# BEFORE THE SCIENCE SUBCOMMITTEE OF THE INDEPENDENT CITIZENS' OVERSIGHT COMMITTEE TO THE

# CALIFORNIA INSTITUTE FOR REGENERATIVE MEDICINE ORGANIZED PURSUANT TO THE CALIFORNIA STEM CELL RESEARCH AND CURES ACT

#### REGULAR MEETING

LOCATION: AS INDICATED ON THE AGENDA

DATE: TUESDAY, APRIL 26, 2011

2 P.M.

REPORTER: BETH C. DRAIN, CSR

CSR. NO. 7152

BRS FILE NO.: 89802

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4. DISCUSSION AND CONSIDERATION OF NOT PROCESS FOR OBTAINING SUPPLEMENTAL INFORMATION OF APPLICANTS FOR CLINICAL TRIAL AND DISEATEAM GRANT ROUNDS, INCLUDING AN OPPORTUNITY TO OBTAIN INFORMATION DURING PEER REVIEW, SUBJECT TO LATER STAFF CONFIRMATION, OF ADDITIONAL DATA NOT PRESENTED IN THE APPLICATION.	ON ASE
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1	TUESDAY, APRIL 26, 2011
2	02:03 P.M.
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3	_
4	CHAIRMAN SHEEHY: I'D LIKE TO CALL THE
5	MEETING TO ORDER. MELISSA, WOULD YOU LIKE TO CALL
6	THE ROLL.
7	MS. KING: I WILL. SUE BRYANT.
8	DR. BRYANT: HERE.
9	MS. KING: MARCY FEIT. MICHAEL FRIEDMAN.
10	DR. FRIEDMAN: HERE.
11	MS. KING: BOB KLEIN.
12	CHAIRMAN KLEIN: HERE.
13	MS. KING: FRANCISCO PRIETO. PHIL PIZZO.
14	DUANE ROTH.
15	MR. ROTH: HERE.
16	MS. KING: JOAN SAMUELSON.
17	MS. SAMUELSON: HERE.
18	MS. KING: JEFF SHEEHY.
19	CHAIRMAN SHEEHY: HERE.
20	MS. KING: OSWALD STEWARD. AND ART
21	TORRES.
22	MR. TORRES: HERE.
23	MS. KING: OKAY. AND, JAMES HARRISON, ARE
24	YOU ON THE LINE?
25	MR. HARRISON: I AM.
	3

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1	MS. KING: JUST TO GO OVER WHO'S HERE WITH
2	ME IN THE ROOM IN SAN FRANCISCO, WE HAVE A MEMBER OF
3	THE PUBLIC, JUDY ROBERSON. WHO JUST JOINED THE
4	CALL?
5	DR. STEWARD: THIS IS OS STEWARD.
6	MS. KING: GREAT. THANK YOU SO MUCH, DR.
7	STEWARD. I'M JUST GOING THROUGH THE ROLL. SO
8	PERFECT TIMING. AND WHO'S WITH ME HERE IN SAN
9	FRANCISCO AT THE CIRM. SO I HAVE JEFF SHEEHY AND
10	ART TORRES WITH ME AS WELL AS LYNN HARWELL, GEOFF
11	LOMAX, PATRICIA OLSON, CYNTHIA SCHAFFER, ELONA BAUM,
12	AGAIN JUDY ROBERSON, MEMBER OF THE PUBLIC, GIL
13	SAMBRANO, ELLEN FIEGAL, AND LAST, BUT NOT LEAST, OUR
14	PRESIDENT, ALAN TROUNSON, I BELIEVE WEARING COWBOY
15	BOOTS. AM I RIGHT? NO. SORRY ABOUT THAT.
16	OKAY. SO THAT'S WHO'S WITH US HERE IN SAN
17	FRANCISCO. DO WE HAVE MEMBERS OF THE PUBLIC AT ANY
18	OF OUR OTHER SITES?
19	MR. THOMAS: THIS IS JOHN THOMAS ON THE
20	LINE.
21	MS. KING: THANK YOU. DO WE HAVE ANY
22	OTHER MEMBERS OF THE PUBLIC?
23	MS. FEIT: THIS IS MARCY FEIT CHECKING IN,
24	AND I HAVE NO ONE HERE.
25	MS. KING: GREAT. THANK YOU SO MUCH,
	4

1	MARCY. AND AS YOU CAN PROBABLY TELL, WE'RE JUST
2	GETTING STARTED. WE JUST WENT THROUGH A ROLL CALL,
3	AND WITH YOU I BELIEVE WE HAVE A QUORUM. SO THANK
4	YOU. I WILL TURN IT OVER TO OUR ILLUSTRIOUS CHAIR,
5	MR. JEFF SHEEHY.
6	CHAIRMAN SHEEHY: SO I'M GOING TO CHANGE
7	THE ORDER OF THE AGENDA, IF THAT'S OKAY. FIRST OF
8	ALL, ITEM NO. 3, DISCUSSION OF EXTRAORDINARY
9	PETITIONS PROCESS, I THINK THAT THAT'S JUST AN
10	ARTIFACT THAT GOT CARRIED FORWARD. NO ONE HAS BEEN
11	ABLE TO IDENTIFY WHAT ISSUES REMAIN. IT SEEMS LIKE
12	THAT THE APPEAL FOR ADDITIONAL ANALYSIS, WE USED
13	THAT PROCESS AT THE LAST MEETING AND THAT WORKED
14	WELL. AND SO I DON'T THINK THAT THERE'S ANY OTHER
15	TWEAKING WE NEED TO DO FOR RIGHT NOW.
16	ITEM NO. 4, STAFF IS STILL WORKING ON
17	THAT. AND SO WE DECIDED TO PUT THAT FORWARD TO
18	HOPEFULLY ANOTHER SCIENCE SUBCOMMITTEE MEETING IN
19	JUNE WHERE WE CAN HAVE A FULL DISCUSSION OF THOSE
20	ISSUES. IT'S A LITTLE BIT PREMATURE TO BE TALKING
21	ABOUT THAT UNTIL WE HAVE ALL THE TILL STAFF IS
22	READY FOR US TO WORK ON SOMETHING.
23	SO WHAT I'D LIKE TO DO IS GO STRAIGHT TO
24	ITEM NO. 6, WHICH IS THE ONE THAT IS PROBABLY MOST
25	TIMELY AND MOST IMMEDIATE, WHICH IS CONSIDERATION OF

1	THE IPSC REPOSITORY. THERE'S A DOCUMENT THAT
2	ACCOMPANIES THIS ITEM YOU SHOULD HAVE. AND I THINK
3	DR. FEIGAL WILL CONDUCT LEAD US THROUGH THIS.
4	DR. FEIGAL: OKAY. THANKS VERY MUCH. AND
5	I HOPE YOU ALL HAD THE OPPORTUNITY TO READ THE
6	DOCUMENT, BUT I WILL TRY AND SUMMARIZE IT REALLY
7	BRIEFLY. SO THANKS FOR THE OPPORTUNITY TO PRESENT
8	THIS, AND THANKS FOR THE MEMBERS OF THE PUBLIC THAT
9	JOINED US TODAY ON THIS IMPORTANT TOPIC.
10	SO CIRM IS REALLY PROPOSING A TWO-STEP
11	CONCEPT TO DEVELOP INDUCED PLURIPOTENT STEM CELL
12	RESEARCH RESOURCES TO FACILITATE THE SCIENTIFIC
13	EVALUATION OF THESE TYPES OF CELLS FOR DRUG
14	DISCOVERY AND DISEASE MODELING PURPOSES.
15	THE FIRST STEP OF THIS CONCEPT IS TO
16	PROVIDE FUNDING IN SUPPORT OF THE INDUCED
17	PLURIPOTENT STEM CELL CONSORTIA THAT IS FOCUSED ON
18	NEURODEGENERATIVE DISEASES THAT'S CONDUCTED BY THE
19	NATIONAL INSTITUTE OF NEUROLOGICAL DISORDERS AND
20	STROKE THROUGH A PUBLIC/PRIVATE PARTNERSHIP
21	COORDINATED BY THE FOUNDATION FOR THE NATIONAL
22	INSTITUTES OF HEALTH. AND THE FOUNDATION IS ALSO A
23	NONPROFIT ORGANIZATION.
24	THE CIRM COMPONENT OF THE NIH FUNDING
25	WOULD BE APPROXIMATELY A HUNDRED FIFTY THOUSAND PER

1	YEAR FOR TWO YEARS OUT OF A TOTAL OF 4.5 MILLION PER
2	YEAR FROM NINDS, WITH THE SECOND YEAR FUNDING
3	DEPENDING ON NINDS APPROPRIATION. AND THIS WOULD
4	BE OUR PART OF THE FUNDING WOULD BE PROVIDED TO
5	MERITORIOUSLY REVIEWED APPLICANTS FROM CALIFORNIA.
6	THE SECOND STEP OF THE CONCEPT WOULD BE
7	FOR CIRM TO PROVIDE SIMILAR RESEARCH RESOURCES IN
8	OTHER DISEASE AREAS POTENTIALLY IN COLLABORATION
9	WITH OTHER PARTNERS. AND THIS SECOND STEP OF THE
10	CONCEPT WILL BE BROUGHT TO THE SCIENCE COMMITTEE,
11	THE GWG, AND THE ICOC SOMETIME LATER THIS YEAR OR
12	EARLY NEXT.
13	SO LET'S GO BACK TO THE FIRST STEP OF THIS
14	PROPOSAL, WHICH IS A PROJECT DESIGN TO DEVELOP
15	WELL-CHARACTERIZED, PUBLICLY AVAILABLE INDUCED
16	PLURIPOTENT STEM CELL LINES FOR SCIENTIFIC
17	EVALUATION OF DISEASE MODELING AND DRUG DISCOVERY IN
18	THE NEURODEGENERATIVE DISEASES OF PARKINSON'S
19	DISEASE, HUNTINGTON'S DISEASE, AND AMYOTROPHIC
20	LATERAL SCLEROSIS.
21	THERE ARE THREE CONSORTIA THAT WERE
22	INITIALLY LAUNCHED AND FUNDED BY NINDS, THE NATIONAL
23	INSTITUTE OF NEUROLOGIC DISORDERS AND STROKE BACK IN
24	2009, AND THEY WERE FOCUSED ON DEVELOPING
25	WELL-CHARACTERIZED, PUBLICLY AVAILABLE IPS CELL
	<u>_</u>

1	LINES FOR THESE DIFFERENT FORMS OF NEURODEGENERATIVE
2	DISEASES. THIS CONSORTIUM APPROACH ENABLED RAPID
3	RESOURCE AND ANALYTICAL TOOL DEVELOPMENT AND THE
4	INITIAL IDENTIFICATION OF CELLULAR PHENOTYPES THAT
5	WERE ASSOCIATED WITH LATE ONSET NEURODEGENERATIVE
6	DISEASE IN IPS CELL-DERIVED NEURONAL CULTURES.
7	ALL THE FIBROBLAST LINES AND THE IPS CELL
8	LINES DEVELOPED THROUGH THIS CONSORTIA WILL BE MADE
9	AVAILABLE THROUGH THE REPOSITORY AT CORIELL.
10	IN 2010, LAST YEAR, NINDS HELD TWO
11	WORKSHOPS FOR THE CONSORTIA, INVESTIGATORS,
12	NONGOVERNMENT ORGANIZATIONS, AND INDUSTRY
13	REPRESENTATIVES TO DISCUSS THE PROGRESS OF THE
14	CONSORTIA IN DEVELOPING IPS CELL DISEASE-SPECIFIC
15	LINES AND PROTOCOLS AND METHODOLOGIES FOR CELL TYPE
16	SPECIFIC DIFFERENTIATION AND LINEAGE ANALYSIS.
17	THE WORKSHOPS ALSO PROVIDED INDUSTRY
18	PERSPECTIVES REGARDING THE CHALLENGES THAT REMAIN
19	FOR THE UTILIZATION OF THE PATIENT-DERIVED IPS CELL
20	LINE IN THE DRUG DEVELOPMENT PROCESS. TODAY THE
21	CONSORTIA HAS DEVELOPED MORE THAN 87 FIBROBLAST
22	LINES IN THESE DIFFERENT NEURODEGENERATIVE DISEASES
23	AND 25 IPS CELL LINES.
24	THIS PROPOSAL WOULD EXTEND THE LIFE OF THE
25	IPS CONSORTIA FOR AN ADDITIONAL TWO YEARS WORKING IN

1	PARTNERSHIP WITH THE FOUNDATION FOR NIH AND OTHER
2	PUBLIC/PRIVATE PARTNERS IN WHICH CIRM WOULD BE ONE
3	OF THOSE PARTNERS IN ORDER TO PROMOTE THE CONTINUED
4	DEVELOPMENT OF THIS RICH RESEARCH RESOURCE, WHICH
5	WILL ENABLE AND ENCOURAGE COLLABORATIVE RESEARCH
6	AMONG ACADEMIC AND INDUSTRY INVESTIGATORS AND
7	ULTIMATELY ACCELERATE BASIC SCIENCE DISCOVERIES AND
8	THERAPEUTIC DEVELOPMENT FOR LATE ONSET
9	NEURODEGENERATIVE DISEASES.
10	CIRM, AS AN INSTITUTE, WE DESIRE TO
11	SUPPORT THIS PROJECT, WHICH IS ALIGNED WITH CIRM'S
12	MISSION AND ALSO RECOMMENDATIONS OF OUR EXTERNAL
13	REVIEW PANEL TO MAINTAIN FOCUS ON MEANINGFUL,
14	TARGETED SCIENTIFIC EXCELLENCE, TO ADOPT A MORE
15	AGGRESSIVE, PROACTIVE APPROACH TO IDENTIFYING
16	INNOVATIVE PROJECTS ACROSS THE STEM CELL THERAPEUTIC
17	LANDSCAPE THAT SHOWS PROMISE FOR MOVING INTO
18	TRANSLATIONAL RESEARCH, CLINICAL TRIALS, AND PRODUCT
19	DEVELOPMENT, AND ALSO TO ENSURE SERIOUS ENGAGEMENT
20	WITH INDUSTRY.
21	POTENTIAL BENEFITS TO CIRM OF BEING PART
22	OF THIS PUBLIC/PRIVATE PARTNERSHIP WITH NINDS IS THE
23	OPPORTUNITY TO PLAY A ROLE IN A NATIONAL EFFORT BY
24	LEVERAGING RESOURCES AND EXPERTISE, ACCESS TO CIRM
25	FUND INVESTIGATORS AS A NATIONAL RESOURCE IN IPS

1	CELL LINES, AS WELL AS THE OPPORTUNITY FOR MORE
2	IN-DEPTH INTERACTIONS WITH INDUSTRY, TRANSLATIONAL
3	SCIENTISTS, AND PATIENT ADVOCACY ENGAGEMENT ALL
4	FOCUSED ON INCREASING THE KNOWLEDGE OF STRATEGIC
5	THERAPEUTIC AREAS AND CATALYZING THE DEVELOPMENT OF
6	MORE ACCURATE AND PREDICTIVE SCREENS FOR DRUG
7	DISCOVERY AND DEVELOPMENT.
8	THAT'S SORT OF A READER'S DIGEST VERSION
9	OF WHAT YOU HAVE AS A DOCUMENT THAT'S ON OUR PUBLIC
10	WEBSITE. MAYBE AT THIS POINT I COULD TAKE TIME TO
11	ANSWER SPECIFIC QUESTIONS.
12	CHAIRMAN KLEIN: JEFF, COULD I ASK A
13	QUESTION OF ELLEN?
14	CHAIRMAN SHEEHY: SURE. PLEASE.
15	CHAIRMAN KLEIN: ELLEN, MY UNDERSTANDING
16	OF THIS IS THAT WE GET THE ADVANTAGE OF SUBSTANTIAL
17	LEVERAGE BECAUSE THE TOTAL AMOUNT BEING PUT UP BY
18	THE PARTNERS IS 4.5 MILLION AND WE'RE PUTTING UP
19	150,000?
20	DR. FEIGAL: THAT'S RIGHT.
21	CHAIRMAN KLEIN: WHAT DOES IT MEAN WHEN IT
22	SAYS THE PROJECT FOR CIRM IS A HUNDRED FIFTY
23	THOUSAND FOR TWO YEARS, AND THE DOLLARS, 300,000
24	TOTAL, THE DOLLARS WOULD BE DIRECTED TO
25	CALIFORNIA-BASED INVESTIGATORS? WHAT DOES THAT
	10

1	MEAN?
2	DR. FEIGAL: WELL, RIGHT NOW THIS IS A
3	COMPETITIVE SUPPLEMENT TO THE ALREADY EXISTING THREE
4	CONSORTIA, ONE FOCUSED ON PARKINSON'S, ONE FOCUSED
5	ON HUNTINGTON'S, ONE FOCUSED ON ALS. TWO OF THOSE
6	THREE CONSORTIA HAVE CALIFORNIA INVESTIGATORS ON
7	THEM. SO IF THE CURRENTLY CONFIGURED CONSORTIA WHO
8	ARE ELIGIBLE TO APPLY DO APPLY, WE WOULD HAVE THE
9	OPPORTUNITY TO TAKE A LOOK AT THE OPPORTUNITY TO
10	FUND AT LEAST PART OF WHAT THEY RECEIVE FROM THE NIH
11	FROM OUR FUNDS.
12	IN ADDITION, BY BEING A PARTNER, WE HAVE
13	THE ABILITY TO SIT ON THE STEERING COMMITTEE AS ONE
14	OF THE PUBLIC/PRIVATE PARTNERS TO TALK ABOUT THE
15	RESEARCH, TO HELP THINK ABOUT WAYS TO LEAD THAT
16	RESEARCH FORWARD. SO THE ONE PART OF IT IS THE
17	FINANCIAL. I THINK THE MORE INTANGIBLE PART OF IT
18	IS THE INTERACTION AND EXPOSURE TO THIS NATIONAL
19	NETWORK AND A CHANCE TO BE ON THE LEADING EDGE OF AN
20	IPS REPOSITORY AND HOW IT'S UTILIZED.
21	DR. BRYANT: COULD I ASK A QUESTION,
22	PLEASE? I'M JUST A LITTLE BIT UNSURE ABOUT WHAT
23	ACCESS WE WOULD HAVE IN CALIFORNIA WITH FUNDING OF
24	EITHER IN NIH OR, WELL, I GUESS FROM CIRM
25	FUNDING. WHAT ACCESS WOULD WE HAVE TO SUCH A

1	CONSORTIUM IF WE DIDN'T PONY UP? I MEAN IS THIS A
2	PRIVATE KIND OF THING THAT ONLY PEOPLE THAT HAVE
3	CONTRIBUTED TO CAN GET INTO?
4	DR. FEIGAL: THE INTENT IS THAT THIS WOULD
5	BE A PUBLICLY AVAILABLE RESEARCH RESOURCE. AND FOR
6	A MODEST FUNDING COMMITMENT COMPARED TO THE TOTAL
7	FUNDING COMMITMENT, WE'D ACTUALLY BE ALLOWED TO HAVE
8	A PART OF A LEADERSHIP ROLE ON THIS STEERING
9	COMMITTEE OF THIS RESEARCH RESOURCE.
10	DR. BRYANT: SO WE'RE BUYING INTO THE
11	MANAGEMENT OR THE DIRECTION AND LEADERSHIP OF THIS,
12	NOT NECESSARILY ACCESS TO IT? I JUST WANTED TO BE
13	CLEAR ABOUT THAT.
14	DR. FEIGAL: WELL, IT WOULD BE A PUBLICLY
15	ACCESSIBLE RESEARCH RESOURCE, YOU'RE RIGHT,
16	REGARDLESS OF WHETHER OR NOT WE PUT DOWN THIS
17	RELATIVELY MODEST AMOUNT OF FUNDING. I THINK IT'S
18	MORE THE LEADERSHIP AND THE INTERACTIONS THAT ARE
19	OTHER ATTRACTANTS TO BE INVOLVED HERE.
20	MR. HARRISON: DR. FEIGAL, JUST TO
21	INTERJECT FOR A MOMENT. DR. BRYANT MAY NOT KNOW
22	THAT UCI IS INVOLVED IN ONE OF THE CONSORTIA. SO
23	SHE SHOULD, THEREFORE, REFRAIN FROM PARTICIPATING IN
24	THIS DISCUSSION.
25	DR. BRYANT: CAN I LISTEN THOUGH?
	12

1	MR. HARRISON: ABSOLUTELY.
2	DR. FEIGAL: CEDARS-SINAI IS ANOTHER
3	INSTITUTION. I DON'T BELIEVE THERE'S ANYBODY THAT
4	WOULD BE INVOLVED WITH THAT.
5	MR. HARRISON: NO ONE WHO'S ON THE PHONE
6	CURRENTLY.
7	CHAIRMAN KLEIN: SO FROM A LEADERSHIP
8	PERSPECTIVE, ARE WE GAINING INFORMATION AND
9	KNOWLEDGE ABOUT HOW TO HAVE OUR OWN IPSC REPOSITORY
10	IN THE FUTURE? IT'S MY UNDERSTANDING THAT'S ONE OF
11	THE GOALS OF THIS PARTICIPATION.
12	DR. FEIGAL: BOB, I THINK YOU'RE
13	ABSOLUTELY CORRECT, AND THAT'S WHY I PRESENTED AS A
14	TWO-STEP CONCEPT. SO THIS WOULD BE PART OF WHOLE.
15	WE'D HAVE THE OPPORTUNITY TO WORK ON THIS IN
16	NEURODEGENERATIVE DISEASES, HAVE LESSONS LEARNED IN
17	TERMS OF HOW THIS COULD BE DEVELOPED, HOW IT COULD
18	BE UTILIZED, THE INTERACTIONS WITH OTHER
19	TRANSLATIONAL SCIENTISTS AND INDUSTRY, AND THEN TAKE
20	SOME OF THESE LESSONS LEARNED TO APPLY IT.
21	NOW, CIRM, AS YOU KNOW, HAS HAD OUR OWN
22	WORKSHOP ON IPS ISSUES AND HOW TO DEVELOP A ROBUST
23	REPOSITORY AND A FRAMEWORK OF ORGANIZATION.
24	SO, YEAH, I WOULD VIEW IT AS WE WOULD BE
25	ABLE TO UTILIZE SOME LESSONS LEARNED FROM THIS FIRST

1	STEP.
2	DR. TROUNSON: I THINK IT'S VERY IMPORTANT
3	TO RECOGNIZE THAT THERE'S NOT A LOT KNOWN ABOUT THE
4	CAUSES, THE BASIC CAUSES, OF SOME OF THESE
5	NEURODEGENERATIVE DISEASES. I THINK THAT THEY'RE
6	PRETTY COMPLEX AND PROBABLY INVOLVE A RAFT OF GENES
7	AND A RAFT OF SITUATIONS, ENVIRONMENTAL EFFECTS. SO
8	TO BE ABLE TO BE INVOLVED IN A VERY MAJOR SCREEN
9	WOULD BE VERY INFORMATIVE, I THINK, AND WOULD HELP
10	US TO BE ABLE TO FEED BACK TO OUR SCIENTISTS SORT OF
11	THE FRONT LINE OF WHAT'S MOVING IN THIS AREA.
12	AND PARTICULARLY BECAUSE THIS IS ONE OF
13	THE THIS IS ONE OF THE DIFFICULT AREAS FOR ALL OF
14	US TO HAVE AN IMPACT. SO WHATEVER WE CAN DO IN
15	THESE AREAS, AND THIS I THINK IS A GREAT
16	OPPORTUNITY, WHATEVER WE CAN DO IN THESE AREAS,
17	WE'LL ACTUALLY MOVE OUR PROGRAM FORWARD.
18	CHAIRMAN SHEEHY: IS THIS OUR FIRST FORMAL
19	COLLABORATION WITH THE NIH?
20	DR. TROUNSON: THERE'S ONE ON THE TABLE,
21	JEFF, WHICH IS STILL HAVING SOME ISSUES ABOUT
22	GETTING SOLVED. WE HAVE AN AGREEMENT THAT HASN'T
23	ACTUALLY YET BEEN SIGNED WITH THE CLINICAL INSTITUTE
24	AT NIH. SO THEY ARE WAITING BECAUSE THE TRANSLATION
25	CENTER IS JUST BEING FORMED, AND THEY WANT THAT TO

1	BE PART OF THE ARRANGEMENT.
2	DR. FEIGAL: THAT'S IN THE WORKS. THIS
3	WOULD ACTUALLY BE THE FIRST.
4	CHAIRMAN SHEEHY: BOB, DID YOU HAVE A
5	QUESTION?
6	CHAIRMAN KLEIN: I CAN WAIT ON MY
7	QUESTIONS IF ANOTHER BOARD MEMBER WOULD LIKE TO ASK
8	A QUESTION FIRST.
9	MS. SAMUELSON: I HAVE A QUESTION. THIS
10	IS JOAN. THANK YOU, ALAN, FOR MENTIONING THE
11	ENVIRONMENTAL CONNECTION. THERE'S A FAIR AMOUNT,
12	ALTHOUGH NOT NEARLY ENOUGH, KNOWN ABOUT THE CAUSE OF
13	PARKINSON'S AS FAR AS THE ENVIRONMENT PLAYS A ROLE.
14	AND I'M WONDERING IF YOU KNOW, ALAN, IF OTHER
15	INSTITUTES AT THE NIH THAT ARE INVOLVED IN SOME OF
16	THESE ISSUES ARE GOING TO BE COLLABORATING OR
17	WHETHER
18	DR. FEIGAL: JOAN, FOR THIS PARTICULAR ONE
19	THAT'S FOCUSED ON NEURODEGENERATIVE DISEASES, THE
20	LEAD INSTITUTE AND I THINK THE ONLY INSTITUTE AT NIH
21	WILL BE THE NINDS, BUT THERE WILL BE OTHER PARTNERS
22	FROM PATIENT FOUNDATIONS AND FROM INDUSTRY THAT WILL
23	WANT TO BE A PART OF THIS PUBLIC/PRIVATE
24	PARTICIPATION.
25	MS. SAMUELSON: WELL, LET ME JUST LET YOU
	15
	Τ.)

Т	KNOW THAT THE NATIONAL INSTITUTE FOR ENVIRONMENTAL
2	HEALTH SCIENCES HAS PLAYED A VERY IMPORTANT ROLE IN
3	FUNDING SOME OF THE KEY DEVELOPMENTS IN UNCOVERING
4	ENVIRONMENTAL CAUSES AND TYING THEM TO GENETIC
5	PREDISPOSITIONS AND SO ON. AND SO IT WOULD BE
6	IMPORTANT THAT ALL OF THAT DATA IS SOMEHOW PART OF
7	THIS.
8	DR. FEIGAL: THAT'S A GOOD POINT. I
9	THOUGHT YOU WERE ASKING ABOUT NIH, BUT THERE MAY BE
10	OTHER AGENCIES.
11	MS. SAMUELSON: WELL, NIEHS IS AT NIH.
12	IT'S ONE OF THE NIH INSTITUTES, ENVIRONMENTAL HEALTH
13	SCIENCES, AND THEY PLAYED A KEY ROLE.
14	YOU MENTIONED LATE ONSET. I WONDERED WHAT
15	THAT MEANT.
16	DR. FEIGAL: JOAN, I THINK THERE WILL BE
17	DIFFERENT STAGES OF DISEASE THAT WILL BE LOOKED AT
18	IN PARKINSON'S. AND SO I THINK THEY'LL WANT TO GET
19	A DIVERSITY. SO AT THIS POINT IN TIME, I CAN'T
20	REALLY COMMENT ON THE EXACT CLINICAL PARAMETERS OR
21	HOW THEY'RE ASSESSING THE LATENESS OF THE DISEASE IN
22	PARKINSON'S. BUT ALL THOSE PARAMETERS IN TERMS OF
23	CLINICAL, GENETIC, LABORATORY WORK THAT WILL BE DONE
24	TO ASSESS THAT, I THINK THEY'LL WANT A DIVERSITY OF
25	THE MATURATION OF THE DISEASE.

1	MS. SAMUELSON: OKAY.
2	DR. FEIGAL: I DON'T HAVE ANY MORE
3	SPECIFICS THAN THAT RIGHT NOW.
4	MS. SAMUELSON: OKAY. GREAT. THANKS.
5	MR. ROTH: ELLEN, I WONDER IF I CAN JUST
6	TALK A LITTLE BIT ABOUT THE BUDGET. THIS INSTITUTE
7	WAS CREATED WITH ARRA FUNDS. AND I SEE THAT THEIR
8	BUDGET NOW HAS TO COME OUT OF APPROPRIATIONS FROM
9	THE NINDS.
10	DR. FEIGAL: CAN I JUST CORRECT THAT?
11	NINDS ALWAYS RECEIVES AN APPROPRIATION FROM THE
12	FEDERAL GOVERNMENT. WHAT THEY GOT WAS STIMULUS
13	FUNDS CALLED ARRA THAT WAS USED IN 2009 TO HELP
14	BASICALLY STIMULATE THE ECONOMY. AND AS PART OF
15	THAT, THIS INITIATIVE WAS PUT FORWARD.
16	MR. ROTH: I UNDERSTAND THAT. BUT I WAS
17	ASKING REALLY ABOUT THE CONFIDENCE THAT THAT
18	CONTINUING BUDGET WILL BE THERE FOR THE OPERATION OF
19	THIS, THAT 4.5 MILLION.
20	DR. FEIGAL: NINDS WILL GET AN
21	APPROPRIATION. I CAN'T FORESEE THE FEDERAL
22	GOVERNMENT NOT PROVIDING FUNDING TO A MAJOR
23	INSTITUTE AT NIH. WHAT THEY CAN'T COMMIT TO IS THE
24	EXACT DOLLAR AMOUNT. SO THAT WILL DEPEND ON WHAT
25	THE FUTURE APPROPRIATION IS IN FISCAL YEAR 2012. SO

1	WHAT THEY'RE COMMITTING TO IS 4.5 MILLION IN FISCAL
2	YEAR 2011 WITH THE SUBSEQUENT YEAR TO BE DETERMINED
3	BY THEIR APPROPRIATION.
4	MR. ROTH: SO THAT WILL BE AN ONGOING
5	BUDGET ITEM THAT THEY'LL HAVE TO MAKE A PRIORITY
6	EACH YEAR, HOPEFULLY. ANYWAY, I'M SUPPORTIVE OF
7	THIS INITIATIVE. I THINK IT WILL GET MORE THAN THE
8	AMOUNT OF MONEY WE'RE PUTTING IN BY ORDERS OF
9	MAGNITUDE BACK IN EXPERIENCE, KNOWLEDGE. I THINK
10	IT'S A GOOD IDEA TO TIE INTO NIH WHEREVER WE CAN
11	LEGALLY TO GET THOSE RELATIONSHIPS GOING.
12	AND, ART, YOU WOULD PROBABLY TELL
13	EVERYBODY THAT WE'RE GOING TO TAKE MY SIDE OF THE
14	AISLE IS GOING TO TAKE BACK ALL THE MONEY ANYWAY.
15	SO YOU MIGHT AS WELL SPEND IT WHILE YOU'VE GOT IT.
16	MR. TORRES: RIGHT. AND WE'LL BE BACK IN
17	2012.
18	MS. FEIT: I HAVE A QUICK QUESTION. YOU
19	MENTIONED THAT WE WOULD GET A SEAT ON THE BOARD.
20	HAVE YOU SAID WHO WOULD BE TAKING THAT SEAT, AND HOW
21	WE ARE GOING TO BE MONITORING THE PROCESS IN THIS?
22	DR. FEIGAL: WELL, NINDS, THE INSTITUTE
23	WOULD NOMINATE THE REPRESENTATIVE FROM THE WHOM
24	WE SELECT TO REPRESENT THE INSTITUTE, AND THAT
25	PERSON WOULD HAVE A SEAT AT THE TABLE. IS THAT WHAT
	10

1	YOU'RE ASKING?
2	MS. FEIT: YES. BUT SO WE WOULD PUT
3	FORWARD A LIST OF NAMES?
4	DR. FEIGAL: NO. WE WOULD PRESUMABLY PUT
5	FORWARD THE NAME THAT WE WOULD WANT TO SIT ON THAT
6	ONE SLOT. AND THEN THEY WOULD I SUPPOSE IF THERE
7	WAS A I CAN'T IMAGINE IF WE RECOMMENDED A SPOT
8	UNLESS THERE WAS SOMETHING INAPPROPRIATE OR
9	SOMETHING ABOUT THE RECOMMENDATION, THAT THEY WOULD
10	PROBABLY HONOR THAT RECOMMENDATION.
11	CHAIRMAN KLEIN: ELLEN, MY UNDERSTANDING
12	IS WE WOULD NOMINATE ONE OF OUR SCIENCE OFFICERS OR
13	SCIENCE LEADERSHIP PERSONNEL TO BE ON THAT SPOT.
14	DR. FEIGAL: YEAH. I MEAN WE CAN DISCUSS
15	THAT, BUT IT'S A SCIENTIFIC PROJECT. AND THERE WILL
16	ALREADY BE OTHER PATIENT ADVOCACY FOUNDATION GROUPS
17	REPRESENTED. SO I THINK THE SCIENTIFIC
18	REPRESENTATION WOULD PROBABLY BE A GOOD
19	RECOMMENDATION TO COME FROM US.
20	CHAIRMAN KLEIN: AND SO WHEN WE DO OUR OWN
21	IPSC REPOSITORY, WOULD WE BE FOLLOWING THE SAME
22	MODEL WITH HAVING SOME PATIENT ADVOCATE
23	REPRESENTATIVES ON THAT BOARD?
24	DR. TROUNSON: I THINK IT'S A LITTLE BIT
25	EARLY TO DECIDE ABOUT THAT AT THIS STAGE. I WOULD

1	HAVE THOUGHT THAT THAT WOULD BE IDEAL. SO WE
2	HAVEN'T TRIED TO FRAME THAT UP AT THIS POINT IN
3	TIME. I WOULD HAVE THOUGHT THAT WOULD BE IDEAL. I
4	THINK THE MORE WE HAVE ADVOCACY IN THIS SPACE THE
5	BETTER.
6	CHAIRMAN KLEIN: IN ANY CASE THIS IS A
7	SCIENTIFIC SPOT BECAUSE WE'RE TRYING TO GLEAN
8	SCIENTIFIC KNOWLEDGE TO BE ABLE TO SET UP OUR OWN
9	REPOSITORY.
10	DR. TROUNSON: I THINK IT WOULD BE GOOD
11	FOR US IF WE HAVE A SENIOR PERSON WHO IS IN THE
12	NEURODEGENERATION AREA IF WE HAVE THE TIME. SO WE
13	WOULD BE I WOULD BE HOPEFULLY PUTTING FORWARD A
14	NAME THAT WOULD THAT THE BOARD WOULD RECOGNIZE AS
15	IMPORTANT FOR US.
16	CHAIRMAN KLEIN: THAT WAS MY
17	UNDERSTANDING. I WAS JUST TRYING TO CLARIFY IT FOR
18	EVERYONE.
19	SO LET ME ASK, IF I COULD, IN THE LAST SIX
20	WEEKS, THERE WAS A MAJOR PUBLICATION COMING OUT OF
21	STANFORD OF AN IPSC-DERIVED MODEL FOR PARKINSON'S IN
22	A DISH. WHAT IS THE PROCESS FOR THAT KIND OF
23	IPSC-DERIVED MODEL? WOULD THAT MODEL THEN BE
24	CONTRIBUTED TO THIS REPOSITORY OR HOW
25	DR. TROUNSON: WELL, IT MIGHT BE, BOB. IT
	20

1	DEPENDS WHETHER THEY'RE PART OF THE ORGANIZATION.
2	AND, OF COURSE, IF WE BECOME PART OF IT, THEN WE
3	WOULD I THINK WE WOULD STRONGLY SUGGEST THAT THAT
4	INFORMATION IS TAKEN ON BOARD BY ALL THE TEAMS. I
5	THINK THAT WAS A VERY GOOD PUBLICATION. I ACTUALLY
6	THINK IT'S REALLY GOOD WORK; BUT, OF COURSE, WE'LL
7	HAVE TO SIT DOWN WITH THE LEADERSHIP OF THE WHOLE
8	EXERCISE TO TRY AND DETERMINE WHAT METHODS THAT THEY
9	WILL USE.
10	IT'S CLEAR TO ME AT THE MOMENT, MAYBE ONE
11	OF THE SLIGHT DEFICIENCIES HERE IS THAT THESE CELLS
12	WILL BE DERIVED BY A VARIETY OF METHODS. AND SO
13	THAT INTRODUCES ONE ELEMENT OF VARIATION, WHICH I
14	THINK IS UNFORTUNATE, BUT IT MAY NOT HAVE A LOT OF
15	EFFECT. BUT I JUST THINK OF IT'S A SOURCE OF
16	VARIATION.
17	AGAIN, THE DIFFERENTIATION PATHWAYS TO
18	CREATE, IF YOU LIKE, DISEASE-IN-A-DISH MODELS IS
19	PRETTY IMPORTANT. SO I THINK WE WOULD BE POINTING
20	THAT OUT TO THE COMMITTEE AND GETTING THEM TO
21	RECOMMEND THAT IF THAT WAS APPROPRIATE.
22	CHAIRMAN KLEIN: AT LEAST HAVING A
23	STRATEGICALLY HAVING A POSITION ON THIS BOARD PUTS
24	US IN A POSITION THAT WE CAN ADVOCATE FOR WORK THAT
25	WE'RE AWARE OF THROUGH OUR FUNDING ON THE WEST COAST

1	THAT MIGHT CONTRIBUTE TO THE RICHNESS OF THIS
2	REPOSITORY.
3	DR. TROUNSON: THAT'S CORRECT. THAT GOES
4	ALSO FOR THE HUNTINGTON'S DISEASE BECAUSE THERE'S A
5	SLIGHTLY DIFFERENT PROGRAM INVOLVED THERE FOR THE
6	DIFFERENTIATION, AND SOME OF THE GRANTEES HERE IN
7	CALIFORNIA HAVE DONE A REALLY GOOD JOB ON THOSE
8	CELLS AS WELL. SO, YOU KNOW, FOR THAT AND FOR
9	ALZHEIMER'S, WE WOULD CERTAINLY BE MAKING THAT
10	INFORMATION AVAILABLE AND MAKING THE ARGUMENT THAT
11	THERE ARE AT LEAST VERY SOUND METHODOLOGIES THAT
12	OUGHT TO BE CONSIDERED OR INCORPORATED AS A PRIORITY
13	IN THE PROGRAM.
14	DR. FEIGAL: I THINK THE OTHER INTERESTING
15	ASPECT, SINCE IT'S A PARTNERSHIP THAT COULD ALSO BE
16	OPEN TO INDUSTRY, IS THAT ANOTHER DESIRED FUNCTION
17	WOULD BE DEVELOPMENT OF CELL-BASED PLATFORMS THAT
18	MEET THE REQUIREMENTS FOR INDUSTRY FOR TARGETED
19	VALIDATION AND SECONDARY SCREENING. THEY'RE VERY
20	INTERESTED IN USING THIS AS A TOOL. SO I THINK
21	THEIR INPUT AND THE REGULATORY NEEDS FOR WHAT MIGHT
22	BE USEFUL TO THEM WILL ALSO BE PART OF THE
23	CONSIDERATION.
24	SO I THINK THE SCIENCE AND RESEARCH WILL
25	BE GREATLY BENEFITED, BUT ALSO AT THE END OF THE
	22

1	DAY, WE'RE INTERESTED IN MOVING US TOWARDS CLINICAL
2	APPLICATION.
3	DR. TROUNSON: AS YOU KNOW, THE GENOMICS
4	AREA IS SOMETHING THAT WE HAVE SOME CONCERN ABOUT,
5	GENOMICS AND EPIGENOMICS, SO WE WILL BE PRESSING FOR
6	SOME SEQUENCING WHERE APPROPRIATE TO ENSURE THAT
7	WE'RE NOT DEALING WITH VARIATION REALLY JUST CAUSED
8	BY MAKING THESE CELLS AND DIFFERENTIATING THEM.
9	CHAIRMAN KLEIN: ALAN, YOU AND ELLEN HAVE
10	BROUGHT UP A NUMBER OF VERY GOOD POINTS ABOUT WHY
11	THE STRATEGIC LEADERSHIP ROLE IS VALUABLE. AND
12	MAYBE AT THE BOARD WE COULD HAVE MORE OF A BULLET
13	POINT SUMMARY OF ALL THOSE STRATEGIC ADVANTAGES,
14	BOTH IN KNOWLEDGE POSITION, ADVOCACY, WORK FROM THE
15	WEST COAST, AND OTHER BENEFITS BECAUSE CERTAINLY THE
16	CONCEPTS ARE ALL LAID OUT WELL HERE, BUT I THINK THE
17	DISCUSSION HAS BROUGHT OUT SOME MORE ARTICULATION OF
18	THIS THAT WOULD BE VALUABLE TO PUT INTO THE BOARD
19	RECORD.
20	DR. FEIGAL: WE CAN DO THAT.
21	CHAIRMAN SHEEHY: ARE THERE ADDITIONAL
22	BOARD QUESTIONS OR COMMENTS?
23	MR. ROTH: JEFF, WOULD YOU LIKE A MOTION
24	TO APPROVE?
25	CHAIRMAN SHEEHY: I THINK SO. I DON'T
	22

1	THINK WE HAVE TO DO IT, BUT I THINK IT'S USEFUL FOR
2	THE BOARD TO HAVE A STRONG RECOMMENDATION FROM THE
3	COMMITTEE. AND THEN PERHAPS IT WILL MAKE IT GO A
4	LITTLE BIT MORE SMOOTHLY AT THE BOARD.
5	MR. ROTH: SO I WOULD BE HAPPY TO MAKE
6	THAT MOTION TO APPROVE THE CONCEPT AS PRESENTED.
7	MS. SAMUELSON: THIS IS JOAN. I'LL SECOND
8	IT.
9	I GUESS I HAVE ONE QUESTION. DO WE HAVE
10	ANY KIND OF CERTAINTY, THAT PROBABLY REALLY SOUNDS
11	LIKE AN OVERSTATEMENT, THAT 150,000 IS THE
12	COMMITMENT JUST BECAUSE WE'RE BASING IT ON THAT, AND
13	YET THE FUTURE FUNDING FOR THE INSTITUTE IS OF
14	CONCERN. I'M WONDERING IF THEY'RE GOING TO COME
15	BACK FOR MORE.
16	DR. FEIGAL: NO. NO. WE'VE ALREADY HAD A
17	DISCUSSION ABOUT WHAT WE WOULD BE WILLING TO
18	PROVIDE, AND WE WILL DOCUMENT THAT IN A MEMORANDUM
19	OF UNDERSTANDING WITH THE FOUNDATION FOR NIH. SO
20	THAT IS WHAT WE SAID WOULD BE AN ACCEPTABLE
21	COMMITMENT.
22	MS. SAMUELSON: AND THEY'VE COMMITTED TO
23	US PLAYING THE ROLE DECIDED UPON FOR AD INFINITUM.
24	DR. FEIGAL: THEY ARE VERY INTERESTED IN
25	HAVING US AS A PARTNER. SO I THINK THEY ALSO SEE IT
	2.4

1	BEYOND THE MONETARY AMOUNT.
2	CHAIRMAN KLEIN: JOAN, I DON'T THINK WE
3	CAN REASONABLY BELIEVE THAT'S AN INDEFINITE
4	COMMITMENT GIVEN THE VOLATILITY IN THE NEAR TERM,
5	BUT IT IS CERTAINLY A COMMITMENT OF INTENT.
6	MS. SAMUELSON: OKAY. I THINK IT'S
7	IMPORTANT WE BE CLEAR WITH OUR VARIOUS
8	CONSTITUENCIES BECAUSE WE WOULDN'T WANT TO BE
9	CRITICIZED OURSELVES.
10	DR. FEIGAL: THE 4.5 MILLION IS ACTUALLY
11	PUBLICLY OUT THERE. THEY'VE MADE THAT COMMITMENT.
12	CHAIRMAN KLEIN: ELLEN, SHE'S TALKING
13	ABOUT YEAR THREE, YEAR FOUR WHAT HAPPENS.
14	DR. FEIGAL: WELL, IT'S ONLY YEAR TWO.
15	OKAY.
16	CHAIRMAN KLEIN: THE POINT IS THAT THE NIH
17	HAS MADE A STRONG COMMITMENT HERE. THEY'VE ALREADY
18	INVESTED A GREAT DEAL OF FUNDS. THEY'VE COMMITTED
19	THE 4.5 THEY HAVE CONTROL OVER, BUT
20	MS. SAMUELSON: IS THAT FISCAL 12 OR
21	SOMETHING?
22	DR. FEIGAL: THAT'S FISCAL YEAR 2011. THE
23	PLAN IS TO FUND THIS BEFORE SOMETIME LATER IN THE
24	SPRING.
25	MS. SAMUELSON: I SEE. GREAT.
	25

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1	CHAIRMAN SHEEHY: SO I THINK WE'RE READY
2	FOR PUBLIC COMMENT. AND I THINK, JUDY ROBERSON.
3	MS. ROBERSON: I'M JUDY ROBERSON,
4	PRESIDENT OF THE HDSA NORTHERN CALIFORNIA CHAPTER.
5	THE HUNTINGTON'S DISEASE COMMUNITY ENTHUSIASTICALLY
6	SUPPORTS CIRM'S IPS CELL REPOSITORY PROPOSAL. OUR
7	COMMUNITY IS ORGANIZED AND EAGER TO PARTICIPATE IN
8	STEM CELL RESEARCH. BECAUSE ALTHOUGH THE GENE FOR
9	HUNTINGTON'S DISEASE WAS FOUND 18 YEARS AGO, A
10	TREATMENT OR CURE FOR HD HAS REMAINED ILLUSIVE. OUR
11	BIG HOPE LIES IN STEM CELL RESEARCH. THANK YOU SO
12	MUCH.
13	CHAIRMAN SHEEHY: THANK YOU, JUDY. SO I
14	THINK WE'RE READY FOR A ROLL CALL VOTE. AND SO,
15	JAMES, COULD YOU REPEAT THE MOTION? AND I THINK
16	WHAT WE'RE TALKING ABOUT IS APPROVAL FOR STEP ONE AS
17	PROPOSED IN THE DOCUMENT WE HAVE.
18	MR. HARRISON: RIGHT, JEFF. AS I
19	UNDERSTAND IT, THE MOTION IS TO RECOMMEND TO THE
20	BOARD APPROVAL OF STEP ONE AS OUTLINED IN THE IPSC
21	REPOSITORY CONCEPT PLAN.
22	CHAIRMAN SHEEHY: GREAT.
23	MS. KING: SUSAN BRYANT.
24	DR. BRYANT: ABSTAIN.
25	MS. KING: AND THE SAME IS TRUE FOR DR.
	26
	26

1	STEWARD; IS THAT CORRECT?
2	MR. HARRISON: CORRECT.
3	MS. KING: MARCY FEIT.
4	MS. FEIT: YES.
5	MS. KING: MICHAEL FRIEDMAN.
6	DR. FRIEDMAN: YES.
7	MS. KING: BOB KLEIN.
8	CHAIRMAN KLEIN: YES.
9	MS. KING: DUANE ROTH.
10	MR. ROTH: YES.
11	MS. KING: JOAN SAMUELSON.
12	MS. SAMUELSON: YES.
13	MS. KING: JEFF SHEEHY.
14	CHAIRMAN SHEEHY: YES.
15	MS. KING: AND ART TORRES.
16	MR. TORRES: AYE.
17	MS. KING: AND FOR THE RECORD, THE MOTION
18	CARRIES. THANK YOU.
19	MS. SAMUELSON: CAN I ASK ONE FINAL
20	QUESTION? AND IT'S PROBABLY VERY MINOR, BUT JUST IN
21	TERMS OF CLARITY FOR THE PUBLIC. WHEN IT'S CALLED A
22	PUBLIC/PRIVATE PARTNERSHIP, I'M NOT SURE QUITE WHO
23	THAT'S REFERRING TO. AND I THINK OF US AND NINDS,
24	IT SEEMS LIKE WE'RE PUBLIC PUBLIC.
25	DR. FEIGAL: JOAN, I TOO NOODLED ON THAT.
	27

1	I TOO WOULD CONSIDER US PUBLIC, BUT THERE ARE
2	PRIVATE PEOPLE. IT'S SORT OF LIKE THE PATIENT
3	FOUNDATIONS ARE
4	MS. SAMUELSON: IT'S THE LARGER GROUPING.
5	DR. FEIGAL: I THINK THE OTHER I
6	THOUGHT THE OTHER THING YOU WERE GOING TO ASK IS
7	GOVERNMENT VERSUS NONGOVERNMENT. AT ANY RATE, I
8	THINK WE CONSIDER OURSELVES IN THE PUBLIC, AND THE
9	INDUSTRY WOULD BE THE PRIVATE SECTOR.
10	MS. SAMUELSON: GOT IT. OKAY. THANK YOU.
11	CHAIRMAN KLEIN: AND, ELLEN, IS THERE A
12	LIST OF THE TOTAL NUMBER OF PARTICIPANTS?
13	DR. FEIGAL: YOU KNOW, I CAN'T SHARE THAT
14	RIGHT NOW BECAUSE I DON'T THINK IT'S PUBLICLY
15	AVAILABLE. I DIDN'T SEE IT ON THE WEBSITE. SO
16	THAT'S WHY I'M NOT SHARING IT.
17	CHAIRMAN KLEIN: IF YOU COULD JUST CONFIRM
18	BEFORE THE BOARD MEETING WHAT THE POLICY IS AS TO
19	THE MEMBERSHIP.
20	DR. FEIGAL: I CAN CONFIRM HOW THE
21	FOUNDATION FOR NIH WORKS TO GET PARTNERS, IF YOU
22	JUST MEANT THE POLICY.
23	CHAIRMAN KLEIN: THE QUESTION IS WHAT IS
24	THE POLICY ON PUBLIC INFORMATION ON THE PARTICIPANTS
25	IN THIS PROGRAM?

1	DR. FEIGAL: OH, IT WILL BE I CAN TELL
2	YOU THAT NOW. ANY PARTNERS WILL BE PUBLICLY THEY
3	HAVE TO BE PUBLIC. SO IF THERE IS A PARTNER THAT'S
4	CONTRIBUTING FUNDING THROUGH THE FOUNDATION FOR NIH,
5	THAT WILL BECOME PUBLIC INFORMATION.
6	CHAIRMAN KLEIN: OKAY. SO THAT WOULD BE
7	GOOD TO RECOUNT AT THE BOARD.
8	DR. FEIGAL: I WILL DO THAT.
9	CHAIRMAN KLEIN: THANK YOU VERY MUCH.
10	MS. SAMUELSON: IS THE WEBSITE AT THE
11	FOUNDATION FOR NIH?
12	DR. FEIGAL: THE WEBSITE WELL, THERE'S
13	A COUPLE WAYS THINGS CAN BE MADE PUBLICLY AVAILABLE.
14	GENERALLY THEY WON'T BE MADE PUBLICLY AVAILABLE
15	UNTIL MOU'S HAVE BEEN EXECUTED. SO I CAN'T PROMISE
16	THAT THAT'S GOING TO HAPPEN BEFORE THE BOARD, AND
17	MOST LIKELY IT WON'T. BUT WHEN IT IS PUBLICLY
18	AVAILABLE, IT WILL BE AVAILABLE ON THE FOUNDATION
19	FOR NIH WEBSITE, AND THEN NINDS IS ALSO WILLING TO
20	ACKNOWLEDGE THE PARTNERS IN ANY PUBLIC ANNOUNCEMENTS
21	THEY MAKE ABOUT THE CONSORTIA.
22	MS. SAMUELSON: OKAY. AND THE WEBSITE
23	THAT WAS REFERRED TO, THAT'S AT THE FOUNDATION FOR
24	NIH?
25	DR. FEIGAL: YEAH. THE ABBREVIATION IS
	29

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1	FNIH.
2	MS. SAMUELSON: OKAY. THANK YOU.
3	DR. FEIGAL: IF YOU CAN'T FIND IT, I'LL
4	SEND IT TO YOU, JOAN.
5	MS. SAMUELSON: GREAT. THANKS.
6	CHAIRMAN SHEEHY: ANY OTHER QUESTIONS?
7	OKAY.
8	AND THEN THE OTHER ITEM, WHICH I DON'T
9	THINK IS QUITE READY FOR PRIME TIME, BUT I DON'T
10	KNOW. YOU MAY HAVE SOME DISCUSSION ON IT, BOB,
11	MAYBE STAFF. BUT ONE OF THE THINGS THAT WE HEARD AT
12	THE LAST GRANTS WORKING GROUP MEETING FROM SEVERAL
13	OF THE SCIENTISTS INVOLVED WAS JUST KIND OF NOODLING
14	AROUND HOW VALUABLE THEY THOUGHT OUR NEW FACULTY
15	AWARDS WERE AND HOW THAT MIGHT BE AN INTERESTING
16	GRANT ROUND FOR US TO REPEAT.
17	SO I THINK WE'RE STILL IN THE EARLY PHASES
18	OF HAVING CONVERSATIONS ABOUT THIS, BUT I KNOW ONE
19	OF THE THINGS WE'RE TALKING ABOUT IS THE SCIENTIFIC
20	STAFF COMING FORWARD WITH SOME SORT OF PRODUCTIVITY
21	FROM THOSE GRANTEES. I KNOW JUST FROM WHAT I'VE
22	SEEN IN TERMS OF FOLLOW-ON GRANTS AND PUBLICATIONS
23	AND IN THE MEDIA, SEVERAL OF THOSE INVESTIGATORS
24	HAVE BEEN EXTRAORDINARILY PRODUCTIVE. AND CERTAINLY
25	IT'S ONE OF THE FEW WAYS WE CAN ACTUALLY CREATE

1	ADDITIONAL CAPACITY IN CLINICAL SPACE, ESPECIALLY
2	WITH THOSE CLINICIAN-SCIENTISTS.
3	I DON'T KNOW IF OTHER MEMBERS OF THE
4	COMMITTEE WOULD LIKE TO OPINE ON THIS. OBVIOUSLY
5	THIS ISN'T SOMETHING WE WOULD DO IN THE NEAR TERM.
6	THERE'S CONSIDERABLE LEAD-TIME THAT WOULD BE
7	REQUIRED, FIRST OF ALL, BECAUSE THESE ARE NOT PREAP
8	PROCESS SCREENED APPLICATIONS. INSTITUTIONS HAVE TO
9	PUT FORTH THE NOMINATIONS. THAT'S HOW WE LIMIT
10	THEM. PLUS INSTITUTION SUPPORT IS A BIG COMPONENT
11	OF THE EVALUATION OF THESE GRANTS. SO THE
12	INSTITUTIONS HAVE TO BE PREPARED TO SUPPORT SOME OF
13	THEIR RISING STARS. I DON'T KNOW, BOB, OR ANYONE
14	ELSE.
15	CHAIRMAN KLEIN: JEFF
16	CHAIRMAN SHEEHY: ALAN HAS A COMMENT.
17	MS. SAMUELSON: I DO TOO.
18	CHAIRMAN KLEIN: I'D LIKE TO ALAN, JUST
19	BRIEFLY, I THINK YOU DESCRIBED TO ME A COUPLE OF
20	OPTIONS. AND MAYBE YOU COULD SUMMARIZE THOSE. AND
21	COULD YOU ALSO COMMENT IN TERMS OF FOCUSED SUPPORT
22	FOR CLINICAL FACULTY IN TERMS OF YOUR APPRAISAL OF
23	WHERE WE ARE AND HAVING SUFFICIENT CLINICAL FACULTY
24	TO REALLY MAKE THE TRANSITION TO THE PATIENTS?
25	DR. TROUNSON: THANKS, BOB. I'M VERY
	21

1	SUPPORTIVE OF THIS IN AT LEAST PART. AS YOU KNOW,
2	WE'RE REALLY TRYING TO GET FOCUSED ON THE
3	TRANSLATION AND MOVING OUR OPPORTUNITY TO THE CLINIC
4	BECAUSE THIS IS REALLY THE PRIMARY GOAL OF THE
5	THINGS THAT WE NEED TO DEMONSTRATE TO CALIFORNIA,
6	THAT THE DISCOVERIES ARE CAPABLE OF ACTUALLY GETTING
7	INTO THE CLINIC. SO ONE OF THE WELL, THERE ARE
8	SEVERAL ISSUES.
9	WITH THE STRAIGHT ACADEMIC FACULTY AWARDS,
10	THERE IS A BIT OF LEAKAGE, IF YOU LIKE. 40 PERCENT
11	OF THE FACULTY DON'T ACTUALLY GET FOCUSED DIRECTLY
12	ON THE THINGS THAT WE'RE SPECIFICALLY INTERESTED IN
13	BECAUSE THEY HAVE FACULTY AWARDS, AND THEY CAN
14	BASICALLY DO PRETTY MUCH WHAT THEY LIKE WITHIN THE
15	GENERAL AREA.
16	BUT IF WE WERE ABLE TO FOCUS SOME OF THIS
17	CAPACITY ON WHERE WE'RE GOING, PARTICULARLY BRINGING
18	M.D./PH.D.S THROUGH TO THE TRANSLATIONAL CLINICAL
19	PART OF THE PROGRAMS, I THINK WE WOULD BE MUCH
20	OUR PROGRAM WOULD BE MUCH MORE EFFECTIVE OR AT LEAST
21	WOULD GIVE US STRONGER DEPTH.
22	ONE OF THE THINGS THAT PAT OLSON HAD
23	SUGGESTED TO US, THAT WE MIGHT BE ABLE TO
24	INCORPORATE THIS PROJECT OR THESE AWARDS WITHIN THE
25	TRANSLATIONAL PROGRAM, FOR EXAMPLE. YOU COULD IF

1	YOU REALLY WANTED TO DO IT IN THE BASIC SCIENCE
2	PROGRAM, IF YOU WISHED AS WELL, BUT I LIKE THE IDEA
3	OF CREATING MORE OF SOME OF THESE YOUNG SCIENTISTS
4	WHO ARE REALLY MAKING THE PACE, WHO ARE REALLY
5	HAVING A BIG IMPACT. YOU CAN LOOK AT REALLY QUITE A
6	NUMBER OF THEM THROUGH CALIFORNIA. AND JUST TO
7	MENTION ONE, CATRIONA JAMIESON, CAME THROUGH ON ONE
8	OF THESE PROGRAMS. IF WE COULD FIND PEOPLE LIKE HER
9	AND OTHERS THAT ARE REALLY VERY PRODUCTIVE AT THAT
10	SPACE, WE'RE LIKELY TO HAVE EVEN MORE IMPACT ON THE
11	DIRECT OPPORTUNITIES FOR CLINICAL DEVELOPMENTS. AND
12	ALSO THOSE PEOPLE CARRY ON TO THE CLINIC, OF COURSE.
13	SO THIS IS THE SPACE IN WHICH WE'RE TRYING
14	TO BE AS ACTIVE AS POSSIBLE. THAT'S WHY WE GOT
15	ELLEN FEIGAL ON THE TEAM HERE. WE'RE ACTUALLY
16	PUSHING HARD IN THIS DIRECTION. SO I LIKE THE IDEA
17	OF THE FOCUS OF THOSE TRANSLATION AWARDS DRAWING THE
18	CAPACITY OF SOME OF THESE YOUNG M.D./PH.D. PEOPLE
19	THROUGH INTO THE PROGRAM.
20	IF YOU LOOK WORLDWIDE AT THE M.D./PH.D.
21	PROGRAMS, THEY'RE THE ONES THAT ARE KIND OF
22	SUFFERING UNFORTUNATELY IN THE FUNDING SHORTFALLS
23	BECAUSE BASIC SCIENCE TENDS TO GET CONTINUED
24	FUNDING, BUT THERE'S NOT NECESSARILY A FOCUS ON
25	HELPING THESE M.D./PH.D. PEOPLE SORT OF DRIVE

1	THROUGH THAT SPACE.
2	SO I THINK WE WOULD OURSELVES END UP BEING
3	MORE COMPETITIVE IN THE SPACE AS WELL, BUT I THINK
4	OUR RESOURCES WOULD BE STRONGER FOR THAT. AND I
5	DON'T THINK WE WOULD HAVE THE SORT OF LEAKAGE OUT
6	INTO STEM CELL AREAS THAT THESE PEOPLE WORK ON, BUT
7	NOT NECESSARILY THE ONES THAT WE'RE REALLY KEEN
8	ABOUT.
9	CHAIRMAN KLEIN: THANK YOU. I APPRECIATE
10	THAT.
11	MS. SAMUELSON: I'D LIKE TO UNDERSTAND
12	THAT JUST A LITTLE BIT BETTER. I DON'T WANT TO TAKE
13	A LOT OF TIME WITH IT. SO THAT MEANS THAT 40
14	PERCENT OF THE FTE, LET'S SAY, OF NEW FACULTY THAT
15	WE'RE FUNDING, CAN YOU GIVE AN EXAMPLE OF WHAT IT IS
16	THEY SPEND THEIR TIME ON THAT WE'RE PAYING FOR?
17	DR. OLSON: LET ME ELABORATE A LITTLE BIT.
18	AS YOU MAY OR MAY NOT RECALL, THE NEW FACULTY
19	AWARDS, THE IDEA WAS ESSENTIALLY THE FOCUS WAS ON
20	CAREER DEVELOPMENT. AND WE DID NOT SPECIFY AT ALL,
21	AS WE DO IN ALMOST, NOT ALMOST, IN EVERY OTHER RFA,
22	THAT THE FOCUS BE HUMAN STEM CELL WORK. SO AS A
23	CONSEQUENCE OF THAT, AND WE HAD BOTH THE
24	PHYSICIAN/SCIENTISTS AND THE SCIENTISTS, BUT WE
25	HAD THE IDEA WAS BRING IN THE PEOPLE AT THE START

1	OF THEIR CAREERS WITHIN THE FIRST FEW YEARS OF
2	INDEPENDENT OF THEIR FIRST INDEPENDENT
3	APPOINTMENTS TO JUST GET THEM ENGAGED IN STEM CELLS.
4	WE DID NOT ASK THAT THEY FOCUS ON HUMAN.
5	SO 15 PERCENT OF OUR AWARDS ARE, STRICTLY
6	SPEAKING, STEM CELL MODEL SYSTEMS. THERE'S NO HUMAN
7	WORK AT ALL. OVER 50 PERCENT OF THE AWARDS ARE A
8	COMBINATION OF HUMAN AND MODEL SYSTEMS, BUT OF THOSE
9	TO DATE, HALFWAY THROUGH THE AWARDS, HALF OF THOSE
10	PEOPLE ARE ONLY FOCUSED ON MODEL SYSTEMS.
11	NOW, THIS WAS PART OF THIS. THIS WAS
12	CONTRIBUTING BASICALLY TO FUNDAMENTAL KNOWLEDGE
13	WITHIN THE STEM CELL FIELD. I THINK THE STRATEGIC
14	QUESTION THAT THIS GROUP NEEDS TO ADDRESS AS WE MOVE
15	FROM ESSENTIALLY OUR PHASE I BUILDING THIS KIND OF
16	INTELLECTUAL INFRASTRUCTURE IS MORE WE'RE MOVING
17	MORE TO A STAGE 2 FOCUS ON MOVING THINGS TO THE
18	CLINIC AND TO THERAPY DEVELOPMENT. AND SO, WHEREAS,
19	I THINK WE ALL APPRECIATE THE IDEA OF ESSENTIALLY
20	PROVIDING AN OPPORTUNITY FOR PEOPLE WHO, BECAUSE OF
21	THEIR TRACK RECORD, THEY HAVEN'T BEEN DOING THIS FOR
22	20 YEARS, WE WANT TO PROVIDE AN OPPORTUNITY TO STILL
23	ACKNOWLEDGE A CHANCE FOR SOME OF THE BEST TO
24	PARTICIPATE IN OUR PROGRAM, IN OUR PROJECT-FOCUSED
25	PROGRAMS THAT ARE ON THE TRACK TO, SAY, THERAPY

1	DEVELOPMENT.
2	SO THIS IS WHERE WE PICK UP THE CONCEPT
3	THAT ALAN INTRODUCED OF THE NOTION OF IN THE CONCEPT
4	IN THE CONTEXT OF, SAY, A BASIC BIOLOGY AWARD, IN
5	THE CONTEXT OF AN EARLY TRANSLATIONAL AWARD, DO WE
6	SPECIFICALLY STATE THAT WE WANT TO ENCOURAGE SOME
7	NEW FACULTY HERE? WE ALLOW FOR CHECK BOXES ON THE
8	APPLICATION THAT STATE THEY'RE WITHIN THIS YEAR. WE
9	ASKED FOR A LETTER FOR THOSE. WE ALLOW PERHAPS
10	IN PROGRAMMATIC DISCUSSION, WE SAY WE WOULD LIKE TO,
11	IF THERE'S SOME GOOD ONES, TO HAVE A PERCENTAGE OF
12	PEOPLE. I THINK WE'VE ALL SAT IN REVIEWS WHERE
13	WE'VE HEARD THIS IS AN INTERESTING IDEA, BUT THIS
14	PERSON IS A LITTLE BIT JUNIOR OR SOMETHING. SO WE'D
15	LIKE TO GIVE THESE PEOPLE A CHANCE, BUT WE'D LIKE TO
16	DO IT IN THE CONTEXT OF OUR FOCUSED PROGRAMS.
17	AND SO THAT'S WHAT I WOULD SUGGEST WE
18	CAN YOU KNOW, I'D BE CURIOUS TO HEAR SOME
19	DISCUSSION ON THIS BECAUSE I THINK IT'S A QUESTION
20	OF CAREER DEVELOPMENT OR THERAPY DEVELOPMENT, OR IS
21	THERE A WAY TO SORT OF MERGE THE CONCEPTS?
22	CHAIRMAN KLEIN: SO, JOAN, THIS IS BOB.
23	YOU MIGHT REMEMBER THAT NICHOLAS WADE CRITICIZED US
24	FOR NOT HAVING THE FORESIGHT TO INVEST SOME IN MODEL
25	SYSTEMS LIKE ZEBRAFISH SPECIFICALLY. HE DIDN'T

1	ACTUALLY REALIZE THAT WE'D ACTUALLY FUNDED STUDIES
2	OF REGENERATIVE NATURE OF ZEBRAFISH AS PART OF THE
3	WORK THAT HAD ALREADY BEEN APPROVED. BUT IN TERMS
4	OF CRITICAL ALLOCATION OF FUNDS, I THINK AT THIS
5	STAGE WE'RE LOOKING AT THE FACT THAT TO MOVE
6	DOWNSTREAM TOWARDS PATIENTS, THAT POTENTIALLY WE
7	HAVE MORE OF A SHORTFALL WHERE WE CAN PLAY A
8	CRITICAL ROLE IN THE M.D./PH.D. PROGRAMS AND THE
9	CLINICAL FELLOWS PROGRAM; WHEREAS, DR. OLSON SAYS
10	PROVIDING AN INCENTIVE WITHIN OTHER GRANT PROGRAMS
11	TO ADVANCE PEOPLE WITH THAT TYPE OF A BACKGROUND
12	COMING UP THROUGH THE RANKS, THEN WE CAN IN ADDING
13	TO WORK THAT THE NIH OR OTHERS CAN ALREADY FUND.
14	CHAIRMAN SHEEHY: DR. OLSON.
15	DR. OLSON: I JUST WANT TO ELABORATE ON
16	THE STATEMENT THAT BOB JUST MADE. I DID LOOK UP THE
16 17	THE STATEMENT THAT BOB JUST MADE. I DID LOOK UP THE NIH FUNDING ALLOCATION FOR 2010 TO DIFFERENT AREAS
17	NIH FUNDING ALLOCATION FOR 2010 TO DIFFERENT AREAS
17 18	NIH FUNDING ALLOCATION FOR 2010 TO DIFFERENT AREAS OF STEM CELL RESEARCH. AND I JUST WANT TO JUST LET
17 18 19	NIH FUNDING ALLOCATION FOR 2010 TO DIFFERENT AREAS OF STEM CELL RESEARCH. AND I JUST WANT TO JUST LET THE COMMITTEE KNOW AND THE PUBLIC WHO IS INTERESTED
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17 18 19 20 21 22	NIH FUNDING ALLOCATION FOR 2010 TO DIFFERENT AREAS OF STEM CELL RESEARCH. AND I JUST WANT TO JUST LET THE COMMITTEE KNOW AND THE PUBLIC WHO IS INTERESTED THAT IN NONEMBRYONIC, NONHUMAN STEM CELL RESEARCH, THAT THE NIH ACTUALLY SPENT LAST YEAR, THIS IS NOT ALLOCATED, THIS IS THEIR BUDGET FOR 2010, THEY SPENT \$570 MILLION ON THAT PARTICULAR CLASS OF STEM CELL

1	ANOTHER \$74 MILLION. SO ESSENTIALLY THERE WAS \$640
2	MILLION SPENT ON NONEMBRYONIC, NONHUMAN.
3	BOB, I TAKE YOUR POINT, THERE'S ALWAYS
4	GOING TO BE GREAT RESEARCH DONE IN ALL SORTS OF
5	AREAS AND ZEBRAFISH. AND THE QUESTION IS WHERE DO
6	WE FOCUS OUR PRIORITIES? WELL, THE FUNDING IS ALL I
7	WOULD SAY. AND SO THESE ARE AREAS THAT DO HAVE SOME
8	FUNDING. NIH SPENDS ACTUALLY QUITE A LOT OF MONEY
9	ON STEM CELL RESEARCH. THAT'S PROBABLY
10	PREDOMINANTLY INVESTIGATOR-INITIATED RESEARCH.
11	DR. BRYANT: COULD I JUST SAY SOMETHING?
12	AM I ALLOWED TO TALK? AS A REGENERATION BIOLOGIST,
13	I WOULD JUST SAY THAT NIH MAY CLASSIFY THE GRANTS
14	THAT THEY HAVE FUNDED IN A WAY THAT MAKES IT SOUND
15	LIKE THEY PUT A LOT OF MONEY INTO THIS, BUT I HAVE
16	TO TELL YOU WORKING ON ONE OF THE ONLY MODEL SYSTEMS
17	THAT REGENERATES PERFECTLY, THEY ARE VERY DERELICT
18	IN FUNDING IN THAT AREA. AND IT'S BECAUSE IT'S A
19	DIFFICULT SYSTEM, AND IT'S NOT AS EASY AS SOMETHING
20	LIKE ZEBRAFISH, WHICH REALLY DOES NOT HAVE THE
21	POWERS THAT SALAMANDERS DO.
22	SO I'M JUST SAYING THAT BECAUSE NIH SAYS
23	IT FUNDS REGENERATION, AND REALLY THE WAY TO
24	UNDERSTAND IT IS TO TAKE A SYSTEM THAT REALLY DOES
25	REGENERATE AND FIGURE IT OUT. AND EVERYTHING ELSE
	20

1	IS BASICALLY DESTROYING YOU'RE JUST THROWING
2	DARTS AT THE PROBLEM BECAUSE IF IT DOESN'T HAPPEN
3	NATURALLY, MAKING IT HAPPEN IS GOING TO BE VERY
4	DIFFICULT. I'M SORRY TO RAMBLE, BUT I JUST FEEL
5	I'VE ACTUALLY FELT THAT CIRM HAS MISSED THE BOAT A
6	LITTLE BIT BY NOT INCLUDING SOME OF THOSE SYSTEMS
7	THAT REALLY COULD SHOW THE WAY TO HOW TO PROCEED
8	WITH HUMANS RATHER THAN JUST FUMBLING IN THE DARK.
9	AND WHICH FUMBLING IN THE DARK MAY GET US THERE, BUT
LO	I THINK UNDERSTANDING SOMETHING DEEPLY IS ALSO A WAY
L1	TO GO.
L2	MS. SAMUELSON: I'D LIKE TO GO ON RECORD
L3	AGAINST FUMBLING IN THE DARK. I'M SO GLAD YOU
L4	SPOKE. I THINK MAYBE WE NEED TO ADDRESS THIS AT A
L5	BOARD MEETING SO THAT WE HAVE A POLICY ON IT BECAUSE
L6	IT SEEMS TO ME IF WE'RE GOING TO TRY TO GET
L7	THERAPEUTIC BREAKTHROUGHS AS SOON AS HUMANLY
L8	POSSIBLE, WE SHOULD BE USING THE BEST MECHANISMS TO
L9	DO IT. IF IT'S ZEBRAFISH, LET'S DO IT.
20	CHAIRMAN KLEIN: I THINK WHAT SHE'S SAYING
21	IS THERE'S ACTUALLY BETTER MODEL SYSTEMS THAN
22	ZEBRAFISH. THE ISSUE IS DO WE MAINTAIN ADEQUATE
23	BASIC SCIENCE RESEARCH THAT COVERS MODEL SYSTEMS AT
24	THE SAME TIME WE'RE PROVIDING SUFFICIENT ALLOCATION
25	TO THE TRANSLATIONAL CRITICAL SHORTAGES, WHICH MAY

1	BE M.D./PH.D. TRAINED SCIENTISTS WHO ARE GOING TO
2	MOVE TRANSLATION TO PATIENTS. THIS IS A FUNDAMENTAL
3	STRATEGIC DISCUSSION THAT I THINK WE'RE GOING TO
4	NEED ANOTHER SCIENCE SUBCOMMITTEE AS WELL AS THEN A
5	BOARD DISCUSSION ON. JEFF, WHAT'S YOUR
6	MS. SAMUELSON: I THINK WE DO NEED THAT.
7	CHAIRMAN SHEEHY: I ACTUALLY THANKS,
8	BOB. THAT'S WHERE I WAS HEADED. I THINK STAFF AT A
9	LATER DATE, WE'VE GOT A LOT GOING ON BEFORE THE NEXT
10	MEETING, IS GOING TO COME TO US WITH AN ANALYSIS OF
11	THE PRODUCTIVITY FOR THIS ROUND. I DO THINK THAT
12	THERE'S SOME EMERGING CONSENSUS THAT THERE'S
13	ENORMOUS UTILITY IN GETTING SOME M.D./PH.D. MAYBE
14	TRYING TO HAVE THAT REPEAT IN SOME FASHION IN THAT I
15	KNOW, AT LEAST IN THE HIV FIELD, THERE'S A CONSTANT
16	SHORTAGE OF CLINICIAN/SCIENTISTS. WE TALKED ABOUT
17	THIS. IT'S KIND OF REPEATING WHAT WE SAID WHEN WE
18	DID THIS ORIGINALLY. CLINICIANS, THERE'S
19	COMPETITION FOR THEIR TIME WITH THEIR CLINICAL WORK.
20	IF YOU CAN'T CREATE SPACE AND CREATE SOME
21	ATTRACTIVENESS FOR THEM TO GET INTO RESEARCH SPACE,
22	A LOT OF REALLY TALENTED PEOPLE REALLY EITHER DON'T
23	GET INTO IT OR DON'T STAY WITH IT.
24	AND CREATING THAT CAPACITY, ESPECIALLY IN
25	THIS EMERGING CLINICAL FIELD, IS OF VALUE TO US. I

1	THINK THERE'S SOME SENSE THAT THAT'S A GOOD IDEA.
2	AND THE MECHANICS FOR DOING THAT CAN EMERGE AS WE
3	HAVE THIS LARGER DISCUSSION ABOUT WHAT'S HAPPENED IN
4	THE FIRST TWO ROUNDS AND HOW WE DO IT, IT CAN ALL
5	KIND OF COME TOGETHER.
6	BUT I THINK IT'S GOOD THAT THIS HAS COME
7	UP BECAUSE I DON'T THINK WE REALLY TALKED ABOUT
8	DOING THIS ROUND AGAIN, AND IT'S KIND OF FALLEN OFF
9	THE RADAR. AND I THINK THAT THAT'S A VALUE FROM THE
10	GRANTS WORKING GROUP THAT THESE REVIEWERS HAVE PUT
11	THIS FORWARD FOR US.
12	ARE THERE ANY OTHER FURTHER COMMENTS FROM
13	BOARD MEMBERS?
14	MS. SAMUELSON: YEAH. THIS IS JOAN. AND
15	MAYBE I'M JUST BEATING A DEAD HORSE. BUT IT SEEMS
16	TO ME THAT IT'S IMPORTANT THAT WE DEVOTE TIME, NOT
17	SO MUCH JUST TO GRANT ROUNDS, BUT TO A STRATEGIC
18	DIRECTION AND THE ELEMENTS OF IT SO THAT WE'RE
19	CONFIDENT AND WE CAN SAY THIS, THAT WE'RE FUNDING
20	WHAT THE SCIENTISTS TELL US ARE THE MECHANISMS THAT
21	ARE GOING TO MOST EFFECTIVELY, MOST SPEEDILY DEVELOP
22	THERAPIES. AND THAT MIGHT NOT NECESSARILY BE MORE
23	MONEY FOR THIS OR THAT KIND OF FUNDING MECHANISM.
24	THOSE ARE TWO DIFFERENT THINGS AS I'M THINKING ABOUT
25	IT.

1	CHAIRMAN SHEEHY: OKAY.
2	CHAIRMAN KLEIN: I'D LIKE TO THANK DR.
3	BRYANT FOR HER COMMENTS. AND I'D LOVE TO GET SOME
4	REFERENCES ON SOME PAPERS ON MODEL SYSTEMS, DR.
5	BRYANT, AS TO WHICH ARE WHERE THE GREATEST PROMISE
6	MIGHT BE AND WHICH MODEL SYSTEMS.
7	DR. BRYANT: BE HAPPY TO DO THAT. I THINK
8	WHAT I'M TRYING TO SAY IS THAT THE APPROACHES THAT
9	ARE GOING STRAIGHT FOR THE DISEASE MODELS DIRECTLY
10	ARE WE MAY GET SOME LOW HANGING FRUIT THERE. BUT
11	IF WE REALLY WANT TO UNDERSTAND HOW TO CREATE
12	REGENERATIVE MEDICINE IN HUMANS, WE HAVE TO BE ABLE
13	TO UNDERSTAND HOW IT'S DONE IN A SIMILAR ORGANISM.
14	SO EVEN ZEBRAFISH DON'T COME AS CLOSE AS THEY SHOULD
15	IN TERMS OF THEIR ABILITIES. SO, YES, I'D BE HAPPY
16	TO. SHALL I JUST SEND THEM TO YOU, BOB, OR WHAT?
17	CHAIRMAN KLEIN: SURE. JEFF, WOULD YOU
18	LIKE TO SEE IT AS WELL?
19	CHAIRMAN SHEEHY: SURE. THAT'D BE GREAT.
20	THANK YOU, DR. BRYANT.
21	MS. SAMUELSON: SO WOULD I.
22	CHAIRMAN SHEEHY: IS THERE ANY OTHER
23	MR. TORRES: I JUST WANT TO MAKE SURE. I
24	KNOW DR. BRYANT DIDN'T MEAN TO SAY THAT, BUT WE ARE
25	NOT FUMBLING IN THE DARK HERE. I DO THINK THAT
	42

1	WE'RE HEADED IN THE RIGHT DIRECTION. HOW WE REFINE
2	THAT DIRECTION, I THINK, IS WHAT DR. BRYANT WAS
3	SAYING. THERE'S NO QUESTION THAT YOU DO NEED
4	CONSISTENT REDIRECTION AND REFINEMENT OF WHERE WE'RE
5	GOING. BUT FROM MY EXPERIENCE, I'VE ONLY BEEN HERE
6	TWO YEARS, I JUST DON'T SEE THAT WE'RE FUMBLING IN
7	THE DARK.
8	DR. BRYANT: NO. WE'RE APPROACHING A
9	PROBLEM IN A WAY THAT IS LET'S SEE IF THIS WORKS
10	RATHER THAN LET'S UNDERSTAND IT AND THEN FIGURE OUT
11	HOW TO FIX IT. I TAKE BACK THE FUMBLING IN THE DARK
12	BECAUSE OBVIOUSLY WE'RE NOT. IT'S JUST DIFFERENT
13	PHILOSOPHIES ABOUT HOW TO APPROACH THINGS. I THINK
14	A COMBINATION OF BOTH IS VERY APPROPRIATE, BUT I
15	SEE MAYBE IT'S APPROPRIATE THAT WE LEAVE BEHIND
16	THE UNDERSTANDING PART BECAUSE WE ONLY HAVE A
17	LIMITED AMOUNT OF TIME AND MONEY, BUT I DO THINK
18	THAT WE CAN GET A LITTLE BIT TOO FAR OUT ON ONE
19	PARTICULAR MODE. OF THE STUDIES THAT HAVE BEEN DONE
20	SO FAR ON ADDING CELLS BACK TO BRAINS AND SO FORTH,
21	ALTHOUGH THEY HAVE A BENEFICIAL EFFECT, IT'S NOT AS
22	IF THEY INTEGRATE INTO THE SYSTEM AND CREATE.
23	SO THERE ARE SOME EFFECTS WE DON'T
24	REALLY CAN'T REALLY ANTICIPATE WHAT THE RESULTS
25	ARE GOING TO BE IS WHAT I'M TRYING TO SAY. AND IF
	43

1	IT'S POSITIVE, THAT'S GREAT. BUT IT'S NOT
2	NECESSARILY WHAT YOU THINK IF YOU DON'T HAVE A
3	DEEPER UNDERSTANDING OF THE SITUATION.
4	CHAIRMAN KLEIN: SO, DR. BRYANT, THIS IS
5	BOB. I THINK, IN FACT, WE HAVE FUNDED BOTH, AND THE
6	ISSUE HERE STRATEGICALLY TO THINK ABOUT IS WHAT
7	ALLOCATION WE KEEP TO THESE MODEL SYSTEMS WHILE
8	WE'RE ALSO TRYING TO FILL CRITICAL SHORTFALLS IN THE
9	CLINICAL POSITIONS THAT WE NEED TO GO FORWARD. AND
10	SO THE STRATEGIC PAPERS THAT YOU CAN GIVE TO US
11	WOULD BE VERY HELPFUL TO INFORM US.
12	MS. SAMUELSON: I THINK THAT'S RIGHT.
13	BECAUSE WE HAVE THE EXTERNAL ADVISORY PANEL TELLING
14	US WE'RE SUPPOSED TO BE MUCH MORE PROACTIVE IN THIS
15	NEXT PHASE, AND I'M NOT SURE WHAT THAT MEANS IN THIS
16	CONTEXT.
17	CHAIRMAN SHEEHY: SO I THINK WE'VE HAD A
18	NICE DISCUSSION ON THIS. AND UNLESS THERE'S
19	REALLY WE ARE GOING TO TALK ABOUT THIS AGAIN.
20	AND UNLESS THERE'S ANYTHING ADDITIONAL, I THINK WE
21	HAVE PRETTY MUCH COVERED OUR AGENDA.
22	MR. ROTH: THANK YOU, JEFF.
23	CHAIRMAN SHEEHY: I THINK WE'RE ADJOURNED.
24	(THE MEETING WAS THEN CONCLUDED AT
25	02:59 P.M.)

#### 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 TELEPHONIC PROCEEDINGS BEFORE THE SCIENCE SUBCOMMITTEE TO THE INDEPENDENT CITIZEN'S OVERSIGHT COMMITTEE OF THE CALIFORNIA INSTITUTE FOR REGENERATIVE MEDICINE IN THE MATTER OF ITS REGULAR MEETING HELD ON APRIL 26, 2011, 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 BARRISTER'S REPORTING SERVICE 1072 BRISTOL STREET SUITE 100 COSTA MESA, CALIFORNIA (714) 444-4100