL HEALING IN THE ELDERLY.
22	WE'RE NOW ON THE FINAL APPROACH TO AN IND. THE
23	PROPOSAL IN FRONT OF YOU HAS COMMERCIAL GRADE
24	MILESTONES. IT'S BEEN VETTED BY A PREMIER
25	CALIFORNIA VENTURE FUND. IT'S BEEN ANALYZED BY
	46

1	EXTERNAL CONSULTANTS ON EVERY LEVEL, AND IT'S GUIDED
2	BY A WORLD-CLASS SCIENTIFIC ADVISORY BOARD.
3	I'M STANDING BEFORE YOU NOW BECAUSE WE'RE
4	ON THE CUSP OF FUNDING. EACH GRANT APPLICATION
5	BRINGS WITH IT NEW REVIEWERS AND A NEW SET OF
6	QUESTIONS. AS A SCIENTIST, I HOLD PEER REVIEW
7	PROCESS IN THE HIGHEST REGARD, BUT BOTH YOU ON THIS
8	BOARD AND I RECOGNIZE THAT THE PROCESS IS NOT
9	PERFECT.
10	IN NEARLY EVERY ICOC MEETING YOU CHOOSE TO
11	FUND SOME TIER II PROJECTS, AND THE CENTRAL
12	RATIONALE HAS BEEN THAT THE MEDIAN SCORE OF THE
13	PROPOSAL IS 75 OR HIGHER AND THAT THE MAJORITY OF
14	REVIEWERS PLACED THE GRANT IN TIER I. OUR PROPOSAL
15	FULFILLS BOTH OF THESE RIGOROUS CRITERIA.
16	THE CRITICISMS OF OUR PROPOSAL WERE NOT
17	ABOUT EFFICACY OR APPROACH, ABOUT CMC, OR MECHANISM
18	OF ACTION. INSTEAD THEY CENTERED AROUND THE NEED TO
19	HAVE A REGULATORY EXPERT AS PART OF OUR TEAM AND THE
20	SUGGESTION THAT WE FOCUS ON A SINGLE LEAD
21	INDICATION.
22	SOME REVIEWERS SIMPLY MISSED THE FACT THAT
23	WE HAVE A REGULATORY COMPANY AS PART OF OUR TEAM.
24	AND AS FOR SELECTING A LEAD INDICATION AND
25	SCHEDULING A PRE-IND MEETING, YOU ALREADY SAW IN THE

1	PRESENTATION BEFORE THIS THAT THEY ARE WITHIN SCOPE
2	OF THIS RFA, AND THEY'RE BOTH A PART OF OUR EXISTING
3	MILESTONE PLAN.
4	THE RFA ALSO STATES THAT PROGRAMS THAT
5	BRING IN 25-PERCENT EXTERNAL CO-FUNDING WILL BE
6	PRIORITIZED. WE HAVE A CO-FUNDER. IT'S BACKED BY
7	AVALON VENTURES, A PREMIER CALIFORNIA VENTURE FUND.
8	AND THEIR COMMITMENT IS WELL IN EXCESS OF 25
9	PERCENT. THEIR CO-FUNDING IS A DIRECT DEMONSTRATION
10	OF CIRM'S MISSION TO COMMERCIALIZE STEM CELL
11	TECHNOLOGIES, AND WE RESPECTFULLY REQUEST THAT THE
12	ICOC APPLY THIS PRIORITY.
13	NOW, I'M FULLY AWARE OF THE RECOMMENDATION
14	THAT WE REAPPLY THROUGH CIRM 2.0, BUT WE'VE DONE THE
15	CALCULATIONS. A PRE-IND MEETING IS REQUIRED FOR
16	THOSE APPLICATIONS, SO IF WE FOLLOW THIS ADVICE,
17	IT'S HITTING PAUSE BUTTON FOR TEN MINUTES.
18	I HAVE A DEEPLY PERSONAL CONNECTION TO
19	DEVELOPING A STEM CELL THERAPY FOR CALIFORNIANS AND
20	THE WORLD. I URGE YOU TO VOTE IN FAVOR OF OUR
21	PROPOSITION.
22	DR. MADIGAN: MY NAME IS SANDY MADIGAN,
23	AND I'M THE FOUNDING CEO OF ANKASA REGENERATIVE
24	THERAPEUTICS. ANKASA IS OSTENSIBLY THE CO-FUNDER
25	FOR DR. HELMS' PROJECT. PERHAPS NOW WE'RE THE ONLY

1	FUNDER. I GUESS WE'LL SEE.
2	SO CO-FUNDING WITH A COMMERCIAL PARTNER
3	HAS BEEN A NOTED INTEREST OF CIRM, AND I'M GOING TO
4	TALK A BIT ABOUT WHY US AND WHY THAT'S IMPORTANT.
5	AFTER FOLLOWING DR. HELMS' WORK FOR ABOUT
6	FOUR YEARS NOW, KEVIN KINSELLA, WHO IS THE FOUNDING
7	AND MANAGING PARTNER OF AVALON VENTURES, DECIDED
8	THIS WAS THE TIME TO STEP IN. IT'S IMPORTANT, AS A
9	LOT OF PEOPLE TELL YOU ALREADY, TO PROVIDE THESE
10	THERAPEUTIC OPTIONS FOR ELDERLY PATIENTS, ESPECIALLY
11	IN LIGHT OF DR. HELM' TECHNOLOGY. SO MR. KINSELLA
12	HAS A SUCCESSFUL HISTORY OF DEVELOPING COMPANIES
13	OVER 30 YEARS IN CALIFORNIA AND BEYOND.
14	HE'S ALSO WORKED WELL WITH NONPROFIT
15	ORGANIZATIONS, AND THIS IS EVIDENCED BY THE
16	SUCCESSFUL DEVELOPMENT OF KALYDECO, A NEW CYSTIC
17	FIBROSIS DRUG THAT WAS WORKED OUT IN PARTNERSHIP
18	WITH THE CYSTIC FIBROSIS FOUNDATION. CALLED THE
19	MOST IMPORTANT NEW DRUG OF 2012 BY FORBES, IT WAS
20	DEVELOPED WITH SUPPORT TO THE TUNE OF \$150 MILLION
21	OVER 15 YEARS FROM THE CF FOUNDATION TO TWO
22	COMPANIES, AURORA BIOSCIENCES AND THEIR ACQUIRER,
23	VEDTEV DUADMACEUTICALS IMPORTANTIV CE
	VERTEX PHARMACEUTICALS. IMPORTANTLY, CF
24	FOUNDATION'S SALE OF THIS ROYALTY STREAM RECENTLY

1	BY WARDEN AS A DEAL THAT WILL CHANGE PHILANTHROPY.
2	SO WHY DO I MENTION THESE COMPANIES?
3	KEVIN KINSELLA WAS PERSONALLY THE FOUNDER AND
4	FINANCIER OF BOTH OF THOSE COMPANIES THAT BENEFITED
5	FROM THE CF FOUNDATION'S WORK. SO CLEARLY HE'S AN
6	EMINENT PARTNER IN THESE AREAS WITH NOT-FOR-PROFIT
7	ORGANIZATIONS.
8	SO IF CIRM IS REALLY SERIOUS ABOUT RAPIDLY
9	DEVELOPING DRUGS TO HELP PATIENTS, THERE'S NO BETTER
10	CO-FUNDER THAN KEVIN KINSELLA AND AVALON VENTURES.
11	SO I'D LIKE TO MOVE ON TO THE REVIEW
12	PROCESS A BIT. IT'S SUGGESTED THAT THE RFA HAS
13	BECOME A LITTLE BIT OF A MOVING TARGET. I THINK
14	WE'VE SEEN PRESENTATIONS THAT SHOWED WHAT THE
15	TARGETS WERE, AND IT SEEMS THAT DURING THE REVIEW
16	THAT TARGET HAS BEEN MOVED. FOR EXAMPLE, DR. HELMS
17	ALREADY NOTED THAT HER APPLICATION WAS CRITICIZED
18	FOR HAVING TWO INDICATIONS RATHER THAN ONE.
19	SELECTED INDICATION IS IN SCOPE; THEREFORE, IT'S
20	UNFAIRLY PUNITIVE TO APPLY CRITERIA THAT DIDN'T
21	EXIST.
22	SIMILARLY, THE APPLICATION WAS CRITICIZED
23	FOR LACK OF FDA ENGAGEMENT. AGAIN, THAT'S IN SCOPE.
24	AND, AGAIN, THAT'S AN APPLIED CRITERIA THAT DID NOT
25	EXIST AT THE TIME OF THE SUBMISSION.

1	MR. SHEEHY: DO WE HAVE ADDITIONAL PUBLIC
2	COMMENT?
3	DR. SCHUBERT: SPEAKING OF ARTHRITIS, HERE
4	COMES A VICTIM RIGHT NOW. TALKING ABOUT ALZHEIMER'S
5	NOW. MY NAME IS DAVE SCHUBERT. I'M ON THE FACULTY
6	AT THE SALK INSTITUTE. I'M SPEAKING IN SUPPORT OF A
7	PRECLINICAL DEVELOPMENT AWARD CALLED "STEM
8	CELL-BASED SMALL MOLECULE FOR ALZHEIMER'S DISEASE."
9	THE GOAL IS TO GET DO THE NECESSARY
10	PRECLINICAL WORK IN ORDER TO GET THIS DRUG INTO THE
11	CLINIC, DRUG CANDIDATE. AND THE BASIS FOR THIS
12	APPEAL IS FUNDING LEVEL. AS MY PREDECESSORS HAVE
13	TALKED ABOUT WITH THEIR SITUATION, WE HAVE NEW DATA
14	WHICH WAS NOT AVAILABLE AT THE TIME OF THE REVIEW
15	PROCESS AND THE MEDICAL NEED.
16	MEDICAL NEED, I THINK, IS OBVIOUS TO
17	EVERYONE. THERE'S NO EFFECTIVE THERAPY FOR
18	ALZHEIMER'S. THIS IS PROBABLY THE WORST MEDICAL
19	SITUATION IN THE UNITED STATES. IT'S THE THIRD
20	LEADING CAUSE OF DEATH. AND THERE ARE ABOUT 600,000
21	CASES IN CALIFORNIA THAT ARE KNOWN AND PROBABLY A
22	MILLION ALTOGETHER INCLUDING THE ONES THAT ARE NOT
23	KNOWN.
24	WE ARE CURRENTLY JUST FINISHING THE THIRD
25	YEAR OF AN EARLY TRANSLATIONAL RESEARCH GRANT WHICH
	רי

1	WAS GENEROUSLY FUNDED BY CIRM, AND THE GOAL OF THIS
2	WAS TO USE HUMAN ES CELLS AS A BASIS, ES-DERIVED
3	NEURAL PRECURSOR CELLS AS A BASIS FOR SCREENING DRUG
4	CANDIDATES THAT STIMULATE THE PRODUCTION OF NERVE
5	CELLS IN THE NEW NERVE CELLS IN THE HUMAN BRAIN
6	AND ALSO IS VERY NEUROPROTECTIVE, WHICH IS REQUIRED.
7	AND SO THIS WAS WE HAVE MADE A GREAT
8	DEAL OF PROGRESS AS DOCUMENTED IN THE PROGRESS
9	REPORTS OVER THE YEARS. WE THINK THIS PROGRAM WAS
10	ENORMOUSLY SUCCESSFUL. WE CAME UP WITH ONE
11	CANDIDATE WHICH STIMULATES THE PRODUCTION OF HUMAN
12	NERVE CELLS IN CULTURE SITUATION. AND IT'S
13	EXTREMELY NEUROPROTECTIVE. THIS WORKS EXCEPTIONALLY
14	WELL IN VARIOUS ANIMAL MODELS OF ALZHEIMER'S
15	DISEASE, AND WE THINK IT'S ONE OF THE BEST
16	CANDIDATES, NOT THE BEST AD CANDIDATE ON THE MARKET.
17	SO AS PREVIOUS SPEAKERS HAVE SAID, WE'RE
18	IN THE SECOND TIER. THE FUNDING WAS MARGINAL. SO
19	THE BASIS OF THIS APPEAL IS NEW DATA. WE HAVE JUST
20	COMPLETED, AS DOCUMENTED IN THE LAST PROGRESS
21	REPORT, WHICH I DON'T THINK WAS AVAILABLE FOR THE
22	REVIEWERS OF THIS APPLICATION TO SEE, WE WERE ABLE
23	TO SHOW THAT THIS DRUG CANDIDATE RESTORES MEMORY IN
24	OLD MICE. IN OLD ALZHEIMER'S MICE, IT RESTORES IT
25	BACK TO THE LEVEL OF THEIR NON-AD CONTROLS AND
	E 2

1	ALSO WORKS IN TWO OTHER MODELS OF AD MICE. AS FAR
2	AS WE KNOW, THIS IS THE ONLY DRUG CANDIDATE THAT
3	ACTUALLY CAN WORK IN THIS TYPE OF MODEL.
4	AND WE'VE ALSO LEARNED IN THE LAST FEW
5	WEEKS ACTUALLY THE MOLECULAR TARGET OF THIS DRUG
6	WHICH IS A BIG HELP IN GETTING IT THROUGH THE IND
7	PHASE. SO WE HAVE SPENT THAT'S A SIGN TO STOP?
8	OKAY.
9	ANYWAY, CAN I GO ON FOR ANOTHER 30
10	SECONDS? NO. IT'S HARD OKAY.
11	SO WE ARE AT THIS MARGINAL LEVEL OF
12	FUNDING. THE MEDIAN SCORE WAS THE SAME AS SOME
13	FUNDED GRANTS. WE HAD THE HIGHEST SINGLE SCORE
14	WITHIN OUR GROUP. WE WERE A LOWER COST THAN OTHER
15	GROUPS, AND I THINK THE NEED IS TREMENDOUS, AND
16	NOBODY CAN ARGUE ABOUT THAT. THANK YOU.
17	DR. CASHMAN: I'M JOHN CASHMAN, PRESIDENT
18	OF HUMAN BIOMOLECULAR RESEARCH INSTITUTE IN SAN
19	DIEGO, A SMALL NONPROFIT RESEARCH INSTITUTE. I'M
20	SPEAKING HERE ON BEHALF OF MY TEAM. DANIEL IS GOING
21	TO SPEAK AFTER ME AND TALK A LITTLE BIT ABOUT THE
22	SCIENCE. NATE QUARRY, WHO IS AN INTERNATIONALLY
23	RECOGNIZED KICK FIGHTER FROM MMA AND IFC, IS GOING
24	TO SPEAK ABOUT HIS PERSONAL INTERACTIONS WITH SPINAL
25	DEGENERATION WHICH IS WHAT THE GRANT WAS ABOUT.

1	THE GRANT WAS ENTITLED "INDUCED STEM CELL
2	IMPLANTS FOR SPINAL FUSION," 08129.
3	I WAS HERE ABOUT A YEAR AGO BECAUSE I
4	RESPONDED TO THE TRANSLATIONAL APPLICATION, AND
5	DUANE ROTH WAS SITTING RIGHT OVER THERE. AND THE
6	GRANT GOT A SCORE OF ABOUT 65, WHICH PUT IT ON THE
7	CUSP. AND DUANE SAID, "WELL, CIRM WILL WORK WITH
8	YOU TO GET THAT OVER THE HUMP SO THAT IT CAN GET
9	FUNDED AND APPROVED." AND SO, IN FACT, ALONG THE
10	SAME LINES AS RANDY HAS TALKED ABOUT IN 2.0, WE
11	WORKED CLOSELY WITH CIRM. WE WORKED WITH GIL AND
12	PAT AND THEIR TEAM TO DESIGN THE EXPERIMENT TO
13	ADDRESS THE ISSUES, AND THERE WERE MAINLY TWO
14	ISSUES.
15	ONE WAS THE REVIEW THOUGHT THAT THERE WAS
16	NO MEDICAL NEED. WELL, THERE ARE 88,000 PEOPLE IN
17	CALIFORNIA THAT SUFFER FROM SPINAL DEGENERATION.
18	IT'S A MAJOR ISSUE. ABOUT HALF OR 44,000 DON'T SEEK
19	TREATMENT BECAUSE OF SOME OF THE REASONS THAT NATE
20	WILL EXPLAIN TO YOU.
21	NONETHELESS, WE WORKED VERY HARD, AND IT'S
22	VERY EXPENSIVE, AS SPEAKERS POINTED OUT BEFORE, TO
23	FUND RESEARCH IN A SMALL NONPROFIT THAT DOESN'T HAVE
24	ANY DISCRETIONARY FUNDS TO GET THE PRELIMINARY
25	RESULTS TO RESPOND TO THE REVIEWERS. NEVERTHELESS,
	r r

1	WE DID THAT, AND THE EXPERIMENTS WERE MASSIVELY
2	SUCCESSFUL. WE GREW HUMAN BONE CELLS IN RATS IN A
3	ROBUST FASHION THAT'S NEVER BEEN REPORTED BEFORE.
4	AS A PRE-IND ENABLING MODEL, THIS WAS A SPECTACULAR
5	RESULT.
6	SO THEN WE WENT BACK IN TO CIRM, AND WE
7	DID NOT GET INTO TIER II. WE GOT INTO TIER III.
8	THIS IS AFTER DILIGENTLY DOING THE EXPERIMENT THAT
9	CIRM AND THE TEAM OUTLINED FOR US TO DO AND THE
10	REVIEWERS REQUESTED.
11	SO THIS IS EXTREMELY FRUSTRATING. THE
12	LAST CONVERSATION I HAD WITH DUANE AT THE OAKLAND
13	AIRPORT ABOUT A YEAR AGO WAS, "JOHN, I REALLY LIKE
14	YOUR PROGRAM. I REALLY LIKE YOUR PRODUCT. IT'S
15	VERY FDA ENABLED. ALL THE COMPONENTS ARE EITHER FDA
16	APPROVED OR FDA CERTIFIED. IT'S ALL READY TO GO.
17	CIRM IS INTERESTED IN DEVELOPING PRODUCTS. THIS IS
18	A NEAR PRODUCT DEVELOPMENT THAT'S GOING TO ADDRESS A
19	MAJOR UNMET NEED IN CALIFORNIA."
20	ARE THOSE THE ANGELES BELLS? SO I'M HERE
21	TO REQUEST THAT THE BOARD RECONSIDER THIS.
22	CERTAINLY THE SYSTEM IS BROKEN. WE WERE TOLD IF WE
23	DID THIS, WE WOULD HAVE OBVIOUSLY A VERY GOOD CHANCE
24	TO GET FUNDED. I REQUEST A RE-REVIEW. THANK YOU.
25	DR. RYAN: GOOD MORNING. DANIEL RYAN,

1	PRINCIPAL SCIENTIST AT HBRI.
2	JUST TO EXPAND ON JOHN'S COMMENTS, AT HBRI
3	WE FEEL WE'VE HAD A BREAKTHROUGH IN USING HUMAN STEM
4	CELL IMPLANTS FOR SPINAL FUSION PROCEDURES. THIS
5	TECHNOLOGY WAS DEVELOPED THROUGH A SERIES OF STEPS.
6	FIRST, WE DISCOVERED THAT ONE OF THE MINOR
7	COMPONENTS OF THE FOOD SPICE TURMERIC WAS POTENTLY
8	ABLE TO PROMOTE HUMAN STEM CELLS TO ADOPT THE BONE
9	LINEAGE. WE ISOLATED THAT MOLECULE, PREPARED IT,
10	AND PROVED THAT OUT.
11	SECONDLY, WE USED THE SMALL MOLECULES IN A
12	NEW WAY IN STEM CELL RESEARCH.
13	MR. SHEEHY: EXCUSE ME. I'M JUST VERY
14	UNCOMFORTABLE WITH I THINK IF THINGS ARE GOING TO
15	BE HANDED TO BOARD MEMBERS, THEY SHOULD BE HANDED TO
16	TEAM MEMBERS TO HAND OUT TO THE BOARD. I JUST FIND
17	IT VERY UNCOMFORTABLE. I JUST DON'T THINK THAT'S
18	GENERALLY HOW WE MAKE THINGS AVAILABLE TO BOARD
19	MEMBERS, AND I THINK IT IS DISCONCERTING TO ME.
20	DR. CASHMAN: MR. CHAIRMAN, WE'RE JUST
21	TRYING TO MAKE IT
22	MR. SHEEHY: THAT'S FINE. WE'LL ACCEPT
23	ANY DOCUMENTS, BUT THEY REALLY SHOULD GO TO TEAM
24	MEMBERS TO HAND TO THE BOARD. YOU CAN PROCEED.
25	DR. RYAN: THE SECOND ASPECT OF THIS
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1	TECHNOLOGY WAS HOW WE USE THESE SMALL MOLECULES FROM
2	TURMERIC. STEM CELLS NORMALLY NEED A CARRIER SO
3	THAT THEY'RE MAINTAINED AT THE SITE OF IMPLANTATION.
4	WE FOUND THAT WHEN WE TREATED THE STEM CELLS ON A
5	BONE MATRIX WITH THE SMALL MOLECULES, THE AMOUNT OF
6	BONE GROWTH FAR EXCEEDED OUR EXPECTATIONS FROM THE
7	COMPOUND ALONE. WE TESTED THIS IN A RAT MODEL OF
8	SPINAL FUSION. THE GOAL IS TO FORM HUMAN BONE IN
9	RAT. AND THE HANDOUTS THAT YOU'VE RECEIVED OR MAY
10	RECEIVE SHOW SOME OF THE RESULTS OF THIS STUDY.
11	USING RADIOGRAPHIC ANALYSIS, WE SAW ROBUST
12	SPINAL FUSION AND ACHIEVED 95 PERCENT OF THE
13	IMPLANTS ACHIEVED FUSION. THIS FAR EXCEEDED OUR
14	EXPECTATIONS IN THAT ALL PRIOR REPORTS OF HUMAN STEM
15	CELL IMPLANTS IN THIS MODEL FORM VERY LITTLE BONE.
16	HISTOLOGY AND MATRICES CONFIRM ROBUST HUMAN BONE
17	GROWTH WITH ADDITIONAL VASCULARIZATION AND IN-GROWTH
18	OF BLOOD VESSELS TO SUPPORT THAT TISSUE. IN EFFECT,
19	BONE GROWTH IS GREATER. AND SINCE WE STIMULATE THE
20	CELLS ON THE MATRIX, PRACTICALLY WE'RE STIMULATING
21	WHAT GOES INTO THE ACTUAL IMPLANT. SO IT'S REDUCED
22	TO PRACTICE VERY EFFICIENTLY.
23	WE FEEL THERE ARE A LOT OF POTENTIAL
24	OPPORTUNITIES FOR ROBUST STEM CELL IMPLANTS FOR
25	SPINAL FUSION PATIENTS. THE LEADERS IN THIS AREA
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1	ARE OVERSEAS, BUT WE HAVE THE TECHNOLOGY WITH YOUR
2	SUPPORT TO DEVELOP A BETTER, FIRST IN CLASS OR BEST
3	IN CLASS THERAPEUTIC RIGHT HERE IN CALIFORNIA. SO
4	WITH THAT, I WOULD LIKE TO ALSO REQUEST A REVIEW OF
5	THE STEM CELL IMPLANT APPLICATION. THANK YOU.
6	MR. SHEEHY: HAVE ADDITIONAL PUBLIC
7	COMMENT?
8	MR. QUARRY: MY NAME IS NATHAN QUARRY. I
9	AM THE PATIENT. A FEW YEARS AGO MY CAREER WAS AS A
10	PROFESSIONAL FIGHTER. I FOUGHT FOR THE UFC, THE
11	ULTIMATE FIGHTING CHAMPIONSHIP. IF YOU'VE EVER RAN
12	FLIPPING THROUGH THE CHANNELS AND YOU SEE TWO
13	KNUCKLEHEADS BEATING THE HELL OUT OF EACH OTHER IN A
14	CAGE, THAT WAS ME. AND I WAS STRUCK DOWN WITH
15	DEGENERATIVE DISK DISEASE IN MY LUMBAR. MY L-2/3
16	HAD COMPLETELY COLLAPSED, AND I WENT FROM FIGHTING
17	FOR THE WORLD TITLE TO NOT BEING ABLE TO PICK UP MY
18	LITTLE GIRL.
19	FORTUNATELY THE TECHNOLOGY HAD PROGRESSED
20	WHERE I COULD LOOK INTO GETTING A SPINAL FUSION. AS
21	MY SURGEON TOLD ME, THE GOLD STANDARD WAS TO HARVEST
22	BONE OFF MY HIP. WELL, JUST A COUPLE WEEKS AFTER MY
23	SURGERY, MY BACK WAS FEELING SO MUCH BETTER, THE
24	INFLAMMATION HAD GONE DOWN, BUT FOR MONTHS MY HIP
25	WAS ACHING TO THE POINT WHERE I HAD TO TAKE EXTENDED
	50

PAIN MEDICATIONS JUST TO DEAL WITH THAT. AND NOW I
REPRESENT THOUSANDS AND MILLIONS OF PEOPLE THAT ARE
SUFFERING FROM BACK PAIN.
AS WE GO THROUGH OUR LIVES, 80 PERCENT OF
US WILL SUFFER FROM BACK PAIN. SO I HEAD UP A
PATIENT SUPPORT PROGRAM CALLED THE BETTER WAY BACK.
I HOPE TO EDUCATE THESE PATIENTS. AND THE FEAR THAT
I HEAR ALL THE TIME, WHAT ABOUT THE PAIN? WHAT
ABOUT THIS HARVESTING OF MY BONE? BECAUSE IT'S
HORRIBLY PAINFUL, AND IT LEADS TO MORE DRUG USE,
MORE TIME IN THE HOSPITAL. WE'RE IN AN EPIDEMIC
RIGHT NOW AS PRESCRIPTION DRUG ABUSE HAS SURPASSED
ILLEGAL DRUG USE AS FAR AS DEATH AND TIME LOST. AND
WHEN I HEAR ABOUT THIS TECHNOLOGY, I GET EXCITED.
I'M EXCITED TO BE HERE REPRESENTING ALL OF THESE
PATIENTS. AND I THINK ABOUT THIS TECHNOLOGY THAT
WILL HELP DRIVE AMERICA BACK TO THE FOREFRONT IN
THESE AREAS.
SO MANY PEOPLE ARE SUFFERING FROM BACK
PAIN, AND WE NEED TO CHANGE THE PUBLIC MISCONCEPTION
THAT THIS IS A LIFE SENTENCE, THAT YOU JUST HAVE TO
LIVE WITH THIS. AS I WAS TOLD, IF YOU GET BACK
SURGERY, YOUR LIFE IS OVER. YOU'LL NEVER FIGHT
AGAIN. YOU'LL NEVER WORK AGAIN. FORGET ABOUT
PICKING UP YOUR LITTLE GIRL. AND I OPTED TO HAVE MY
60

1	SURGERY.
2	NOW KNOWING THAT I HAVE DEGENERATIVE DISK
3	DISEASE, OTHER DISKS OF MINE ARE COLLAPSING, AND I'M
4	NOT LOOKING FORWARD TO THOSE DAYS WHEN THE DOCTOR
5	SAYS WE CAN HARVEST BONE OFF OF YOUR HIP TO GET YOU
6	BACK, GIVE YOU THAT GOLD STANDARD. NO. I PREFER TO
7	WAIT FOR THIS TECHNOLOGY TO TAKE PLACE AS EVEN WITH
8	THAT, WITH THE BONE GRAFTED OFF MY HIP, THAT'S A
9	SIX-MONTH PROCESS FOR THE BONE TO COME TOGETHER AND
10	HEAL MY BACK; WHEREAS, WITH THIS TECHNOLOGY SPEEDING
11	UP THAT PROCESS, LETTING ME GET BACK TO WORK,
12	LETTING ALL OF AMERICA GET OVER THESE ISSUES, GET
13	BACK TO WORK, TAKE CARE OF THEIR FAMILIES, GET OFF
14	OF THESE PAIN MEDICATIONS THAT ARE KILLING OUR
15	COUNTRY. THAT'S WHAT I'M EXCITED TO SEE, AND
16	HOPEFULLY THAT THIS TECHNOLOGY WILL CONTINUE MOVING
17	FORWARD. THANK YOU.
18	MR. SHEEHY: THANK YOU. DO WE HAVE
19	ADDITIONAL PUBLIC COMMENT? OKAY. SO I THINK, DR.
20	SAMBRANO, WILL YOU REITERATE THE TEAM
21	RECOMMENDATIONS?
22	DR. SAMBRANO: SURE. I WANTED TO JUST
23	POINT OUT THAT WE HAD SOME FORMAL APPEALS THAT WERE
24	SUBMITTED, SO THERE WERE THREE. THERE'S A MEMO THAT
25	I PROVIDED THAT SUMMARIZES OUR FINDINGS ON THOSE.

1	TWO OF THOSE WERE DENIED. ONE APPLICATION, WHICH IS
2	PC1-08132, WHICH IS IN TIER III, IS BEING DEFERRED.
3	THAT'S THE LAST ONE LISTED ON THE MEMO. SO THAT ONE
4	WILL NOT BE CONSIDERED AT TODAY'S BOARD MEETING.
5	MR. SHEEHY: THANK YOU, DR. SAMBRANO.
6	PERHAPS IF WE COULD GET JUST A REITERATION WHAT THE
7	TEAM'S RECOMMENDATIONS WERE FOR TIER II. I THINK
8	SOME OF IT WAS KIND OF ALLUDED TO IN WHAT WE HEARD
9	FROM THE PUBLIC COMMENT IS THAT IN SOME WAYS THERE
10	WERE FLAWS THAT WERE IN OUR PREVIOUS SYSTEM THAT
11	HAVE NOW BEEN ADDRESSED THROUGH CIRM 2.0 AND THE
12	OPPORTUNITY TO COME BACK IN IN CIRM 2.0 IN
13	SITUATIONS WHERE IF YOU DO END UP IN TIER II, YOU
14	GET CLEAR DIRECTION ON WHAT NEEDS TO BE FIXED IN
15	YOUR PROJECT, YOU FIX THOSE, AND YOU CAN COME BACK
16	IN A REALLY SHORT TIME. WE GET A BETTER TURNAROUND,
17	WE GET BETTER INFORMATION GOING BACK AND FORTH
18	BETWEEN THE REVIEW TEAM AND THE APPLICANTS.
19	AND I THINK SOME OF THE THINGS THAT HAVE
20	BEEN ALLUDED TO LIKE EVERY TIME YOU GO THROUGH A
21	REVIEW, IT'S A NEW SET OF REVIEWERS. THERE'S GOING
22	TO BE MORE CONSISTENCY AND COHERENCE BETWEEN THE
23	REVIEWS. THAT'S JUST PART OF THE FEATURES, BUT I'LL
24	HAND IT TO DR. MILLS.
25	DR. MILLS: SO THAT'S CORRECT. COUPLE OF
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1	THINGS TO POINT OUT ABOUT THE REVIEW PROCESS. ONE,
2	THE REVIEW PROCESS IN ITSELF IS SUBOPTIMAL IN ITS
3	CURRENT FORM, WHICH IS WHY WE MADE A LOT OF THOSE
4	CHANGES. I THINK THAT'S AN IMPORTANT PART OF IT TO
5	POINT OUT.
6	THE SECOND THING IS THE FEEDBACK MECHANISM
7	FROM THE REVIEW PROCESS IS ALSO SUBOPTIMAL. SO IT'S
8	THE BEST UNDER THIS CURRENT SYSTEM, IT'S THE BEST
9	WE COULD DO TO PROVIDE FEEDBACK, BUT THE TOTALITY OF
10	THE DECISIONS THAT THE REVIEWERS THE TOTALITY OF
11	THE INFORMATION THE REVIEWERS CONSIDERED AND THEIR
12	REASONS FOR WHY THEY VOTED THE WAY THEY VOTED ISN'T
13	ONE-TO-ONE COMMUNICATED BACK TO THE APPLICANTS. SO
14	IT WOULD BE A FALSE STATEMENT TO SAY THE ONLY
15	PROBLEMS THE REVIEWERS HAD WITH THE APPLICATION WERE
16	X, Y, OR Z. WE DON'T UNDER THE CURRENT SYSTEM HAVE
17	A WAY OF PROVIDING ALL OF THAT INFORMATION BACK.
18	THAT SAID, THE RECOMMENDATION TO NOT FUND
19	ANYTHING NOT IN TIER I WAS BASED ON TWO
20	CONSIDERATIONS. ONE IS NOT FUND ANYTHING NOT IN
21	TIER I WERE BASED ON TWO CONSIDERATIONS. ONE, ALL
22	OF THE THINGS NOT IN TIER I CLEARLY HAD, WHILE SOME
23	OF THEM ARE PROMISING, HAD THINGS THAT COULD BE MADE
24	BETTER ABOUT THEM. THEY WERE NOT OPTIMAL GRANTS.
25	IT WAS NOT AN UNANSWERABLE CALL. THERE WERE THINGS
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1	IN TIER I THAT DID VERY WELL. AND THESE
2	APPLICATIONS DID NOT HAVE THAT.
3	SECONDLY, THE PROXIMITY OF THE NEXT
4	OPPORTUNITY TO APPLY IS RELATIVELY CLOSE COMPARED TO
5	PREVIOUSLY WHERE THE NEXT OPPORTUNITY MIGHT NOT COME
6	AROUND FOR 18 MONTHS OR SO. SINCE OUR GOAL HERE IS
7	TO LAUNCH PROJECTS THAT HAVE THE BEST OPPORTUNITY
8	ULTIMATELY TO GO ON AND SUCCESSFULLY IMPACT A
9	PATIENT AND NOT JUST TO GET THINGS LAUNCHED AS
10	QUICKLY AS WE CAN, OUR RECOMMENDATION WAS TO NOT
11	FUND THESE APPLICATIONS AND THEN TO GIVE THEM
12	CONSIDERATION FOR REAPPLICATION IN THE TWO
13	MECHANISMS WHICH WERE STATED.
14	AND LASTLY, UNDER WHICH A PROCESS WE HOPE
15	IS MUCH BETTER IN THAT IF A PROGRAM ENDS UP IN WHAT
16	IS TODAY TIER II, THEY WOULDN'T BE FACED WITH AN
17	UP-OR-DOWN DECISION, BUT THEY'D BE FACED WITH MAKING
18	THESE CORRECTIONS QUICKLY AND LET'S GET IT BACK INTO
19	THE PROCESS FOR ANOTHER RE-REVIEW IN 30 DAYS. WE'RE
20	LOOKING AT WHAT TODAY MIGHT BE A 73 IN 30 DAYS COULD
21	BE A 95, WE COULD ALL BE HAPPY WITH IT AND MOVE ON.
22	MR. SHEEHY: THANK YOU, DR. MILLS.
23	DR. JUELSGAARD: YES. DR. MILLS, JUST A
24	COUPLE OF THINGS THAT I'VE NOTED IN KIND OF LOOKING
24 25	COUPLE OF THINGS THAT I'VE NOTED IN KIND OF LOOKING THROUGH THESE MATERIALS AND LISTENING TO THE

1	COMMENTS ABOUT OUR PROCESS. THESE ARE PROCESS
2	QUESTIONS, I GUESS.
3	SO ONE OF THEM IS THE SCORING AND USING AN
4	AVERAGE SCORE VERSUS A MEDIAN SCORE. SO MEDIAN IS
5	THE MIDDLE POINT, RIGHT; WHEREAS, AVERAGE TAKES
6	EVERYTHING INTO ACCOUNT. AND IN PARTICULAR WHEN YOU
7	SEE THIS KIND OF STANDARD DEVIATION GOING ON IN THE
8	TIER II, WITH TWO OF THEM THERE WAS A STANDARD
9	DEVIATION OF TEN AND A COUPLE THAT WAS A STANDARD
10	DEVIATION OF NINE, SUGGESTING THERE'S THIS BROAD
11	RANGE OF OPINION GOING ON WITH A LOW SCORE, FOR
12	EXAMPLE, 45 IN ONE OF THEM VERSUS A HIGHER SCORE OF
13	85.
14	AND SO I DON'T KNOW QUITE HOW THAT HAPPENS
14 15	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT
15	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT
15 16	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE
15 16 17	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE  THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M
15 16 17 18	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE  THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M  WONDERING JUST ABOUT THE WAY WE DO THINGS. DO WE
15 16 17 18 19	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE  THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M  WONDERING JUST ABOUT THE WAY WE DO THINGS. DO WE  USE AN AVERAGE SCORE, STAY WITH AN AVERAGE SCORE, DO
15 16 17 18 19 20	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE  THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M  WONDERING JUST ABOUT THE WAY WE DO THINGS. DO WE  USE AN AVERAGE SCORE, STAY WITH AN AVERAGE SCORE, DO  WE USE A MEDIAN SCORE? DO WE THROW OUT THE TOP
15 16 17 18 19 20 21	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE  THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M  WONDERING JUST ABOUT THE WAY WE DO THINGS. DO WE  USE AN AVERAGE SCORE, STAY WITH AN AVERAGE SCORE, DO  WE USE A MEDIAN SCORE? DO WE THROW OUT THE TOP  SCORE AND THE BOTTOM SCORE AND USE THE AVERAGE OR
15 16 17 18 19 20 21	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT  APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE  THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M  WONDERING JUST ABOUT THE WAY WE DO THINGS. DO WE  USE AN AVERAGE SCORE, STAY WITH AN AVERAGE SCORE, DO  WE USE A MEDIAN SCORE? DO WE THROW OUT THE TOP  SCORE AND THE BOTTOM SCORE AND USE THE AVERAGE OR  MEDIAN OF THE REMAINING SCORES TO TRY AND CREATE A
15 16 17 18 19 20 21 22	IN A REVIEW LIKE THIS. THERE ARE SOME PEOPLE THAT APPARENTLY REALLY LIKE THE PROJECT AND SOME PEOPLE THAT REALLY DON'T LIKE THE PROJECT. AND SO I'M WONDERING JUST ABOUT THE WAY WE DO THINGS. DO WE USE AN AVERAGE SCORE, STAY WITH AN AVERAGE SCORE, DO WE USE A MEDIAN SCORE? DO WE THROW OUT THE TOP SCORE AND THE BOTTOM SCORE AND USE THE AVERAGE OR MEDIAN OF THE REMAINING SCORES TO TRY AND CREATE A LITTLE MORE UNIFORMITY? I DON'T KNOW. SO THAT'S

1	AMPLIFIED FOR ME IN WHAT I SEE LINED IN GRAY HERE.
2	THE SECOND IS A COMMENT ON ONE OF THE
3	FROM GIL TO US, AND IT HAD TO DO WITH ONE OF THE
4	PROJECTS. AND AT THE VERY END OF THE LITTLE
5	PARAGRAPH, THIS IS ON 081000, IT SAYS, "AN APPEAL
6	POLICY DEFINES MATERIAL NEW INFORMATION," AND THEN
7	IT SAYS, "AS A MANUSCRIPT THAT HAS BEEN PEER
8	REVIEWED AND PUBLISHED OR ACCEPTED FOR PUBLICATION."
9	SO I CAN UNDERSTAND THAT PERHAPS IN THE ACADEMIC
10	WORLD. I HAVE A MUCH HARDER TIME UNDERSTANDING THAT
11	IN THE COMMERCIAL WORLD.
12	THERE'S A LOT OF WORK THAT GETS DONE THAT
13	COMMERCIAL ENTITIES COULD HAVE VERY MATERIAL NEW
14	INFORMATION AND THAT THEY WOULD NOT PUBLISH AND
15	WOULD NOT CHOOSE TO PUBLISH BECAUSE IT'S OF
16	COMPETITIVE ADVANTAGE. AND SO I JUST WOULD ASK US
17	TO THINK ABOUT WHETHER THAT SHOULD BE A PART OF OUR
18	POLICY AROUND MATERIAL NEW INFORMATION.
19	DR. MILLS: I AGREE WITH EVERYTHING YOU'VE
20	SAID. AND WHAT I CAN TELL YOU IS THAT THIS IS THE
21	LAST REVIEW WE'RE HOLDING UNDER THE 1.0 PROCESS FOR
22	THAT REASON. THE PROCESS ITSELF IS SUBOPTIMAL. ALL
23	OF THE ISSUES YOU RAISED WITH REGARDS TO SCORING,
24	THERE WOULDN'T EVEN BE A NEED FOR AN APPEAL UNDER
25	THE 2.0 PROCESS. AND SO I AGREE WITH ALL OF THOSE

1	THINGS AND SAY OUR RESPONSE TO THAT WAS 2.0, WHICH
2	STARTS ITS FIRST REVIEW ON MONDAY.
3	I COULD HAVE GIL TAKE YOU THROUGH THE
4	SCORING, BUT THE WAY WE'VE DONE SCORING HISTORICALLY
5	HAS USED THE MEAN SCORE. THAT'S WHAT IT'S BEEN.
6	DR. SAMBRANO: I MAY ALSO POINT OUT THAT,
7	REGARDING YOUR LATTER POINT, WE DO CONSIDER FOR
8	FOR-PROFIT ENTITIES DATA THAT IS CONFIDENTIAL OR
9	PROPRIETARY THAT DOES NOT NEED TO BE PUBLISHED.
10	THAT'S PART OF THE POLICY IN THIS PARTICULAR CASE.
11	IT IS AN ACADEMIC INSTITUTION; THEREFORE, THE
12	REQUIREMENT IS THAT IT'S PUBLISHED.
13	MR. SHEEHY: SO DO WE HAVE ANY OTHER
14	QUESTIONS BEFORE WE MOVE INTO CONSIDERATION OF THESE
15	GRANTS?
16	SO THE FIRST MOTION I WILL TAKE WILL BE TO
17	MOVE ANY APPLICATION FROM TIER III TO TIER I. SO
18	THAT WOULD BE FROM THE WHITE PART INTO THE GREEN
19	PART. OKAY. I SEE NO MOTIONS.
20	THE NEXT MOTION I WILL TAKE WILL BE TO
21	MOVE AN APPLICATION FROM TIER I TO TIER III. SO
22	SOMETHING OUT OF THE FUNDABLE CATEGORY, WHICH IS THE
23	GREEN BOX, DOWN INTO THE WHITE BOX, THE UNFUNDABLE.
24	NO MOTIONS THAT I SEE. SO THE NEXT STEP
25	IS TO CONSIDER MOTIONS THAT WOULD MOVE SOMETHING
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1	FROM TIER II, WHICH IS THE GRAY AREA, INTO TIER I.
2	DO I HAVE ANY MOTIONS?
3	DR. PRIETO: YES. I'D LIKE TO MAKE A
4	MOTION TO MOVE PC1-08128 FROM TIER II INTO TIER I.
5	MR. TORRES: WHICH ONE IS THAT?
6	DR. PRIETO: I'M SORRY. I MISREAD.
7	PC1-08086, "HUMAN STEM CELL-BASED DEVELOPMENT OF A
8	POTENT ALZHEIMER'S DRUG CANDIDATE" FROM TIER II INTO
9	TIER I.
10	MR. SHEEHY: DO I HAVE A SECOND FOR THAT
11	MOTION?
12	MS. MILLER: SECOND.
13	MR. SHEEHY: MS. MILLER SECONDS.
14	I THINK THE NEXT STEP NOW, COULD WE JUST
15	GET A BRIEF OVERVIEW OF THAT PARTICULAR APPLICATION
16	BY A MEMBER OF THE CIRM TEAM?
17	DR. KADYK: THIS IS APPLICATION PC1-08086.
18	AND THIS PROPOSAL IS FOCUSED ON THE DEVELOPMENT OF A
19	SMALL MOLECULE DRUG FOR ALZHEIMER'S DISEASE. THIS
20	CANDIDATE SMALL MOLECULE WAS IDENTIFIED THROUGH
21	SCREENING OF HUMAN EMBRYONIC STEM CELL-DERIVED
22	NEURAL PRECURSOR CELLS FOR COMPOUNDS THAT ARE BOTH
23	NEUROGENIC AND NEUROPROTECTIVE. AND AS YOU KNOW,
24	ALZHEIMER'S DISEASE IS A PROGRESSIVE
25	NEURODEGENERATIVE DISEASE, AND IT'S A MAJOR UNMET
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1	NEED.
2	FOR THIS PARTICULAR APPLICATION, THE
3	APPLICANT PROPOSES THAT THE CANDIDATE SMALL MOLECULE
4	COULD POTENTIALLY PROTECT EXISTING NEURONS IN THE AD
5	PATIENT BRAIN AND STIMULATE THE PRODUCTION OF NEW
6	NEURONS, THEREBY IMPROVING MEMORY AND COGNITION.
7	AND THE ACTIVITIES PROPOSED IN THE
8	APPLICATION INCLUDE IDENTIFICATION OF METABOLITES OF
9	THE CANDIDATE SMALL MOLECULE, PK STUDIES,
10	PHARMACOKINETIC STUDIES. AGAIN, THAT'S THE EFFECT
11	OF THE BODY ON THE DRUG. IDENTIFICATION OF THE DRUG
12	TARGET OR PATHWAY, DRUG SYNTHESIS, OFF-TARGET AND
13	SAFETY SCREENS, AND DOING AN EFFICACY STUDY IN AN
14	ANIMAL MODEL IN PREPARATION FOR A PRE-IND MEETING
15	WITH THE FDA.
16	WOULD YOU LIKE TO HEAR A SUMMARY OF THE
17	REVIEWERS' COMMENTS?
18	MR. SHEEHY: I THINK WOULD THAT BE HELPFUL
19	TO THE MAKERS.
20	DR. KADYK: SO UNDER SIGNIFICANCE AND
21	IMPACT, THE REVIEWERS AGREED THAT THIS PROPOSAL
22	ADDRESSES A SERIOUS UNMET MEDICAL NEED AND THAT, IF
23	IT'S SUCCESSFUL, COULD HAVE AN ENORMOUS IMPACT ON
24	THE TREATMENT OF ALZHEIMER'S DISEASE. AND THEY FELT
25	THAT THE TARGET PRODUCT PROFILE WAS HIGHLY

69

1	APPROPRIATE AND CONSISTENT WITH THE OBJECTIVES OF
2	THIS RFA AND OTHER ALZHEIMER'S DISEASE DRUG
3	DEVELOPMENT EFFORTS.
4	THE SCIENTIFIC RATIONALE, THE REVIEWERS
5	AGREED THAT THE PROPOSED THERAPEUTIC CANDIDATE HAS
6	DESIRABLE DRUG FEATURES BASED ON BOTH IN VITRO AND
7	IN VIVO DRUG STUDIES. FOR EXAMPLE, THEY APPRECIATED
8	IT'S PHARMACOLOGIC PROPERTIES AND THAT IT CAN
9	PENETRATE THE BLOOD-BRAIN BARRIER, WHICH IS
10	OBVIOUSLY NECESSARY FOR ALZHEIMER'S TREATMENT.
11	THEY THOUGHT THE SCIENTIFIC RATIONALE WAS
12	GENERALLY SOUND. EVEN THOUGH THE DRUG CANDIDATE HAD
13	BEEN IDENTIFIED USING A PHENOTYPIC SCREEN RATHER
14	THAN BY UNDERSTANDING OF A BIOCHEMICAL TARGET, WHICH
15	IS JUST A DIFFERENT APPROACH THAT'S NOT TYPICAL OF
16	CURRENT DRUG DEVELOPMENT EFFORTS THESE DAYS, BUT IN
17	FACT THERE'S CERTAINLY PRECEDENT FOR THAT TYPE OF
18	DRUG DEVELOPMENT.
19	LET'S SEE. THE REVIEWERS WERE SOMEWHAT
20	CONCERNED ABOUT THE ABSENCE OF INFORMATION REGARDING
21	THE DRUG TARGET AND THE MECHANISM OF ACTION BECAUSE
22	THAT DOES MAKE IT SOMEWHAT MORE COMPLICATED TO
23	DEVELOP THE DRUG WITHOUT UNDERSTANDING EXACTLY WHAT
24	THE TARGET IS.
25	SO I THINK ONE OF THE MAJOR CRITICISMS WAS

1	THAT IN SMALL MOLECULE DRUG DEVELOPMENT UNACCEPTABLE
2	TOXICITY IS A MAJOR REASON FOR FAILURE. AND SO THEY
3	FELT THAT IT WOULD MAKE SENSE TO PRIORITIZE, IF THIS
4	AWARD WERE FUNDED, EVALUATING THE TOXICITY IN ANIMAL
5	MODELS INSTEAD OF FOCUSING ON MECHANISM OF ACTION
6	STUDIES.
7	THEY FELT THAT THE PLANNED ACTIVITIES,
8	HOWEVER, COULD LEAD TO A ROBUST PACKAGE FOR A
9	PRE-IND MEETING AND FOUND THAT THE PRECLINICAL
10	DEVELOPMENT PLAN WAS FEASIBLE.
11	WITH RESPECT TO THE PI AND DEVELOPMENT
12	TEAM, THERE WAS SOME CONCERN THAT THERE WAS SOME
13	LACK OF DRUG DEVELOPMENT EXPERIENCE IN REGULATORY
14	AND CMC EXPERTISE AND FELT THAT WAS A SIGNIFICANT
15	WEAKNESS THAT SHOULD BE ADDRESSED. BUT ALSO AT THE
16	SAME TIME NOTICED THAT THE PI IS A VERY WELL
17	ESTABLISHED AND ACCOMPLISHED INVESTIGATOR AND HAS
18	ASSEMBLED A STRONG TEAM OF SCIENTISTS.
19	MR. SHEEHY: SO ARE THERE QUESTIONS,
20	COMMENTS ANY BOARD MEMBER WOULD LIKE TO SPEAK TO
21	THIS? ANY QUESTIONS FOR CIRM TEAM MEMBERS?
22	DR. PRIETO: I DON'T KNOW IF WE WANT TO
23	GET THIS INTO THE WEEDS ABOUT THE SCORING, BUT CAN I
24	ASK HOW MANY REVIEWERS SCORED THIS BELOW THE
25	FUNDABLE RANGE AND HOW MANY IN THE FUNDABLE RANGE?
	7.1

1	MR. SHEEHY: IF YOU LOOK AT THE CHART IN
2	FRONT OF YOU, YOU GET TIER I SCORES AND TIER II
3	SCORES.
4	DR. PRIETO: I SEE. I'M SORRY.
5	DR. KADYK: SO EIGHT SCORED IN TIER I AND
6	THEN SEVEN SCORED IN EITHER TIER II OR TIER III.
7	MR. SHEEHY: IS THERE ADDITIONAL
8	DISCUSSION? IS THERE PUBLIC COMMENT ON THIS
9	APPLICATION? WE DID HEAR FROM FOLKS A FEW MINUTES
10	AGO.
11	I THINK THE NEXT STEP IS TO MOVE TO A ROLL
12	CALL IF THERE'S NO PUBLIC COMMENT.
13	DR. PRIETO: OR BOARD COMMENT.
14	MR. SHEEHY: OR BOARD COMMENT.
15	MR. TORRES: YOU HAVE TO ADDRESS THE MIC.
16	OTHERWISE YOUR COMMENTS WILL NOT BE RECORDED.
17	MR. SHEEHY: AND, AGAIN, PLEASE INTRODUCE
18	YOURSELF.
19	DR. SCHUBERT: I'M DAVE SCHUBERT. I'M THE
20	PI OF THE GRANT THAT WAS JUST DISCUSSED. AND ONE OF
21	THE I WAS REQUESTED NOT TO PUT ANYTHING IN THE
22	APPEAL LETTER THAT I WROTE ABOUT THE SCIENTIFIC
23	REVIEW PROCESS ITSELF. AND SO THAT WAS NOT DELETED,
24	BUT I CAN MAKE SOME COMMENT ABOUT SOMETHING THAT WAS
25	JUST MENTIONED IN THE CONTEXT OF DISCOVERY.
	70
	72

1	WE HAVE A DRUG NOW WHICH IS ACTUALLY JUST
2	FINISHING THE IND PROCESS. AND WE HAVE A TEAM OF
3	CONSULTANTS AND WE HAVE PEOPLE WE WORK WITH AS FAR
4	AS ALL THE CLINICAL DEVELOPMENT.
5	AND WITHIN THE GRANT APPLICATION ITSELF, I
6	SHOULD MENTION THIS APPLICATION IS A VERY, VERY
7	DIFFICULT FORM TO DEAL WITH. I MEAN THE GRANTS
8	PEOPLE HAVE A HORRIBLE TIME WITH THIS, BUT THAT'S
9	BESIDE THE POINT. BUT WITHIN THE GRANT APPLICATION,
10	THERE WAS A VERY DETAILED CLINICAL DEVELOPMENT PHASE
11	I DESCRIPTION OF THE TRIAL TO IND PROCESS. AND FOR
12	SMALL MOLECULES, THE FDA DOES NOT REQUIRE, IN FACT,
13	THEY DISCOURAGE AND DO NOT ALLOW IN MOST CASES A
14	PRECLINICAL MEETING WITH THE FDA. SO THAT WAS AN
15	ASSUMPTION.
16	THIS IS DIFFERENT WITH STEM CELL-BASED
17	THERAPIES, BUT WITH SMALL MOLECULES, IF IT'S A
18	STANDARD MOLECULE THERAPY, THE FDA DOESN'T REQUIRE
19	THAT. SO THAT SHOULD NOT BE CONSIDERED AS A WEAK
20	POINT.
21	MR. SHEEHY: THANK YOU.
22	DR. HELMS: CAN I MAKE A COMMENT ABOUT
23	THIS PROCESS. JILL HELMS, STANFORD. WE'VE HEARD
24	ABOUT THE VAGARIES OF THE SCORING, AND I THINK THAT
25	YOU'RE ALL AWARE OF THAT AS VOTING MEMBERS. AND
	7.7

1	WE'VE ALSO HEARD FROM THE PRESIDENT THAT CIRM 2.0 IS
2	GOING TO TRY TO CORRECT THESE. BUT TO TELL YOU
3	STANDING HERE I HAD TO WORK UNDER THE RFA I WAS
4	GIVEN. I TOTALLY BELIEVE IN RAISING THE BAR, BUT I
5	DON'T THINK YOU CAN MOVE THE GOALPOST. IF YOU ARE
6	GOING TO USE THE GOALPOST OF 2.0, APPLY THEM HERE
7	NOW TO THESE GRANTS.
8	MR. SHEEHY: THANK YOU. I THINK IN SOME
9	WAY THAT'S BEEN THE STAFF RECOMMENDATION WHICH IS
10	FOR THOSE GRANTS IN TIER II, WE'RE GOING TO HAVE A
11	THREE-POINT SCORING SYSTEM. ONE, FUND; TWO, RETURN,
12	LOOKING AT THE RECOMMENDATIONS FROM THE WORKING
13	GROUP AND FIX ACCORDING TO THOSE RECOMMENDATIONS;
14	AND, THREE, WE DON'T THINK THAT THIS IS A GRANT THAT
15	WILL HAVE SUCCESS AT CIRM. SO THAT'S KIND OF WHAT
16	THE STAFF HAS RECOMMENDED, THAT WE FUND TIER I,
17	WHICH WOULD BE EQUIVALENT TO ONE IN THE NEW SYSTEM,
18	AND IN TIER II, THAT INDIVIDUALS TAKE THE COMMENTS
19	THAT WERE MADE BY THE REVIEWERS, RETOOL THEIR GRANTS
20	TO MAKE THEM BETTER GRANTS, AND RETURN AT THE LATEST
21	IN SIX MONTHS WHEN WE HAVE THE CONCEPT OUT FOR OUR
22	TRANSLATIONAL PROGRAM. OR IF YOU'RE LATER DOWN THE
23	ROAD, YOU CAN RETURN TO APPLY TO CIRM WITHIN THE
24	NEXT COUPLE OF MONTHS.
25	SO I ACTUALLY THINK THAT THAT'S WHAT STAFF
	7.4

1	WAS RECOMMENDING. ANYWAY, TO GO TO WHERE WE ARE, IF
2	THERE'S NO MORE PUBLIC COMMENT, I THINK WE'RE READY
3	TO CALL THE ROLL.
4	DR. JUELSGAARD: SO TO SPEAK TO THAT
5	LATTER POINT THAT YOU JUST MADE, JEFF, AND I WANT TO
6	ASK RANDY OR GIL OR WHOEVER. SO IF WE SAID TO ALL
7	THE TIER II PEOPLE WHAT WE WOULD RECOMMEND YOU DO IS
8	JUST RESUBMIT UNDER A NEW AND DIFFERENT PROCESS, HOW
9	QUICKLY COULD THAT HAPPEN IN YOUR MIND FOR ANY OF
10	THESE INDIVIDUAL PROJECTS THAT HAVE YOU CAN
11	COMMENT LATER, BUT I'M SPEAKING WITH OUR PEOPLE
12	FIRST. SO HOW LONG WOULD THAT TAKE IN YOUR
13	ESTIMATION?
14	DR. MILLS: I'M NOT GOING TO COMMENT AND
15	SAY ANY SPECIFIC APPLICATION BECAUSE I HAD A
16	CONFLICT WITH ONE OF THE APPLICATIONS. BUT IN
17	GENERAL TERMS, IF THEY WERE RESPONSIVE TODAY TO
18	15-01, WHICH IS AT AN ADVANCED STAGE, BUT IT
19	APPEARED THAT SOME OF THE APPLICATIONS WERE THERE OR
20	VERY CLOSE TO THERE, THEN THEY CAN APPLY TODAY.
21	THEY WOULD BE REVIEWED LITERALLY THEY WOULD HAVE
22	THE GWG RECOMMENDATION IN 60 DAYS AND AN AWARD
23	DECISION IN 81 DAYS, SO VERY, VERY QUICKLY.
24	IF THEY WERE NOT, THEN THEY WOULD HAVE TO
25	WAIT FOR THE CONCEPT PLAN AND THE RFA'S COMES OUT
	75

1	WHICH GO TO THE BOARD MEETING IN JULY.
2	DR. JUELSGAARD: OKAY.
3	MR. SHEEHY: COULD I JUST ASK ONE QUESTION
4	ABOUT THE PROCESS? AND I'M NOT SURE. WOULD THEY BE
5	FUNDED ACTUALLY FASTER UNDER 2.0 BECAUSE WE'VE
6	STREAMLINED THE FUNDING PROCESS TO SUCH A DEGREE
7	THAT YOU MIGHT ACTUALLY RECEIVE YOUR FUNDS FASTER
8	UNDER THE FRAMEWORK OF 2.0 THAN THE PROCESSES WE'RE
9	DEPLOYING RIGHT NOW UNDER 1.0?
10	DR. MILLS: SO YOU WOULD HAVE TO BE ABLE
11	TO GET YOUR APPLICATION IN IMMEDIATELY FOR THIS
12	MONTH'S FUNDING CYCLE. AND THEN IT WOULD PROBABLY
13	BE CLOSE BECAUSE HISTORICALLY CONTRACTING AND
14	AWARDING HAS TAKEN ABOUT, IF YOU JUST LOOK AT THE
15	AVERAGE, IT'S BEEN ABOUT SEVEN MONTHS UNDER THE
16	CURRENT PROCESS. BUT IF YOU CAN CONTRACT MORE
17	EFFICIENTLY, THEN ANSWER WOULD BE NO. IT WOULD BE
18	FASTER THIS WAY. YOU COULDN'T GET IT CONTRACTED
19	QUICKLY, THEN IT THEORETICALLY COULD BE, BUT ONLY
20	FOR THOSE APPLICATIONS THAT ARE READY TO GO IN THE
21	LATER STAGE.
22	MR. SHEEHY: THANK YOU.
23	DR. HELMS: JILL HELMS FROM STANFORD.
24	WITH REGARDS TO APPLYING FOR CIRM 2.0, A
25	PREREQUISITE FOR THE EXISTING PROGRAM IS THAT YOU

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1	HAVE HAD A PRE-IND MEETING. WE HAVE NOT. THE
2	PROGRAM THAT YOU TALK ABOUT COMING DOWN THE
3	PIPELINE, IT'S NOT YET ON THE WEBSITE. SO LET'S SAY
4	EARLIEST IT BECOMES AVAILABLE IN JULY. WE'VE DONE
5	THE CALCULATIONS. THAT'S THE ONLY PROGRAM WE CAN
6	APPLY TO. IF YOU DON'T FUND US NOW, THAT'S TEN
7	MONTHS AT THE MINIMUM, AND IT'S TO APPLY FOR A
8	PROGRAM THAT IS STILL IN THE PROCESS, AS WE'VE HEARD
9	FROM THE PRESIDENT, OF BEING DEVELOPED. SO THAT
10	MEANS THE TEAM DISINTEGRATES. WORK STOPS. THAT'S
11	WHAT I MEAN WHEN I SAID HIT THE PAUSE BUTTON.
12	MR. SHEEHY: THANK YOU.
13	DR. SCHUBERT: THIS IS DAVE SCHUBERT FROM
14	THE SALK INSTITUTE. I SAID THIS BEFORE, I BELIEVE,
15	BUT WE HAVE BEEN THROUGH THIS PROCESS BEFORE. AND I
16	CAN ASSURE YOU THAT FOR SMALL MOLECULE DRUG
17	DEVELOPMENT, IF YOU LOOK AT THE FDA WEBSITE FOR THE
18	GUIDANCE TO INDUSTRY FOR SMALL MOLECULES, THEY WILL
19	SAY THAT A PRE-IND MEETING IS NOT ALLOWED IN MOST
20	CASES. AND SO THIS IS A REQUIREMENT FOR THIS CIRM
21	2.0, AND I THINK THAT ALSO WOULD STOP US FROM
22	APPLYING FOR THAT.
23	AND THE OTHER ASPECT, WHICH SHOULD BE
24	POINTED OUT TO THE MEMBERS OF THIS PANEL IS THAT FOR
25	THE IND PROCESS ITSELF, IT HAS TO BE DONE IN AN FDA

1	APPROVED FACILITY BY CRO'S. AND SO AS A CONSEQUENCE
2	THERE'S NO LAB FUNDING FOR THAT. SO ONCE THE
3	PROJECT GOES INTO THE CRO'S GROUP THAT'S DOING THE
4	IND, THE LAB FUNDING BASICALLY IS ELIMINATED.
5	SO THIS IS ANOTHER PROBLEM THAT ACADEMIC
6	SCIENTISTS FACE. WE HAVE TO MAINTAIN SOME COHERENCE
7	IN OUR PROGRAM. AND IN MY CASE PARTICULARLY, IT WAS
8	WE THOUGHT THERE WAS A SEQUENCE OF GRANTS THAT WAS
9	DETERMINED BEFORE WE APPLIED FOR THE INITIAL GRANT,
10	AND SO IT'S A DIFFICULT PROBLEM.
11	MR. SHEEHY: THANK YOU.
12	DR. MILLS: I WANT TO CLARIFY SOMETHING
13	THAT'S JUST FACTUALLY INCORRECT. SO WE RECOGNIZE AT
14	CIRM THAT CERTAIN CATEGORIES OF DRUGS EITHER DO NOT
15	REQUIRE OR IN SOME INSTANCES ARE NOT EVEN ABLE TO
16	OBTAIN A PRE-IND MEETING WITH THE FDA. PRE-IND
17	MEETING IS NOT A REQUIREMENT IF YOUR PRODUCT IS NOT
18	A BIOLOGIC WHERE A PRE-IND MEETING WOULD BE
19	NECESSARY. SO IT'S NOT A REQUIREMENT UNDER CIRM 2.0
20	TO HAVE A PRE-IND MEETING IF YOU HAVE A THERAPEUTIC
21	CANDIDATE THAT DOES NOT REQUIRE ONE.
22	MR. REED: THIS IS JUST A QUICK STRUCTURAL
23	QUESTION. THE ROLL CALL THAT YOU'RE JUST ABOUT TO
24	TAKE, AND THIS IS JUST ON THE ONE PROJECT, RIGHT?
25	MR. SHEEHY: YES.

1	MR. REED: NOT ON THE WHOLE BLOCK?
2	MR. SHEEHY: JUST ON THIS ONE PROJECT.
3	CHAIRMAN THOMAS: SO JUST WANT TO REMIND
4	THE BOARD THE CRITICAL REASON FOR HAVING 2.0 IS WE
5	HAVE PROJECTS THAT UP TO THIS POINT AND INCLUDING
6	THIS ROUND FALL INTO TIER II WHICH MEANS THEY DO
7	HAVE SOME CONSIDERABLE PROMISE, BUT THEY ALSO HAVE
8	ISSUES. AND THE REASON TO HAVE 2.0 IN PLACE AND A
9	FREQUENT OPPORTUNITY TO REAPPLY IS TO ACTUALLY
10	REFINE GOOD PROJECTS TO HOPEFULLY MAKE THEM GREAT
11	AND GIVE THEM ULTIMATELY A BETTER CHANCE OF SUCCESS
12	THAN THEY MIGHT OTHERWISE HAVE IF THEY STILL HAVE
13	FLAWS IN THEM AS PERCEIVED BY OUR PEER REVIEWERS.
14	SO I THINK WE ALL ARE INTERESTED IN
15	GETTING THERAPIES THROUGH TO PATIENTS AS BEST WE
16	CAN; BUT TO GIVE US THE BEST SHOT AT DOING THAT,
17	WE'D LIKE THE PROJECTS TO BE REFINED TO A STAGE
18	WHERE THEY ACTUALLY HAVE THE GREATEST OPPORTUNITY
19	FOR SUCCESS. THAT'S THE ESSENCE OF 2.0. DR. MILLS,
20	AM I CORRECT ON THAT STATEMENT?
21	DR. MILLS: EXACTLY.
22	MR. SHEEHY: SO ANY MORE BOARD SENATOR
23	TORRES.
24	MR. TORRES: THANK YOU FOR CHAIRING THIS
25	INITIAL SESSION. WE'RE FINDING OUT HOW WE'RE
	79

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1
     DEALING WITH THIS PROCESS. SO I NO LONGER HAVE A
 2
     CONFLICT, BECAUSE MY SURGERY IS OVER, ON
 3
     OSTEOARTHRITIS, BUT I AM CONCERNED ABOUT THE FUTURE
 4
     WITH MY AGE GROUP, WHICH WE ARE CHRONOLOGICALLY
 5
     GIFTED, AND WE ARE INCREASING IN NUMBERS.
               MR. JUELSGAARD: I THOUGHT WE WERE TALKING
 6
 7
     ABOUT ALZHEIMER'S.
               MR. TORRES: I'M SORRY. I THOUGHT WE
 8
 9
     VOTED. WE HAVEN'T VOTED YET. WELL, LET'S GET TO
10
     IT. CALL THE QUESTION.
               MR. SHEEHY: I WAS HEADED THAT DIRECTION.
11
12
     OKAY. COULD WE CALL THE ROLL, PLEASE.
13
               MS. BONNEVILLE: ANNE-MARIE DULIEGE.
14
               DR. DULIEGE: NO.
15
               MS. BONNEVILLE: DAVID HIGGINS.
16
               DR. HIGGINS: CAN YOU CLARIFY EXACTLY WHAT
17
     A YES VOTE AND A NO VOTE IS?
               MR. SHEEHY: A YES VOTE WOULD BE TO MOVE
18
19
     THIS APPLICATION INTO TIER I AND FUND IT. A NO VOTE
     WOULD BE TO LEAVE IT WHERE IT IS.
20
21
               DR. HIGGINS: GOT YOU. YES.
22
               MS. BONNEVILLE: STEPHEN JUELSGAARD.
23
               MR. JUELSGAARD: YES.
24
               MS. BONNEVILLE: KATHY LAPORTE.
25
               DR. LAPORTE: YES.
                               80
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1	MS. BONNEVILLE: LAUREN MILLER.
2	MS. MILLER: YES.
3	MS. BONNEVILLE: FRANCISCO PRIETO.
4	DR. PRIETO: AYE.
5	MS. BONNEVILLE: ROBERT QUINT.
6	DR. QUINT: YES.
7	MS. BONNEVILLE: AL ROWLETT.
8	MR. ROWLETT: YES.
9	MS. BONNEVILLE: JEFF SHEEHY.
10	MR. SHEEHY: ABSTAIN.
11	MS. BONNEVILLE: JONATHAN THOMAS.
12	CHAIRMAN THOMAS: YES.
13	MS. BONNEVILLE: ART TORRES.
14	MR. TORRES: AYE.
15	MS. BONNEVILLE: DIANE WINOKUR.
16	MS. WINOKUR: I ABSTAIN.
17	MS. BONNEVILLE: MOTION CARRIES.
18	MR. SHEEHY: OKAY. THANK YOU. SO THE
19	NEXT ARE THERE ADDITIONAL MOTIONS OR ANY MOTIONS
20	TOWARDS ANOTHER APPLICATION?
21	MR. TORRES: YES. A POINT OF INFORMATION
22	TO THE PRESIDENT. MR. PRESIDENT, HOW ARE YOU? WHAT
23	HAPPENS IF WE DO NOT MOVE AN ITEM, AND I'M
24	SPECIFICALLY LOOKING AT PC1-08128, TO THE FIRST
25	TIER? WHAT'S THE STATUS OF THAT PROJECT THEN AT
	0.1
	81

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1	THIS POINT UNDER 2.0?
2	DR. MILLS: IT'S NOT UNDER 2.0.
3	MR. TORRES: SO REAPPLICATION WOULD HAVE
4	TO OCCUR?
5	DR. MILLS: YES, ABSOLUTELY.
6	MR. TORRES: MR. CHAIRMAN, I MOVE THAT WE
7	MOVE FROM TIER II TO TIER I PC1-08128.
8	DR. JUELSGAARD: I SECOND THE MOTION.
9	MR. SHEEHY: MOTION BY SENATOR TORRES,
10	SECOND BY MR. JUELSGAARD. COULD WE HEAR FROM CIRM
11	TEAM ABOUT THIS APPLICATION?
12	DR. KADYK: THIS IS PC1-08128. THIS IS A
13	PROJECT THAT AIMS TO DEVELOP A NOVEL CELL-BASED
14	THERAPY FOR REPAIR OF CARTILAGE DEFECTS, AS WE HAVE
15	HEARD, CAUSED BY TRAUMA OR DISEASE.
16	THE PROPOSED APPROACH WOULD COMBINE
17	ALLOGENEIC EMBRYONIC STEM CELL-DERIVED CARDIAC
18	PROGENITOR CELLS THAT HAVE THE ABILITY TO
19	DIFFERENTIATE AND DEVELOP INTO MATURE CARTILAGE WITH
20	A SCAFFOLDING MATERIAL WHICH IS THEN FACILITATING
21	TISSUE REPAIR. THIS CELL SCAFFOLD COMBINATION
22	PRODUCT WOULD BE SURGICALLY IMPLANTED TO A CARTILAGE
23	DEFECT.
24	THE PROPOSED ACTIVITIES INCLUDE MAKING AND
25	TESTING GOOD MANUFACTURING PRACTICES COMPLIANT
	9.7

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1	MASTER AND WORKING CELL BANKS, TECHNOLOGY TRANSFER,
2	SCALE-UP, PRODUCTION AND CHARACTERIZATION OF GMP
3	PRODUCT CANDIDATE, DOSE FINDING, AND PILOT
4	PRECLINICAL STUDIES IN ORDER TO REACH THE OVERALL
5	OBJECTIVE OF CONDUCTING WELL-SUPPORTED PRE-IND
6	MEETING WITH THE FDA.
7	SO DURING THE REVIEW, UNDER SIGNIFICANCE,
8	THE REVIEWERS HAD SOMEWHAT MIXED OPINIONS AS TO THE
9	LEVEL WHICH THIS WAS AN UNMET NEED. SOME FELT IT
10	WAS ONLY A MODERATE UNMET NEED WHERE OTHERS THOUGHT
11	IT WAS A VERY CLEAR UNMET NEED AND THAT THERE WAS A
12	COMMERCIAL OPPORTUNITY IN THIS TARGETED SUBGROUP OF
13	YOUNGER PATIENTS IN PARTICULAR BECAUSE THERE'S
14	VARIABLE REPORTS OF THE EFFECTIVENESS AND DURABILITY
15	OF CURRENTLY AVAILABLE THERAPIES. SO FOR A YOUNGER
16	PATIENT, DURABILITY IS PARTICULARLY IMPORTANT.
17	REVIEWERS COMMENTED THAT THE CARTILAGE
18	REPAIR FIELD IS COMPETITIVE WITH SOME OTHER
19	THERAPEUTIC OPTIONS THAT ARE ALREADY AVAILABLE OR IN
20	DEVELOPMENT, INCLUDING SOME CELL THERAPIES.
21	HOWEVER, SOME REVIEWERS DID BELIEVE THERE'S ROOM FOR
22	A NEW TREATMENT OPTION AND FELT THAT THIS PROPOSED
23	CELL SCAFFOLD COMBINATION HAD SOME FEATURES THAT
24	DIFFERENTIATES IT FROM OTHER PROPOSED OR OTHER
25	PROCEDURES THAT ARE IN DEVELOPMENT AND HAS THE
	0.7

1	POTENTIAL TO BE AN OFF-THE-SHELF PRODUCT AND WOULD
2	HAVE AVAILABLE A CONSISTENT PROGENITOR CELL
3	POPULATION AND A NOVEL SCAFFOLD THAT COULD
4	CONTRIBUTE TO PRODUCT SAFETY AND ACTIVITY.
5	THE REVIEWERS DID NOTE THE RELATIVELY
6	SHORT HALF-LIFE OF THE CURRENT PRODUCT AND JUST
7	ENCOURAGED THEM TO DEVELOP A FROZEN FORMULATION
8	FURTHER DOWN THE ROAD SINCE THAT STABILITY AND SHELF
9	LIFE AS IT CURRENTLY EXISTS MIGHT LIMIT WIDESPREAD
10	USE. SO THEY MADE THAT SUGGESTION.
11	IN TERMS OF SCIENTIFIC RATIONALE AND
12	PRECLINICAL DEVELOPMENT READINESS, THEY FELT THAT
13	THE PRELIMINARY DATA WERE DEFINITELY SUPPORTIVE OF
14	PRECLINICAL DEVELOPMENT READINESS. AND, IN FACT,
15	SOME REVIEWERS FELT THAT THE PRODUCT CANDIDATE IS
16	CLOSE TO THE STAGE WHERE THEY'D BE ABLE TO HOLD A
17	PRE-IND MEETING, AND THEY MAY BE ABLE TO SHORTEN THE
18	TERM OF THE AWARD IF THEY WERE TO RECEIVE IT.
19	WHILE THE RATIONALE WAS VIEWED AS SOUND,
20	SOME REVIEWERS FELT THAT THE PRELIMINARY DATA COULD
21	HAVE BEEN MORE COMPELLING IF THEY HAD DEMONSTRATED A
22	STATISTICALLY SIGNIFICANT DIFFERENCE BETWEEN THEIR
23	APPROACH AND COMPETING TECHNOLOGIES.
24	FOR DESIGN AND FEASIBILITY, REVIEWERS
25	FOUND THAT THE PROPOSED STUDIES WERE WELL DESIGNED

1	AND FEASIBLE AND HAD ALL NECESSARY TECHNIQUES IN
2	PLACE. ALTHOUGH THEY DID COMMENT THAT THERE WERE
3	SOME POSSIBLE OPPORTUNITIES TO STREAMLINE THE PLAN.
4	AND SOME OF THE PROPOSED PRECLINICAL STUDIES COULD
5	BE COMBINED AND PERHAPS MORE APPROPRIATELY CONDUCTED
6	AFTER THE PRE-IND MEETING.
7	THEY DID SUGGEST THAT IT WOULD BE
8	IMPORTANT TO SEEK OUT EXPERT REGULATORY INPUT TO
9	IDENTIFY THE CRITICAL PATH ACTIVITIES NEEDED TO GET
10	TO THE PRE-IND MEETING UNDER THIS AWARD.
11	REVIEWERS COMMENTED ON THE IMPORTANCE OF
12	HAVING AN ACCEPTABLE RISK PROFILE FOR THIS
13	PARTICULAR PRODUCT BECAUSE IT IS AN EMBRYONIC STEM
14	CELL-DERIVED PRODUCT.
15	THEY SUGGESTED THAT USING A SINGLE MARKER
16	TO TEST FOR UNDIFFERENTIATED STEM CELLS MIGHT NOT BE
17	SUFFICIENT AND THAT ADDITIONAL MARKERS SHOULD ALSO
18	BE EXPLORED.
19	REVIEWERS WERE SUPPORTIVE OF THE
20	ALLOGENEIC STEM CELL SOURCE WITH THE CUSTOM SCAFFOLD
21	AS AN OFF-THE-SHELF PRODUCT, BUT THEY WEREN'T AS
22	CLEAR TO WHETHER THE PROPOSED THERAPY WAS GOING TO
23	BE COMPARED AGAINST THE CURRENT SURGICAL STANDARD OF
24	CARE WHICH IS MICROFRACTURE. AND THEY FELT THAT
25	DIRECT COMPARISON SHOULD BE MADE.
	ΟΓ

1	THE PROPOSED TIMELINES FOR THE PRE-IND
2	MEETING WERE APPROPRIATE, AND, AS I MENTIONED
3	BEFORE, COULD POTENTIALLY EVEN BE ACCELERATED.
4	THEY FELT IN TERMS OF THE TEAM AND THE PI,
5	THE TEAM IS EXCELLENT AND HAS STRONG EXPERTISE IN
6	CARTILAGE DEVELOPMENT, AND REVIEWERS WERE VERY
7	APPRECIATIVE OF THE PLANNED ADDITION OF A SECOND
8	PROJECT MANAGER THAT IS CURRENTLY BEING SOUGHT.
9	THEY, AGAIN, AS I MENTIONED EARLIER, THE
10	TEAM COULD USE SOME ADDITIONAL EXPERTISE IN
11	REGULATORY AFFAIRS TO HELP DEFINE THE CRITICAL PATH
12	ACTIVITIES. AND THERE WERE NO CONCERNS ABOUT THEIR
13	COLLABORATIONS, ASSETS, RESOURCES, AND ENVIRONMENT.
14	MR. SHEEHY: OTHER QUESTIONS OR DISCUSSION
15	BY BOARD MEMBERS?
16	DR. JUELSGAARD: SO ONE OF THE THINGS
17	STRUCK ME, SOME OF THE CRITICISM OF THIS PARTICULAR
18	APPLICATION WAS THAT IT SEEMED AS IF THE REVIEWERS
19	FELT THAT THERE WERE ALTERNATIVE THERAPIES AVAILABLE
20	AND, THEREFORE, THERE WAS A STANDARD OF CARE. AND
21	THE QUESTION WAS HOW DID THIS RELATE TO THE STANDARD
22	OF CARE. AND THAT THERE WEREN'T STUDIES DONE TO
23	SHOW THE SUPERIORITY OF THIS PARTICULAR TYPE OF
24	TREATMENT, THE SUPERIORITY POTENTIAL AGAINST
25	STANDARD OF CARE.
	96

1	FIRST OF ALL, I THINK THAT'S A
2	PROGRAMMATIC ISSUE MORE THAN ANYTHING. ARE WE GOING
3	TO FUND SOMETHING WHERE WE THINK THERE'S ALREADY A
4	LOT OF STUFF GOING ON OUT THERE IN OTHER AREAS. I
5	APPRECIATE THEIR VIEWS, BUT I THINK THAT THAT'S
6	REALLY SOMETHING WE NEED TO CONSIDER.
7	ABOVE AND BEYOND THAT, THERE'S ALWAYS THE
8	QUESTION OF WHETHER YOU NEED SUPERIORITY OR
9	NONINFERIORITY, SO THE TWO DIFFERENT STANDARDS THAT
10	ARE USED FOR FDA APPROVAL. NONINFERIORITY MEANS
11	THAT YOU HAVE TO PROVE THAT, IN ESSENCE, YOU'RE NOT
12	INFERIOR TO AN EXISTING STANDARD OF TREATMENT. SO
13	WHY WOULD PEOPLE DO THAT? WHY ARE THERE SOMETIMES
14	NONINFERIORITY STANDARDS USED THAT ARE, AT THE END
15	OF THE DAY, APPROVED? AND ONE OF THE REASONS IS
16	THAT THE PRODUCT IS SAFER OR THAT THE PRODUCT IS
17	EASIER TO ADMINISTER AND CREATES LESS TREATMENT DOWN
18	THE ROAD, HOSPITALIZATION, ETC.
19	AND ONE THING THE FDA DOES NOT TAKE INTO
20	ACCOUNT, BUT I THINK FOR US IS SOMETHING THAT WE
21	HAVE TO THINK ABOUT, IS COST. IF WE HAVE A
22	TREATMENT THAT POTENTIALLY COULD BE GIVEN IN THE
23	FACE OF EXISTING TREATMENTS BUT IS SUBSTANTIALLY
24	LESS IN COST, THEN I THINK NONINFERIORITY IS AN
25	ACCEPTABLE STANDARD.
	0.7

1	SO I JUST I DON'T KNOW THAT WE KNOW
2	ENOUGH AT THIS POINT ABOUT, WELL, THERE'S JUST
3	PLENTY OF OTHER STUFF OUT THERE, SO WE REALLY
4	SHOULDN'T CONSIDER THIS, IF WE ARE REALLY AT THAT
5	POINT WHERE THAT DECISION COULD BE MADE. SO TO THE
6	EXTENT THAT THOSE CRITICISMS WERE LEVELED AT THIS
7	APPLICATION, I, FOR ONE, TEND TO, I GUESS, TEND TO
8	NOT TAKE THEM TOO SERIOUSLY IN MY OWN MIND. THAT'S
9	OBVIOUSLY OPINIONS OF OTHER PEOPLE, BUT I THINK WE
10	NEED MORE INFORMATION BEFORE WE CAN FIGURE OUT
11	WHETHER THIS IS A BETTER THERAPY OR SIMPLY A MORE
12	USEFUL THERAPY THAN WHAT ALREADY EXISTS.
13	MR. SHEEHY: SO CAN I JUST ASK ARE YOU
14	ARGUING IN FAVOR?
15	DR. JUELSGAARD: YES, I'M ARGUING IN
16	FAVOR.
17	MR. TORRES: WAIT A MINUTE. HE WAS MY
18	SECOND.
19	DR. JUELSGAARD: NO. I'M ARGUING IN
20	FAVOR. I THINK THERE'S A LOT MORE WORK THAT NEEDS
21	TO BE DONE BEFORE YOU CAN GET TO THE POINT OF
22	SAYING, WELL, WAIT A MINUTE. YOU CAN ALWAYS REPLACE
23	A KNEE USING HARDWARE, RIGHT? AND THAT'S JUST AS
24	GOOD OR BETTER THAN WHAT YOU'RE TALKING ABOUT. I
25	DON'T THINK WE KNOW ENOUGH ABOUT THAT RIGHT NOW TO
	88

USE THAT AS A REALISTIC CRITERIA FOR SAYING THAT WE
SHOULDN'T FUND THIS.
MR. SHEEHY: ARE THERE OTHER QUESTIONS OR
COMMENTS FROM BOARD MEMBERS? ANY PUBLIC COMMENT?
DR. D'LIMA: DARRYL D'LIMA AGAIN. I'M THE
PI ON THE GRANT THAT'S BEING REVIEWED AT THE MOMENT.
AND I'D JUST LIKE TO MAKE A COMMENT ABOUT THE
CONCERNS ABOUT REGULATORY ISSUES. WE HAVE THREE
MEMBERS ON OUR LEADERSHIP TEAM WHO ARE ACTUALLY ON
THE FDA ADVISORY PANEL FOR CELL THERAPY AND GENE
THERAPY. WE HAVE CONTRACTED WITH AN INTERNATIONAL
COMPANY. IT'S IN THE APPLICATION. SO FOR A
REVIEWER TO SAY THAT WE NEED MORE REGULATORY SUPPORT
IS DISINGENUOUS.
THE CRITICISM ABOUT THE SINGLE MARKER,
THAT'S WHAT THE FDA RECOMMENDED. SO WE'RE USING
WHAT THE FDA HAS TOLD US TO PUT IN OUR PRE-PRE-IND
MEETING THAT THEY WOULD ACCEPT TODAY. THEY MIGHT
CHANGE THEIR MIND LATER, BUT THAT'S WHAT THEY WOULD
ACCEPT TODAY AS A MARKER OF LACK OF PLURIPOTENCY OR
LACK OF TERATOGENESIS.
TO THE POINT ABOUT ALTERNATIVE THERAPIES,
THEY MENTION MICROFRACTURE, WHICH IS A MARROW
STIMULATION PROCEDURE. THAT'S WHAT WE PUT IN OUR
STIMULATION PROCEDURE. THAT S WHAT WE PUT IN OUR
PRECLINICAL. WE KNOW THAT WE HAVE TO DO BETTER THAN

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1
     THE EXISTING CELL THERAPIES. WE KNOW THAT THE
 2
     EXISTING CELL THERAPIES DON'T WORK. THE PEOPLE WHO
 3
     KNOW WHAT THE UNMET NEEDS ARE ARE THE PATIENTS AND
 4
     THE PHYSICIANS WHO TREAT THEM. AND I THINK THAT THE
 5
     DATA THAT WE'VE SEEN TODAY IS VERY COMPELLING. AND
     THE TESTING WE PROPOSED, WE'VE TESTED AGAINST ADULT
 6
 7
     STEM CELLS, WE'VE TESTED AGAINST BONE MARROW STEM
 8
     CELLS, AND WE'VE TESTED AGAINST THE SO-CALLED
 9
     MICROFRACTURE, WHICH SOME CLAIM IS STANDARD OF CARE,
     AND IT'S FAR BETTER THAN ALL THREE OF THOSE IN
10
11
     ANIMALS.
               THANK YOU.
12
               MR. SHEEHY: ANY OTHER PUBLIC COMMENT?
13
     THEN, MS. BONNEVILLE, COULD YOU CALL THE ROLL,
14
     PLEASE.
               MS. BONNEVILLE: ANNE-MARIE DULIEGE.
15
16
                DR. DULIEGE: YES.
17
               MS. BONNEVILLE: STEPHEN JUELSGAARD.
18
               MR. JUELSGAARD: YES.
19
               MS. BONNEVILLE: KATHY LAPORTE.
20
                DR. LAPORTE: YES.
21
               MS. BONNEVILLE: LAUREN MILLER.
22
               MS. MILLER: YES.
23
               MS. BONNEVILLE: FRANCISCO PRIETO.
24
                DR. PRIETO: ABSTAIN.
25
                MS. BONNEVILLE: ROBERT QUINT.
                               90
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1	DR. QUINT: YES.
2	MS. BONNEVILLE: AL ROWLETT.
3	MR. ROWLETT: YES.
4	MS. BONNEVILLE: JEFF SHEEHY.
5	MR. SHEEHY: ABSTAIN.
6	MS. BONNEVILLE: JONATHAN THOMAS.
7	CHAIRMAN THOMAS: YES.
8	MS. BONNEVILLE: ART TORRES.
9	MR. TORRES: AYE.
10	MS. BONNEVILLE: DIANE WINOKUR.
11	MS. WINOKUR: YES.
12	MR. HARRISON: MOTION CARRIES.
13	MR. SHEEHY: DO WE HAVE ANY ADDITIONAL
14	MOTIONS? MR. JUELSGAARD.
15	DR. JUELSGAARD: YES. I MOVE THAT WE MOVE
16	APPLICATION PC1-08105 FROM TIER II TO TIER I.
17	MR. SHEEHY: DO WE HAVE A SECOND FOR THAT
18	MOTION?
19	MR. TORRES: SECOND.
20	MR. SHEEHY: DISCUSSION FROM STAFF, FROM
21	TEAM MEMBERS.
22	DR. KADYK: OKAY. THIS IS PC1-08105.
23	THIS APPLICATION SEEKS TO DEVELOP AN IMPROVED
24	THERAPY FOR BONE REGENERATION BY ENHANCING OR
25	RESTORING THE OSTEOGENIC POTENTIAL OF A PATIENT'S
	91
	31

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1	OWN STEM CELLS AS PART OF A SURGICAL BONE GRAFTING
2	TECHNIQUE CALLED AUTOGRAFTING. SO THE PROPOSED
3	DEVELOPMENT CANDIDATE WOULD COMPRISE BONE-DERIVED
4	CELLS THAT ARE HARVESTED FROM A PATIENT, TREATED EX
5	VIVO WITH A PROPRIETARY FORMULATION WNT3A, WHICH IS
6	A PROTEIN INVOLVED IN STEM CELL SELF-RENEWAL AND
7	TISSUE REGENERATION. AND THEN THOSE TREATED CELLS
8	WOULD THEN BE TRANSPLANTED INTO SITES REQUIRING BONE
9	REPAIR.
10	FOR THIS APPLICATION THE MAJOR PROJECT
11	ACTIVITIES INCLUDE COMPLETION OF PRECLINICAL
12	STUDIES, DETERMINING EFFECTIVE DOSE RANGE,
13	DEVELOPING AND VALIDATING APPROPRIATE MANUFACTURING
14	PROCESSES AND ASSAYS, SELECTING THE TARGET
15	INDICATION, AND DEVELOPING THE CLINICAL PLAN WITH
16	THE ULTIMATE GOAL OF CONDUCTING A WELL-PREPARED
17	PRE-IND MEETING.
18	SO UNDER SIGNIFICANCE AND IMPACT,
19	REVIEWERS FELT THAT THE TARGET PRODUCT PROFILE MIGHT
20	BE TOO BROAD AND SHOULD BE FOCUSED AROUND A SINGLE
21	INDICATION. HOWEVER, THEY ALSO STATED THAT, IF
22	SUCCESSFULLY DEVELOPED, THE PROPOSED APPROACH COULD
23	IMPROVE UPON THE STANDARD OF CARE, POTENTIALLY
24	EXTENDING THE USE OF AUTOGRAFT PROCEDURES IN OLDER
25	PATIENTS.

1	THERE WAS SOME CONCERN ABOUT THE POTENTIAL
2	VARIABILITY OF AN AUTOGRAFT THAT WAS EXPRESSED BY
3	SOME OF THE REVIEWERS.
4	AND OF THE TWO LEAD INDICATIONS THAT WERE
5	MENTIONED IN THE DRAFT CLINICAL SYNOPSES, REVIEWERS
6	CONSIDERED THE OSTEONECROSIS OF THE HIP TO BE A MUCH
7	MORE COMPELLING UNMET NEED THAN THE SPINAL FUSION.
8	UNDER SCIENTIFIC RATIONALE AND PRECLINICAL
9	DEVELOPMENT READINESS, THEY FELT THAT THE RATIONALE
10	FOR EXPLOITING THE WNT3A PATHWAY IN BONE REPAIR WAS
11	VERY WELL SUPPORTED BY PRELIMINARY DATA AS WELL AS
12	BY THE FIELD IN GENERAL.
13	REVIEWERS BELIEVED THE DATA PRESENTED WERE
14	SUFFICIENT TO SUPPORT THE READINESS OF THE PROPOSED
15	CANDIDATE FOR PRECLINICAL DEVELOPMENT.
16	UNDER DESIGN AND FEASIBILITY, REVIEWERS
17	HAD SOMEWHAT MIXED VIEWS ABOUT THE OVERALL
18	FEASIBILITY DUE TO REGULATORY QUESTIONS AND HOW THE
19	FDA WOULD CLASSIFY AND REGULATE THE THERAPEUTIC
20	CANDIDATE. SOME BELIEVE THAT THAT COULD BE VERY
21	STRAIGHTFORWARD WHILE OTHERS FELT IT MIGHT BE
22	CHALLENGING. AND THE CONSENSUS WAS THAT EARLIER FDA
23	ENGAGEMENT AND ADVICE WOULD BE CRITICAL TO CLEARLY
24	ESTABLISH THE PATH FORWARD TO A PRE-IND AND IND
25	MEETING. SO THEY SHOULD CONSULT THE FDA AS SOON AS
	93

1	POSSIBLE.
2	THE REVIEWERS FELT THAT THE PROPOSED
3	STUDIES REQUIRE SOME REFINEMENT INCLUDING A FOCUS ON
4	A SINGLE INDICATION, WHICH IS, I THINK, MENTIONED
5	EARLIER. AND SO THEY SHOULD DISCUSS WITH THE FDA IN
6	HOW TO GET ALIGNMENT FOR A WELL-PREPARED PRE-IND
7	MEETING.
8	THE PROPOSAL ITSELF HAD VERY CLEAR
9	GO/NO-GO DECISION POINTS AND MILESTONES. THE
10	APPLICANTS DO HAVE EXTENSIVE EXPERIENCE WITH THE
11	PROPOSED ANIMAL MODELS AND SHOULD HAVE LITTLE
12	TROUBLE ACHIEVING THEIR TECHNICAL MILESTONES.
13	THE REVIEWERS FELT THAT THE PROPOSED
14	SURGICAL APPROACH IS ACHIEVABLE AND, HOWEVER, IT
15	COULD BE FURTHER STREAMLINED PERHAPS TO MINIMIZE THE
16	TIME SPENT UNDER ANESTHESIA. AND SO THIS WAS
17	SPECULATION THAT THEY FELT SHOULD BE DISCUSSED
18	FURTHER WITH INPUT FROM SURGEONS.
19	REVIEWERS COMMENTED THAT AN ALTERNATIVE TO
20	AN AUTOGRAFT, SUCH AS A SYNTHETIC CARRIER FOR WNT3A,
21	MIGHT ALLOW FOR MORE EXPERIMENTAL REPRODUCIBILITY
22	AND PRESENT AN EASIER REGULATORY PATH.
23	THE PI AND TEAM, THEY FELT THEY SAID
24	THE PI HAS LARGELY PIONEERED AND DEVELOPED THIS
25	PROPOSAL AND IS VERY DEDICATED TO ITS CLINICAL

1	APPLICATION AND PLAYS AN IMPORTANT ROLE ON THE
2	INVESTIGATIVE TEAM. THEY DID SUGGEST THAT THE TEAM
3	WOULD BENEFIT FROM A QUALIFIED REGULATORY EXPERT WHO
4	WOULD WORK AS AN INTEGRAL PART OF THE TEAM.
5	THE PI HAS ASSEMBLED AN OUTSTANDING TEAM
6	OF COLLABORATORS WITH PREMIER EXPERTISE IN THE AREAS
7	OF BOTH WNT SIGNALING AND BONE BIOLOGY. AND THERE
8	WERE NO CONCERNS REGARDING THE ASSETS OR RESOURCES
9	AVAILABLE.
10	MR. SHEEHY: SO ARE THERE ANY QUESTIONS
11	FOR DR. KADYK?
12	MS. WINOKUR: I HAVE A COMMENT, NOT A
13	QUESTION. THERE ARE FIVE PROPOSALS IN TIER II, AND
14	WE ARE ABOUT TO VOTE ON MOVING THE THIRD ONE TO TIER
15	I. I AM UNCOMFORTABLE WITH CHANGING THAT MUCH OF
16	THE PEER REVIEW DECISIONS HERE TODAY.
17	DR. PRIETO: IF IT'S APPROPRIATE TO ASK,
18	SORT OF IN LIGHT OF DIANE'S COMMENT, GIL OR HIS TEAM
19	WHERE WE ARE ON THE BUDGET WITH THE TWO APPLICATIONS
20	THAT WE'VE ALREADY MOVED UP.
21	DR. SAMBRANO: CERTAINLY. SO I PUT UP ON
22	THE SCREEN, IT MAY BE VERY DIFFICULT TO SEE, WE'RE
23	AT 25.2 MILLION CURRENTLY.
24	DR. PRIETO: THAT'S WITH THE CHANGES?
25	DR. SAMBRANO: THAT'S WITH THE TWO THAT
	95

1	WERE ADDED.
2	MR. SHEEHY: WHAT WAS THE BUDGET FOR THIS
3	ROUND?
4	DR. SAMBRANO: IT WAS 40 MILLION.
5	DR. WESTON: I SHARE DIANE'S CONCERN. AND
6	I WONDER IF THERE'S AN ISSUE WITH THE INSTRUCTION,
7	MAYBE I DON'T UNDERSTAND, COMING FROM THE GRANTS
8	WORKING GROUP OR GIVEN TO THEM ABOUT WHAT SHOULD BE
9	FUNDED IF SO MANY DECISIONS ARE OVERTURNED HERE.
10	MR. SHEEHY: THANK YOU.
11	DR. LEVIN: I'M NOT SUPPOSED TO SPEAK ON
12	THIS. IT'S A POINT OF CLARIFICATION. TIER II IS
13	FUND IF FUNDS ARE AVAILABLE.
14	MR. SHEEHY: YOU CAN SPEAK TO THIS, I
15	THINK.
16	DR. LEVIN: THIS IS NOT OVERTURNING PEER
17	REVIEW. IT WAS THE CIRM STAFF RECOMMENDATION THAT
18	TIER II NOT BE FUNDED. BUT THE PEER REVIEW SAYS
19	FUND IF FUNDS ARE AVAILABLE, WHICH IS STILL
20	CONSISTENT WITH THE \$40 MILLION CAP.
21	DR. MILLS: THAT'S JUST WRONG. TIER II IS
22	IT WAS OF MODERATE QUALITY OR IT BIFURCATED AS IN
23	THERE WAS NO CONSENSUS. MEANING SOME PEOPLE
24	DR. LEVIN: I THOUGHT THAT WAS ONLY FOR
25	CIRM 2.0.
	96

1	DR. MILLS: THAT'S THE INSTRUCTIONS THAT
2	ARE GIVEN NOW TO THEM IS VOTE USE THE ENTIRE
3	SPECTRUM AND VOTE IT WHEREVER YOU THINK ALONG THAT
4	SPECTRUM IT IS. BUT IT CAN ONLY BE IN TIER II IF
5	YOU THINK IT'S OF MODERATE QUALITY.
6	DR. JUELSGAARD: WELL, THERE'S A SECOND
7	PART TO TIER II, RANDY, WHICH IS OR CONSENSUS ON
8	MERIT WAS NOT REACHED. IT MAY BE
9	DR. MILLS: THAT'S AN EFFECT OF HOW THEY
10	VOTED, NOT AN INSTRUCTION ON HOW TO VOTE. THEY
11	WOULDN'T KNOW THERE WAS NOT A CONSENSUS.
12	DR. JUELSGAARD: WELL, I'M GOING TO FOCUS
13	ON THAT FOR A MOMENT BECAUSE ONE OF THE THINGS, AND
14	I SAID THIS EARLIER, IT'S A PROCESS ISSUE FOR ME, IS
15	WHEN YOU HAVE THESE LARGE STANDARD DEVIATIONS GOING
16	ON, AND HERE YOU HAVE A LOW SCORE OF 45 LAID UP
17	AGAINST A HIGH SCORE OF 85 AND YOU HAD A MEDIAN
18	SCORE OF 75. AND SO THERE'S THIS HUGE RANGE GOING
19	ON THROUGHOUT ALL OF THIS, AND THAT'S A LITTLE
20	BOTHERSOME TO ME THAT THERE ARE SUCH DIVERGENT
21	POINTS OF VIEW. THAT'S WHY I FOCUS ON CONSENSUS ON
22	MERIT NOT REACHED. IN OTHER WORDS, SOME PEOPLE
23	THOUGHT IT WAS VERY MERITORIOUS AND SOME PEOPLE
24	THOUGHT THAT IT WASN'T.
25	SO THEN IF YOU GO BACK AND LOOK AT THE

1	COMMENTS THAT WERE MADE, FROM MY POINT OF VIEW,
2	THERE ARE TWO PRIMARY COMMENTS. ONE IS THAT YOU
3	SHOULD BE PURSUING ONE INDICATION, NOT TWO. AND
4	THAT'S ALWAYS A DEBATE THAT GOES ON WHEN YOU'RE
5	FIRST DEVELOPING A DRUG. DO YOU FASHION ON ONE
6	INDICATION, OR DO YOU FASHION ON MORE THAN ONE
7	INDICATION? AND IT'S NOT AN INAPPROPRIATE DECISION
8	AT THE VERY BEGINNING TO ACTUALLY LOOK AT MORE THAN
9	ONE AND THEN NARROW OVER TIME.
10	THE SECOND CRITICISM WAS AROUND FDA
11	INVOLVEMENT AND WHERE WAS THE FDA EXPERTISE. I
12	DON'T REALLY KNOW THE ANSWER TO THAT QUESTION. I DO
13	KNOW THAT FDA EXPERTISE CAN BE BROUGHT INTO THESE
14	PROJECTS TO MAKE SURE THAT YOU DO THE RIGHT THINGS
15	ALONG THE WAY FROM AN FDA POINT OF VIEW. BUT I
16	DIDN'T SEE ANY OTHER SUBSTANTIAL CRITICISMS AT LEAST
17	OF THE SCIENCE THAT'S GOING ON HERE.
18	SO TO THE POINT THAT YOU FEEL CONCERNED
19	ABOUT, IN ESSENCE, OVERRULING, SO TO SPEAK, THE
20	GRANTS WORKING GROUP, I THINK, PARTICULARLY IN A
21	CASE LIKE THIS WHERE THERE'S SUCH A WIDE RANGE OF
22	OPINION, THAT REALLY IS OUR JOB. OUR JOB IS TO
23	THINK ABOUT WHETHER OR NOT WHAT THE GRANTS WORKING
24	GROUP DID WAS A GOOD, FULL, FAIR ANALYSIS
25	PARTICULARLY WHEN YOU'VE GOT SUCH DIVERGENT POINTS

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1
     OF VIEW ON THE GRANTS WORKING GROUP. SO I'M NOT --
 2
     I GUESS THE PROCESS IS ONE THAT BOTHERS ME, AND I
 3
     DON'T QUITE KNOW HOW WE GET TO THAT POINT OF VIEW.
 4
     AND THEN I WONDER ABOUT WHO WAS ON THE GRANTS
 5
     WORKING GROUP THAT REVIEWED THIS. WHERE WAS THE
 6
     BREADTH OF EXPERTISE, ETC.? SO ANYWAY, I'LL BE
 7
     QUIET.
 8
                MR. SHEEHY: SO I HAVE DR. DULIEGE AND
 9
     THEN I THINK DR. PRIETO. I HOPE I'M NOT NEGLECTING
10
     PEOPLE OVER HERE.
11
               MS. WINOKUR: MAY I SPEAK TO MY ORIGINAL
12
     COMMENT?
13
               MR. SHEEHY: IF EVERYONE IS OKAY BECAUSE
14
     I'M GETTING A CUE HERE.
15
                MS. WINOKUR: MY ORIGINAL COMMENT HAD
16
     NOTHING TO DO WITH WHETHER WE HAVE ENOUGH MONEY TO
17
     DO THIS. IT WAS A COMMENT REGARDING OUR PROCESS
     HERE TODAY AND IF WE REALIZE THAT WHAT WE ARE DOING
18
19
     IS OVERTURNING MOST OF THE DECISIONS THAT WERE MADE
20
     TO PLACE FIVE PROJECTS IN TIER II.
21
                DR. DULIEGE: I JUST WANT TO COME BACK AND
22
     ADD A LITTLE BIT TO WHAT YOU WERE SAYING, STEVE.
23
     I'M TALKING ABOUT THE GENERAL PROCESS, NOT ABOUT
24
     THIS PARTICULAR APPLICATION FOR WHICH I'M
25
     CONFLICTED. I THINK IN GENERAL WE HAVE EXACTLY TO
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1	USE OUR JUDGMENT ABOUT THE CRITERIA FOR WHICH A
2	PARTICULAR APPLICATION WAS PUT IN TIER II. AND IF
3	IT'S SCIENTIFIC MERIT, IT'S A REALLY IMPORTANT
4	CONCERN. IF IT'S ABOUT ACQUIRING EXPERTISE, AND
5	PARTICULARLY, AS YOU SAID, STEPHEN, IN REGULATORY,
6	THERE ARE MANY, MANY PEOPLE THAT CAN COME. THIS IS
7	VERY EASILY MANAGEABLE. THIS IS WHY I THINK WE
8	SHOULD
9	MR. SHEEHY: I THINK WE'RE SPEAKING TO THE
10	GRANT, AREN'T WE?
11	DR. DULIEGE: ABOUT THE PROCESS IN
12	GENERAL.
13	DR. MILLS: I JUST WANT TO MAKE TWO
14	COMMENTS. ONE, I STRONGLY, STRONGLY CAUTION YOU ON
15	READING TOO MUCH INTO THE SUMMARIES ON WHAT THE GWG
16	THOUGHT AND DELIBERATED ABOUT. IT WAS OUR BEST
17	EFFORT TO TRY TO FIGURE OUT WHAT THEY THOUGHT WAS
18	IMPORTANT, BUT I CAN TELL YOU AS A GWG MEMBER
19	HISTORICALLY, HAVING LOOKED BACK ON WHAT MADE OR NOT
20	MADE IT IN THE SUMMARY, THEY DO NOT CORRELATE UNDER
21	THE CURRENT PROCESS VERY WELL WITH WHY SOMEBODY
22	FEELS LIKE THEY MAKE A DECISION OR DON'T FEEL.
23	HOPEFULLY WE'RE GOING TO ADDRESS THAT IN 2.0 BY
24	ASKING THE REVIEWERS DIRECTLY TO COMMENT AND THEN
25	GIVING YOU THOSE COMMENTS DIRECTLY. THAT IS NOT
	100

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1	WHAT WE HAVE NOW.
2	THIS HAPPENS ALL THE TIME WHERE YOU SEE A
3	GRANT AND SAY IF BUT NOT FOR THAT, THIS THING WOULD
4	HAVE BEEN PERFECT. AND HAVING GONE THROUGH THAT
5	EXPERIENCE NOW ON GRANTS I HAD REVIEWED, I WILL JUST
6	SAY IN THE CASES WHERE IT'S INVOLVED ME, THE ANSWER
7	WAS NO. IT WASN'T JUST THAT ONE THING.
8	THE SECOND PART GOES TO THIS COMMENT ABOUT
9	WHEN THERE'S NOT CONSENSUS. I AGREE WITH DR.
10	JUELSGAARD. IT'S EXACTLY WHAT YOU SHOULD DO. THESE
11	DO HAPPEN TO BE BIFURCATED RESULTS FROM THE GWG
12	WHERE THERE CLEARLY WASN'T CONSENSUS. SOME THOUGHT
13	IT WAS GOOD; SOME CLEARLY THOUGHT IT WASN'T GOOD.
14	AND SO I THINK THAT'S EXACTLY THE KIND OF THING YOU
15	SHOULD BE DOING. I WANT TO MAKE SURE EVERYONE
16	UNDERSTANDS THAT MAKING A DECISION THAT'S CONTRARY
17	TO THE RECOMMENDATION FROM CIRM IS NOT TAKEN IN ANY
18	WAY BY US NEGATIVELY OR OUT OF CONTEXT. THIS IS
19	YOUR JOB TO DO THAT. AND WE LAID OUT A RATIONALE
20	FOR WHY WE MADE THE RECOMMENDATIONS WE MADE, BUT IT
21	IS ABSOLUTELY APPROPRIATE FOR YOU GUYS TO DO THAT, I
22	THINK.
23	MR. SHEEHY: DR. PRIETO, I THINK, AND
24	THEN
25	DR. PRIETO: MY QUESTION WAS ANSWERED.
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1	DR. HIGGINS: I THINK WE'RE IN A UNIQUE
2	SITUATION TODAY IN THAT WE'VE GOT ONE FOOT IN CIRM
3	1.0 AND WE'VE GOT ONE FOOT IN CIRM 2.0. AND I THINK
4	IF YOU LOOK AT THE SCORES AND YOU JUDGE THEM BY ONE
5	CRITERIA, YOU GET ONE RESULT. YOU JUDGE THEM BY
6	ANOTHER CRITERIA IN 2.0, YOU GET A DIFFERENT RESULT.
7	SO I THINK WE NEED TO BE A LITTLE BIT SENSITIVE TO
8	THE PAST AND THE FUTURE.
9	AND I THINK THIS ACTIVISM AS IT SEEMS ON
10	THE BOARD IS LOOKING AT SOME OF THESE SCORES FROM
11	THE 2.0 PERSPECTIVE, AND I SUPPORT DOING THAT. I
12	DON'T KNOW IF I'M BEING OBSCURE OR NOT. I'M NOT
13	TRYING TO BE, BUT I JUST THINK TODAY IS A SPECIAL
14	CASE THAT'S A LITTLE BIT COMPLICATED BECAUSE WE'RE
15	ALL SORT OF THINKING 2.0, BUT WHAT WE'VE GOT IN
16	FRONT OF US IS 1.
17	MR. SHEEHY: ARE THERE FURTHER BOARD
18	COMMENTS OR QUESTIONS?
19	MR. TORRES: THANK YOU, DAVID. I THINK
20	THAT WAS A VERY PERCEPTIVE COMMENT. AND ALSO THAT
21	DOESN'T MEAN IF WE VOTE IN FAVOR OF MOVING THIS
22	PROJECT TO TIER I THAT THAT'S THE END OF IT BECAUSE
23	WHAT I HAVE EXPERIENCED IN THE PAST, AND I THINK
24	THAT WILL CONTINUE TO BE THE CASE, IF STAFF FEELS
25	THAT THIS PROJECT IS NOT MOVING TOWARDS ITS
	102

1	BENCHMARKS, IF IT'S NOT BEING SUCCESSFUL, IT WILL BE
2	DEFUNDED; IS THAT CORRECT?
3	DR. MILLS: YEAH. IF IT DOESN'T HIT
4	GO/NO-GO MILESTONES, OBVIOUSLY WE WOULD STOP A
5	PROGRAM. OUR RATIONALE, AGAIN, FROM 2.0 WOULD BE
6	LET'S LAUNCH A PROGRAM WITH THE GREATEST CHANCE OF
7	SUCCESS THAN LAUNCH A PROJECT THAT HAS FLAWS
8	ASSOCIATED WITH IT AND THEN TRY TO FIX THEM AS WE'RE
9	ALSO TRYING TO MOVE THE PROGRAM ALONG.
10	MR. SHEEHY: ANY OTHER BOARD COMMENT OR
11	QUESTIONS? ANY PUBLIC COMMENT?
12	DR. MADIGAN: THANK YOU. SANDY MADIGAN
13	AGAIN, CEO OF ANKASA IN SUPPORT OF JILL HELMS AND IN
14	SOME REGARD TO EVERYBODY ELSE THAT'S IN TIER II WITH
15	SOME OF MY COMMENTS.
16	SO FIRST, I'M SORRY MY OPTOMETRIST SETS
17	MY VISION SO I CAN READ AT COMPUTER DISTANCE. SO UP
18	CLOSE DISTANCE, I HAVE TROUBLE. I CAN'T QUITE GET
19	THIS GENTLEMAN'S NAME ON MY RIGHT, BUT CAN I JUST
20	HAND YOU MY NOTES AND YOU TAKE OVER BECAUSE YOU DID
21	A FABULOUS JOB OF SUPPORTING THE PROJECT. I
22	APPRECIATE THAT.
23	SO FIRST OF ALL, I'D LIKE TO ADDRESS. SO
24	LET'S TALK ABOUT CIRM 2.0. YES, IT'S THE NEW KID ON
25	THE BLOCK AND WE CERTAINLY RECOGNIZE THAT THE ADMIN
	103
	TOO

1	WOULD LOVE TO MOVE ALL THE TIER IIS INTO CIRM 2.0
2	BECAUSE THEY'D LIKELY BE EARLY SUCCESSES FOR THE
3	ADMINISTRATION.
4	SECOND, LET'S JUST STOP WITH SEMANTICS
5	ABOUT SCORING. THERE ARE SIX GRANTS LISTED UP THERE
6	WITH A MEAN OF 75. HOW DO YOU DIFFERENTIATE BETWEEN
7	THE TWO THAT ARE IN TIER I AND THE FOUR THAT ARE IN
8	TIER II? YOU CANNOT. YOU CAN'T DO IT. WHAT
9	SEPARATES THOSE? I THINK IN TERMS OF THIS
10	PARTICULAR RFA, THIS RFA WAS ANNOUNCED A YEAR AGO.
11	GRANTS WERE SUBMITTED IN NOVEMBER. THERE WAS NO
12	GUIDANCE ABOUT ANY CIRM 2.0, BUT NOW WE'RE APPLYING
13	THOSE CRITERIA. THAT'S TACITLY UNFAIR.
14	FINALLY, IN TERMS OF SOME OF THE SPECIFIC
15	CRITICISMS OF THE REVIEWERS, TOO MANY USES, IT'S THE
16	FIRST TIME AS A BUSINESS PERSON I'VE EVER BEEN
17	CRITICIZED FOR HAVING TOO MANY APPLICATIONS FOR MY
18	PRODUCT.
19	A SYNTHETIC CARRIER, I BROUGHT THIS UP IN
20	MY THREE-MINUTE TALK. THAT IS A PATENTLY
21	MISUNDERSTOOD APPLICATION OF OUR TECHNOLOGY. IT MAY
22	HAVE LED TO THAT 45, IT MAY HAVE BEEN A SEPARATE
23	REVIEWER, IMPOSSIBLE FOR ME TO KNOW, BUT THERE WAS A
24	REVIEWER WHO DID NOT UNDERSTAND THE PROCESS, PERIOD.
25	TWO DIFFERENT REVIEWERS COMMENTED ON
	104
	I TO4

1	AUTOGRAFT. ONE SAID TOO MUCH RELIANCE ON AUTOGRAFT.
2	ONE SAID, OH, AUTOGRAFT IS GOING TO BE BETTER THAN
3	STANDARD OF CARE. WHICH IS IT? MAKE A DECISION.
4	AND FINALLY, COMMENT ABOUT MINIMIZING TIME
5	UNDER ANESTHESIA, IRRELEVANT. WE ARE APPLYING INTO
6	A BONA FIDE ONGOING SURGICAL PROCEDURE THAT DOCS DO
7	ON A DAILY BASIS. WE DON'T EITHER EXTEND OR
8	DIMINISH TIME UNDER ANESTHESIA. THAT'S NOT OUR
9	PROBLEM. IT'S NOT EVEN PART OF WHAT WE'RE TALKING
10	ABOUT HERE.
11	AND THEN FINALLY, I'M VERY DISTURBED, JUST
12	AS A CALIFORNIA CITIZEN AND VOTER, THAT WE HAVE
13	BOARD MEMBERS WHO ARE UNWILLING TO DO THEIR JOB AND
14	MOVE THESE APPLICATIONS AS THEY SHOULD BE MOVED IF
15	THEY'RE ELIGIBLE AND PEOPLE BELIEVE THEY SHOULD BE
16	MOVED. AND RESPECTFULLY SUBMIT IF THERE ARE PEOPLE
17	THAT ARE NOT WILLING TO DO THAT AND TAKE ON THAT
18	RESPONSIBILITY, WE SHOULD FIND PEOPLE THAT ARE
19	WILLING TO DO SO. THANK YOU.
20	MR. SHEEHY: I JUST WANT TO TAKE OFFENSE
21	AT YOUR LAST COMMENT. WE WERE APPOINTED TO DO OUR
22	JOBS AND NOT TO FUND YOU. AND, PLEASE, YOUR PUBLIC
23	COMMENT IS OVER.
24	DR. MADIGAN: I DIDN'T SUGGEST THAT YOU
25	FUND ME. I SAID IF THERE ARE PEOPLE THAT ARE
	105

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1
     UNCOMFORTABLE MOVING GRANTS, WHICH IS PART OF THE
 2
     BODY OF THIS JOB --
 3
               MR. SHEEHY: I DID NOT RECOGNIZE YOU FOR
 4
     PUBLIC COMMENT.
 5
                DR. MADIGAN: I DIDN'T ASK YOU FOR PUBLIC
 6
     COMMENT.
 7
               MR. TORRES: AT THIS POINT I WOULD CALL
     THE SERGEANT AT ARMS, BUT I DON'T HAVE ONE.
 8
 9
                MR. SHEEHY: I'M SURPRISED.
10
                DR. HELMS: JILL HELMS FROM STANFORD. I
11
     FEEL AS PASSIONATELY AS SANDY DOES ABOUT THIS, ONLY
12
     I'VE BEEN BROUGHT UP IN ACADEMICS. SO, OF COURSE,
13
     WE DON'T SAY THESE THINGS. I MUST SAY THAT IT'S
14
     BEEN VERY DIFFICULT TO UNDERSTAND THE PROCESS BY
15
     WHICH MEDIAN AND AVERAGE SCORES ARE JUDGED. I HOPE
16
     THAT YOU THINK ABOUT THE SCIENCE. I THINK IT'S
17
     VERY, VERY STRONG, AND I HOPE THAT YOU THINK ABOUT
18
     THAT IN YOUR VOTE. THANK YOU.
19
                DR. MILLS: I'VE JUST GOT TO SAY ONE
20
     THING. THERE'S SOME SPEAKING TO THE BOARD, AND I
21
     HOPE THE BOARD KNOWS THIS, UNDER NO CIRCUMSTANCES
22
     WOULD I OR ANY MEMBER OF MY TEAM EVER MAKE A
     RECOMMENDATION TO DO ANYTHING THAT WASN'T WHAT WE
23
24
     FELT WAS IN THE BEST INTEREST OF OUR MISSION, WHICH
25
     IS TO ACCELERATE STEM CELL TREATMENTS TO PATIENTS
                               106
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1	WITH UNMET MEDICAL NEEDS. IT IS WHY I START EVERY
2	PRESENTATION WITH A REMINDER THAT IT'S THE PATIENTS
3	THAT COME FIRST. AND UNDER NO CIRCUMSTANCES WOULD
4	WE DO SOMETHING TO SECURE AN EARLY WIN FOR THE
5	ADMINISTRATION.
6	CHAIRMAN THOMAS: I WANT TO COMMENT JUST
7	AS A GENERAL REMARK THAT TESTIMONY THAT WE JUST
8	HEARD TWO SPEAKERS AGO I VIEW AS ENTIRELY
9	NONPRODUCTIVE. THERE'S A PROTOCOL AND A SENSE OF
10	DECORUM THAT ONE HAS TO FOLLOW IN DEALING IN A
11	PUBLIC AGENCY SETTING LIKE THIS. AND I JOIN MR.
12	SHEEHY IN FINDING THE TENOR OF THOSE COMMENTS TO BE
13	VERY OFFENSIVE AND COUNTERPRODUCTIVE TO THE POINTS
14	THAT YOU WERE MAKING.
15	MR. TORRES: HERE. HERE.
16	MR. SHEEHY: THIS IS NOT TIME FOR PUBLIC
17	COMMENT.
18	DR. MADIGAN: I JUST WANTED TO APOLOGIZE.
19	YOU'RE ABSOLUTELY RIGHT, MR. CHAIRMAN. I APOLOGIZE.
20	I APOLOGIZE FOR BEING OUT OF BOUNDS, BUT I AM VERY
21	PASSIONATE ABOUT THIS PROJECT. SO PLEASE ACCEPT MY
22	APOLOGY.
23	MR. SHEEHY: I JUST THINK IT'S IMPORTANT
24	TO REMEMBER THAT THE PEOPLE WHO ARE SERVING HERE ARE
25	SERVING AS VOLUNTEERS. AND AS SOMEONE WHO'S BEEN ON
	107
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1	THIS BOARD SINCE THE BEGINNING FOR TEN YEARS NOW,
2	YOU KNOW, THE TIME AND EFFORT WE PUT INTO TRYING TO
3	UNDERSTAND THE SCIENCE AND DO THE BEST BY THE PEOPLE
4	OF CALIFORNIA HAS BEEN UNMATCHED, I THINK. THE
5	DEDICATION OF THE BOARD MEMBERS I'VE SERVED WITH,
6	THE COMPLETE ABSENCE OF PERSONAL INTEREST, AND
7	DEDICATION TO DOING THE BEST FOR THE PATIENTS OF
8	CALIFORNIA, I THINK, HAS BEEN IN EVIDENCE FOR THE
9	LAST TEN YEARS.
10	(APPLAUSE.)
11	MR. SHEEHY: AND I WANT TO REITERATE THAT
12	TO CHALLENGE THAT DOESN'T REALLY GET TO THE HEART OF
13	WHY WE'RE HERE. WE'RE HERE TO BE CAREFUL STEWARDS
14	OF THE STATE'S MONEY. THIS IS NOT OUR MONEY. THIS
15	BELONGS TO THE CITIZENS OF CALIFORNIA. AND IT'S TO
16	BALANCE THAT STEWARDSHIP WITH THE NECESSITY OF
17	GETTING TREATMENTS TO PATIENTS AS QUICKLY AND
18	EFFICIENTLY AS WE CAN. SO I'LL JUST LEAVE IT AT
19	THAT. I THINK WE'RE READY TO CALL THE ROLL.
20	DR. JUELSGAARD: JUST TWO QUICK COMMENTS.
21	FIRST OF ALL, I COMPLETELY AGREE WITH EVERYTHING
22	THAT YOU JUST SAID, AND I THINK IT'S VERY
23	UNFORTUNATE THE COMMENTS THAT WERE PREVIOUSLY MADE
24	EVEN THOUGH THERE'S BEEN AN APOLOGY ISSUED.
25	BUT FOR ME THIS VOTE IS NOT SO MUCH ABOUT
	100

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1	THAT. IN FACT, IT'S NOT ALL ABOUT THAT, BUT IT'S ON
2	WHAT I PERSONALLY THINK ABOUT THE VALUE OF THIS
3	PROGRAM AND WHETHER OR NOT IT'S SOMETHING THAT WE
4	SHOULD FUND, AND IT'S NOT ABOUT WHETHER WE'RE IN THE
5	OLD SYSTEM OR THE NEW SYSTEM OR WHATEVER, BUT BASED
6	ON WHAT I READ AS THE COMMENTS, DR. MILLS, THIS IS
7	ALL I HAVE TO GO ON WHAT ARE WRITTEN DOWN. SO IF
8	THERE WERE STRONGER FEELINGS THAT JUST DIDN'T MAKE
9	IT INTO THIS, THERE'S NO WAY I CAN JUDGE. BUT FOR
10	ME IT'S ABOUT WHETHER OR NOT THIS IS A PROGRAM THAT
11	WOULD FIT WELL WITHIN OUR PORTFOLIO OF PROJECTS WITH
12	A STRONG SCIENTIFIC MERIT AND THAT HAS A GREAT TEAM
13	BEHIND IT AND WHETHER OR NOT THE PROCESS THAT WAS
14	USED BY THE GWG WAS ONE WHICH YOU COULD SAY WAS LESS
15	THAN PERFECT BECAUSE I THINK IT WAS. IF YOU LOOK AT
16	THE VOTES, YOU HAVE EIGHT PEOPLE PUTTING THIS IN
17	TIER I, THREE IN TIER II, AND THEN FOUR IN TIER III.
18	IT'S JUST TO ME NOT VERY UNDERSTANDABLE HOW YOU CAN
19	HAVE THOSE KIND OF OUTLIERS GOING ON.
20	SO, ANYWAY, JUST WANTED TO ISSUE MY VIEW
21	OF THIS PROPOSAL.
22	MS. WINOKUR: I THINK THAT THIS DISCUSSION
23	IS A VERY GOOD ONE TO SUPPORT THE CHANGE TO 2.0.
24	MR. SHEEHY: THANK YOU. I THINK WE'RE
25	READY TO CALL THE ROLL.
	109
	109

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1	MS. BONNEVILLE: DAVID HIGGINS.
2	DR. HIGGINS: YES.
3	MS. BONNEVILLE: STEPHEN JUELSGAARD.
4	MR. JUELSGAARD: YES.
5	MS. BONNEVILLE: KATHY LAPORTE.
6	DR. LAPORTE: YES.
7	MS. BONNEVILLE: LAUREN MILLER.
8	MS. MILLER: YES.
9	MS. BONNEVILLE: FRANCISCO PRIETO.
10	DR. PRIETO: ABSTAIN.
11	MS. BONNEVILLE: ROBERT QUINT.
12	DR. QUINT: ABSTAIN.
13	MS. BONNEVILLE: AL ROWLETT.
14	MR. ROWLETT: ABSTAIN.
15	MS. BONNEVILLE: JEFF SHEEHY.
16	MR. SHEEHY: ABSTAIN.
17	MS. BONNEVILLE: JONATHAN THOMAS.
18	CHAIRMAN THOMAS: NO.
19	MS. BONNEVILLE: ART TORRES.
20	MR. TORRES: AYE.
21	MS. BONNEVILLE: DIANE WINOKUR.
22	MS. WINOKUR: ABSTAIN.
23	MR. HARRISON: THE MOTION FAILS WITH FIVE
24	YES VOTES, FIVE ABSTENTIONS, AND ONE NO VOTE.
25	MR. SHEEHY: NOW I WILL TAKE A MOTION TO
	110
	_== <del>*</del>

1	FUND ALL THE APPLICATIONS AND NOT FUND THE REMAINING
2	APPLICATIONS, AND THIS SHOULD BE SOMEONE WITHOUT A
3	CONFLICT.
4	MR. TORRES: SO MOVED.
5	MR. SHEEHY: SENATOR TORRES. DO I HAVE A
6	SECOND?
7	MS. MILLER: SECOND.
8	MR. SHEEHY: SECONDED BY MS. MILLER. ANY
9	BOARD COMMENT? ANY PUBLIC COMMENT? CALL THE ROLL.
10	AND COULD YOU INFORM US ABOUT THE FORM THAT THIS
11	SHOULD TAKE?
12	MR. HARRISON: YES. PLEASE REMEMBER, IF
13	YOU HAVE A CONFLICT WITH RESPECT TO A PARTICULAR
14	APPLICATION, TO VOTE EITHER A YES OR NO EXCEPT WITH
15	RESPECT TO THOSE APPLICATIONS FOR WHICH YOU HAVE A
16	CONFLICT.
17	MS. BONNEVILLE: ANNE-MARIE DULIEGE.
18	DR. DULIEGE: YES, EXCEPT FOR THOSE WITH
19	WHICH I HAVE A CONFLICT.
20	MS. BONNEVILLE: DAVID HIGGINS.
21	DR. HIGGINS: YES, EXCEPT FOR THOSE WITH
22	WHICH I HAVE A CONFLICT.
23	MS. BONNEVILLE: STEPHEN JUELSGAARD.
24	MR. JUELSGAARD: YES.
25	MS. BONNEVILLE: KATHY LAPORTE.
	111

1	DR. LAPORTE: YES.
2	MS. BONNEVILLE: LAUREN MILLER.
3	MS. MILLER: YES.
4	MS. BONNEVILLE: FRANCISCO PRIETO.
5	DR. PRIETO: YES, EXCEPT FOR THOSE WITH
6	WHICH I HAVE A CONFLICT.
7	MS. BONNEVILLE: ROBERT QUINT.
8	DR. QUINT: YES.
9	MS. BONNEVILLE: AL ROWLETT.
10	MR. ROWLETT: YES, EXCEPT FOR THOSE WITH
11	WHICH I HAVE A CONFLICT.
12	MS. BONNEVILLE: JEFF SHEEHY.
13	MR. SHEEHY: YES, EXCEPT FOR THOSE WITH
14	WHICH I HAVE A CONFLICT.
15	MS. BONNEVILLE: JONATHAN THOMAS.
16	CHAIRMAN THOMAS: YES.
17	MS. BONNEVILLE: ART TORRES.
18	MR. TORRES: AYE.
19	MS. BONNEVILLE: DIANE WINOKUR.
20	MS. WINOKUR: YES.
21	MR. HARRISON: MOTION CARRIES.
22	MR. TORRES: WELL DONE, MR. CHAIRMAN
23	SHEEHY.
24	CHAIRMAN THOMAS: I KNOW YOU NEED A BREAK
25	DESPERATELY. WE'RE GOING TO TAKE A FIVE-MINUTE
	112
	±± <u>¢</u>

1	BREAK. WE HAVE A NUMBER OF ITEMS OF BUSINESS TO GET
2	TO HERE, SO PLEASE BE BACK IN FIVE. DON, I THINK
3	WE'RE GOING TO TAKE YOUR PICTURE LATER THAN NOW.
4	(A RECESS WAS TAKEN.)
5	CHAIRMAN THOMAS: OKAY. WOULD EVERYBODY
6	PLEASE TAKE YOUR SEATS. COULD THOSE IN THE AUDIENCE
7	EITHER PLEASE TAKE YOUR SEATS OR CONTINUE THEIR
8	CONVERSATIONS IN THE HALLWAY? THANK YOU.
9	WE ARE NOW GOING TO PROCEED TO ITEM NO. 8,
10	CONSIDERATION OF DEFERRED APPLICATIONS RT THEY'RE
11	LONG NUMBERS. I'LL LET GIL TALK ABOUT THEM FOR
12	RFA 13-05, WHICH IS A CIRM TOOLS AND TECHNOLOGIES
13	III ROUND.
14	DR. SAMBRANO: THANK YOU, MR. CHAIRMAN,
15	MEMBERS OF THE BOARD. SO YOU MIGHT RECALL THAT AT
16	OUR LAST MEETING IN JANUARY, THERE WERE TWO
17	APPLICATIONS FROM THE TOOLS AND TECHNOLOGY II, THE
18	
	RFA 13-05 THAT WERE DEFERRED FROM CONSIDERATION IN
	RFA 13-05 THAT WERE DEFERRED FROM CONSIDERATION IN ORDER TO ASSESS APPEALS THAT WERE MADE BY THE
19	
19 20	ORDER TO ASSESS APPEALS THAT WERE MADE BY THE
19 20 21	ORDER TO ASSESS APPEALS THAT WERE MADE BY THE APPLICANTS. AND THESE WERE BOTH BASED ON A MATERIAL
19 20 21 22	ORDER TO ASSESS APPEALS THAT WERE MADE BY THE APPLICANTS. AND THESE WERE BOTH BASED ON A MATERIAL DISPUTE OF FACT. BOTH HAD SET FORTH ADEQUATE
19 20 21 22 23	ORDER TO ASSESS APPEALS THAT WERE MADE BY THE APPLICANTS. AND THESE WERE BOTH BASED ON A MATERIAL DISPUTE OF FACT. BOTH HAD SET FORTH ADEQUATE GROUNDS FOR CONSIDERATION, SO WHAT WE DO IS WE PUT
19 20 21 22 23 24 25	ORDER TO ASSESS APPEALS THAT WERE MADE BY THE APPLICANTS. AND THESE WERE BOTH BASED ON A MATERIAL DISPUTE OF FACT. BOTH HAD SET FORTH ADEQUATE GROUNDS FOR CONSIDERATION, SO WHAT WE DO IS WE PUT TOGETHER A SUBCOMMITTEE OF THE GRANTS WORKING GROUP

1	WORKING GROUP REVIEW, THREE SCIENTIFIC REVIEWERS,
2	AND ONE OR TWO PATIENT ADVOCATE MEMBERS.
3	NOW, IN BOTH OF THESE CASES, THE GRANTS
4	WORKING GROUP REVIEWERS AGREED THAT THE
5	CLARIFICATION ABOUT THE DISPUTED FACTS DID NOT
6	IMPACT THEIR SCORES AND WOULD NOT HAVE CHANGED THEIR
7	RECOMMENDATION. THEREFORE, BOTH APPLICATIONS ARE TO
8	BE CONSIDERED BY THE ICOC'S APPLICATION REVIEW
9	SUBCOMMITTEE WITH NO CHANGE FROM THE GRANTS WORKING
10	GROUP IN TERMS OF THE RECOMMENDATION.
11	SO I DON'T KNOW THAT YOU CAN SEE THIS ON
12	THE SCREEN. I THINK YOU MAY HAVE A COPY OF IT.
13	THIS IS THE TABLE IN RANK ORDER FROM THE TOOLS AND
14	TECHNOLOGY APPLICATIONS THAT WERE RECEIVED. AND
15	IT'S HERE JUST REALLY TO SERVE AS A REMINDER, AND
16	I'LL GO OVER THIS BRIEFLY.
17	SO THE TWO APPLICATIONS IN QUESTION, THE
18	FIRST ONE IS RT3-07836. THAT ONE RECEIVED A SCORE
19	OF 64, WHICH PLACED IT IN TIER III, AND THE
20	RECOMMENDATION FROM THE WORKING GROUP WAS NOT
21	RECOMMENDED FOR FUNDING. SO, THEREFORE, IT
22	CONTINUES TO BE IN TIER III. AND THE CIRM TEAM
23	RECOMMENDATION CONCURS WITH THAT WORKING GROUP
24	RECOMMENDATION.
25	THE OTHER APPLICATION THAT WAS CONSIDERED
	114

1	WAS RT3-07678. THAT ONE HAD A SCORE OF 74. THE
2	GRANTS WORKING GROUP RECOMMENDATION WAS A TIER II
3	THAT WAS MODERATE QUALITY OR NO CONSENSUS. IT ALSO
4	HAD A CIRM TEAM RECOMMENDATION TO FUND, AND WE
5	CONTINUE TO SUPPORT THAT RECOMMENDATION.
6	OUR RATIONALE FOR CHOOSING TO RECOMMEND
7	THIS ONE FOR FUNDING IS BASED ON THAT THIS PROPOSAL
8	ADDRESSES THE SAFETY OF HUMAN PLURIPOTENT STEM
9	CELL-DERIVED CELLS FOR TRANSPLANTATION. IT'S A
10	CRITICAL BOTTLENECK FOR CLINICAL APPLICATION OF STEM
11	CELL-DERIVED THERAPIES. WE CURRENTLY DO NOT HAVE
12	ANY ACTIVE GRANTS IN THE CIRM PORTFOLIO THAT ADDRESS
13	THIS SPECIFIC BOTTLENECK. THIS IS A SMALL MOLECULE
14	APPROACH THAT WE THINK IS A POTENTIAL COST-EFFECTIVE
15	APPROACH THAT CAN ADDRESS THIS BOTTLENECK.
16	AND THEN YOU MAY ALSO NOTICE IN THAT TIER
17	II GROUP THAT WAS CONSIDERED LAST TIME THERE WERE
18	TWO OTHER TOOLS AND TECH APPLICATIONS THAT WERE
19	RECOMMENDED FOR FUNDING BY THE CIRM TEAM AND
20	APPROVED FOR FUNDING BY THE ICOC. THIS APPLICATION,
21	LIKE THOSE OTHER TWO, HAD A MEDIAN SCORE OF 75 AND A
22	MAJORITY OF THE VOTING MEMBERS OF THE GRANTS WORKING
23	GROUP SCORED THE APPLICATION WITH 75 OR ABOVE.
24	SO WE ARE THEN OPEN TO YOUR CONSIDERATION
25	FOR THESE TWO PROPOSALS.
	115

1	CHAIRMAN THOMAS: TURN THIS OVER TO
2	MR. SHEEHY FOR PROGRAMMATIC.
3	MR. SHEEHY: SURE. AND I'M WONDERING I
4	THINK WE'RE MISSING A COUPLE OF FOLKS, DR. PRIETO
5	AND MR. JUELSGAARD. DO WE NEED THEIR VOTES IN ORDER
6	TO CONSIDER THIS? I JUST WANT TO MAKE SURE THAT WE
7	DON'T I DON'T KNOW WHAT OUR QUORUM CONSTRAINTS
8	ARE.
9	MR. HARRISON: WE HAVE A QUORUM IN THE
10	ROOM.
11	MR. SHEEHY: THEN LET'S TAKE UP THE STAFF
12	RECOMMENDATION TO FUND FIRST, I BELIEVE, BECAUSE THE
13	SECOND ONE I BELIEVE I HAVE A CONFLICT, SO SOMEONE
14	WILL ELSE WILL HAVE TO CARRY THAT ONE. CHAIRMAN
15	THOMAS.
16	SO DO WE HAVE A MOTION TO ACCEPT THE STAFF
17	RECOMMENDATION FOR THIS ONE, WHICH IS TO FUND?
18	MR. TORRES: SO MOVED.
19	MR. SHEEHY: MOVED BY SENATOR TORRES. DO
20	I HAVE A SECOND?
21	DR. JUELSGAARD: SECOND.
22	MR. SHEEHY: DO WE HAVE DISCUSSION OF
23	THIS? DO YOU WANT ADDITIONAL INFORMATION? I THINK
24	DR. SAMBRANO GAVE A BIT OF A SYNOPSIS; BUT IF MORE
25	INFORMATION IS NEEDED, OTHERWISE WE CAN GO TO PUBLIC
	116
	110

1	COMMENT AND A VOTE. IS THERE ANY PUBLIC COMMENT ON
2	THIS? YES, PLEASE.
3	DR. KOEHLER: I'M DR. CARLA KOEHLER. I'M
4	THE PI ON 7678, THE ONE THAT HAS THE LITTLE WHITE
5	BAR UP THERE.
6	THIS IS A GRANT THAT HAS BORNE OUT OF A
7	CIRM BASIC BIOLOGY AND A SEED GRANT, AND WE WOULD
8	ANOTHER POINT OF COMMENT IN THE REVIEWERS WAS THAT
9	WE WERE ONLY GOING TO WORK ON NEURONAL CELLS. BUT
10	AS WE STATED IN SEVERAL PLACES IN THE GRANT, WE ARE
11	GOING TO WORK ON MORE THAN NEURONAL CELLS. WE THINK
12	THIS IS A REALLY IMPORTANT AREA TO DEVELOP. THERE'S
13	A LARGE NUMBER OF STEM CELLS THAT FAIL TO
14	DIFFERENTIATE IN STEM CELL THERAPIES WHEN WE IMPLANT
15	THEM INTO PATIENTS. IF THEY HAVE NOT COMPLETELY
16	DIFFERENTIATED, THEY CAN TURN INTO TERATOMAS. AS
17	GIL SAID, THERE'S REALLY NOTHING THAT'S BEING DONE
18	IN THIS SCOPE, AND SO WE THINK THAT THIS IS REALLY
19	AN IMPORTANT AREA TO DEVELOP AND SOMETHING THAT WE
20	WOULD LIKE TO CONTINUE WORKING ON.
21	MR. SHEEHY: THANK YOU. MS. BONNEVILLE,
22	COULD WE CALL THE ROLL, PLEASE.
23	DR. DULIEGE: CAN WE REPEAT THE MOTION?
24	MR. SHEEHY: THE MOTION IS TO MOVE THIS
25	INTO TIER I.
	117

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1		MR. HARRISON: IT'S APPLICATION RT3-07678.
2		MS. BONNEVILLE: ANNE-MARIE DULIEGE.
3		DR. DULIEGE: AYE.
4		MS. BONNEVILLE: DAVID HIGGINS.
5		DR. HIGGINS: YES.
6		MS. BONNEVILLE: STEPHEN JUELSGAARD.
7		MR. JUELSGAARD: YES.
8		MS. BONNEVILLE: KATHY LAPORTE.
9		DR. LAPORTE: YES.
10		MS. BONNEVILLE: LAUREN MILLER.
11		MS. MILLER: YES.
12		MS. BONNEVILLE: FRANCISCO PRIETO. ROBERT
13	QUINT.	
14		DR. QUINT: YES.
15		MS. BONNEVILLE: AL ROWLETT.
16		MR. ROWLETT: YES.
17		MS. BONNEVILLE: JEFF SHEEHY.
18		MR. SHEEHY: YES.
19		MS. BONNEVILLE: JONATHAN THOMAS.
20		CHAIRMAN THOMAS: YES.
21		MS. BONNEVILLE: ART TORRES.
22		MR. TORRES: AYE.
23		MS. BONNEVILLE: DIANE WINOKUR.
24		MS. WINOKUR: YES.
25		MR. HARRISON: THE MOTION CARRIES WITH A
		118
		110

1	11 YES VOTES.
2	MR. SHEEHY: AND THEN COULD I HAND THE
3	CONSIDERATION OF THE NEXT APPLICATION OVER TO
4	CHAIRMAN THOMAS.
5	CHAIRMAN THOMAS: CERTAINLY, MR. SHEEHY.
6	WE'RE NOW CONSIDERING GRANT RT3-07836; IS THAT
7	CORRECT, DR. SAMBRANO?
8	DR. SAMBRANO: CORRECT.
9	CHAIRMAN THOMAS: AND HAVING HEARD THE
10	RECOMMENDATION, UPON FURTHER WORK, THAT WE CONTINUE
11	TO KEEP THIS IN TIER III AND NOT MOVE IT UP FOR
12	FUNDING, THAT IS THE RECOMMENDATION OF THE TEAM. DO
13	I HEAR A MOTION TO MOVE IT FROM TIER III UP TO NO.
14	i? HEARING NONE, MR. HARRISON, DO WE REQUIRE PUBLIC
15	COMMENT IF WE HAVE NO MOTION?
16	MR. HARRISON: NO.
17	CHAIRMAN THOMAS: SO THAT WILL COMPLETE
18	CONSIDERATION OF THAT PARTICULAR GRANT.
19	MR. HARRISON: CHAIRMAN THOMAS, WE
20	ACTUALLY NEED TO TAKE ANOTHER MOTION NOT TO FUND
21	THAT APPLICATION.
22	CHAIRMAN THOMAS: DO I HEAR A MOTION
23	MR. TORRES: SO MOVED.
24	CHAIRMAN THOMAS: MOVED BY SENATOR TORRES.
25	MR. JUELSGAARD: SECOND.
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CHAIRMAN THOMAS: ANY DISCUSSION BY
MEMBERS OF THE BOARD? MR. HARRISON, DOES THAT
MOTION REQUIRE PUBLIC COMMENT?
MR. HARRISON: YES.
CHAIRMAN THOMAS: IS THERE ANY PUBLIC
COMMENT ON THAT MOTION? SEEING AND HEARING NONE,
MARIA, WILL YOU PLEASE CALL THE ROLL.
MS. BONNEVILLE: ANNE-MARIE DULIEGE.
DR. DULIEGE: AYE.
MS. BONNEVILLE: DAVID HIGGINS.
DR. HIGGINS: YES.
MS. BONNEVILLE: STEPHEN JUELSGAARD.
MR. JUELSGAARD: YES.
MS. BONNEVILLE: KATHY LAPORTE.
DR. LAPORTE: YES.
MS. BONNEVILLE: LAUREN MILLER.
MS. MILLER: YES.
MS. BONNEVILLE: FRANCISCO PRIETO.
DR. PRIETO: YES.
MS. BONNEVILLE: ROBERT QUINT.
DR. QUINT: YES.
MS. BONNEVILLE: AL ROWLETT.
MR. ROWLETT: YES.
MS. BONNEVILLE: JONATHAN THOMAS.
CHAIRMAN THOMAS: YES.
120

MS. BONNEVILLE: ART TORRES.
MR. TORRES: AYE.
MS. BONNEVILLE: DIANE WINOKUR.
MS. WINOKUR: YES.
CHAIRMAN THOMAS: DO WE NEED ANY FURTHER
MOTIONS ON THIS MATTER? SO THIS WILL BE THE END OF
THAT TOPIC THEN; IS THAT CORRECT?
MR. HARRISON: CORRECT. AND THAT MOTION
PASSED WITH 11 YES VOTES.
CHAIRMAN THOMAS: THANK YOU VERY MUCH,
MR. SHEEHY. THANK YOU, DR. SAMBRANO.
ON TO ITEM NO. 9, CONSIDERATION OF
ADOPTION OF THE INTERIM GRANTS ADMINISTRATION POLICY
FOR THE CLINICAL STAGE PROGRAMS. GABE THOMPSON WILL
PRESENT.
MR. SHEEHY: I JUST WANTED TO MAKE THE
COMMENT THAT THE NEXT TWO ITEMS, I BELIEVE, WERE
DISCUSSED IN THE SCIENCE SUBCOMMITTEE. AND THOSE
WERE MOVED THROUGH THE SCIENCE SUBCOMMITTEE
UNANIMOUSLY.
MR. THOMPSON: CHAIRMAN THOMAS, MEMBERS OF
THE BOARD, MEMBERS OF THE PUBLIC, AND CIRM TEAM, I'M
GABRIEL THOMPSON, DIRECTOR OF GRANTS MANAGEMENT AT
CIRM AND OFFERING FOR YOUR CONSIDERATION PROPOSED
INTERIM GRANTS ADMINISTRATION POLICY REGULATIONS FOR
121

1	THE LATE STAGE PROJECTS. THIS WOULD BE THE GRANTS
2	ADMINISTRATION POLICY THAT WOULD APPLY TO NEW AWARDS
3	UNDER PROGRAM ANNOUNCEMENTS 15-01, 15-02, AND 15-03.
4	THE PROPOSED INTERIM GRANTS ADMINISTRATION
5	POLICY FOLLOWS THE SAME BASIC TEMPLATE AS THE
6	EXISTING POLICY. THAT APPLIES TO CURRENT ACTIVE AND
7	FORMER AWARDS. THE POLICY IS CHRONOLOGICALLY
8	ORGANIZED TO GOVERN THE FULL GRANT-MAKING PROCESS,
9	INCLUDING APPLICATION IN THE REVIEW PROCESS, THE
10	PRE-AWARD AND AWARD REQUIREMENTS, AND RULES
11	GOVERNING PAYMENT AND USE OF FUNDS WHEN CIRM GRANTS
12	ARE FUNDED.
13	THE CIRM TEAM REVIEWED EACH ASPECT OF THE
14	EXISTING POLICY THROUGH THE LENS OF THE CLINICAL
15	STAGE PROGRAM AND ITS REQUIREMENTS. AND THE RESULT
16	IS A POLICY DESIGNED TO ATTRACT MORE HIGH QUALITY
17	APPLICATIONS, REDUCE CYCLE TIME FROM APPLICATION TO
18	PROJECT START, ACCELERATE PROGRESSION OF FUNDED
19	PROJECTS, AND PROVIDE A MORE EFFICIENT
20	ADMINISTRATION OF THE PROJECTS.
21	THE GRANTS ADMINISTRATION POLICY WAS
22	PREVIEWED AT THE LAST BOARD MEETING, AND WE WENT
23	OVER THOSE MAJOR AREAS THAT WE ADDRESSED IN THE
24	POLICY AND ASKED THE BOARD FOR ADDITIONAL TIME TO
25	PROVIDE A LITTLE BIT MORE CLARIFICATION IN THE
	122

1	POLICY ITSELF.
2	THOSE ITEMS WE'VE CLARIFIED SINCE THE LAST
3	BOARD MEETING INCLUDES LANGUAGE REGARDING THE NEW
4	SCORING SYSTEM. SO WE EXPLAINED THE SCORING SYSTEM
5	AS YOU WILL HEAR IN THE NEXT AGENDA ITEM. WE'VE
6	CLARIFIED THE REPORTING REQUIREMENTS FOR AWARDS
7	FUNDED UNDER THESE NEW PROGRAM ANNOUNCEMENTS. SO
8	WHEN ARE REPORTS REQUIRED? UPON ACHIEVING
9	OPERATIONAL MILESTONES OR HITTING SUSPENSION EVENTS.
10	AND WHAT ARE THE CONTENT OF THOSE REPORTS?
11	WE CLARIFY WHEN PROTOCOL APPROVALS ARE
12	REQUIRED. THAT WOULD BE IRB, IACUC, AND SCRO
13	PROTOCOL APPROVALS. SO WE CLARIFY WHEN WE WANT TO
14	SEE THOSE DOCUMENTS AND WHEN SELF-CERTIFICATION IS
15	SUFFICIENT.
16	AND THEN WE INCLUDE LANGUAGE ABOUT THE
17	COMMUNICATIONS PLAN UNDER THESE AWARDS. AND WHAT WE
18	MEAN BY COMMUNICATIONS PLAN IS WHEN IS THE AWARDEE
19	REQUIRED TO COMMUNICATE TO CIRM THE STATUS OF A
20	PROJECT, FOR INSTANCE, IF A TRIAL GOES ON A CLINICAL
21	HOLD. SO THEY WOULD HAVE TO REPORT THAT TO CIRM.
22	IF THE AWARDEE WAS SUBJECT TO AN FDA AUDIT, THOSE
23	KINDS OF THINGS, WE SPELL OUT IN MORE DETAIL WHEN
24	THEY'RE REQUIRED TO COMMUNICATE TO CIRM.
25	SO WE ARE ASKING THE BOARD TO FORMALLY
	123

1	APPROVE THESE INTERIM REGULATIONS WHICH WOULD THEN
2	START THE PROCESS BY WHICH WE WOULD GO TO THE
3	STATE'S OFFICE OF ADMINISTRATIVE LAW TO SEEK FORMAL
4	APPROVAL. AND WE WOULD HAVE 270 DAYS TO DO SO. AND
5	WE WOULD IN THAT PERIOD BE ABLE TO CONTINUE TO TAKE
6	INPUT FROM ALL OF OUR STAKEHOLDERS, INCLUDING THE
7	PUBLIC, ON THE REGULATIONS AND CONTINUE TO REFINE
8	THOSE THROUGHOUT THIS PROCESS UNTIL WE GET FORMAL
9	APPROVAL.
10	SO THAT SAID, WE'RE ASKING THE BOARD FOR
11	APPROVAL OF THESE INTERIM REGULATIONS.
12	CHAIRMAN THOMAS: DO I HEAR A MOTION TO
13	APPROVE THESE INTERIM REGULATIONS?
14	MR. SHEEHY: SO MOVED.
15	MR. TORRES: SECOND.
16	CHAIRMAN THOMAS: MOVED BY MR. SHEEHY,
17	SECONDED BY SENATOR TORRES. DISCUSSION BY MEMBERS
18	OF THE BOARD? AS MR. SHEEHY SAID, WE HAD
19	CONSIDERABLE DISCUSSION ON ALL THESE POINTS AT THE
20	SCIENCE SUBCOMMITTEE, AND IT WAS PASSED THROUGH
21	UNANIMOUSLY. SO ALL OF THESE THINGS HAVE BEEN
22	CAREFULLY LOOKED AT AND VETTED.
23	ANY COMMENTS FROM MEMBERS OF THE PUBLIC?
24	MR. HARRISON, DOES THIS REQUIRE A ROLL CALL?
25	MR. HARRISON: VOICE VOTE IS FINE EXCEPT
	124

FOR THOSE ON THE PHONE.
CHAIRMAN THOMAS: SO WE'RE GOING TO DO
FIRST VOICE VOTE IN THE ROOM AND THEN THOSE ON THE
PHONE. ALL THOSE IN FAVOR IN THE ROOM PLEASE SAY
AYE. OPPOSED? ANY ABSTENTIONS? MARIA, WILL YOU
PLEASE CALL THOSE ON THE PHONE?
MS. BONNEVILLE: KATHY LAPORTE.
DR. LAPORTE: AYE.
MS. BONNEVILLE: ELIZABETH FINI.
DR. FINI: YES.
MS. BONNEVILLE: KRISTINA VUORI.
DR. VUORI: YES.
CHAIRMAN THOMAS: MR. HARRISON, I PRESUME
THAT PASSED WITH FLYING COLORS.
MR. HARRISON: MOTION CARRIES.
CHAIRMAN THOMAS: THANK YOU. WE'RE GOING
TO GO ON NOW TO ITEM NO. 10, CONSIDERATION OF
ADOPTION OF AMENDMENTS TO GRANTS WORKING GROUP
BYLAWS. WHEN WE GET THROUGH THAT, WE WILL THEN
DISCUSS LUNCH PLANS. MR. HARRISON.
MR. TORRES: ARE WE GOING TO GET TO ITEM
12 BY 1 0'CLOCK?
CHAIRMAN THOMAS: ABSOLUTELY. WE'RE
ACTUALLY GOING TO WORK THROUGH LUNCH, SO WE WILL
DEFINITELY GET TO ITEM 12 BEFORE 1 O'CLOCK.
125

125

1	MR. HARRISON: THANK YOU, CHAIRMAN THOMAS.
2	AS JEFF SHEEHY EXPLAINED, THE SCIENCE SUBCOMMITTEE
3	MET EARLIER THIS WEEK AND CONSIDERED PROPOSED
4	AMENDMENTS TO THE GWG BYLAWS. AND WE WANT TO
5	BRIEFLY TAKE YOU THROUGH THE SIGNIFICANT CHANGES
6	THAT WE PROPOSE TO MAKE.
7	OUR GOAL IN REVIEWING THE BYLAWS WAS,
8	FIRST, TO UPDATE THEM. WE HAVE NOT AMENDED THE
9	BYLAWS SINCE THE BEGINNING OF 2013, AND THERE HAVE
10	BEEN CHANGES TO PRACTICES AND POLICIES SINCE THEN.
11	SO WE HAVE INCLUDED AMENDMENTS TO BRING THE BYLAWS
12	UP TO DATE AND CONSISTENT WITH OUR CURRENT PRACTICES
13	AND POLICIES.
14	WE'VE ALSO PROPOSED AMENDMENTS TO CONFORM
15	THE BYLAWS TO THE CIRM 2.0 PROCESS, INCLUDING THE
16	SUBJECT OF SCORING, WHICH THE BOARD TALKED ABOUT AT
17	LENGTH THIS MORNING.
18	SO LET ME BRIEFLY DESCRIBE SOME OF THE
19	SIGNIFICANT CHANGES. FIRST, WE HAVE CLARIFIED THE
20	GRANTS WORKING GROUP'S ROLE IN OVERSEEING THE
21	PROGRESS OF FUNDED PROJECTS. UNDER PROP 71 THE
22	GRANTS WORKING GROUP HAS A ROLE TO PLAY IN THE
23	OVERSIGHT OF FUNDED AWARDS. WE INTEND, AS PART OF
24	CIRM 2.0, TO USE THE GRANTS WORKING GROUP IN ITS
25	OVERSIGHT CAPACITY ON A MORE REGULAR BASIS. AS YOU
	126

1	KNOW, AS PART OF CIRM 2.0, A CLINICAL ADVISORY PANEL
2	WILL BE ESTABLISHED FOR EACH CLINICAL STAGE PROJECT.
3	AND THESE CAP'S WILL REPORT ON AN ANNUAL BASIS TO
4	THE GRANTS WORKING GROUP.
5	THE SECOND SIGNIFICANT CHANGE THAT WE
6	WOULD PROPOSE IS TO MORE ACTIVELY ENGAGE THE PATIENT
7	ADVOCATE MEMBERS OF THE GRANTS WORKING GROUP IN THE
8	REVIEW PROCESS. CURRENTLY THE PATIENT ADVOCATE
9	MEMBERS OF THE GRANTS WORKING GROUP PARTICIPATE IN
10	THE GRANTS WORKING GROUP REVIEW, BUT DO NOT SCORE
11	APPLICATIONS AND ARE NOT ASSIGNED AS REVIEWERS. WE
12	PROPOSE TO ENGAGE THE PATIENT ADVOCATE MEMBERS MORE
13	ACTIVELY BY INVITING A PATIENT ADVOCATE MEMBER OF
14	THE GWG TO SERVE AS A REVIEWER ON EACH APPLICATION.
15	ALTHOUGH THE PATIENT ADVOCATE MEMBER WOULDN'T SCORE
16	THE APPLICATIONS, ONLY THE SCIENTIFIC REVIEWERS
17	ASSIGN A SCORE, THE PATIENT ADVOCATE MEMBER WOULD
18	HAVE AN OPPORTUNITY TO PROVIDE HIS OR HER VIEWS ON
19	THE MERITS OF THE APPLICATION DURING THE REVIEW
20	PROCESS AND BEFORE THE SCIENTISTS SCORE THE
21	APPLICATION.
22	THE LAST SIGNIFICANT CHANGE IS A PROPOSED
23	MODIFICATION TO THE SCORING SYSTEM FOR CLINICAL
24	STAGE APPLICATIONS; THAT IS, THOSE APPLICATIONS
25	SUBMITTED IN RESPONSE TO PROGRAM ANNOUNCEMENTS
	127

1	15-01, 15-02, AND 15-03. THE GOAL OF THIS SYSTEM IS
2	REALLY TO ADDRESS SOME OF THE CONCERNS THAT WERE
3	RAISED BY MEMBERS EARLIER TODAY. AND THAT IS TO
4	REALLY OBTAIN CLEAR DIRECTION ABOUT WHETHER TO FUND
5	A PROPOSAL, SEND IT BACK TO THE APPLICANT FOR
6	REFINEMENT AND RESUBMISSION, OR RECOMMEND AGAINST
7	FUNDING AND AGAINST RESUBMITTING THE SAME PROJECT IN
8	THE SAME FORM.
9	SO PURSUANT TO THE SYSTEM, RATHER THAN
10	ASKING THE SCIENTIFIC MEMBERS TO USE A RANGE FROM 1
11	TO 100 TO ASSIGN A SCORE TO THE APPLICATION, WE'D
12	ASK THEM, INSTEAD, TO ASSIGN A SCORE OF ONE, TWO, OR
13	THREE. A SCORE OF ONE WOULD SIGNIFY THAT THE
14	APPLICATION HAS EXCEPTIONAL MERIT AND SHOULD BE
15	FUNDED. A SCORE OF TWO WOULD MEAN THAT THE
16	APPLICATION NEEDS IMPROVEMENT AND DOESN'T WARRANT
17	FUNDING AT THIS TIME, BUT COULD BE RESUBMITTED TO
18	ADDRESS AREAS OF IMPROVEMENT NOTED BY THE GWG. AND
19	FINALLY, A SCORE OF THREE WOULD SIGNIFY THAT THE
20	APPLICATION IS SUFFICIENTLY FLAWED THAT IT DOESN'T
21	WARRANT FUNDING AND SHOULDN'T BE RESUBMITTED FOR
22	REVIEW IN THE SAME FORM.
23	AS PART OF THIS PROCESS, THE CIRM TEAM
24	WOULD TALLY THE NUMBER OF SCIENTIFIC MEMBERS WHO
25	ASSIGNED A SCORE OF ONE, TWO, AND THREE

1	RESPECTIVELY, AND THEN WOULD PRESENT THAT
2	INFORMATION FOR EACH APPLICATION TO THE ENTIRE GWG.
3	IF A PLURALITY OF MEMBERS ASSIGNED A SCORE OF ONE OR
4	TWO, THEN THAT SCORE WOULD CONSTITUTE THE
5	RECOMMENDATION OF THE GWG. SO, FOR EXAMPLE, IF
6	EIGHT SCIENTISTS ASSIGN A SCORE OF ONE, SIX ASSIGNED
7	A SCORE OF TWO, AND ONE ASSIGNED A SCORE OF THREE,
8	THEN THAT APPLICATION WOULD BE PLACED IN TIER I AND
9	RECOMMENDED TO THE APPLICATION REVIEW SUBCOMMITTEE
10	FOR FUNDING.
11	WITH RESPECT TO TIER III, OR A SCORE OF
12	THREE, SINCE THE EFFECT OF ASSIGNING AN APPLICATION
13	TO TIER III WOULD BE THAT THE APPLICANT COULD NOT
14	RESUBMIT THE SAME PROJECT, WE WANTED TO HAVE A
15	SLIGHTLY HIGHER THRESHOLD THERE. SO RATHER THAN A
16	PLURALITY, WE'D REQUIRE A MAJORITY. SO EIGHT OR
17	MORE MEMBERS WOULD HAVE TO ASSIGN A SCORE OF THREE
18	BEFORE THAT APPLICATION WOULD BE ASSIGNED TO TIER
19	III, NOT RECOMMENDED FOR FUNDING.
20	IF THERE'S NO PLURALITY AND THERE'S A
21	NUMERICAL TIE BETWEEN TWO OR MORE SCORES, THEN ANY
22	MEMBER OF THE GWG COULD MAKE A MOTION TO BREAK THAT
23	TIE BY ASSIGNING THE APPLICATION TO TIER I, II OR
24	III. SO AS THE EXAMPLE SUGGESTS, IF YOU HAD SEVEN
25	VOTES IN TIER I AND SEVEN IN TIER II, A MEMBER OF
	100

THE GWG, FOR EXAMPLE, COULD MOVE THAT THE
APPLICATION BE ASSIGNED TO TIER II. AND IF THE
MAJORITY OF MEMBERS APPROVED THAT MOTION, THE
APPLICATION WOULD BE ASSIGNED TO TIER II. IF IT
FAILED, THEN THERE WOULD BE AN OPPORTUNITY FOR A NEW
MOTION.
WITH RESPECT TO TIER III, IF A PLURALITY
OF SCIENTIFIC MEMBERS ASSIGNED AN APPLICATION A
SCORE OF THREE, BUT LESS THAN A MAJORITY, THEN ANY
MEMBER OF THE GWG COULD MAKE A MOTION TO EITHER
ASSIGN THAT APPLICATION TO TIER II OR TIER III.
AGAIN, IF A MOTION WAS MADE TO ASSIGN IT TO TIER II
AND A MAJORITY OF THE MEMBERS VOTED IN FAVOR OF
THAT, THAT WOULD REPRESENT THE RECOMMENDATION OF THE
GRANTS WORKING GROUP.
WITH RESPECT TO WHAT'S PRESENTED TO THE
APPLICATION REVIEW SUBCOMMITTEE, THE CIRM TEAM WOULD
PRESENT THE GWG RECOMMENDATIONS TO THE APPLICATION
REVIEW SUBCOMMITTEE ALONG WITH THE DISTRIBUTION OF
SCORES AMONG THE THREE TIERS.
WE THINK THIS SYSTEM WILL PRODUCE MORE
CLEAR DIRECTION AND GUIDANCE FROM THE GWG AND IS
ALIGNED WITH THE NEW DIRECTION UNDER CIRM 2.0. SO
WE WOULD ASK FOR YOUR APPROVAL. I'D BE HAPPY TO
ANSWER ANY QUESTIONS.
130

1	CHAIRMAN THOMAS: MR. JUELSGAARD.
2	DR. JUELSGAARD: MR. HARRISON, I'D LIKE TO
3	FOCUS ON SLIDES 5 AND 7. I'D LIKE TO FOCUS ON THE
4	USE OF THE WORD "PLURALITY" BECAUSE I'M NOT QUITE
5	SURE I UNDERSTAND. SO SLIDE 5 SAYS, "IF A PLURALITY
6	OF MEMBERS HAS ASSIGNED A SCORE OF ONE OR TWO, THEN
7	THAT SCORE CONSTITUTES THE RECOMMENDATION OF THE
8	GWG." AND THEN YOU SHOW IT AS BASICALLY GOING TO
9	TIER I.
10	MR. HARRISON: CORRECT.
11	DR. JUELSGAARD: SO WHAT DO YOU MEAN BY
12	PLURALITY IN THAT CASE?
13	MR. HARRISON: PLURALITY MEANS THAT THE
14	SCORE THAT HAS RECEIVED THE GREATEST NUMBER OF
15	VOTES.
16	DR. JUELSGAARD: SO DO YOU REALLY MEAN IF
17	A PLURALITY OF MEMBERS HAS ASSIGNED A SCORE OF ONE,
18	AND THEN LEAVE OUT THE "OR TWO," THEN THAT'S THE
19	RESULT?
20	MR. HARRISON: CORRECT.
21	DR. JUELSGAARD: SO THE "OR TWO" ACTUALLY
22	SHOULDN'T BE IN THERE?
23	DR. MILLS: HE MEANS THE PLURALITY RULE
24	APPLIES TO TIER I OR TIER II. THE PLURALITY DOES
25	NOT APPLY TO TIER III. YOU NEED A MAJORITY TO BE
	121
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1	ASSIGNED TO TIER III. A NON-MAJORITY HIS EXAMPLE
2	UP HERE HAPPENS TO ALSO BE A MAJORITY, BUT YOU COULD
3	HAVE A SITUATION WHERE YOU HAD A PLURALITY TIER II
4	OR A PLURALITY TIER I THAT WASN'T A MAJORITY.
5	DR. JUELSGAARD: SO I'M STILL CONFUSED.
6	DR. LEVIN: 6-4-4 IS TIER I, 4-6-4 IS TIER
7	II, 4-4-6 IS NOT TIER II.
8	DR. JUELSGAARD: GO TO SLIDE 7. SO,
9	AGAIN, YOU SAY IF THERE'S A PLURALITY OF IN THIS
10	CASE THERE ISN'T A PLURALITY EVEN THOUGH WE HAVE
11	MR. HARRISON: THIS IS THE TIE BREAKER
12	SITUATION.
13	DR. JUELSGAARD: SO FIRST OF ALL, DO YOU
14	HAVE MORE VOTES IN ONE AND TWO, RIGHT?
15	MR. HARRISON: CORRECT.
16	DR. JUELSGAARD: THAT'S STEP 1. AND THEN
17	STEP 2 IS IF YOU HAVE MORE VOTES IN ONE AND TWO,
18	THEN HOW DO YOU DECIDE BETWEEN A ONE OR A TWO IN
19	THIS CASE TO BE A TIE BREAKER?
20	MR. HARRISON: CORRECT. IF THERE IS A TIE
21	VOTE BETWEEN ONE AND TWO, THEN YOU'D HAVE A MOTION
22	TO BREAK THE TIE. OTHERWISE, IF YOU HAD A PLURALITY
23	OF VOTES IN TIER II, THEN THAT WOULD BE THE
24	RECOMMENDATION.
25	DR. JUELSGAARD: SO IN THIS CASE IF SCORE
	132

1	ONE WAS SIX VOTES, SCORE TWO WAS EIGHT, SCORE THREE
2	WAS ONE, THEN IT WOULD BE AUTOMATICALLY IN TIER II.
3	MR. HARRISON: IN TIER II. EXACTLY.
4	DR. MILLS: BUT SIMILARLY, AND I THINK THE
5	PURPOSE FOR USING PLURALITY IS THAT IF IT WAS 6-7-2,
6	YOU WOULDN'T HAVE A MAJORITY, BUT YOU WOULD HAVE A
7	PLURALITY, AND SO TIER II WOULD GO.
8	DR. LEVIN: CAN I GET A CLARIFICATION ON
9	SOMETHING WE WERE DISCUSSING EARLIER? WHEN IN THE
10	PROCESS ARE THESE SCORES ASSIGNED? AND MORE
11	IMPORTANTLY, IS THERE AN OPPORTUNITY AFTER
12	DISCUSSION AMONG GRANT WORKING GROUP MEMBERS TO
13	ADJUST THEM?
14	DR. MILLS: ABSOLUTELY. THE SCORES ARE
15	NOT FINALIZED UNTIL ALL OF THE DISCUSSION TAKES
16	PLACE AND ALL OF THE IDEAS ARE VETTED. I'LL JUST
17	SAY MY EXPERIENCE AS A GWG MEMBER, ONE OF THE THINGS
18	WHEN I WAS A MEMBER OF THAT BODY, ONE OF THE THINGS
19	I FOUND MOST IMPRESSIVE WAS HOW THE MEMBERS OF THE
20	GROUP WOULD LISTEN TO EACH OTHER AND NOT JUST DRIVE
21	AN OPINION AND SAY, WOW, I DIDN'T REALIZE THAT AND
22	ADJUST THEIR SCORES ACCORDINGLY. I DON'T KNOW IF
23	ANY OTHER MEMBERS THAT SIT ON THE GWG WANT TO
24	COMMENT TO THAT, BUT IT'S AN IMPRESSIVE
25	MR. ROWLETT: I WOULD DITTO THAT. IN
	133
	±33

1	FACT, I THINK THIS IS A TREMENDOUS IMPROVEMENT FROM
2	THE PATIENT ADVOCATE PERSPECTIVE BECAUSE IT DOES
3	CHARGE US WITH REALLY BEING INFLUENTIAL IN THAT
4	PROCESS BECAUSE THE DISCUSSION REQUIRES US TO,
5	BECAUSE EVEN THOUGH WE'RE NOT SCORING, WE ARE
6	PARTICIPATING MORE ACTIVELY IN THE DISCUSSION. AND
7	SO I THINK THIS IS AN ADVANCEMENT FORWARD. I FOUND
8	MYSELF TRYING TO INFLUENCE THE PEOPLE IN MY SEATING
9	AREA RELATED TO THEIR SCORING INFORMALLY. PERHAPS I
10	SHOULDN'T SAY THAT, BUT THAT WAS MY MOTIVATION
11	BECAUSE, AS A PATIENT ADVOCATE, YOU MIGHT NOT ALWAYS
12	BE ABLE TO SPEAK TO ALL OF THE CLINICAL
13	APPLICATIONS, BUT YOU CERTAINLY UNDERSTAND THE
14	POTENTIAL IMPLICATIONS FOR ALL THE CITIZENS. SO
15	YOU'RE THINKING ABOUT THINGS LIKE THE DISTRIBUTION
16	SYSTEM AND HOW INCLUSIVE IT'S GOING TO BE, AND SO
17	YOU'RE TRYING TO INFLUENCE THAT. AND THAT
18	CONVERSATION THE PATIENT ADVOCATES, FROM THE
19	PERSPECTIVE OF ONE, WILL INJECT INTO THIS PROCESS.
20	SO DITTO. I THINK THIS IS A TREMENDOUS
21	ADVANCEMENT FORWARD. NOT THAT THE PROCESS WAS
22	FLAWED COMPLETELY, BUT YOU KNOW WHERE I'M GOING.
23	DR. HIGGINS: THIS IS A DETAIL, BUT I
24	WOULD JUST LIKE THE BOARD TO KNOW AND APPRECIATE THE
25	AMOUNT OF WORK THAT WENT INTO CREATING THIS SYSTEM
	124

1	BY THE STAFF. IF YOU'VE SEEN THE NUTS AND BOLTS OF
2	HOW THIS WORKS, THIS IS AN INCREDIBLE SYSTEM,
3	INCREDIBLY WELL AUTOMATED, WELL DESIGNED, WELL
4	THOUGHT OUT. AND I JUST WANT TO MAKE SURE THAT
5	EVERYBODY APPRECIATES THAT.
6	MR. TORRES: BRAVO.
7	(APPLAUSE.)
8	CHAIRMAN THOMAS: WELL SAID, DR. HIGGINS.
9	OTHER COMMENTS BY MEMBERS OF THE BOARD? DO I HEAR A
10	MOTION TO APPROVE?
11	MR. ROWLETT: I'LL MOVE.
12	CHAIRMAN THOMAS: SO MOVED BY MR. ROWLETT,
13	VERY EMPHATICALLY, I MIGHT ADD, AND NOT INDIRECTLY.
14	MR. TORRES: SECOND.
15	CHAIRMAN THOMAS: ANY DISCUSSION, FURTHER
16	DISCUSSION BY MEMBERS OF THE BOARD?
17	DR. LAPORTE: I JUST HAVE A QUESTION.
18	THERE WAS A LOT OF GOOD DISCUSSION EARLIER TODAY
19	ABOUT THE PROCESS PROBLEMS AND ISSUES, THE PRIOR
20	SCORING AT THE GWG. IS THE EXPECTATION HERE THAT
21	WITH THIS NEW SCORING SYSTEM, WE SHOULD SEE LESS OF
22	THAT? WONDERING HOW TO THINK ABOUT THIS GOING
23	FORWARD.
24	CHAIRMAN THOMAS: DR. MILLS WILL ADDRESS
25	THAT QUESTION.
	125

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1	DR. MILLS: I THINK WITH REGARDS TO HOW
2	THE SCORING SYSTEM IS ONE PIECE TO IT, AND THEN
3	WHAT'S DONE WITH THOSE SCORES IS AS IMPORTANT AS THE
4	SCORING SYSTEM ITSELF. AND SO BASICALLY THE CHOICE
5	THAT WAS GIVEN BEFORE WAS RECOMMENDED FOR FUNDING OR
6	DON'T RECOMMEND IT FOR FUNDING. IT WAS BINARY. IT
7	WAS UP OR DOWN.
8	WHAT WE'VE DONE HERE IS WE'VE ADDED THIS
9	TIER II OR SCORE TWO, AND TIER II IS IT'S GOOD, BUT
10	THERE ARE SOME THINGS THAT CAN BE FIXED. SO, AGAIN,
11	THE GOAL ISN'T TO SEE HOW MANY THINGS WE CAN GET
12	THROUGH OR PUT TO THE BOARD VERY DIFFICULT CHOICES
13	ABOUT THINGS THAT ARE BORDERLINE. MY REAL GOAL IS
14	TO PUT TO THE BOARD THINGS THAT WE'RE ALL
15	ENTHUSIASTICALLY SUPPORTIVE OF BECAUSE WE'VE MADE
16	THEM AS GOOD AS WE POSSIBLY CAN MAKE THEM BEFORE
17	THEY COME BEFORE YOU FOR A DECISION.
18	THAT IS NOT TO SAY THERE ARE NOT SOME
19	THINGS WHERE WE JUST CAN'T MAKE IT ANY BETTER AND
20	IT'S STILL BORDERLINE, AND THOSE ARE THE THINGS THAT
21	YOU'LL BE FACED TO BREAK THE TIE ON. BUT THE
22	SCORING SYSTEM ITSELF IS ONE PIECE OF IT.
23	HOW THE SCORING SYSTEM IS USED IS THE
24	OTHER PART. SO I REALLY DO THINK, FOR THE MOST
25	PART, THE APPLICATIONS YOU'LL BE SEEING WILL BE MADE
	136

1	AS WELL AS THEY POSSIBLY CAN BE MADE BEFORE THEY
2	COME BEFORE YOU. AND I THINK THAT WILL CLEAN UP A
3	LOT OF THIS NOISE THAT WE HAD EARLIER.
4	CHAIRMAN THOMAS: TO SUMMARIZE, YES.
5	ANY FURTHER DISCUSSION OR QUESTIONS FROM
6	MEMBERS OF THE BOARD? ANY PUBLIC COMMENT? HEARING
7	NONE, MR. HARRISON, SIMILAR VOICE VOTE PROTOCOL
8	HERE? FIRST IN THE ROOM, THEN ON THE PHONE. ALL
9	THOSE IN FAVOR PLEASE SAY AYE. OPPOSED? ANY
10	ABSTENTIONS? MARIA, PLEASE CALL THE ROLL OF THOSE
11	ON THE PHONE.
12	MS. BONNEVILLE: KATHY LAPORTE.
13	DR. LAPORTE: ABSOLUTELY.
14	MS. BONNEVILLE: DR. FINI.
15	DR. FINI: YES.
16	MS. BONNEVILLE: KRISTINA VUORI.
17	DR. VUORI: OF COURSE.
18	CHAIRMAN THOMAS: EXCELLENT NEW
19	TERMINOLOGY USED BY MEMBERS ON THE PHONE. MR.
20	HARRISON, I THINK THAT ONE PASSES WITH FLYING COLORS
21	AS WELL.
22	SO WE ARE NOW GOING TO DO A WORKING LUNCH
23	BECAUSE WE WANT TO MAKE SURE WE MAINTAIN QUORUM FOR
24	ALL ITEMS THAT WE NEED TO GET THROUGH BEFORE WE LOSE
25	ANYBODY. SO, MARIA, THE FOOD IS IN THE BACK OF THE
	127
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1	ROOM. SO IF EVERYBODY COULD PLEASE GET THEIR LUNCH
2	AND PROMPTLY RETURN TO YOUR SEATS AND WE WILL
3	CONTINUE. THOSE ON THE PHONE, I HOPE YOU HAVE
4	SOMETHING TASTY TO EAT AS WELL.
5	(A RECESS WAS TAKEN.)
6	CHAIRMAN THOMAS: OKAY. WE ARE BACK LIVE
7	IN THE SONOMA ROOM. WE ARE NOW GOING TO GO TO
8	ITEM 11 IS POSTPONED ITEM 12, WHICH HAS A VERY
9	LONG TITLE, BUT BASICALLY IS ABOUT OFFICE SPACE.
10	GOING TO TURN THAT OVER TO SENATOR TORRES.
11	MR. TORRES: THANK YOU. THANK YOU VERY
12	MUCH, MR. CHAIRMAN. THANK YOU VERY MUCH, RECORDING
13	SECRETARY.
14	IN 2005 CIRM ISSUED A REQUEST FOR
15	APPLICATION FROM MAJOR CITIES AROUND THE STATE TO
16	SEE WHICH CITY WOULD LIKE TO BE HOME TO CIRM. AS A
17	RESULT OF THAT, THERE WERE NO EMPLOYEES AT THE TIME,
18	SO EVERY CITY WAS AVAILABLE TO US. SO WE HAD
19	APPLICATIONS FROM SAN DIEGO, LOS ANGELES,
20	SACRAMENTO, EMERYVILLE, OAKLAND, AND SAN FRANCISCO.
21	AS A RESULT OF THAT, THOSE PRESENTATIONS WERE MADE
22	TO A MEETING OF THE BOARD OF CIRM IN FRESNO,
23	CALIFORNIA. CORRECT SO FAR BECAUSE I WAS NOT PART
24	OF THE BOARD AT THAT TIME?
25	AT THAT MEETING THEN MAYOR GAVIN NEWSOM
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T38

1	MADE THE MOST GENEROUS OFFER BY OFFERING TO PROVIDE
2	CIRM WITH FREE RENT FOR TEN YEARS, FREE PARKING FOR
3	TEN YEARS, FREE MAINTENANCE COSTS FOR TEN YEARS, AND
4	FREE HOTEL ROOMS TO THE TUNE OF 2400 HOTEL ROOMS AND
5	14,000 HOTEL ROOMS AT A DISCOUNT IN SAN FRANCISCO
6	FOR MANY OF THE SPEAKERS THAT WE WOULD INVITE AND
7	OBVIOUSLY THE REVIEWERS THAT WE WOULD HAVE FOR GRANT
8	REVIEW ISSUES.
9	WELL, THAT DAY IS OVER, AND THE LEASE RUNS
10	OUT OCTOBER 31ST, 2015. AS A RESULT, I MADE THE
11	FIRST INITIATIVE TO CONTACT A REAL ESTATE AGENT WITH
12	COLDWELL BANKER ON APRIL 14, 2014, TO BEGIN THE
13	PROCESS TO LOOK FOR POTENTIAL OFFICE SPACE. THE
14	FIRST PRIORITY, AS MR. JUELSGAARD AND I DISCUSSED,
15	LET'S GET SOMETHING FREE AND MOVE FROM THERE. I
16	DON'T THINK THAT, MUCH TO OUR MUTUAL WISHES, WE'RE
17	GOING TO GET ANYTHING FOR FREE AGAIN.
18	ALSO, WE HAVE EMPLOYEES NOW THAT WE NEED
19	TO BE CONCERNED ABOUT IN TERMS OF WHERE THEY LIVE,
20	HOW THEY GET TO WORK, THEIR SAFETY, AND THEIR
21	ABILITY TO LIVE IN AN ENVIRONMENT OF WORK WHICH IS
22	ACCESSIBLE TO DISABLED AND THE HANDICAPPED WHICH IS
23	SAFE AND WHICH IS AN ENVIRONMENT THAT THEY CAN
24	CONTINUE TO DO THE GREAT WORK THAT ARE OUR STAFF
25	DOES.

1	THE SUBCOMMITTEE OF OUR GOVERNANCE
2	COMMITTEE HELD A HEARING A FEW MONTHS AGO AND
3	OUTLINED A SERIES OF CRITERIA THAT WE SHOULD UTILIZE
4	IN LOOKING FOR NEW OFFICE SPACE. AND THAT CRITERIA
5	WAS ESTABLISHED IN CONJUNCTION WITH THE GOVERNANCE
6	SUBCOMMITTEE MEMBERS AS WELL AS WITH OUR CHAIR AND
7	PRESIDENT AND SOME CIRM TEAM MEMBERS, MANY OF WHOM
8	CONTRIBUTED THEIR IDEAS AND THEIR PERSPECTIVES.
9	AGAIN, I WANT TO THANK A NUMBER OF THE BOARD
10	MEMBERS, ESPECIALLY BOARD MEMBER JUELSGAARD, FOR
11	GIVING US SOME PERSPECTIVES OF WHAT WE SHOULD BE
12	LOOKING FOR.
13	AS A RESULT OF THAT, THIS WILL ALLOW US, I
14	THINK, TO MOVE EFFICIENTLY AND OPPORTUNISTICALLY
15	GUIDED BY THE CRITERIA, WHICH IS ITEM NO. 2 IN YOUR
16	PAMPHLET, ITEM NO. 12, BUT PAGE 3 OF YOUR BOARD
17	PAMPHLET.
18	THE RECOMMENDATION BY THE BOARD BY THE
19	SUBCOMMITTEE WAS TO DELEGATE AUTHORITY FOR THE
20	NEGOTIATION AND EXECUTION OF A LEASE FOR NEW OFFICE
21	SPACE IN THE BAY AREA ALONG WITH NEGOTIATION AND
22	EXECUTION OF OTHER CONTRACTS NECESSARY FOR OUR
23	RELOCATION.
24	I CANNOT EMPHASIZE ENOUGH THAT WE HAVE
25	SAVED THE TAXPAYERS A LITTLE OVER 12 MILLION OVER
	140

THIS TEN-YEAR PERIOD BECAUSE WE DID NOT PAY RENT, WE
DID NOT PAY FOR MAINTENANCE COSTS, AND WE DID NOT
PAY FOR PARKING. AND I BELIEVE THAT THIS
INSTITUTION OUGHT TO BE GIVEN CREDIT FOR THAT
BECAUSE NO OTHER STATE AGENCY CAN CLAIM, NO. 1, THAT
IT HAS CREATED 38,000 NEW JOBS, THAT IT BRINGS IN A
POTENTIAL OF 214 MILLION IN NEW REVENUE TO THE STATE
OF CALIFORNIA, AND SAVED THE STATE 12 MILLION
BECAUSE OF THE GENEROSITY AND THE VISION OF OUR
CURRENT LIEUTENANT GOVERNOR WHEN HE WAS MAYOR OF SAN
FRANCISCO.
SO THIS RECOMMENDATION IS PRETTY
STRAIGHTFORWARD, AND I OPEN IT UP TO DISCUSSION, MR.
CHAIRMAN.
CHAIRMAN THOMAS: MS. WESTON.
DR. WESTON: DO YOU HAVE ANY IDEA WHAT THE
RANGE OF RENTS WOULD LIKELY BE?
MR. TORRES: WHEN WE STARTED OUT, THE
RANGE WAS AT ONE LEVEL, WHICH WAS 56 TO \$60 A SQUARE
FOOT IN SAN FRANCISCO BECAUSE THAT WAS AT LEAST MY
PREFERENCE, AND IN WORKING WITH THE MAYOR AND OTHER
PHILANTHROPISTS IN SAN FRANCISCO TO SEE IF WE COULD
FULFILL MR. JUELSGAARD'S WISHES AND DREAMS OF FREE
RENTAL SPACE AND PARKING. AS WE PROCEEDED THROUGH
THE PROCESS, THOSE RENTS INCREASED IN SAN FRANCISCO.
1/1

1	IN OAKLAND THE VARIABLES WERE \$46 A SQUARE
2	FOOT AND ABOUT \$280 A MONTH FOR EMPLOYEE PARKING.
3	IN THE CITY OF EMERYVILLE, WHICH SURPASSED LOS
4	ANGELES TEN YEARS AGO WHEN THEY'RE PRESENTED THEIR
5	PROPOSAL, THEY ARE STILL COMING BACK TO US WITH AN
6	AMOUNT PER SQUARE FOOT, AND I'M SURE IT WILL BE LESS
7	THAN OAKLAND, BUT THEIR PROPOSAL FOR PARKING WAS
8	ANYWHERE BETWEEN \$75 TO \$90 A MONTH IN EMERYVILLE.
9	WE ALSO LOOKED AT PROPERTY IN SOUTH SAN FRANCISCO,
10	AND THAT PROPERTY WAS, AGAIN, COMPETITIVE WITH THE
11	RATES THAT WE SAW IN OAKLAND AND EMERYVILLE.
12	THE OTHER FACTORS THAT WE TOOK INTO MIND
13	WAS TRANSPORTATION ISSUES. HOW CLOSE ARE THESE
14	OFFICES GOING TO BE TO BART? RIGHT NOW WE'RE PRETTY
15	WELL SITUATED WITH CAL. TRAIN STOPPING AT 4TH
16	STREET, AND IT'S ABOUT A THREE-MINUTE WALK TO OUR
17	HEADQUARTERS, PEOPLE BEING ABLE TO CATCH BART BY
18	MUNI EXTENSION JUST ACROSS THE STREET AND GETTING TO
19	THE EAST BAY AND OTHER PARTS OF THE BART SYSTEM;
20	WHEREAS, THOSE PEOPLE WHO LIVE IN MARIN HAVE A MORE
21	DIFFICULT TIME IN ACCESSING PUBLIC TRANSPORTATION
22	UNLESS THEY GET ON A FERRY OR DRIVE IN.
23	SO THOSE WERE THE RATES THAT WE WERE
24	LOOKING AT, AND WE STILL HAVEN'T RECEIVED FINAL
25	COUNTEROFFERS TO ANY OF THESE PROPOSALS THAT I'VE
	142

CITED, AND THAT'S WHAT WE'RE WAITING FOR TO
DETERMINE JUST WHAT THOSE RATES WILL BE ON PARKING,
MAINTENANCE COSTS.
THE OTHER FACTOR, WHICH MR. JUELSGAARD AND
I DISCUSSED AS WELL AS WITH THE CHAIRMAN AND WITH
OUR PRESIDENT, WAS CAN WE FIND OFFICE SPACE THAT
WOULD ALLOW US TO HOLD OUR BOARD MEETINGS IN THAT
OFFICE SPACE? CAN WE SAVE MONEY THAT WAY AS WELL BY
NOT HAVING TO PAY FOR HOTEL SPACE AS WELL AS FOR OUR
GRANT REVIEW SESSIONS AS WELL AS FOR OUR BOARD. AND
THOSE ARE FACTORS THAT WE'VE PUT FORWARD IN OUR
DISCUSSIONS WITH THE THREE CITIES THAT WE'RE TALKING
ABOUT AND TO SEE WHAT THEY COME BACK WITH.
SO WE MET PRELIMINARILY. MY FIRST VENTURE
SINCE MY SURGERY WAS MONDAY, AND WE HAD AN EXCELLENT
MEETING WITH THE NEW MAYOR OF OAKLAND AND HER STAFF,
AND BERT LUBIN, ONE OF OUR BOARD MEMBERS, WAS THERE
AS WELL WHO KNOWS THE NEW MAYOR VERY WELL. OUR
PRESIDENT, MARIA BONNEVILLE, AS WELL AS OUR CHAIR,
J.T., AND OUR REAL ESTATE BROKER WERE ALL PRESENT AT
THAT MEETING.
WE THEN HAD A MEETING WITH THE CITY
MANAGER OF EMERYVILLE TO DISCUSS WHAT POTENTIAL WAS
THERE. EMERYVILLE WAS JUST CREATED INTO A CHARTER
CITY IN THE NOVEMBER ELECTION 2014, WHICH MEANS THAT
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1	THE CITY MANAGER IS ESSENTIALLY THE MAYOR AND MAKES
2	ALL ADMINISTRATIVE DECISIONS. SO WE SAT DOWN WITH
3	HER, AND THEN WE FOUND OUT SHE'S JUST BEEN RECRUITED
4	TO BECOME THE CITY MANAGER FOR OAKLAND. SO WE'LL
5	SEE WHAT KIND OF ALTERNATIVE OFFER HE OR SHE MAY
6	BRING FORWARD.
7	SO WE EXPLORED EVERY SUGGESTION THAT WAS
8	GIVEN TO ME BY STAFF IN THE HALLWAYS, IN THE
9	KITCHEN, AND SOMETIMES IN THE MEN'S RESTROOM, HAVE
10	YOU CHECKED OUT THIS SPACE, HAVE YOU CHECKED OUT
11	THAT SPACE, TO THE POINT WHERE I ALMOST WANT TO GET
12	A REAL ESTATE LICENSE. BUT THE FACT OF THE MATTER
13	IS THIS IS WHERE WE'RE AT IN TERMS OF THE POTENTIAL
14	SPACE. AND NOW WE'RE AWAITING TO GET RESPONSES BACK
15	FROM THESE CITIES TO SEE WHAT ELSE THEY CAN PROVIDE.
16	THE MAYOR OF OAKLAND TOOK INTO
17	CONSIDERATION HOTEL ROOMS AND WHAT KIND OF SPACE IS
18	THERE. AND I DON'T WANT TO GET INTO TOO MANY OTHER
19	DETAILS OTHER THAN TO SAY THEY'RE LOOKING AT
20	WHATEVER ALTERNATIVES THEY CAN COME UP WITH THAT
21	WILL HELP US THROUGH THIS PROCESS. THERE'S NO
22	QUESTION I'M SURE THE GOVERNOR WOULD WANT US TO BE
23	IN OAKLAND SINCE THAT'S HIS HOME CITY, BUT THAT
24	STILL REMAINS TO BE SEEN IN TERMS OF WHAT IS THE
25	BEST OFFER THAT WE CAN LOOK AT IN TERMS OF AN
	144

1	AGENCY.
2	MR. SHEEHY: I'D LIKE TO MAKE MOTION TO
3	ADOPT THIS, TO MOVE THIS, BUT WITH A MINOR
4	AMENDMENT. I WOULD ALSO LIKE TO TASK THE PRESIDENT
5	WITH CONSIDERING A COMPENSATION, A TEAMWIDE
6	COMPENSATION BECAUSE WHAT'S GOING TO HAPPEN LET
7	ME EXPLAIN. IF WE MOVE, SOME OF THESE FOLKS WHO
8	HAVE BEEN JUST ASTONISHING IN THE AMOUNT OF
9	CHANGE WE'VE GONE THROUGH, THE WAY THE TEAM MEMBERS
10	HAVE ADAPTED, AND THE WAY THAT MANY OF THEM HAVE
11	CARRIED US FORWARD FOR OUR ENTIRE MISSION, SOME
12	PEOPLE COULD END UP BEING IN REAL TERMS RECEIVING
13	LESS MONEY BECAUSE THEIR COMMUTE CHANGES. THEY HAVE
14	TO GO OVER A BRIDGE. THEY MAY HAVE TO TAKE A
15	DIFFERENT PATH.
16	AND SO THAT NO MEMBER OF OUR TEAM ENDS UP
17	MAKING LESS TOMORROW THAN TODAY BECAUSE OF SOMETHING
18	THEY HAD NO CONTROL OVER, IF THE PRESIDENT CAN
19	CONSIDER SOME SORT OF POLICY, INDEPENDENT OF MERIT
20	OR COST OF LIVING RAISES, TO MAKE SURE THAT FOLKS
21	ARE NOT HARMED. AND TO BE FAIR, I THINK IT PROBABLY
22	SHOULD BE ACROSS THE BOARD SO THAT WE'RE NOT PAYING
23	FOR PEOPLE'S COMMUTES DIRECTLY, BUT MORE OR LESS AS
24	A RECOGNITION OF JUST THE INCREDIBLE WORK THIS TEAM
25	HAS DONE, THAT THEY HAVE PUT UP WITH A LOT OF NEW

```
1
     THINGS. AND NOT ONLY HAVE THEY PUT UP WITH IT,
 2
     THEY'VE ACCEPTED IT, THEY'VE THRIVED, AND THEY'RE
 3
     LEADING US INTO THE CIRM 2.0, AND THEY'RE
 4
     IRREPLACEABLE.
 5
               AND I JUST, AS SOMEBODY WHO WORKS FOR
 6
     PEOPLE, THE IDEA THAT I WOULD WAKE UP ON MONDAY AND
 7
     HAVE A SUBSTANTIAL -- A BIT LESS MONEY IN MY POCKET
     FOR DOING THE SAME JOB JUST DOESN'T REALLY STRIKE ME
 8
 9
     AS THE WAY WE SHOULD BE REWARDING THIS INCREDIBLE
10
     TEAM.
11
                MR. TORRES: I TOTALLY AGREE, NO. 1. NO.
12
     2, THAT'S WHY WE TOOK INTO CONSIDERATION, BECAUSE
13
     YOU AND I TALKED ABOUT THIS AS WELL, JEFF, HOW TO
     MAKE SURE THAT WE LOOK AT POTENTIAL OFFICE SPACE
14
15
     THAT IS ACCESSIBLE TO BART OR SOME OTHER SHUTTLES
16
     THAT WON'T INCREASE THE COST OF GETTING TO THE
17
     OFFICE.
               THE ISSUE OF PARKING IS GOING TO BE A
18
19
     MAJOR ISSUE, BUT I'D LIKE TO ASK MR. HARRISON, IF I
20
     MAY, JUST WHAT IS THE AUTHORITY THAT WE HAVE AS A
21
     STATE AGENCY TO PROVIDE FOR THE TYPE OF
22
     COMPENSATION -- NOT COMPENSATION, BUT THE TYPE OF
     RELIEF THAT JEFF IS PROPOSING.
23
24
               MR. HARRISON: SO I THINK YOU'RE RIGHT
25
     ACTUALLY TO THINK OF IT AS COMPENSATION. AND
                               146
```

1	ULTIMATELY THE BOARD HAS THE AUTHORITY TO SET
2	COMPENSATION, WHICH INCLUDES BENEFITS, AT LEVELS
3	SIMILAR TO THOSE PAID FOR EMPLOYEES IN EQUIVALENT
4	POSITIONS AT THE NONPROFIT INSTITUTIONS THAT ARE
5	REPRESENTED ON THIS BOARD, INCLUDING THE UNIVERSITY
6	OF CALIFORNIA AND THE PRIVATE NONPROFIT RESEARCH
7	INSTITUTIONS. SO THOSE ARE COMPARATORS, AND THAT'S
8	WHAT WE'RE CHARGED WITH EVALUATING.
9	MR. TORRES: SO WE COULD LEGALLY PROVIDE
10	THAT IRRESPECTIVE OF MERIT INCREASE AND THE OTHER
11	USUAL PANOPLY OF COMPENSATION INCREASES?
12	MR. HARRISON: SO THERE ARE TWO DIFFERENT
13	WAYS TO LOOK AT IT. GENERALLY THE BOARD HAS
14	ESTABLISHED A RANGE OF COMPENSATION FOR EACH
15	POSITION AT CIRM BASED ON THE COMPENSATION OFFERED
16	FOR PEOPLE IN SIMILAR POSITIONS AT OUR COMPARATOR
17	INSTITUTIONS. WE USUALLY TRY TO TARGET ABOUT 80
18	PERCENT OF THE RANGE FOR SALARY. SO WE MAY HAVE
19	SOME ROOM THERE. I THINK WE HAVE TO TAKE A LOOK AT
20	THAT. AND THEN ALSO TO THE EXTENT WE'RE LOOKING AT
21	PARKING SEPARATE, THE DEGREE TO WHICH OUR COMPARATOR
22	INSTITUTIONS OFFER PARKING AS PART OF A BENEFIT
23	PACKAGE
24	MR. TORRES: AT THIS POINT NO STATE AGENCY
25	OFFERS PARKING.

1	MR. HARRISON: THAT'S CORRECT.
2	MR. TORRES: MR. CHAIRMAN, YOU'RE THE
3	CHAIR IN THIS.
4	CHAIRMAN THOMAS: JUST SPEAKING TO MS.
5	WINOKUR.
6	MR. TORRES: MR. JUELSGAARD.
7	DR. JUELSGAARD: SO, SENATOR TORRES, WHEN
8	YOU SAY NO STATE AGENCY OFFERS PARKING, LET'S JUST
9	TAKE THE UC SYSTEM FOR A MOMENT. SO WHEN PROFESSORS
10	OR INSTRUCTORS COME TO TEACH AT THAT INSTITUTION ON
11	A DAILY BASIS AND THEY DRIVE IN FROM SOMEWHERE,
12	ISN'T PARKING PROVIDED? AREN'T THERE ON-SITE
13	PARKING STRUCTURES?
14	MR. TORRES: NO.
15	DR. JUELSGAARD: OH, LINDA, HOW LUCKY YOU
16	HAVE IT.
17	MR. TORRES: I MIGHT WANT TO REMIND YOU
18	NOT UNTIL THE LATE 1980S WAS THE LEGISLATURE
19	PROVIDED HEALTHCARE BENEFITS. I WAS COVERED UNDER
20	MY FORMER WIFE'S CONTRACT FOR HEALTHCARE BENEFITS.
21	SO IT WASN'T UNTIL THE LATE 1980S THAT EVEN THE
22	LEGISLATURE WAS PROVIDED HEALTH BENEFITS, AND
23	CLEARLY STATE EMPLOYEES WERE NOT PROVIDED PARKING.
24	CHAIRMAN THOMAS: DR. LEVIN.
25	DR. LEVIN: THANK YOU. CAN I JUST SAY I
	1.10
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1	THINK THIS IS A GREAT THING TO POINT OUT TO
2	CONSIDER; BUT AS WE'VE DISCUSSED IN PRIOR YEARS JUST
3	IN TERMS OF WHAT THE BOARD'S AUTHORITY IS OR SHOULD
4	BE AND PUBLIC PERCEPTION, THAT THIS KIND OF GETS
5	DOWN INTO THE WEEDS OF MANDATING POLICY FOR CIRM
6	ITSELF. AND I THINK WE SHOULD PROBABLY LEAVE IT AS
7	A SUGGESTION TO THE PRESIDENT AND LET HIM DEAL WITH
8	THE WORKINGS OF HIS OFFICE BECAUSE THIS REALLY IS A
9	STAFF
10	MR. TORRES: THAT WAS THE NATURE OF MR.
11	SHEEHY'S RECOMMENDATION.
12	DR. LEVIN: WE STARTED TALKING ABOUT WHAT
13	OUR AUTHORITY IS TO SET THINGS. IT SEEMS TO ME
14	OVERREACHING TO GO DOWN AND TO STIPULATE THAT CIRM
15	SHOULD PROVIDE THIS COMPENSATION FOR THIS THING OR
16	THIS FOR THAT FOR ITS STAFF.
17	MR. SHEEHY: THAT WAS NOT ACTUALLY WHAT I
18	WAS SAYING. I WAS JUST ASKING THE PRESIDENT TO
19	CONSIDER IF I DON'T KNOW WHERE WE'RE GOING TO END
20	UP; BUT IF THIS ENDS UP BECOMING SOMETHING THAT
21	HAPPENS, WITHOUT SPECIFYING OR I THINK THE WORD I
22	USED WAS CONSIDER, SO THAT'S ALL. AND WHAT I WANTED
23	TO DO IS GIVE SUPPORT TO THE PRESIDENT IF HE FELT
24	THAT THIS WAS SOMETHING WE MAY HAVE RETENTION
25	ISSUES. IF YOU'RE TALKING GOING FROM SAN FRANCISCO
	140

1	TO I MEAN SOME OF THESE PLACES, GETTING AROUND
2	THE CITY CAN BE A CHALLENGE. AND THESE ARE
3	EXTREMELY TALENTED PEOPLE WE HAVE. IT'S NOT LIKE
4	THERE ARE OTHER PEOPLE WHO WOULDN'T WANT TO HIRE
5	THEM. AND WE HAVE PEOPLE WITH FAMILIES. IT'S JUST
6	NOT EASY TO GET AROUND THE BAY AREA.
7	SO WHAT I'M TRYING TO DO IS, FIRST AS A
8	BOARD MEMBER, IS RECOGNIZE THE INCREDIBLE THIS IS
9	JUST ONE THING. OUR TEAM HAS ADAPTED SO GRACEFULLY
10	AND WITH SUCH EFFORT, I THINK THAT WE SHOULD
11	RECOGNIZE THAT INDEPENDENTLY IN MY OWN MIND, BUT I
12	WANT TO GIVE DIRECTION GIVE SUPPORT TO THE
13	PRESIDENT IF THIS IS SOMETHING THAT HE FEELS LIKE IS
14	USEFUL AS WE MAKE THIS TRANSITION. THAT'S THE
15	LIMIT, JUST TO PUT SOMETHING ON AS A BOARD TO MAKE
16	THAT STATEMENT.
17	DR. MILLS: THAT'S APPRECIATED, NOT JUST
18	BY ME, THE SUPPORT TO BE ABLE TO DO IT AND TO TAKE
19	CARE OF THE TEAM AND THE RECOGNITION OF THEIR WORK
20	ON BEHALF OF THEM. THANK YOU.
21	CHAIRMAN THOMAS: OKAY. SO, MR. SHEEHY,
22	PERHAPS COULD YOU RESTATE YOUR MOTION? I DON'T
23	THINK WE'VE HAD A SECOND ON IT YET.
24	MR. HARRISON: SO THE MOTION IS TO
25	DELEGATE AUTHORITY FOR THE NEGOTIATING AND EXECUTION

150

1	OF A LEASE FOR NEW OFFICE SPACE IN THE BAY AREA
2	ALONG WITH OTHER CONTRACTS NECESSARY FOR CIRM'S
3	RELOCATION TO THE CIRM PRESIDENT IN CONSULTATION
4	WITH THE CHAIR AND VICE CHAIR AND TO TASK THE
5	PRESIDENT WITH CONSIDERING A POLICY TO ENSURE THAT
6	TEAM MEMBERS ARE NOT FINANCIALLY HARMED BY THE
7	RELOCATION.
8	MR. SHEEHY: AND THIS WOULD BE INDEPENDENT
9	OF MERIT OR COST OF LIVING RAISES THAT MAY OR MAY
10	NOT BE IN THE WORKS.
11	CHAIRMAN THOMAS: OKAY. SO THAT'S THE
12	MOTION. DO WE HAVE A SECOND?
13	MR. ROWLETT: SECOND.
14	CHAIRMAN THOMAS: SECONDED BY MR. ROWLETT.
15	FURTHER DISCUSSION BY MEMBERS OF THE BOARD?
16	DR. PRIETO: WELL, I THINK THIS IS I
17	GUESS THERE'S NOT A PRECEDENT, FROM WHAT I
18	UNDERSTAND, FOR AN EMPLOYER TO OFFER FREE OR REDUCED
19	TRANSIT TO EMPLOYEES. CERTAINLY THAT EXISTS IN THE
20	PRIVATE SECTOR. I GUESS THERE'S NOT A PRECEDENT IN
21	STATE GOVERNMENT FOR THAT. MAYBE MR. HARRISON CAN
22	CORRECT ME.
23	MR. HARRISON: CIRM DOES HAVE A PUBLIC
24	TRANSIT PROGRAM.
25	DR. PRIETO: THAT WE PROVIDE?
	151
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160 S. OLD SPRINGS ROAD, SUITE 270, ANAHEIM, CALIFORNIA 92808 1-800-622-6092 1-714-444-4100 EMAIL: DEPO@DEPO1.COM

1	MR. TORRES: PUBLIC TRANSIT.
2	DR. PRIETO: WE FUND IT. BUT NOT PARKING.
3	MR. HARRISON: THAT'S RIGHT. PARKING WAS
4	OFFERED, AS SENATOR TORRES SAID, AS PART OF THE FREE
5	OFFICE SPACE.
6	CHAIRMAN THOMAS: OTHER COMMENTS,
7	QUESTIONS FROM MEMBERS OF THE BOARD? ANY PUBLIC
8	COMMENT? MR. REED. HOLD ON ONE SECOND, MR. REED.
9	MS. MILLER.
10	MS. MILLER: A CLARIFICATION ON IT. SO A
11	YES VOTE WOULD MEAN THAT YOU WOULD CONSIDER FUNDING
12	A PROGRAM LIKE THAT, NOT NECESSARILY THAT YOU WOULD
13	BE FUNDING IT?
14	MR. HARRISON: CORRECT.
15	MS. MILLER: OKAY.
16	MR. REED: TWO SUGGESTIONS ABOUT WHICH I
17	KNOW ALMOST NOTHING.
18	MR. TORRES: NOT THE FIRST TIME, DON.
19	MR. REED: I WAS GOING TO SAY THAT. FIRST
20	OFF, I WOULD LOVE TO SEE SOME KIND OF A STATUE TO
21	MEMORIALIZE THE TREMENDOUS EFFORT OF THE CALIFORNIA
22	STEM CELL PROGRAM. I'D LOVE TO SEE A STATUE, I
23	DON'T KNOW, SOMETHING WONDERFUL, SOMEBODY HELPING A
24	CHILD OUT OF A WHEELCHAIR. IF PEOPLE COULD GO AND
25	SEE AND TOUCH, EVEN IF THEY'VE NEVER BEEN TO A
	4-0
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1
     MEETING, BUT THEY CAN SEE AND TOUCH THIS. I'D LOVE
 2
     TO SEE -- MAYBE HAVE A LOCAL ARTIST CONTEST,
 3
     SOMETHING WHICH SUMMARIZES THE DREAM OF CIRM. I'D
 4
     LOVE TO SEE THAT.
               SECONDLY, I WONDER IF IT'S POSSIBLE, AS WE
 5
     LOOK FOR NEW OFFICE SPACE, TO CONSIDER A MILITARY
 6
 7
     BASE. THEY MAY HAVE BUILDINGS THAT WORK FOR US, BUT
 8
     ARE NOT DOING ANYTHING. MAY BE A POSSIBILITY.
 9
               CHAIRMAN THOMAS: THANK YOU FOR YOUR
10
     COMMENTS, MR. REED. OTHER COMMENTS FROM MEMBERS OF
     THE PUBLIC? MR. HARRISON, THIS IS ANOTHER VOICE
11
12
     VOTE, I PRESUME. FIRST IN THE ROOM, ALL THOSE IN
13
     FAVOR PLEASE SAY AYE. OPPOSED? ABSTENTIONS?
14
     MARIA, ROLL OF THOSE ON THE PHONE.
               MS. BONNEVILLE: KATHY LAPORTE.
15
16
               DR. LAPORTE: AYE.
17
               MS. BONNEVILLE: ELIZABETH FINI. KRISTINA
18
     VUORI.
19
               DR. VUORI: YES.
20
               CHAIRMAN THOMAS: THAT ITEM PASSES. ON TO
21
     THE NEXT ITEM, ITEM 13, CONSIDERATION OF APPOINTMENT
22
     OF NEW SCIENTIFIC MEMBERS AND PATIENT ADVOCATE
     MEMBER TO THE GRANTS WORKING GROUP. DR. SAMBRANO
23
24
     FOLLOWED BY J.T.
25
               DR. SAMBRANO: THANK YOU, MR. CHAIRMAN.
                              153
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160 S. OLD SPRINGS ROAD, SUITE 270, ANAHEIM, CALIFORNIA 92808 1-800-622-6092 1-714-444-4100 EMAIL: DEPO@DEPO1.COM

1	SO WE'RE BRINGING FOR YOUR CONSIDERATION TWO
2	NOMINEES FOR MEMBERSHIP INTO THE GRANTS WORKING
3	GROUP. THESE ARE THE SCIENTIFIC MEMBERS WHO ARE
4	BRINGING EXPERTISE IN CLINICAL OPERATIONS, STEM CELL
5	DEVELOPMENT, IMMUNOLOGY, AND CANCER. THE BRIEF
6	BIOGRAPHIES OF THE TWO INDIVIDUALS FOR CONSIDERATION
7	HAVE BEEN MADE AVAILABLE TO YOU. THEY ARE DR. LINDA
8	CUSTER AND DR. CASSIAN YEE.
9	CHAIRMAN THOMAS: DO I HEAR A MOTION TO
10	APPROVE?
11	DR. PRIETO: SO MOVED.
12	CHAIRMAN THOMAS: MOVED BY DR. PRIETO.
13	MR. JUELSGAARD: SECOND.
14	CHAIRMAN THOMAS: SECONDED BY MR.
15	JUELSGAARD. DISCUSSION FROM MEMBERS OF THE BOARD?
16	PUBLIC COMMENT? ALL THOSE IN FAVOR PLEASE SAY AYE.
17	OPPOSED? ABSTENTIONS? MARIA.
18	MS. BONNEVILLE: KATHY LAPORTE.
19	DR. LAPORTE: YES.
20	MS. BONNEVILLE: ELIZABETH FINI.
21	DR. FINI: YES.
22	MS. BONNEVILLE: KRISTINA VUORI.
23	DR. VUORI: YES.
24	CHAIRMAN THOMAS: THANK YOU. THE
25	ADDITIONAL WE NEED TO CONSIDER IS PATIENT ADVOCATE
	4-4
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	DARKESTERS REPORTERS SERVICE
1	LAUREN MILLER TO JOIN THIS AUGUST GROUP. DO I HEAR
2	A MOTION THAT SHE BE APPOINTED AS WELL?
3	DR. HIGGINS: SO MOVED.
4	DR. BURTIS: SECOND.
5	CHAIRMAN THOMAS: MOVED BY DR. HIGGINS,
6	SECONDED BY DR. BURTIS. DISCUSSION BY MEMBERS OF
7	THE BOARD? PUBLIC COMMENT? ROLL CALL VOTE. ALL IN
8	THE ROOM IN FAVOR PLEASE SAY AYE. OPPOSED?
9	ABSTENTIONS? MARIA.
10	MS. BONNEVILLE: KATHY LAPORTE.
11	DR. LAPORTE: YES.
12	MS. BONNEVILLE: ELIZABETH FINI.
13	DR. FINI: YES.
14	MS. BONNEVILLE: KRISTINA VUORI.
15	DR. VUORI: YES.
16	CHAIRMAN THOMAS: OKAY. THAT PASSES AS
17	WELL. THANK YOU.
18	ON TO ITEM 14, CONSIDERATION OF POLICY TO
19	EXTEND WORKER'S COMP COVERAGE TO CIRM VOLUNTEERS.
20	MR. HARRISON.
21	MR. HARRISON: SO OVER THE YEARS CIRM HAS
22	BENEFITED FROM THE SERVICES OF VOLUNTEERS RANGING
23	FROM SCIENTISTS TO ATTORNEYS TO STUDENT INTERNS. IN
24	TOTAL WE'VE HAD 27 INDIVIDUALS WHO VOLUNTEERED THEIR
25	TIME TO THE AGENCY. HISTORICALLY WE HAVE OFFERED TO
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1	PROVIDE WORKERS' COMPENSATION COVERAGE TO THESE
2	INDIVIDUALS IN RECOGNITION OF THE FACT THAT THEY ARE
3	HELPING TO ADVANCE CIRM'S MISSION. AND THAT AS AN
4	AGENCY THAT CARES ABOUT PATIENTS, IT'S IMPORTANT FOR
5	US TO PROTECT THEM IN THE EVENT THAT THEY'RE INJURED
6	WHILE VOLUNTEERING ON OUR BEHALF.
7	THE CALIFORNIA DEPARTMENT OF HUMAN
8	RESOURCES, HOWEVER, RECENTLY ADVISED US THAT IN
9	ORDER TO CONTINUE EXTENDING WORKERS' COMPENSATION
10	COVERAGE TO OUR VOLUNTEERS, WE NEEDED YOUR APPROVAL
11	IN ORDER TO DO SO.
12	SO WE WOULD ASK FOR YOUR VOTE TODAY TO
13	EXTEND WORKERS' COMPENSATION COVERAGE TO CIRM
14	VOLUNTEERS. I'D BE HAPPY TO ANSWER ANY QUESTIONS.
15	DR. JUELSGAARD: SO, JAMES, AT THE STATE
16	LEVEL IS WORKMAN'S COMPENSATION PROVIDED THROUGH AN
17	INSURANCE ARRANGEMENT?
18	MR. HARRISON: CORRECT. IT'S PROVIDED
19	THROUGH THE STATE COMPENSATION INSURANCE FUND.
20	DR. JUELSGAARD: SO IS IT THAT THE MORE
21	EMPLOYEES OR THE MORE PEOPLE YOU COVER, THE GREATER
22	THE AMOUNT OF PREMIUM THAT YOU WIND UP PAYING?
23	MR. HARRISON: I'M LOOKING FOR CHILA
24	SILVA-MARTIN. I ACTUALLY DON'T BELIEVE THAT IT WILL
25	HAVE AN EFFECT ON CIRM'S TOTAL PAYMENTS UNLESS WE
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1	ACTUALLY HAVE A VOLUNTEER WHO TAKES ADVANTAGE OF THE
2	POLICY. TO DATE WE HAVE NOT.
3	CHAIRMAN THOMAS: THERE'S CHILA COMING
4	RIGHT NOW, JAMES.
5	MR. TORRES: MR. CHAIRMAN, THIS IS A TREND
6	THAT IS OCCURRING WITH A LOT OF NON-PROFITS AND
7	VOLUNTEER ORGANIZATIONS WHERE THE NONPROFIT HAS TO
8	PROTECT ITSELF IN TERMS OF VOLUNTEERS WHO MAY BE
9	INJURED AS THEY'RE GIVING TIME. WE'RE DOING THAT
10	NOW WITH ONE LEGACY, THE OTHER BOARD THAT I SIT ON
11	FOR ORGAN TRANSPLANTATION WHERE WE HAVE WHAT'S
12	CALLED AMBASSADORS, AND A LOT OF THEM HELP IN
13	DEVELOPING THE ROSE PARADE FLOAT. WE'VE HAD A
14	NUMBER OF ACCIDENTS, AND NOW WE REALIZE WE'VE GOT TO
15	HAVE THEM COVERED. OTHERWISE WE'RE GOING TO BE IN
16	TREMENDOUS LIABILITY.
17	MR. HARRISON: MR. JUELSGAARD, TO CLARIFY,
18	WE DON'T ACTUALLY PAY ANY PREMIUM. THE AGENCY WOULD
19	ONLY PAY IF AND WHEN A VOLUNTEER IS INJURED.
20	DR. JUELSGAARD: THEN IT ISN'T INSURANCE.
21	MR. HARRISON: WELL, IT'S SORT OF A FORM
22	OF SELF-INSURANCE, I GUESS.
23	MR. ROWLETT: QUICK QUESTION. THE
24	POTENTIAL EXPOSURE IF WE DON'T HAVE PEOPLE COVERED
25	COULD BE FAR GREATER. SO CONSEQUENTLY YOU HAVE
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1	PEOPLE COVERED GIVEN THAT THE ORGANIZATION USES A
2	LARGE NUMBER OF VOLUNTEERS, CORRECT?
3	MR. HARRISON: THAT'S CORRECT. THEY COULD
4	GO TO CIVIL COURT TO SEEK DAMAGES OTHERWISE.
5	CHAIRMAN THOMAS: DO I HEAR A MOTION? IS
6	THIS A COMMENT, DR. LEVIN?
7	DR. LEVIN: I WAS JUST GOING TO SAY I
8	THINK IT'S GREAT. THIS BOARD HASN'T PAID ENOUGH
9	ATTENTION TO THE DEVASTATING EFFECTS OF GRANT REVIEW
10	INJURY IN THE PAST.
11	MR. TORRES: WE ALMOST HAD SUCH AN INJURY
12	EARLIER TODAY.
13	CHAIRMAN THOMAS: I TAKE THAT DOES THAT
14	MEAN YOU MOVE THE ITEM?
15	DR. LEVIN: I MOVE TO ACCEPT.
16	DR. GASSON: SECOND.
17	CHAIRMAN THOMAS: SECONDED BY DR. GASSON.
18	ANY OTHER BOARD DISCUSSION? PUBLIC COMMENT? VOICE
19	VOTE. ALL IN THE ROOM PLEASE SAY AYE IF YOU LIKE
20	THIS. THANK YOU. OPPOSED? ABSTENTIONS?
21	MS. BONNEVILLE: KATHY LAPORTE.
22	DR. LAPORTE: YES.
23	MS. BONNEVILLE: ELIZABETH FINI.
24	DR. FINI: YES.
25	MS. BONNEVILLE: KRISTINA VUORI.
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1	DR. VUORI: YES.
2	MS. BONNEVILLE: THANK YOU.
3	CHAIRMAN THOMAS: THAT ITEMS PASSES. ON
4	TO ITEM NO. 15, CONSIDERATION OF AUGMENTATION TO THE
5	REMCHO, JOHANSEN & PURCELL CONTRACT. AND THIS IS
6	ALWAYS THE ITEM WHERE JAMES FEELS VERY ANTSY AND
7	STARTS LOOKING AT HIS KEYBOARD.
8	MR. TORRES: WITH A FLUSHED FACE.
9	CHAIRMAN THOMAS: IN JULY OF 2014, MR.
10	HARRISON, WHO TO THAT POINT HAD BEEN COUNSEL TO THE
11	BOARD, WAS GIVEN THE LARGER RESPONSIBILITY OF
12	GENERAL COUNSEL TO THE AGENCY. AND THAT ADDITIONAL
13	AMOUNT OF WORK, WHICH HAS BEEN CONSIDERABLE, HAS
14	BROUGHT TO BEAR FURTHER HOURS THAT MR. HARRISON
15	NEEDS TO WORK ON BEHALF OF THE AGENCY AND THE PEOPLE
16	OF CALIFORNIA AND REQUIRED THAT THE \$500,000 AMOUNT
17	THAT WE HAD PREVIOUSLY BUDGETED FOR HIS FIRM'S
18	COMPENSATION BE INCREASED ACCORDINGLY. AND WE FEEL
19	THAT, INSTEAD OF 500, THAT 600,000 IS THE
20	APPROPRIATE AMOUNT TO COVER THE TOTAL AMOUNT OF WORK
21	HE'S DOING.
22	SO I WOULD LIKE TO ENTERTAIN A MOTION TO
23	INCREASE MR. HARRISON AND REMCHO'S SALARY BY
24	\$100,000 FOR THIS CALENDAR YEAR.
25	MR. TORRES: SO MOVED.
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1	DR. GASSON: SECOND.
2	CHAIRMAN THOMAS: SO MOVED BY SENATOR
3	TORRES, SECONDED BY DR. GASSON. DR. MILLS HAS A
4	COMMENT.
5	DR. MILLS: IF ANYONE HAS ANY QUESTIONS
6	ABOUT IT, I CAN PROVIDE COLOR. OTHERWISE NO.
7	CHAIRMAN THOMAS: AS ALWAYS, JUST TO
8	FURTHER THE EMBARRASSMENT OF MR. HARRISON, THIS IS
9	ALWAYS THE POINT IN THIS PARTICULAR AGENDA TOPIC
10	WHERE I POINT OUT HOW
11	MR. TORRES: YOU'RE ENJOYING IT A LITTLE
12	TOO MUCH.
13	CHAIRMAN THOMAS: HE'S DOING SUCH AN
14	EXEMPLARY JOB AS ALWAYS, AND WE ARE VERY FORTUNATE
15	TO HAVE HIM AS OUR GENERAL COUNSEL.
16	(APPLAUSE.)
17	DR. MILLS: I ACTUALLY WILL JUST POINT OUT
18	TWO THINGS BECAUSE I THINK THEY'RE FAIRLY MATERIAL.
19	ONE IS THE INCREASE WE'RE TALKING ABOUT IS THE
20	MAXIMUM AMOUNT WE WOULD PAY HIM, NOT THE AMOUNT WE
21	WILL PAY HIM. HE'S FEE FOR SERVICE. SO IF HE WORKS
22	THAT TIME, THEN HE WOULD GET PAID. BUT IF HE
23	DOESN'T, THEY DON'T.
24	SECONDLY, THE INCREASE THAT WE'RE TALKING
25	ABOUT OVER HIS PREVIOUS IS \$100,000. SINCE HE CAME
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1	ON TO THE ROLE OF GENERAL COUNSEL, WE'VE ACTUALLY
2	REMOVED \$400,000 IN LEGAL EXPENSE. SO I JUST WANT
3	TO MAKE IT CLEAR. IT'S A POTENTIAL INCREASE OF A
4	HUNDRED THOUSAND ONLY IF THE WORK IS THERE FOR AN
5	ALREADY GAINED REDUCTION OF 400,000.
6	MR. TORRES: THAT'S AN IMPORTANT POINT,
7	MR. CHAIRMAN, BECAUSE THE REDUCTION IS SUBSTANTIAL.
8	AND I THINK THAT THAT REALLY LENDS ITSELF TO THE
9	OVERVIEW THAT THE PRESIDENT JUST ARTICULATED, THAT
10	THIS AGENCY, ESPECIALLY WHEN THIS NEW PRESIDENT CAME
11	ON BOARD, HAS ALWAYS BEEN TRYING TO FIND WHERE CAN
12	WE SAVE MONEY, WHERE CAN WE CUT COSTS, AND I APPLAUD
13	HIM FOR DOING THAT.
14	CHAIRMAN THOMAS: THANK YOU, MR. SENATOR.
15	THANK YOU, DR. MILLS. OTHER COMMENTS OR QUESTIONS
16	BY MEMBERS OF THE BOARD?
17	DR. LAPORTE: I HAPPEN TO HAVE LEGAL THAT
18	REPORTS TO ME IN A COMPANY OF ABOUT THE SAME SIZE
19	BUDGET AS THIS AGENCY. (INAUDIBLE) HIGHER. ONE
20	COULD CONSIDER INTERNAL COUNSEL. THAT'S ALWAYS A
21	TRADE-OFF, BUT THAT'S IN LINE WITH WHAT I'M SEEING.
22	CHAIRMAN THOMAS: ANY OTHER COMMENTS? WE
23	WON'T EVEN ASK MR. HARRISON IF THIS IS A VOICE VOTE
24	OR NOT SINCE HE'S CONFLICTED, BUT WE WILL GO TO DO
25	ONE ANYWAY. SO ALL THOSE IN FAVOR OF THIS MOTION
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1	PLEASE SAY AYE. OPPOSED? ABSTENTIONS? MARIA.
2	MS. BONNEVILLE: KATHY LAPORTE.
3	
	DR. LAPORTE: YES.
4	MS. BONNEVILLE: ELIZABETH FINI.
5	DR. FINI: YES.
6	MS. BONNEVILLE: KRISTINA VUORI.
7	DR. VUORI: YES.
8	CHAIRMAN THOMAS: OKAY. THAT PASSES.
9	CONGRATULATIONS, MR. HARRISON. AND THANK YOU ONCE
10	AGAIN FOR YOUR EXTREMELY EXEMPLARY WORK.
11	OKAY. WE GET TO GO NOW TO CLOSED SESSION.
12	MR. HARRISON, WOULD YOU INSTRUCT US ON TODAY'S
13	TOPICS?
14	MR. HARRISON: THE BOARD WILL BE MEETING
15	IN CLOSED SESSION PURSUANT TO HEALTH AND SAFETY CODE
16	SECTION $125290.30(f)(3)(D)$ TO CONSIDER THE
17	EVALUATION OF THE CHAIR.
18	CHAIRMAN THOMAS: OKAY. MARIA, WOULD YOU
19	LIKE TO DESCRIBE THE LOGISTICS OF THIS?
20	MS. BONNEVILLE: FOR THOSE OF YOU ON THE
21	PHONE, I'VE SENT YOU A CLOSED SESSION DIAL-IN
22	NUMBER. YOU CAN DIAL IN AND WE'LL BE OVER THERE IN
23	A COUPLE OF MINUTES. FOR PEOPLE WHO ARE HERE, IT'S
24	JUST RIGHT ACROSS THE HALL. SO THE BOARD MEMBERS
25	SHOULD FEEL FREE TO WANDER ACROSS THE HALL.
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1	CHAIRMAN THOMAS: WE ARE GOING TO GO OUT
2	FOR A WHILE FOR CLOSED SESSION. WE WILL THEN
3	RECONVENE WHEN CLOSED SESSION IS FINISHED TO COME
4	BACK TO COMPLETE TODAY'S AGENDA.
5	BEFORE ANYBODY LEAVES, JUDY, DID YOU WANT
6	TO ADDRESS THE BOARD? NO REASON FOR YOU TO HANG
7	AROUND.
8	MS. ROBERSON: I'M JUDY ROBERSON FROM
9	SACRAMENTO. AND CHAIRMAN THOMAS AND ICOC MEMBERS,
10	THE HUNTINGTON'S DISEASE FAMILIES OF CALIFORNIA
11	THANK YOU FOR FUNDING UC IRVINE'S DR. LESLIE
12	THOMPSON'S GRANT TODAY. BECAUSE OF CIRM,
13	HUNTINGTON'S FAMILIES HAVE HOPE FOR A FUTURE
14	TREATMENT FOR THIS HEREDITARY AND ALWAYS FATAL BRAIN
15	DISEASE THAT AFFECTS BOTH ADULTS AND CHILDREN
16	GENERATION AFTER GENERATION, FAMILIES LIKE MINE.
17	THANK YOU SO MUCH.
18	CHAIRMAN THOMAS: THANK YOU, JUDY. AND
19	THANK YOU FOR MAKING THE TRIP AS ALWAYS.
20	MEMBERS OF THE BOARD, IF YOU CAN JUST
21	PROCEED ACROSS THE HALL AT THIS TIME.
22	(THE BOARD THEN MET IN CLOSED
23	SESSION, NOT REPORTED NOR HEREIN TRANSCRIBED. AT
24	THE CONCLUSION OF THE CLOSED SESSION, THE FOLLOWING
25	WAS HEARD IN OPEN SESSION:)
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1	CHAIRMAN THOMAS: FOR THOSE OF YOU WHO
2	STUCK IT OUT ON THE PHONE, WE ARE RECONVENING. DOWN
3	TO HOME STRETCH HERE.
4	NO. 1, WE'RE GOING TO HAVE A
5	COMMUNICATIONS UPDATE. ANN HOLDEN, COULD YOU PLEASE
6	COME AND PRESENT?
7	KEVIN, AM I INTRODUCING YOU AND YOU'RE
8	INTRODUCING ANN?
9	MR. MC CORMACK: I'M GOING TO GIVE A
10	PRESENTATION, AND THEN ANN WILL FOLLOW ME.
11	CHAIRMAN THOMAS: I WILL SAY YOU'RE
12	LOOKING QUITE SARTORIAL TODAY, IF I DO SAY SO.
13	MR. MC CORMACK: IT'S SHOW BIZ, DARLING.
14	CHAIRMAN THOMAS: THERE YOU GO.
15	MR. MC CORMACK: DARLING CHAIRMAN THOMAS,
16	MEMBERS OF THE BOARD, AND JEANNE BEING THE ONLY
17	MEMBER OF THE PUBLIC HERE, AND COLLEAGUES, AND DON,
18	OF COURSE. I'M JUST GOING TO GIVE A BRIEF
19	PRESENTATION ABOUT SOME OF THE MEDIA THAT WE'VE HAD
20	OVER THE LAST COUPLE OF MONTHS SINCE WE LAST MET IN
21	JANUARY. AND FOLLOWING ON FROM THE PRESIDENT, I'M
22	GOING TO TRY AND BE PITHY AND MAKE WAY AFTER THAT
23	FOR ANN HOLDEN WHO'S GOING TO TALK ABOUT SOMETHING
24	THAT I THINK IS REALLY IMPORTANT WHICH IS THE
25	COMPLETE REDO OF OUR WEBSITE.
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1	SO WE'VE HAD OVER THE LAST COUPLE OF
2	MONTHS SOME PRETTY HEAVY MEDIA COVERAGE, STARTING
3	MOSTLY WITH THE NEWS FROM CITY OF HOPE, THAT THEY'VE
4	BEEN APPROVED FOR A CLINICAL TRIAL IN HIV/AIDS. AND
5	THIS PROJECT, WHICH ALSO INVOLVES USC AND SANGAMO
6	BIOSCIENCES, WAS FEATURED IN A FAIRLY GOOD ARTICLE
7	IN BUZZFEED, WHICH IS AN INTERNATIONAL GLOBAL MEDIA
8	WEBSITE. AND THAT GOT A LITTLE PICKUP AROUND THE
9	INTERNET. SO WE GOT A LOT OF REALLY GOOD REACH AND
10	A LOT OF REALLY GOOD RESPONSE FROM THAT ONE PIECE.
11	IT WAS ALSO FEATURED IN AN ARTICLE IN THE
12	SAN FRANCISCO BUSINESS TIMES AND ALSO KQED RADIO IN
13	THE "FORUM" RADIO SHOW, WHICH IS A BROADCAST
14	STATEWIDE. SO THAT WAS GREAT.
15	JOHN ZAIA, WHO'S THE PRINCIPAL
16	INVESTIGATOR FROM CITY OF HOPE ON THIS PROJECT, AND
17	OUR COLLEAGUE ON THE BOARD, JEFF SHEEHY, WERE THE
18	GUESTS ON THAT. AND THEY BOTH DID A WONDERFUL JOB
19	OF TALKING ABOUT THE PROJECT, THE IMPORTANCE OF THIS
20	WORK IN HIV/AIDS, AND ALSO THE IMPORTANCE OF THE
21	STEM CELL AGENCY IN HELPING TO FUND IT. SO THAT WAS
22	REALLY GOOD. WE GOT A REALLY GREAT RESPONSE FROM
23	THAT PARTICULAR PIECE.
24	AND, AGAIN, DR. ZAIA DID AN INTERVIEW WITH
25	24/7 NEWS, WHICH IS A RADIO SYNDICATION SERVICE THAT
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1	FEEDS STORIES LIKE THIS TO SOME 250 RADIO STATIONS
2	AROUND THE COUNTRY. SO, AGAIN, AN AWFUL LOT OF
3	EXPOSURE FROM ONE INTERVIEW. SO IT WAS REALLY GOOD
4	BE TO BE ABLE TO GET THAT NEWS OUT.
5	THERE WAS ANOTHER ONE WITH ONE OF OUR
6	OTHER CLINICAL TRIAL PARTNERS, ASTERIAS. THIS IS
7	SOMETHING THAT DR. MILLS WILL KNOW ALL ABOUT. THEY
8	WERE GIVEN THE OPPORTUNITY TO RING THE OPENING BELL
9	AT THE NEW YORK STOCK EXCHANGE. FOR THEM IT WAS A
10	GREAT THRILL OBVIOUSLY, AND IT ALSO GAVE US A CHANCE
11	TO KIND OF GET SOME STORIES IN THE MEDIA,
12	PARTICULARLY ABOUT THE SPINAL CORD INJURY TRIAL THAT
13	THEY'RE DOING, THEY'RE CARRYING OUT AND HAVE JUST
14	BEGUN AND THAT WE'RE FUNDING. SO, AGAIN, WE
15	SHAMELESSLY EXPLORE EVERY OPPORTUNITY TO GET OUR
16	NAME IN THE MEDIA.
17	AT THE LAST MEETING YOU VOTED ON THE TOOLS
18	AND TECHNOLOGY AWARDS. AND, AGAIN, THAT GOT SOME
19	REALLY GOOD COVERAGE IN THE SACRAMENTO BUSINESS
20	TIMES JOURNAL, VARIOUS OTHER NEWS ORGANIZATIONS, AND
21	SOME OF OUR OTHER RESEARCH PROJECTS AS WELL GOT SOME
22	NATIONAL COVERAGE AND THINGS LIKE THE SCIENTISTS AND
23	MEDICAL NEWS TODAY.
24	AND, FINALLY, XCONOMY, WHICH IS A BIOTECH
25	WEBSITE, DID A REALLY NICE PROFILE OF DR. MILLS AND
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1	CIRM 2.0. I PARTICULARLY LIKED THE OPENING
2	PARAGRAPH. IT SAYS, "HOME REMODELING SHOWS ARE A
3	REALITY TV STAPLE, BUT NO PARK AVENUE MANSION OR
4	COUNTRY ESTATE CAN TOP THE ONE BILLION PRICE TAG C.
5	RANDALL MILLS IS TRYING TO RENOVATE ON THE FLY IN
6	CALIFORNIA," WHICH MADE US SOUND LIKE SOME KIND OF
7	REAL HOUSEWIVES OF ORLANDO SHOW. IT WAS GREAT.
8	BUT, REALLY, I THINK ONE OF THE MOST
9	IMPORTANT THINGS THAT'S HAPPENED LATELY IS THE REDO
10	OF OUR WEBSITE. AND DR. ANNE HOLDEN, WHO'S OUR
11	WEBSITE MANAGER AND OUR SOCIAL MEDIA GURU, HAS BEEN
12	WORKING WITH A TEAM OF PEOPLE FOR THE LAST FEW
13	MONTHS ON DOING THIS. AND SO I'D LIKE YOU TO HEAR
14	FROM ANNE.
15	DR. HOLDEN: THANKS, EVERYONE. I WANT TO
16	TAKE A FEW MINUTES TO UPDATE THE BOARD AND EVERYONE
17	ELSE HERE ON THE CREATION, DEVELOPMENT, AND
18	EXECUTION OF OUR NEW WEBSITE THAT HOPEFULLY YOU ALL
19	HAVE TAKEN A LOOK AT IN THE LAST FEW DAYS.
20	SO AS SOME OF YOU MAY KNOW, OUR WEBSITE
21	WAS ACTUALLY REDESIGNED A FEW YEARS AGO, BUT THERE
22	ARE SEVERAL IMPORTANT REASONS WHY WE FELT ANOTHER
23	REDESIGN AT THIS POINT IN TIME IN CIRM'S LIFETIME
24	WAS NECESSARY. WITH THE LAUNCH OF CIRM 2.0, WE
25	WANTED TO OFFER AN IMPROVED USER EXPERIENCE TO TWO

1	OF OUR VERY IMPORTANT CONSTITUENTS; THAT IS, CURRENT
2	AND POTENTIAL GRANTEES, PATIENTS, AND PATIENT
3	ADVOCATES, OR REALLY ANYONE IN THE GENERAL PUBLIC
4	THAT'S INTERESTED IN LEARNING MORE ABOUT CIRM.
5	SO REDESIGNING OUR WEBSITE WITH THOSE TWO
6	GROUPS IN MIND, WE BELIEVED IT WOULD HELP TO ATTRACT
7	TOP QUALITY APPLICANTS FROM INSTITUTIONS AND FROM
8	INDUSTRY AND INCREASE AWARENESS OF CIRM-FUNDED
9	RESEARCH, ULTIMATELY HELPING TO ACHIEVE OUR MISSION
10	OF ACCELERATING STEM CELL TREATMENTS TO PATIENTS IN
11	NEED.
12	WHEN WE BEGAN LOOKING AT HOW TO IMPROVE
13	THE SITE TO MEET THESE GOALS, WE FOCUSED ON THE
14	MAJOR PAIN POINTS WITH OUR CURRENT SITE. AS YOU CAN
15	SEE HERE FROM NOW OUR OLD SITE, THERE WAS
16	SIGNIFICANT UNUSED WHITE SPACE. FOR THE NEW VISITOR
17	IT WAS NOT CAPTIVATING. IT ALSO CAUSED PROBLEMS
18	WITH WHERE THE USER SHOULD GO TO LEARN MORE ABOUT
19	WHO WE ARE AND WHAT WE DO.
20	IN ADDITION, WHEN YOU DRILL DOWN TO THE
21	INDIVIDUAL SUBPAGES, THE HORIZONTAL DROP-DOWN WAS
22	DIFFICULT TO USE, AS YOU CAN SEE UP AT THE TOP. WE
23	ACTUALLY RECEIVED MANY COMPLAINTS ABOUT THIS. ME
24	PERSONALLY ALMOST EVERY DAY SINCE I STARTED WORKING
25	HERE. ALSO, INDIVIDUAL SUBPAGES WERE ALMOST
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1	ENTIRELY RELIANT ON TEXT, AS YOU CAN SEE HERE. THIS
2	IS DESPITE THE FACT THAT WE HAVE SEVERAL OR WE HAVE
3	MANY BEAUTIFUL SCIENTIFIC IMAGES AS WELL AS IMAGES
4	OF OUR PATIENTS AND ADVOCATES THAT REALLY WEREN'T
5	BEING PUT TO GOOD USE.
6	AND, FINALLY, A VERY IMPORTANT POINT, THE
7	MOBILE EXPERIENCE ON OUR SITE WAS AT BEST UNPLEASANT
8	AND IN MANY CASES MUCH WORSE. I'LL REFRAIN FROM
9	USING WORDS THAT I'VE HEARD ABOUT IT HERE. OUR
10	ANALYTICS ACTUALLY SHOW THAT 40 PERCENT OF USERS
11	WERE VISITING OUR SITE FROM A MOBILE OR TABLET
12	DEVICE, AND WE BELIEVED THAT THEY DESERVE TO HAVE AN
13	ENJOYABLE USER EXPERIENCE.
	SO WITH THAT IN MIND, WE BEGAN TO DEVELOP
14	
	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE
15	, and the second se
15 16	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE
15 16 17	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE THE USER EXPERIENCE EFFECTIVELY AND EFFICIENTLY. WE
15 16 17 18	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE THE USER EXPERIENCE EFFECTIVELY AND EFFICIENTLY. WE CONTRACTED WITH A BRAND-NEW UP AND COMING SMALL
15 16 17 18 19	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE THE USER EXPERIENCE EFFECTIVELY AND EFFICIENTLY. WE CONTRACTED WITH A BRAND-NEW UP AND COMING SMALL BOUTIQUE FIRM CALLED RADIANT DIGITAL THAT ARE BASED
15 16 17 18 19 20	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE THE USER EXPERIENCE EFFECTIVELY AND EFFICIENTLY. WE CONTRACTED WITH A BRAND-NEW UP AND COMING SMALL BOUTIQUE FIRM CALLED RADIANT DIGITAL THAT ARE BASED BOTH IN NEWPORT, CALIFORNIA, AND IN A NEW YORK TO
15 16 17 18 19 20	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE THE USER EXPERIENCE EFFECTIVELY AND EFFICIENTLY. WE CONTRACTED WITH A BRAND-NEW UP AND COMING SMALL BOUTIQUE FIRM CALLED RADIANT DIGITAL THAT ARE BASED BOTH IN NEWPORT, CALIFORNIA, AND IN A NEW YORK TO HELP WITH DESIGN AND DRUPAL DEVELOPMENT. WE ALSO
15 16 17 18 19 20 21	OUR PLAN LAST FALL AROUND OCTOBER, BEING TO IMPROVE THE USER EXPERIENCE EFFECTIVELY AND EFFICIENTLY. WE CONTRACTED WITH A BRAND-NEW UP AND COMING SMALL BOUTIQUE FIRM CALLED RADIANT DIGITAL THAT ARE BASED BOTH IN NEWPORT, CALIFORNIA, AND IN A NEW YORK TO HELP WITH DESIGN AND DRUPAL DEVELOPMENT. WE ALSO RELIED HEAVILY ON IN-HOUSE EXPERTISE FOR DRUPAL
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1	IMPROVED CIRM.CA.GOV. AS YOU CAN SEE THE VISUAL,
2	WE'VE MADE A NUMBER OF VISUAL IMPROVEMENTS TO THE
3	SITE, INCLUDING FEATURING ONE OF OUR PATIENT
4	ADVOCATES, DIANA SOUZA, ON OUR HOME PAGE. JUST
5	BELOW THE IMAGES, YOU WILL SEE CURRENT FUNDING
6	OPPORTUNITIES AND LEARN MORE ABOUT CIRM. SO THERE
7	ARE TWO MAIN AUDIENCES, CURRENT AND POTENTIAL
8	GRANTEES, AS WELL AS INTERESTED MEMBERS OF THE
9	PUBLIC CAN DISCOVER RIGHT AWAY WHO WE ARE AND WHAT
10	WE DO AND HOW THEY CAN APPLY FOR FUNDING.
11	YOU ALSO SEE A SPECIAL BANNER DEDICATED TO
12	CIRM 2.0 SO VISITORS CAN EXPLORE AND UNDERSTAND HOW
13	CIRM 2.0 WORKS.
14	WE'VE ALSO IMPROVED THE MAIN MENU
15	NAVIGATION TO A VERTICAL DROP-DOWN AS OPPOSED TO THE
16	HORIZONTAL FROM PREVIOUSLY. THIS SHOULD VASTLY
17	IMPROVE THE USER EXPERIENCE IN NAVIGATING THROUGHOUT
18	THE SITE.
19	WE MADE A NUMBER OF IMPROVEMENTS TO THE
20	EXISTING PAGES, AGAIN, TAKING ADVANTAGE OF THE VAST
21	ARRAY OF IMAGES AT OUR DISPOSAL. FOR EXAMPLE,
22	HERE'S OUR REVISED STEM CELL BASICS PAGE WHICH, WHEN
23	YOU COMPARE TO THE PREVIOUS ITERATION, OFFERS A MUCH
24	RICHER AND MORE ENGAGING USER EXPERIENCE.
25	WE'VE ALSO IMPROVED OUR FUNDING
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1	OPPORTUNITIES PAGE TO REFLECT THE CHANGES IN HOW
2	RESEARCHERS ARE APPLYING FOR GRANTS. WITH THE
3	ASSISTANCE OF BOB DEMETRIUS AND HIS TEAM IN LA'S
4	LSHD ADVERTISING, WHO ALSO WERE THE PEOPLE BEHIND
5	OUR NEW SUITE OF LOGOS, WE CREATED AN INTERACTIVE
6	IMAGE MAP THAT ALLOWS POTENTIAL GRANTEES TO CLICK ON
7	VARIOUS AWARD CATEGORIES. AT PRESENT, CLICKING ON
8	THE CLINICAL TAB HERE HIGHLIGHTED IN RED WILL TAKE
9	THE VISITOR TO THE CLINICAL STAGE PROGRAM
10	ANNOUNCEMENTS. AND WHEN THE ANNOUNCEMENTS FOR
11	DISCOVERY AND TRANSLATIONAL GO LIVE, THE BUTTONS
12	WILL TAKE THE GRANTEE OR POTENTIAL GRANTEE TO THOSE
13	PAGES AS WELL.
14	WE'VE ALSO USED THIS OPPORTUNITY TO CREATE
14	
15	PAGES THAT DID NOT EXIST ON THE OLD WEBSITE, BUT
	PAGES THAT DID NOT EXIST ON THE OLD WEBSITE, BUT THAT WE FEEL ARE VERY IMPORTANT FOR THE USER
15	
15 16	THAT WE FEEL ARE VERY IMPORTANT FOR THE USER
15 16 17	THAT WE FEEL ARE VERY IMPORTANT FOR THE USER EXPERIENCE. FOR EXAMPLE, WE CREATED A DEDICATED
15 16 17 18	THAT WE FEEL ARE VERY IMPORTANT FOR THE USER  EXPERIENCE. FOR EXAMPLE, WE CREATED A DEDICATED  LANDING PAGE FOR OUR NEW ALPHA CLINICS STEM CELL
15 16 17 18 19	THAT WE FEEL ARE VERY IMPORTANT FOR THE USER  EXPERIENCE. FOR EXAMPLE, WE CREATED A DEDICATED  LANDING PAGE FOR OUR NEW ALPHA CLINICS STEM CELL  NETWORK AND AS WELL A SERIES OF SUBPAGES THAT WILL
15 16 17 18 19	THAT WE FEEL ARE VERY IMPORTANT FOR THE USER EXPERIENCE. FOR EXAMPLE, WE CREATED A DEDICATED LANDING PAGE FOR OUR NEW ALPHA CLINICS STEM CELL NETWORK AND AS WELL A SERIES OF SUBPAGES THAT WILL HELP DESCRIBE THE PURPOSE AND MISSION OF THE ALPHA
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15 16 17 18 19 20 21 22	THAT WE FEEL ARE VERY IMPORTANT FOR THE USER  EXPERIENCE. FOR EXAMPLE, WE CREATED A DEDICATED  LANDING PAGE FOR OUR NEW ALPHA CLINICS STEM CELL  NETWORK AND AS WELL A SERIES OF SUBPAGES THAT WILL  HELP DESCRIBE THE PURPOSE AND MISSION OF THE ALPHA  CLINICS ALONG WITH LIVE LINKS TO EACH CENTER'S HOME  PAGE, CITY OF HOPE, UCSD, AND SO ON.  IN COORDINATION WITH THESE CENTERS, WE ARE

1	WEEKS.
2	WE'VE ALSO CREATED A PATIENT TOOLBOX,
3	HOUSING CONTENT FOR PATIENTS AND ADVOCATES TO GIVE
4	THEM THE TOOLS THEY NEED WHEN SPEAKING ABOUT CIRM IN
5	PUBLIC, SUCH AS DURING MEDIA INTERVIEWS OR
6	MODERATING A PANEL.
7	IMPORTANTLY, WE'VE CREATED A FULLY
8	RESPONSIVE MOBILE AND TABLET FAMILY VERSION OF OUR
9	SITE SO THAT NO MATTER WHERE YOU ARE AND WHAT YOUR
10	DEVICE, I'M INCLUDING IPADS, KINDLE FIRE, ANDROID
11	TABLETS AND IPHONES, YOU CAN HAVE AN ENJOYABLE USER
12	EXPERIENCE, WE HOPE.
13	AND FINALLY, I THINK THIS IS IMPORTANT
14	ESPECIALLY FOR THE PEOPLE IN THIS ROOM. THE NEW
15	WAYS TO NAVIGATE TO ANYTHING RELATED TO THE BOARD,
16	GOVERNING BOARD, MEETINGS, THINGS LIKE THAT, CAN
17	EITHER BE THROUGH THE EVENTS TAB, CIRCLED HERE, OR
18	UNDER GOVERNANCE AT THE TOP OF THE PAGE.
19	BUT OUR WORK IS NOT DONE. THERE ARE A
20	NUMBER OF ADDITIONAL IMPROVEMENTS THAT WE'RE ALREADY
21	WORKING ON IN THE NEXT FEW MONTHS, INCLUDING OUR TOP
22	PRIORITY, WHICH IS AN IMPROVED SEARCH. STAY TUNED
23	FOR THAT. WE'RE ALSO GOING TO BE INCREASING THE USE
24	OF DATA VISUALIZATION INFOGRAPHICS AND DECREASING
25	THE RELIANCE ON TEXT EVEN FURTHER TO HELP INFORM

1	PATIENTS, GRANTEES, EDUCATORS, AND OTHERS. WE ARE
2	ALSO WORKING ON SOME BEHIND-THE-SCENES FUNCTIONALITY
3	IMPROVEMENTS THAT YOU MAY NOT NOTICE, BUT WILL MAKE
4	THE OVERALL EXPERIENCE FASTER AND MORE EFFICIENT.
5	AND FINALLY IN THE BRIEF TIME I HAVE LEFT,
6	I WOULD LIKE TO THANK EVERYONE WHO PARTICIPATED IN
7	THIS DESIGN. IT REALLY WAS A TEAM EFFORT FROM
8	EVERYONE AT CIRM WHO'S LISTED HERE, ESPECIALLY THE
9	ONES LISTED HERE, RADIANT DIGITAL, ESPECIALLY TIM
10	HOBERT, WHO WORKED WITH OUR TEAM TO DEVELOP THE
11	PROJECT, AND LSHD ADVERTISING FOR THE BEAUTIFUL
12	IMAGE AND GRAPHICS.
13	I'M READY TO TAKE ANY QUESTIONS. GREAT.
14	CHAIRMAN THOMAS: ANNE, I THINK WE SPOKE
15	IN THE OFFICE A COUPLE DAYS AGO. I THINK IT'S A
16	WONDERFUL WEBSITE. YOU TOOK WHAT WAS ALREADY A
17	GREAT WEBSITE AND MADE IT EVEN BETTER IN KEEPING
18	WITH THE 2.0 THEME. AND CONGRATULATIONS ON A VERY
19	FINE PIECE OF WORK.
20	DR. HOLDEN: THANK YOU VERY MUCH. THANKS,
21	EVERYONE.
22	MR. MC CORMACK: I JUST WANTED TO ADD ONE
23	MORE NOTE, WHICH IS THAT ANNE HAS ONLY BEEN WITH US
24	FOR ABOUT A YEAR. IN THAT TIME SHE'S COMPLETELY
25	CHANGED THE WAY WE LOOK ONLINE. SHE, FIRST OF ALL,
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1	COMPLETELY UPDATED OUR BLOG TO MAKE IT LOOK
2	WONDERFUL, AND SHE'S DONE THE SAME WITH OUR WEBSITE.
3	SO SHE'S DONE AN EXTRAORDINARY JOB IN A VERY SHORT
4	TIME. AND SHE SHOWS GREAT SHE'S ALWAYS VERY CALM
5	DESPITE ALL THE FLURRY OF SUGGESTIONS THAT EVERYONE
6	IS COMING BY. EVERY TIME THEY GO BY HER DESK, THEY
7	SAY COULD WE DO, COULD WE, AND SHE NEVER GETS UPSET
8	OR ANYTHING. SHE'S JUST GREAT. SHE'S A JOY TO WORK
9	WITH.
10	I ALSO WANTED TO KIND OF EMPHASIZE THE
11	POINT SHE TALKED ABOUT, THE MOBILE DEVICE, BECAUSE
12	ABOUT 40 PERCENT OF OUR TRAFFIC RIGHT NOW COMES FROM
13	MOBILE DEVICES. AND THAT'S ONLY GOING TO GET MORE
14	IN THE FUTURE. SO TO BE ABLE TO LOOK AT OUR WEBSITE
15	IN A WAY THAT MAKES SENSE AND IS REALLY EASY TO
16	NAVIGATE ON A CELL PHONE OR A TABLET IS SUCH AN
17	IMPORTANT ADVANCE. SO I REALLY WANT TO AGAIN THANK
18	ANNE AND THE REST OF THE TEAM FOR THAT WORK.
19	MR. SHEEHY: THIS IS A LITTLE OFF TOPIC,
20	BUT IN THE LINE OF MEDIA, ONE OF THE THINGS THAT
21	I'VE NOTICED A LOT OF, JUST INSTANTLY IN THE LAST
22	TWO WEEKS, AN ISSUE THAT'S EXPLODED IS THE GENE
23	MODIFICATION OF GERM LINES WITH BOTH THE PUBLICATION
24	IN NATURE AND IN SCIENCE LED BY DR. BALTIMORE, OUR
25	FORMER COLLEAGUE, AND THE OTHER BY THE ALLIANCE FOR

1	REGENERATIVE MEDICINE. AND I NOTICE THAT WE HAVE A
2	STANDARDS WORKING GROUP COMING UP, AND WE HAVE BEEN
3	SUCH LEADERS, THOUGHT LEADERS, ACTUALLY I THINK WE
4	HAD THE FIRST OPERATIVE POLICIES IN PLACE FOR
5	EMBRYONIC STEM CELL RESEARCH WHEN WE FIRST DID OUR
6	STANDARDS WHEN WE FIRST STARTED TEN YEARS AGO.
7	SO I WONDER IF THERE'S A WAY OBVIOUSLY
8	WE DON'T HAVE AN ITEM ON OUR AGENDA, SO WE CAN'T
9	TAKE ANY FORMAL ACTION, BUT TO COMMUNICATE TO THE
10	CIRM TEAM, TO THE STANDARDS WORKING GROUP CHAIRS
11	THAT IT WOULD BE HELPFUL TO HEAR WHAT OUR POLICY IS
12	BECAUSE THIS IS BECOMING A BIG SUBJECT. I THINK
13	CLARITY ON HOW WE ARE WHAT OUR RULES ARE AROUND
14	THIS WOULD BE GOOD. SEE IF THERE ARE FURTHER POINTS
15	FOR DISCUSSION AND SEE IF WE NEED TO A TAKE
16	LEADERSHIP ON THIS BECAUSE I THINK, IF YOU'VE READ
17	THIS STUFF, THIS IS A HOT ISSUE.
18	ONE OF THE THINGS I'M FINDING IS THERE'S A
19	LOT OF CONFUSION IN THE COMMUNITY. SO A LOT OF
20	PEOPLE THINK THAT THIS IMPLIES THAT ALL GENETIC
21	MODIFICATION, INCLUDING THE GENE THERAPY STUDIES
22	THAT WE HAVE IN OUR PORTFOLIO, ARE NOW SUSPECT
23	BECAUSE THEY HAVE THE POTENTIAL TO PERMANENTLY ALTER
24	THE HUMAN GENOME. SO I THINK HAVING A DIALOGUE AND
25	DISCUSSION AT THE STANDARDS WORKING GROUP, IF
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1	THERE'S STILL TIME TO GET THAT ON THEIR AGENDA, I
2	THINK WOULD BE HELPFUL. AND WITH THE ETHICISTS WE
3	HAVE WHO HAVE BEEN PARTICIPATING IN THAT EFFORT
4	ENABLE US TO BE PART OF THE CONVERSATION AS IT MOVES
5	FORWARD.
6	I THINK THE PIECES THAT WERE WRITTEN ARE
7	VERY INTERESTING. I THINK IT'S AN IMPORTANT SOCIAL
8	TOPIC AS WE MOVE FORWARD WITH THESE NEW
9	TECHNOLOGIES. AND IF THERE'S SOME WAY WE CAN KIND
10	OF EXPRESS THAT, IF THE PRESIDENT AND WHOEVER I'M
11	TRYING TO PERSUADE TO GET THAT TO MOVE FORWARD, I
12	THINK IT WOULD BE AN INTERESTING THING TO HAVE AN
13	INTERESTING ROLE THAT CIRM CAN CONTINUE TO PLAY AND
14	HAS PLAYED IN THE PAST.
15	CHAIRMAN THOMAS: THANK YOU, JEFF. ANY
16	OTHER COMMENTS?
17	MR. ROWLETT: JUST IN GENERAL, THE
18	WEBSITE, REGARDING THAT, I FIND IT TO BE A
19	TREMENDOUS WORK IN A POSITIVE DIRECTION, USABILITY
20	AND JUST IN TERMS OF THE FLEXIBILITY OF THE WEBSITE.
21	I FIND IT TO BE FAR MORE INTUITIVE, AND THAT'S, I
22	THINK, IMPORTANT WHEN YOU SPEAK TO CONSTITUENTS.
23	AND THE OTHER THING THAT I WANT TO SAY IS
24	I APPRECIATE THE DIVERSITY THAT'S REFLECTED IN OUR
25	WEBSITE. I THINK THAT YOU TRIED TO INCORPORATE ALL
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OF CALIFORNIA, INCLUDING THE GREAT STATE OF LOS ANGELES. AND THOSE OF YOU WHO WERE IN THE OTHER ROOM KNOW WHAT I'M TALKING ABOUT. CHAIRMAN THOMAS: IS THAT ANOTHER THINLY VEILED DODGER/GIANT LINE? MR. ROWLETT: NO, MR. PRESIDENT, IT'S NOT. SO CONGRATULATIONS ON THE POSITIVE WORK IN PROGRESS. KEEP IT UP. AND I APPLAUD WHAT YOU'VE DONE SO FAR. MR. MC CORMACK: THANK YOU. CHAIRMAN THOMAS: OKAY. NEXT THIS IS, I THINK, A VERY IMPORTANT UPDATE. YOU MAY RECALL WE HAD A VERY LARGE INITIATIVE TWO OR THREE YEARS AGO IN THE INDUCED PLURIPOTENT STEM CELL BANKING SPACE. AND WE HAVE A PROGRAM THAT'S IN MIDSTRIDE RIGHT NOW. AND I ASKED DR. GRISHAMMER IF SHE WOULD SPEAK TO US AND GIVE US AN UPDATE ON WHAT'S GOING ON BECAUSE I'M SURE THERE ARE A LOT OF NEAT THINGS. SO DOCTOR. DR. GRISHAMMER: GOOD AFTERNOON AND THANKS. YES, I'M HERE TO UPDATE YOU ON THE HUMAN INDUCED PLURIPOTENT STEM CELL OR IPS CELL INITIATIVE. IN LINE WITH CIRM'S MISSION TO ACCELERATE STEM CELL TREATMENTS TO PATIENTS WITH UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS		
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AND WE HAVE A PROGRAM THAT'S IN MIDSTRIDE RIGHT NOW.  AND I ASKED DR. GRISHAMMER IF SHE WOULD SPEAK TO US  AND GIVE US AN UPDATE ON WHAT'S GOING ON BECAUSE I'M  SURE THERE ARE A LOT OF NEAT THINGS. SO DOCTOR.  DR. GRISHAMMER: GOOD AFTERNOON AND  THANKS. YES, I'M HERE TO UPDATE YOU ON THE HUMAN  INDUCED PLURIPOTENT STEM CELL OR IPS CELL  INITIATIVE. IN LINE WITH CIRM'S MISSION TO  ACCELERATE STEM CELL TREATMENTS TO PATIENTS WITH  UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS	12	HAD A VERY LARGE INITIATIVE TWO OR THREE YEARS AGO
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THANKS. YES, I'M HERE TO UPDATE YOU ON THE HUMAN INDUCED PLURIPOTENT STEM CELL OR IPS CELL INITIATIVE. IN LINE WITH CIRM'S MISSION TO ACCELERATE STEM CELL TREATMENTS TO PATIENTS WITH UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS	17	SURE THERE ARE A LOT OF NEAT THINGS. SO DOCTOR.
20 INDUCED PLURIPOTENT STEM CELL OR IPS CELL 21 INITIATIVE. IN LINE WITH CIRM'S MISSION TO 22 ACCELERATE STEM CELL TREATMENTS TO PATIENTS WITH 23 UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS	18	DR. GRISHAMMER: GOOD AFTERNOON AND
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22 ACCELERATE STEM CELL TREATMENTS TO PATIENTS WITH 23 UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS	20	INDUCED PLURIPOTENT STEM CELL OR IPS CELL
UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS	21	INITIATIVE. IN LINE WITH CIRM'S MISSION TO
, and the second	22	ACCELERATE STEM CELL TREATMENTS TO PATIENTS WITH
24 TO CREATE AN TRE CELL RANK WHICH IS IN ESSENCE A	23	UNMET MEDICAL NEEDS, THE GOAL OF THIS INITIATIVE IS
ZI TO CICATE AN ITO CLLE DANK, WITCH IS, IN ESSENCE, A	24	TO CREATE AN IPS CELL BANK, WHICH IS, IN ESSENCE, A
25 LIBRARY WHERE STEM CELLS ARE STORED AND DISTRIBUTED	25	LIBRARY WHERE STEM CELLS ARE STORED AND DISTRIBUTED
177		177

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1	TO USE THEM TO ACCELERATE THERAPEUTIC DISCOVERY.
2	THE BUDGET FOR THIS INITIATIVE IS A LITTLE
3	OVER \$32 MILLION. IN MARCH 2013 YOU APPROVED THE
4	GRANTS THAT MAKE UP THIS INITIATIVE. AND THEN IN
5	NOVEMBER 2013 ALL THE GRANTS THAT ARE PART OF THIS
6	INITIATIVE WERE LAUNCHED. SO THE FIRST FEW MINUTES
7	I'LL GO OVER AND REMIND YOU WHAT THIS INITIATIVE IS
8	ABOUT.
9	SO THERE ARE THREE DIFFERENT DISTINCT
10	ACTIVITIES THAT ARE FUNDED UNDER THIS INITIATIVE.
11	AND THESE ACTIVITIES ARE THE STEPS THAT ARE NEEDED
12	TO CREATE THIS RESOURCE. THE FIRST STEP IS CALLED
13	TISSUE COLLECTION WHERE PHYSICIAN SCIENTISTS RECRUIT
14	TISSUE DONORS WITH VARIOUS DISEASES WHO DONATE SOME
15	BLOOD OR A SMALL PIECE OF SKIN. THESE TISSUES ARE
16	THEN TRANSFERRED TO THE IPS CELL GENERATION FACILITY
17	WHERE THE IPS CELLS ARE GENERATED. AND THEN,
18	FINALLY, THE CELLS ARE STORED AND DISTRIBUTED FROM
19	THE IPS CELL BANK FOR RESEARCHERS AND DRUG
20	DEVELOPERS TO USE THESE CELLS TO MODEL DISEASES TO
21	BETTER UNDERSTAND HUMAN DISEASE AND TO ENGAGE IN
22	DRUG DISCOVERY AND DEVELOPMENT.
23	SO WHY CREATE SUCH AN IPS CELL BANK? THE
24	INTENDED USES OF THESE IPS CELLS FROM THIS BANK ARE
25	ILLUSTRATED ON THE SLIDE. SO AS AN EXAMPLE, IF YOU
	170

1	HAVE A PATIENT WITH NEURODEGENERATIVE DISEASE SUCH
2	AS ALZHEIMER'S OR PARKINSON'S DISEASE, YOU CAN ASK
3	THEM TO DONATE SOME BLOOD. THE BLOOD CELLS ARE
4	CONVERTED TO IPS CELLS, THEN STORED IN THE BANK.
5	AND THEN THE RESEARCHERS WHO THEN OBTAIN THESE CELLS
6	FROM THE IPS CELL BANK WILL EXPAND THEM FURTHER IN
7	THEIR LABORATORIES AND INDUCE THEM TO BECOME BRAIN
8	CELLS IN THIS CASE. AND SINCE THESE CELLS WERE
9	ORIGINALLY OBTAINED FROM A PATIENT WITH
10	NEURODEGENERATIVE DISEASE, THESE BRAIN CELLS IN THE
11	DISH MAY NOW DISPLAY SOME DEFECTS THAT RESEMBLE
12	THOSE SEEN IN THE PATIENT WITH NEURODEGENERATIVE
13	DISEASE AND CAN, THEREFORE, BE USED BY THE
14	RESEARCHER TO STUDY THE DISEASE.
15	IN ADDITION, DRUG DEVELOPERS CAN NOW USE
16	THESE SICK BRAIN CELLS IN THE DISH AND SCREEN LARGE
17	LIBRARIES OF POTENTIAL DRUG COMPOUNDS TO IDENTIFY
18	CANDIDATES THAT REVERT THE DISEASED CELLS BACK TO
19	NORMAL. IF YOU FIND A COMPOUND THAT DOES THAT, IT
20	CAN THEN BECOME A CANDIDATE DRUG THAT CAN MOVE
21	TOWARD AND INTO CLINICAL TRIAL.
22	NOW, IPS CELLS HAVE THREE COMPELLING
23	CHARACTERISTICS THAT MAKE THEM SO BROADLY APPLICABLE
24	TO STUDYING MANY DIFFERENT HUMAN DISEASES. THE
25	FIRST IS THAT THEY'RE DERIVED FROM TISSUES THAT CAN
	1=0

1	EASILY BE OBTAINED FROM LIVING PEOPLE, SUCH AS BLOOD
2	OR SMALL PIECE OF SKIN. THE IPS CELLS THEMSELVES
3	HAVE WHAT APPEAR TO BE AN UNLIMITED CAPACITY, THE
4	SECOND CHARACTERISTIC, AN UNLIMITED CAPACITY TO
5	DIVIDE AND PROLIFERATE IN CULTURE, WHICH MEANS THAT
6	HUGE AMOUNTS OF CELLS CAN BE EASILY PRODUCED. AND
7	THEN, FINALLY, THE THIRD CHARACTERISTIC IS THAT,
8	LIKE EMBRYONIC STEM CELLS, IPS CELLS ARE
9	PLURIPOTENT. THAT'S WHERE THEIR NAME COMES FROM.
10	PLURIPOTENT MEANS THAT THEY HAVE THE ABILITY TO FORM
11	THE CELLS OF ANY ORGAN OR TISSUE IN YOUR BODY, SUCH
12	AS HEART, GUT, PANCREAS, LIVER, OR BRAIN.
13	SO, THEREFORE, THIS TECHNOLOGY, THIS IPS
14	CELL TECHNOLOGY, MAKES IT POSSIBLE FOR RESEARCHERS
15	AND DRUG DEVELOPERS TO STUDY HEART CELLS FROM
16	PATIENTS WITH HEART DISEASE OR BRAIN CELLS FROM
17	PATIENTS BRAIN DISEASE; IN OTHER WORDS, THIS
18	TECHNOLOGY, THE IPS CELL GENERATION, IS A MEANS TO
19	BE ABLE TO ACCESS PREVIOUSLY INACCESSIBLE OR HARD TO
20	ACCESS TISSUES FROM CELL TYPES FROM PATIENTS TO
21	CREATE LARGE QUANTITIES OF THESE CELLS FOR STUDY.
22	NOW, IN THE IPS CELL INITIATIVE THAT IS
23	FUNDED BY CIRM, WE ARE INCLUDING DISEASES OF THE
24	BRAIN, THE HEART, THE LIVER, THE LUNG, AND THE EYE.
25	THE SPECIFIC CONDITIONS ARE LISTED HERE AND SO ARE
	100

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1	THE CLINICIAN SCIENTISTS WHO ARE RECRUITING THE
2	PATIENTS TO PARTICIPATE IN THIS STUDY. THEY ALSO
3	RECRUIT HEALTHY INDIVIDUALS WHOSE IPS CELLS WILL
4	SERVE AS NORMAL CONTROLS. AND SO THOSE ARE THE
5	INDIVIDUALS WHO COLLECT THE BLOOD OR SKIN PIECES.
6	THE TISSUES ARE THEN TRANSFERRED TO THE
7	IPS CELL GENERATION FACILITY WHICH IS OPERATED BY A
8	COMPANY CALLED CELLULAR DYNAMICS INTERNATIONAL OR
9	CDI. THEY ARE A COMPANY THAT PROVIDES IPS CELLS AND
10	MATURE CELLS DERIVED FROM THESE IPS CELLS TO
11	RESEARCHERS AND THE PHARMACEUTICAL INDUSTRY. SO
12	THEY'RE WELL-SUITED FOR THIS TASK OF GENERATING
13	INDEED IPS CELLS FROM 3,000 TISSUE DONORS, WHICH IS
14	WHAT THEY HAVE SIGNED UP TO DO.
15	ONCE THE IPS CELLS ARE GENERATED, THEY ARE
16	TRANSFERRED TO THE BANK, WHICH IS OPERATED BY
17	CORYELL. THAT'S A NONPROFIT ORGANIZATION THAT HAS
18	DECADES OF EXPERIENCE IN OPERATING CELL BANKS.
19	NOW, CDI AND CORYELL ARE NOT LOCATED IN
20	CALIFORNIA, BUT THEY ESTABLISHED FACILITIES HERE IN
21	CALIFORNIA AT THE BUCK INSTITUTE UP IN NOVATO TO
22	EXECUTE THEIR GRANTS.
23	SO ON MY LAST SLIDE, I WANT TO TELL YOU
24	WHERE THIS INITIATIVE STANDS. AS I MENTIONED ON MY
25	FIRST SLIDE, THE AWARDS WERE ALL LAUNCHED BY
	181

1	NOVEMBER 2013. THE FIRST FEW MONTHS WERE SPENT FOR
2	THE PROCESSES THAT WERE NECESSARY TO COORDINATE THE
3	DIFFERENT ACTIVITIES TO BE WORKED OUT AND TO
4	ESTABLISH INDEED THE IPS CELL GENERATION AND BANKING
5	OPERATIONS.
6	THE TISSUE COLLECTION FROM PATIENTS
7	STARTED IN MARCH 2014. AND AS OF THIS MONTH MORE
8	THAN 1,000 SAMPLES FROM PATIENTS HAVE BEEN COLLECTED
9	AND TRANSFERRED TO THE BUCK INSTITUTE. AND IPS
10	CELLS FROM MORE THAN 350 INDIVIDUALS HAVE BEEN
11	GENERATED AND VALIDATED.
12	THE NEAR-TERM MILESTONES ARE TO HAVE IPS
13	CELLS FROM 750 DONORS GENERATED BY THE END OF MAY,
14	AND THEN THE BANK IS SLATED TO OPEN FOR BUSINESS BY
15	THE END OF AUGUST WHEN THE IPS CELLS FROM THE FIRST
16	300 DONORS WILL BECOME AVAILABLE FROM THE BANK.
17	THE LAST THREE TIME POINTS SHOW THE SLATED
18	ENDPOINTS FOR THE THREE ACTIVITIES I WAS DESCRIBING.
19	AND WITH THE IPS CELLS FROM ALL DONORS BECOMING
20	AVAILABLE BY THE END OF 2017.
21	SO I WANT TO STOP HERE. I FORGOT TO
22	MENTION THE ACTIVITIES THAT ARE INVOLVED HERE NEED
23	TO BE COORDINATED, AND CIRM IS ACTUALLY ACTIVELY
24	INVOLVED IN HELPING COORDINATE THESE VARIOUS
25	ACTIVITIES. I WANT TO THANK MICHAEL YAFFE WHO HAS
	182

1	BEEN IN CHARGE OF THIS INITIATIVE UNTIL RECENTLY.
2	AND DUE TO THE REORGANIZATION WE RECENTLY HAD AT
3	CIRM, I AM NOW IN CHARGE OF THIS INITIATIVE. AND
4	I'M HAPPY TO TAKE ANY QUESTIONS.
5	MR. SHEEHY: SURE. SO THIS LOOKS GREAT.
6	I HAD A QUESTION. AND THIS ACTUALLY CAME UP IN A
7	PRESENTATION THAT I WAS AT WITH SOMEONE WHO IS
8	HEAVILY INVOLVED IN THE FORMATION OF THE EUROPEAN
9	BANK FOR INDUCED PLURIPOTENT STEM CELLS. AND ONE
10	FEATURE THAT THEY HAD THAT WE DID NOT INCLUDE, WHICH
11	I'M ACTUALLY BEGINNING TO THINK IS A GAP, BECAUSE
12	THEY ACTUALLY PARTNERED WITH A BIG PHARMA, WITH
13	PFIZER. AND WHAT THAT DID OBVIOUSLY WAS STRETCH
14	THEIR RESOURCES MUCH FURTHER, BUT IT ALSO BROUGHT
15	CONCRETE EXPERTISE IN TARGETING AND DRUG
16	DEVELOPMENT. THE TWO PIECES OF IT, OF THAT FIRST
17	SCHEME OF IT, ARE KIND OF LAYING OUT THERE.
18	I WONDER IF ANY THOUGHT HAS BEEN GIVEN TO
19	ACTUALLY SEEKING AN INDUSTRY PARTNER IN ORDER BOTH
20	TO MAKE THIS MORE ECONOMICALLY RATIONAL FOR US, BUT
21	ALSO TO MAKE IT MORE EFFICIENT IN THAT YOU BRING IN
22	EXPERTISE THAT REALLY ALREADY EXISTS FOR TARGETING
23	AND DRUG DEVELOPMENT.
24	IT JUST SEEMS LIKE WE'RE MISSING A PIECE
25	ON THAT END OF THE OUR PROPOSAL LOOKS GREAT.
	100

1	WE'RE GOING TO HAVE ALL THESE LINES, BUT REALLY WHAT
2	HAPPENS TO THEM, WHO'S VESTED IN REALLY DEVELOPING
3	THE PRODUCTS OUT OF THOSE LINES? AND, AGAIN, THE
4	EUROPEAN EXAMPLE, I HAVE THEIR WEBSITE UP, WORKING
5	CLOSELY WITH A BIG PHARMA COMPANY MAY ADVANTAGE THEM
6	GREATLY.
7	DR. GRISHAMMER: IF I CAN MAKE TWO
8	COMMENTS. ONE IS, FIRST, YES, I ACKNOWLEDGE THAT
9	THERE WAS NO MOVE TO INCLUDE PHARMACEUTICAL FUNDING
10	IN THIS INITIATIVE. BUT TWO COMMENTS. ONE THING
11	THAT WE DID DO WITH FORESIGHT OVER THE FACT WE DO
12	WANT THESE IPS CELLS NOT ONLY TO BE USED FOR
13	ACADEMIC RESEARCHERS, BUT ALSO BY THE PHARMACEUTICAL
14	INDUSTRY FOR DRUG DISCOVERY, IT ACTUALLY WAS ELONA
15	BAUM WHO REALLY DROVE THAT AT THE TIME, WAS THINK
16	THROUGH HOW WE COULD ENSURE THAT THE CELLS THAT ARE
17	DEPOSITED HERE WOULD BE AS WIDELY AVAILABLE AND
18	EASILY AVAILABLE TO EVERYBODY, INCLUDING THE
19	PHARMACEUTICAL INDUSTRY.
20	AND SO ACTUALLY, AS PART OF THE RFA AND
21	REVIEW PROCESS, WE ASKED ALL THE APPLICANTS TO TELL
22	US ABOUT THEIR IP POSITION IN TERMS OF THEIR OWN
23	PATENTS AND LICENSES THEY HAD THAT WOULD ALLOW
24	THAT ARE FOR THE COMMERCIAL USE OF THESE CELL LINES.
25	AND SO WHEN CDI WAS SELECTED, THEY DO HAVE
	18/

1	A VERY STRONG PORTFOLIO OF PATENTS SURROUNDING IPS
2	CELL TECHNOLOGY. AND WHAT WE WILL BE ABLE TO OFFER
3	TO THE PHARMACEUTICAL INDUSTRY IS CELL LINES FROM
4	3,000 DONORS WHICH HAVE CLARITY AROUND THE IP THAT
5	EXISTS FOR THE VERY CELL LINES THAT THEY'RE BUYING,
6	WHICH MEANS THAT AT THAT POINT THEY CAN EITHER BUY
7	THOSE LICENSES THAT HAVE ALREADY BEEN NEGOTIATED AND
8	THEY KNOW EXACTLY WHAT THE PACKAGE LOOKS LIKE, BUT
9	THEY DON'T HAVE TO. THEY CAN GO AND NEGOTIATE THEIR
10	OWN LICENSES. SO IT'S A VERY DIFFERENT SUBJECT
11	MATTER THAN YOU SAID, BUT WE WERE DEFINITELY TRYING
12	TO FORESEE AND MAKE EASY THE USE OF THESE CELLS BY
13	INDUSTRY.
14	AND THE SECOND POINT I WANT TO MAKE IS
15	THAT ACTUALLY CDI, CORYELL, AND CIRM ARE GETTING
16	TOGETHER RIGHT NOW TO DESIGN OUR ADVERTISING
17	ACTIVITY AROUND THIS ACTIVITY WITH THE BANK OPENING
18	IN AUGUST. SO THAT OBVIOUSLY BOTH CDI AND CORYELL
19	ARE VERY INTERESTED THAT THESE CELLS WILL BE WIDELY
20	USED, INCLUDING BY INDUSTRY. SO THERE IS AN EFFORT
21	TO HAVE AN ADVERTISING CAMPAIGN FOR THE EXISTENCE OF
22	THIS BANK.
23	MR. SHEEHY: BUT WOULDN'T IT BE AN
24	INTERESTING THOUGHT PROCESS TO CONSIDER WHETHER IT
25	MAKES SENSE LOOKING FOR A SPECIFIC INDUSTRY PARTNER
	105

1	THAT WOULD COMMIT TO DEVELOPING PRODUCTS BASED ON
2	THESE LINES? BECAUSE WE WILL HAVE ALREADY DONE THE
3	FRONT-END INVESTMENT. I WONDER IF YOU MAKE THEM
4	AVAILABLE TO A WHOLE HOST OF PEOPLE WHAT BECOMES
5	ARE THEY GOING TO COMPETE WITH EACH OTHER USING
6	THESE CELLS? IS THAT SOMETHING WE ANTICIPATE
7	HAPPENING, DIFFERENT PHARMACEUTICAL COMPANIES USING
8	THESE CELLS AND THEN RUNNING IT THROUGH THEIR
9	TARGETING? IT'S NOT CLEAR TO ME. I'M NOT A
10	BUSINESS PERSON, SO I DON'T KNOW HOW THE BUSINESS
11	MODEL SYNCS UP.
12	EUROPEANS THE PERSON WHO WAS DISCUSSING
13	EUROPEAN EXPERIENCE THOUGHT THAT THEY WOULD GET A
14	BETTER YIELD BY PARTNERING DIRECTLY WITH ONE COMPANY
15	WITH AN INTEREST IN DEVELOPING PRODUCTS IN THE AREAS
16	THAT THEY WERE BANKING.
17	DR. GRISHAMMER: THAT'S CERTAINLY
18	SOMETHING WE CAN TALK ABOUT AND THINK THROUGH.
19	CHAIRMAN THOMAS: UTA, HOW DOES OUR IPS
20	BANK STACK UP AGAINST OTHER IPS BANKS IN TERMS OF
21	SIZE, COMPREHENSIVENESS, AND ALL THAT SORT OF THING?
22	DR. GRISHAMMER: SO IN TERMS OF SIZE, IT
23	DEPENDS A LITTLE WHICH PUBLICATION YOU READ OR WHO
24	YOU TALK TO. THERE ARE SEVERAL EFFORTS OUT THERE TO
25	BE CREATING AN IPS CELL BANK FOR VARIOUS DIFFERENT
	186

1	DISEASES. AND I WON'T THROW OUT NUMBERS JUST
2	BECAUSE I DO HEAR DIFFERENT NUMBERS FROM DIFFERENT
3	PEOPLE, AND I DON'T KNOW WHAT THE REALITY ULTIMATELY
4	WILL BE FOR THE VARIOUS BANKS. SO THIS IS NOT THE
5	ONLY LARGE BANK. THREE THOUSAND TISSUE DONORS
6	INCLUDED IN A BANK LIKE THIS IS CONSIDERED
7	DEFINITELY A LARGE-SCALE BANK. THE NIH, THE
8	WELLCOME TRUST, THE EUROPEAN EFFORTS, THE NEW YORK
9	STEM CELL FOUNDATION, JAPAN ALL HAVE ACTIVITIES IN
10	THIS REGARD AS WELL.
11	IN TERMS OF THE DISEASES THAT ARE COVERED,
12	WE TRY TO ACTUALLY MAKE AN EFFORT TO MAKE SURE THAT
13	THERE WOULDN'T BE TOO MUCH REPETITION. WE ACTUALLY
14	EXCLUDED CERTAIN DISEASES COMING IN FROM THE GET-GO
15	BECAUSE CIRM WAS ALREADY PARTICIPATING IN AN
16	NIH-FUNDED PROCESS FOR FUNDING IPS CELL BANKING FOR
17	SEVERAL NEUROLOGICAL DISEASES.
18	IN ADDITION, WE ASKED OUR APPLICANTS TO
19	TALK ABOUT THE COMPETITIVE OF THEIR DISEASE AS BEING
20	PART OF AN IPS CELL BANK IN TERMS OF WHAT WAS KNOWN
21	AT THE TIME OF WHAT EVERYBODY WAS GENERATING, BUT IT
22	WAS REALLY NOT THAT KNOWN. SO REALLY THE COMMENT I
23	WANT TO MAKE IS THAT THERE WILL BE PROBABLY OVERLAP
24	BETWEEN THE CIRM BANK AND SOME OTHER BANKS IN TERMS
25	OF AN EXACT TYPE OF DISEASE THAT IS GOING TO BE

1	INCLUDED, BUT I THINK ACTUALLY THAT COULD BE TO THE
2	BENEFIT FOR THE RESEARCH COMMUNITY WHERE PEOPLE OF
3	DIFFERENT ETHNICITIES, FOR INSTANCE, WOULD BE
4	INCLUDED IN ONE BANK, LET'S SAY, WITH ALZHEIMER'S
5	DISEASE VERSUS OUR BANK WITH ALZHEIMER'S DISEASE.
6	CHAIRMAN THOMAS: THANK YOU. ANY OTHER
7	COMMENTS BY MEMBERS OF THE BOARD? THANK YOU VERY
8	MUCH FOR THAT UPDATE.
9	DO WE HAVE ANY REMAINING PUBLIC COMMENT ON
10	ANY PARTICULAR TOPICS? OKAY.
11	HEARING NONE, COUPLE OF CLOSING COMMENTS.
12	NO. 1, I'D LIKE TO CONGRATULATE SCOTT TOCHER FOR
13	YESTERDAY GETTING HIS PILOT'S LICENSE.
14	(APPLAUSE.)
15	CHAIRMAN THOMAS: HE JOINS DR. MILLS AS, I
16	BELIEVE, THE ONLY TWO PILOTS INVOLVED IN THIS
17	OPERATION.
18	DR. MILLS: WE'RE STARTING AN AIRLINE.
19	CHAIRMAN THOMAS: AND SHOULD YOU HAVE
20	PROBLEMS, AS YOU ALWAYS DO, GETTING IN OR OUT OF
21	SFO, YOU MIGHT CHOOSE TO GO FLY AIR TOCHER AND AIR
22	MILLS.
23	MY SECOND COMMENT IS AND CLOSING COMMENT,
24	WITH OPENING DAY TWO WEEKS AGO OR TWO WEEKS FROM NOW
25	AND HAVING WEATHERED AND SUFFERED THROUGH THREE
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GIANTS WORLD CHAMPIONSHIPS IN THE LAST FIVE YEARS,
 1
 2
      ENOUGH ALREADY. GO DODGERS. WE STAND ADJOURNED.
 3
                MS. BONNEVILLE: OUR MAY BOARD MEETING
     WILL NOT BE IN SAN DIEGO AS IT SAYS ON THE SCHEDULE.
 4
 5
     THAT WILL BE IN SEPTEMBER. SO THE NEXT BOARD
 6
     MEETING WILL BE IN THE BAY AREA.
 7
                CHAIRMAN THOMAS: THANK YOU, EVERYBODY.
 8
     WE WILL SEE YOU IN MAY.
 9
                     (THE MEETING WAS THEN CONCLUDED AT
10
     02:49 P.M.)
11
12
13
14
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16
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#### REPORTER'S CERTIFICATE

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 THE INDEPENDENT CITIZEN'S OVERSIGHT COMMITTEE OF THE CALIFORNIA INSTITUTE FOR REGENERATIVE MEDICINE IN THE MATTER OF ITS REGULAR MEETING HELD AT THE LOCATION INDICATED BELOW

THE CLAREMONT HOTEL
44 TUNNEL ROAD
BERKELEY, CALIFORNIA
ON
MARCH 26, 2015

WAS HELD AS HEREIN APPEARS AND THAT THIS IS THE ORIGINAL TRANSCRIPT THEREOF AND THAT THE STATEMENTS THAT APPEAR IN THIS TRANSCRIPT WERE REPORTED STENOGRAPHICALLY BY ME AND TRANSCRIBED BY ME. I ALSO CERTIFY THAT THIS TRANSCRIPT IS A TRUE AND ACCURATE RECORD OF THE PROCEEDING.

BETH C. DRAIN, CSR 7152 BARRISTERS' REPORTING SERVICE 160 S. OLD SPRINGS ROAD SUITE 270 ANAHEIM, CALIFORNIA (714) 444-4100