

BEFORE THE
IP TASK FORCE
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: UCSD
9500 GILMAN DRIVE
EUCALYPTUS POINT
LA JOLLA, CALIFORNIA

DATE: THURSDAY, APRIL 27, 2006
8:15 A.M.

REPORTER: BETH C. DRAIN, CSR
CSR. NO. 7152

BRS FILE NO.: 75296

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1 THURSDAY, APRIL 27, 2006

2

3 MS. KING: WE'LL CALL THE MEETING TO ORDER
4 THEN.

5 CHAIRMAN PENHOET: WE HAVE A TRANSCRIBER AT
6 WORK, SO WE CAN START THE MEETING. THIS MEETING IS
7 BEING CONDUCTED IN FIVE PLACES. WE'RE AT UC SAN DIEGO,
8 AND WE THANK OUR HOST FOR PROVIDING THIS SPACE, AT
9 NUVELO CORPORATION, IN LOS ANGELES, AND IN CHICO.
10 WE'LL DO A ROLL CALL SO WE KNOW EXACTLY WHO FROM THE
11 CIRM IS PRESENT.

12 MS. KING: SUSAN BRYANT.

13 DR. BRYANT: HERE.

14 MS. KING: MICHAEL GOLDBERG.

15 MR. GOLDBERG: HERE.

16 MS. KING: SHERRY LANSING. TED LOVE.

17 UNIDENTIFIED SPEAKER: HE STEPPED OUT. HE'LL
18 BE RIGHT BACK.

19 MS. KING: ED PENHOET.

20 CHAIRMAN PENHOET: HERE.

21 MS. KING: PILL PIZZO. JOHN REED.

22 DR. REED: HERE.

23 MS. KING: JEFF SHEEHY. OSWALD STEWARD.

24 DR. STEWARD: HERE.

25 MS. KING: JANET WRIGHT.

1 DR. WRIGHT: HERE.

2 CHAIRMAN PENHOET: GOOD. SO THANK YOU ALL.
3 SORRY WE'RE LATE IN GETTING STARTED, BUT WE WILL
4 PROCEED WITH TODAY'S SESSION. AS ALL OF YOU KNOW, I
5 THINK THIS IS OUR SECOND MEETING IN OUR ATTEMPTS TO
6 DEFINE AN INTELLECTUAL PROPERTY POLICY FOR GRANTS,
7 CONTRACTS, LOANS, WHATEVER FORM OF PAYMENTS WE MIGHT
8 MAKE TO INDUSTRIAL ORGANIZATIONS.

9 AND WE HAD A SORT OF VERY, I THINK,
10 PRODUCTIVE FIRST MEETING AT STANFORD NOW SIX WEEKS AGO,
11 I GUESS IT WAS. AND THIS ROUND WITH THE SECOND MEETING
12 IN SAN DIEGO. WE PLAN TO HAVE A THIRD MEETING PROBABLY
13 IN SACRAMENTO SOMETIME LATER IN THE SUMMER. SO WE'RE
14 MOVING THIS PROCESS ALONG. WE'RE VERY INDEBTED TO THE
15 PEOPLE WHO HAVE AGREED TO COME AND SHARE THEIR
16 PERSPECTIVES ON THIS ISSUE WITH US TODAY.

17 WE HAVE A GROUP OF PRESENTERS, FIVE IN TOTAL,
18 AS YOU KNOW. AND THE FIRST OF THOSE IS JOHN SIMPSON.
19 JOHN IS THE STEM CELL PROJECT DIRECTOR FOR THE
20 FOUNDATION FOR TAXPAYER AND CONSUMER RIGHTS. SO WE'LL
21 JUST GET RIGHT INTO JOHN'S PRESENTATION. JOHN, THANK
22 YOU FOR JOINING US TODAY AND FOR PUTTING YOUR
23 PRESENTATION TOGETHER.

24 A REMINDER, WE HAD TO WAIT FOR A TRANSCRIBER
25 BECAUSE ALL THESE MEETINGS ARE TRANSCRIBED, SO WE GET

1 ALL OF THE COMMENTS AND WE KEEP ALL OF THE POWERPOINT
2 PRESENTATIONS AS WELL. WE HAVE FIVE, SIX PEOPLE IN THE
3 AUDIENCE HERE THIS MORNING, ONE OF WHOM IS OUR NEWEST
4 ICOC MEMBER, DUANE ROTH, WHO MOST OF THE LOCAL PEOPLE
5 KNOW THANK YOU FOR JOINING US TODAY.

6 MR. SIMPSON: THANK YOU, VERY MUCH, DR.
7 PENHOET. IS THIS MICROPHONE WORKING LOUD ENOUGH SO
8 THAT EVERYONE CAN HEAR IT OKAY? I'M WIRELESSLY HOOKED
9 UP HERE. I'M NOT QUITE SURE. WHAT ABOUT THE REMOTE
10 SITES? ARE YOU ABLE TO HEAR ON THE TELEPHONES?

11 (ALL SITES RESPOND AFFIRMATIVELY.)

12 MR. SIMPSON: WELL, FIRST OF ALL, I REALLY
13 WANTED TO THANK THE CHAIRMAN AND THE FIRST DEPUTY FOR
14 GIVING ME THE OPPORTUNITY TO PRESENT OUR VIEWS. WE DID
15 ISSUE A REPORT BACK IN JANUARY WHICH WAS CALLED
16 "AFFORDABILITY, ACCESSIBILITY, AND ACCOUNTABILITY FOR
17 CALIFORNIA STEM CELL RESEARCH." THAT WAS AIMED MORE
18 TOWARDS THE NONPROFITS. WE HAVE COPIES OF THAT HERE
19 AVAILABLE TO THE BOARD MEMBERS. WHAT I ALSO THEN
20 PREPARED IS A WHITE PAPER TODAY THAT IS ALSO AVAILABLE.
21 IT'S THE BASIS FOR THIS PRESENTATION. I THINK YOU'VE
22 GOT THAT AS WELL.

23 I JUST WOULD POINT OUT IF YOU HAVE STRONG
24 AGREEMENTS OR DISAGREEMENTS OR WHATEVER WITH WHAT I
25 MIGHT SAY, MY COORDINATES ARE THERE ON THE FIRST SLIDE.

1 I CAN BE REACHED AT JOHN@CONSUMERWATCHDOG.ORG. OUR
2 WEBSITE IS WWW.CONSUMERWATCHDOG.ORG, AND WE HAVE OUR
3 OWN SPECIAL STEM CELL PAGES AT STEMCELLWATCH.ORG.

4 DR. WRIGHT: JOHN, THIS IS JANET WRIGHT. IS
5 THE WHITE PAPER THAT YOU HAVE THERE TODAY ACCESSIBLE
6 THROUGH THE WEBSITE, OR ARE YOU GOING TO MAIL THAT OUT
7 TO THOSE OF US WHO AREN'T THERE? HOW DOES THAT WORK?

8 MR. SIMPSON: THAT WAS E-MAILED, I THINK, WAS
9 IT? IN TERMS OF THE SLIDES. I CAN SEND IT. IT WILL
10 BE AVAILABLE ON OUR WEBSITE. I HAVE SENT A COPY OF IT
11 TO CIRM.

12 DR. MAXON: I CAN DISTRIBUTE IT AGAIN.

13 DR. WRIGHT: GREAT. THANKS.

14 CHAIRMAN PENHOET: -- HERE EVERYBODY SHOULD
15 HAVE A COPY.

16 MR. SIMPSON: THE THING I FIRST WANT TO TALK
17 ABOUT WHO WE ARE. WE'RE A NONPROFIT, NONPARTISAN
18 CONSUMER WATCHDOG ORGANIZATION. WE WERE KNOWN FOR THE
19 PROPOSITION 103. OUR FOUNDER, HARVEY ROSENFELD, WROTE
20 THAT AND REFORMED THE INSURANCE INDUSTRY. WE'VE BEEN
21 INVOLVED RECENTLY IN HEALTHCARE, INSURANCE, ACCESS,
22 FIGHTING EXCESSIVE OIL PROFITS, CAMPAIGN FINANCE
23 REFORM, THAT SORT OF THING.

24 I THINK IT'S IMPORTANT TO UNDERSTAND THAT WE
25 DID NOT HAVE A POSITION ON PROPOSITION 71 WHEN IT WAS

1 PASSED. WHEN IT WAS PASSED, WE FELT THAT OUR ROLE WAS
2 TO MAKE SURE THAT THE PUBLIC BENEFIT PROMISES ARE KEPT
3 AND ENACTED. SO SOME ORGANIZATIONS HAVE BEEN AGAINST
4 IT FROM THE BEGINNING. WE STEPPED INTO IT ONCE
5 ESSENTIALLY IT WAS OVERWHELMINGLY APPROVED BY THE
6 PEOPLE.

7 AND A LITTLE BIT ABOUT MYSELF. I'M A FORMER
8 NEWSPAPERMAN, A LONG TIME WITH THE GANETT COMPANY IN
9 PLACES IN UPSTATE NEW YORK, BINGHAMTON, ITHACA, GUAM,
10 AND THEN MOST OF MY CAREER AT U.S.A. TODAY WHERE I
11 ENDED UP BEING DEPUTY EDITOR, WHICH WAS A GLORIFIED WAY
12 OF SAYING I RAN THE INTERNATIONAL EDITION. I DID A
13 LITTLE BIT OF CONSULTING WITH THE IRISH TIMES, THE
14 GLEANER IN JAMAICA, TAUGHT JOURNALISM IN IRELAND, ENDED
15 UP AT A SYNDICATE COMPANY IN L.A., AND NOW I'M WORKING
16 FOR THE FOUNDATION. I SHOULD STRESS I'M NOT A LAWYER.
17 SO I DON'T KNOW WHETHER THAT HELPS MY VIEW OF IP OR
18 HINDERS IT, BUT THAT IS A DISCLAIMER.

19 AT THE LAST HEARING WE HEARD WHAT I THOUGHT
20 WAS SORT OF AN EXAMPLE OF, YOU KNOW, DO IT OUR WAY OR
21 WE'RE GOING TO TAKE THE PETRIE DISHES AND GO HOME.
22 GENENTECH, ALTHOUGH THEY'RE NOT PROBABLY GOING TO BE
23 INVOLVED IN STEM CELL RESEARCH, TALKED ABOUT THE
24 BIOTECH INDUSTRY AND SAID, LOOK, WE'RE NOT GOING TO
25 ENGAGE IF THERE ARE APPROPRIATE PUBLIC BENEFIT

1 REQUIREMENTS ATTACHED TO THESE GRANTS.

2 BRAD MARGUS FROM PERLEGEN SCIENCES BASICALLY
3 SAID COMPANIES AREN'T GOING TO COME TO YOU ON OUR
4 KNEES. YOU WANT TO ENCOURAGE THE BEST CLASS OF PLAYERS
5 TO PARTICIPATE, NOT JUST THE FINANCIALLY DESPERATE.
6 WELL, I THINK THERE'S SOME FACTS THAT NEED TO BE
7 EXAMINED. LAST YEAR \$5.9 BILLION WENT OUT TO THE
8 BIOTECH INDUSTRY. A 120 MILLION OF THAT WAS FOR STEM
9 CELL RESEARCH. THERE WAS \$30 MILLION OF FEDERAL STEM
10 CELL MONEY. AND WHAT THAT REALLY MEANS IS, I THINK,
11 THAT WHEN CIRM STARTS PUTTING OUT \$300 MILLION A YEAR,
12 YOU'RE GOING TO BE THE BIGGEST FUNDER OF STEM CELL
13 RESEARCH IN THE WORLD PROBABLY. SO TO ME THAT MEANS,
14 YES, THE BIOTECH INDUSTRY REALLY IS KEENLY INTERESTED
15 IN THIS PUBLIC MONEY AND WANTS IT.

16 THE OTHER THING I THINK I JUST WANT TO ADD
17 ABOUT THE WHOLE VENTURE CAPITAL PICTURE, WHICH IS
18 IMPORTANT, IS THAT LOOMING OVER THE WHOLE INDUSTRY AND
19 CAUSING A GREAT AMOUNT OF UNCERTAINTY ON THE PART OF
20 VENTURE CAPITALISTS IN STEM CELLS ARE THE PATENTS ON
21 HUMAN EMBRYONIC STEM CELLS ON ALL OF THEM IN THE UNITED
22 STATES, THE PATENTS ONLY RECOGNIZED IN THE UNITED
23 STATES THAT ARE HELD BY THE WISCONSIN AREA
24 ALUMNI -- I'M SORRY -- WISCONSIN ALUMNI RESEARCH
25 FOUNDATION. THE WARF PATENTS REALLY ARE OUTRAGEOUS. I

1 MEAN THEY MEAN, FOR INSTANCE, THAT IF THE STEM CELL
2 RESEARCHER, LIKE JEANNE LORING, WHO'S IN THE AUDIENCE,
3 COMES UP WITH 20 NEW STEM CELL LINES, SHE CAN'T PROVIDE
4 THEM TO ANYBODY ELSE UNLESS WARF SAYS SO. AT LEAST
5 THAT'S LITERALLY WHAT IT MEANS.

6 PROPOSITION 71, THE PROMISES OF THAT. WELL,
7 WE ALL KNOW ABOUT THE PROMISES OF CURES AND HOPE AND
8 THAT, BUT THE OTHER THING WE'VE GOT TO REMEMBER WAS
9 THAT IN THE PROPOSITION ITSELF WERE SPECIFIC PROVISIONS
10 TO BENEFIT THE PEOPLE. I QUOTE HERE FROM SECTION III
11 OF THE PURPOSE AND INTENT OF PROP 71. ONE OF THEM WAS
12 TO PROTECT THE BENEFIT TO THE CALIFORNIA BUDGET BY
13 PROVIDING AN OPPORTUNITY FOR THE STATE TO BENEFIT FROM
14 ROYALTIES, PATENTS, AND LICENSING FEES THAT RESULT FROM
15 RESEARCH.

16 THE VOTE OVERWHELMINGLY PROVED THE CONCEPT OF
17 PROP 71. FIFTY-NINE PERCENT OF THE PEOPLE VOTED FOR
18 IT. THEY DID IT, THOUGH, WITH THE DIRECT UNDERSTANDING
19 AND BELIEF BOTH IN THAT SPECIFIC LANGUAGE AND IN THE
20 LANGUAGE OF VARIOUS ANALYSES THAT THE SUPPORTERS OF THE
21 PROPOSITION USED AT THE TIME THAT ESTIMATED THAT AN
22 AMOUNT FROM 6.4 BILLION TO 12.6 BILLION WOULD CAN BACK
23 TO THE STATE. SO WHAT I THINK IS THAT IT WAS AN
24 OVERWHELMING SUPPORT FOR PROP 71, BUT NOT A BLANK CHECK
25 FOR BIOTECH.

1 THE OTHER THING I THINK THAT'S KEY IN ALL OF
2 THIS IS THAT THE WAY THAT YOU CAN MAKE SURE THAT THE
3 PUBLIC BENEFIT IS MET IS THROUGH IP POLICY. IP POLICY
4 IS THE MECHANISM TO ASSURE THE PROMISES ARE KEPT.

5 NOW, YOU CAN TALK ABOUT PATENTS AND
6 COMMISSIONS AND ROYALTIES, AND YOUR EYES WILL GLAZE
7 OVER WHEN YOU START THINKING ABOUT HOW TO APPROACH IT
8 IN THAT DIRECTION. I LIKE TO USE SOME SIMPLE BUSINESS
9 MODELS, SOME SIMPLE SCENARIOS TO KIND OF UNDERSTAND
10 WHAT I THINK ARE AT STAKE. AND TO ME THERE'S ONLY FOUR
11 WAYS THAT CIRM MONEY CAN GO TO A BUSINESS. AND I CALL
12 SCENARIO 1 LIKE BUILDING A HOUSE. BASICALLY YOU PAY A
13 COMPANY TO BUILD A HOUSE. WHEN IT'S FINISHED, YOU OWN
14 THE HOUSE. YOU MIGHT LIVE IN IT OR SELL IT, AND THE
15 COMPANY THAT RECEIVED YOUR MONEY MADE A PROFIT AND HAS
16 AN INCENTIVE TO BUILD MORE HOUSES. I THINK IN THE SAME
17 WAY IF YOU GIVE A GRANT TO A BIOTECH COMPANY TO DO
18 RESEARCH AND THE TAXPAYERS HAVE PUT UP THE MONEY, ANY
19 DISCOVERIES THAT COME OUT OF THAT GRANT THAT LEAD TO
20 SOMETHING THAT'S PATENTABLE, TAXPAYERS SHOULD OWN THAT
21 PATENT.

22 WE WOULD URGE THAT THAT PATENT GO INTO A
23 PATENT POOL TO MAXIMIZE ACCESS TO IT FOR THE MAXIMUM
24 NUMBER OF PEOPLE. WE WOULD ALSO SAY THAT IF THERE WERE
25 CASES WHERE FOR COMMERCIALIZATION OF THE PRODUCT THAT

1 WERE NECESSARY TO HAVE AN EXCLUSIVE LICENSE, THE STATE
2 WOULD LICENSE IT BACK EXCLUSIVELY TO THAT GRANTEE
3 ORGANIZATION IF IT WERE ABSOLUTELY NECESSARY TO
4 COMMERCIALIZE.

5 THE SECOND SCENARIO IS WHAT I CALL THE
6 PARTNERSHIP MODEL. I KIND OF LIKE THIS NOTION OF A
7 HOUSE, BUT HERE SUPPOSE SOMEBODY HAS GOT LAND AND THEY
8 COME TO YOU AND SAY GIVE US SOME MONEY. WE'LL BUILD A
9 HOUSE ON THAT LAND. YOU PUT UP THE MONEY, WHEN THE
10 HOUSE IS SOLD, YOU GET A RETURN ON YOUR INVESTMENT. IN
11 THE WORLD OF PROPOSITION 71 AND BIOTECH, THIS WOULD BE
12 ANALOGOUS TO A SITUATION WHERE A COMPANY ALREADY HAS A
13 LICENSE OR A PATENT, AND THEY WANT THE MONEY TO FURTHER
14 DEVELOP IT AND BRING WHATEVER THE PRODUCT IS USING THE
15 EXISTING PATENT THAT THEY OWN OR CONTROL, BRINGING THAT
16 TO MARKET.

17 SO IN THIS CASE I WOULD ARGUE THAT THE
18 TAXPAYER SHOULD DESERVE SOME SORT OF RETURN ON THAT
19 INVESTMENT THAT THEY'VE PUT IN. AND THAT WOULD BE SOME
20 KIND OF A COMMISSION.

21 SCENARIO 3 IS MY BANK LOAN MODEL. COMPANY
22 COMES AND SAYS WE'D LIKE TO BORROW X MILLIONS OF
23 DOLLARS TO DO Y PROJECT. EVERYONE SITS DOWN AND AGREES
24 THAT, OKAY, YOU GET IT AT THIS INTEREST RATE, YOU PAY
25 IT BACK OVER THIS PERIOD OF TIME. I WOULD ASSERT THAT

1 IN THAT SITUATION YOU PROBABLY WOULD NOT HAVE A CLAIM
2 THAT THE STATE COULD PUT ON THE INTELLECTUAL PROPERTY;
3 BUT JUST LIKE WHEN A BANK MAKES A MORTGAGE, THE
4 MORTGAGEE OFTEN HAS REQUIREMENTS PLACED ON THEM THAT
5 THEY HAVE TO ENSURE THE PROPERTY OR THINGS LIKE THAT.
6 I THINK THAT IT'S PERFECTLY APPROPRIATE IF YOU ARE
7 GOING TO GET A LOAN FROM CIRM, THAT THERE BE PUBLIC
8 BENEFIT REQUIREMENTS ATTACHED TO THE LOAN.

9 FINALLY, SCENARIO NO. 4 IS CONTRACT SERVICE.
10 I COMPARE THAT TO THE SAME SORT OF THING AS HIRING
11 SOMEBODY TO COME IN AND CLEAN YOUR HOUSE. FOR PROP 71
12 MONEY, IT COULD BE SOMETHING WHERE CIRM WOULD CONTRACT
13 TO PROVIDE A PARTICULAR SERVICE, MAYBE MAINTAIN A STEM
14 CELL BANK, MAYBE PROVIDING SOME SORT OF WIDELY
15 NECESSARY RESEARCH TOOL THAT THE RESEARCHERS DON'T
16 REALLY WANT TO MAKE, BUT IS NECESSARY FOR THE WORK.
17 CIRM COULD CONTRACT TO DO THAT.

18 AGAIN, I DON'T SEE THAT THERE WOULD BE
19 INTELLECTUAL PROPERTY CREATED THAT THE STATE WOULD
20 NECESSARILY HAVE A ROLE IN, BUT I DO THINK THAT, JUST
21 AS IF I GO HIRE A CONTRACTOR, IT'S SUGGESTED TO ME THAT
22 I GET THREE BIDS AND GET THE BEST PERSON IN TO DO THE
23 JOB, I THINK IT WOULD BE APPROPRIATE FOR CIRM TO HAVE
24 PUBLIC BENEFIT REQUIREMENTS ATTACHED TO WHATEVER
25 CONTRACT WAS AWARDED.

1 SO THOSE ARE SOME BROAD KINDS OF THEORIES.
2 WE BELIEVE THAT THE IP POLICY NEEDS TO BE GROUNDED IN
3 THREE PRINCIPLES. WE SAY AFFORDABILITY, ACCESSIBILITY,
4 ACCOUNTABILITY. WE MADE THAT POINT IN OUR ORIGINAL
5 PAPER THAT WE PUT OUT. AND WHAT THAT BASICALLY MEANS
6 VERY QUICKLY IS THE TREATMENTS HAVE TO BE PRICED SO
7 THEY CAN BE SOLD -- SO THAT ALL CALIFORNIANS CAN AFFORD
8 AND BENEFIT THEM. ACCESSIBILITY MEANS NOT JUST FOR
9 POTENTIAL PATIENTS THAT CAN USE THE THERAPIES. IT ALSO
10 MEANS THAT RESEARCHERS NEED TO HAVE ACCESS TO EACH
11 OTHER'S RESEARCH, WHICH I THINK IS ONE OF THE MAJOR
12 IMPORTANT POSITIONS OF THE IP POLICY FOR NONPROFITS.
13 AND FINALLY, ACCOUNTABILITY. POLICIES ARE NO GOOD
14 UNLESS THERE'S A MECHANISM TO ENSURE ENFORCEMENT, AND
15 WE THINK THAT NEEDS TO BE ATTORNEY GENERAL.

16 SO WITH THAT SORT OF BROAD FRAMEWORK, I'VE
17 GOT SOME SPECIFIC SORT OF BULLET POINTS OF THINGS THAT
18 WE WOULD LIKE TO SEE INCORPORATED IN WHATEVER IP POLICY
19 FINALLY COMES OUT, AND IT'S IN GREATER DETAIL IN THE
20 WHITE PAPER, BUT JUST TO HIT THE HIGHLIGHTS VERY
21 QUICKLY, AND WE THINK THIS ONE IS CRITICAL. THIS IS
22 NOT NOW IN THE NONPROFIT ONES. WE THINK IT NEEDS TO BE
23 ADDED. WE THINK THAT THAT'S THE PRINCIPLE THAT ANY
24 THERAPIES OR DIAGNOSTICS THAT ARE DEVELOPED WITH
25 PROPOSITION 71 MONEY HAVE TO BE MADE AVAILABLE AT A

1 REASONABLE PRICE. WE THINK A REASONABLE PRICE IS ONE
2 THAT REFLECTS THE TRUE COST OF DEVELOPMENT OF THE DRUG
3 OR THERAPY AND ALSO THE PUBLIC'S INVESTMENT.

4 WE THINK THAT IN THE SITUATIONS WHERE PERHAPS
5 A COMPANY IS GETTING LICENSING REVENUE, ROYALTY REVENUE
6 IN FOR SOMETHING THAT IT'S DEVELOPED WITH PROP 71
7 MONEY, WE THINK THAT, LIKE WITH THE NONPROFITS, THERE
8 SHOULD BE A DIRECT PAYBACK TO THE STATE, AND WE THINK
9 25 PERCENT OF THE ROYALTIES WOULD BE APPROPRIATE.

10 NEXT SLIDE. WE WOULD BE THINKING ABOUT
11 SITUATIONS WHERE THERE WOULD BE A COMMISSION OR A
12 LICENSE FEE COMING BACK TO THE STATE. WE THINK A
13 ROYALTY RATE WHEN A ROYALTY COMES INTO PLAY OUGHT TO BE
14 THE SAME SORT OF ROYALTY RATE THAT WOULD APPLY TO
15 SIMILAR SITUATIONS IF THE UNIVERSITY OF CALIFORNIA HAD
16 DONE IT.

17 OTHER POINTS THERE, WE ALWAYS THINK IT'S
18 IMPORTANT TO HAVE THE BUSINESSES EXPLAIN HOW WHAT
19 THEY'RE GOING TO DO IS GOING TO BENEFIT ALL OF
20 CALIFORNIANS. AND AGAIN, SAME AS THE NONPROFITS, WE
21 THINK THERE NEEDS TO BE THE PROVISION THAT ANY
22 DIAGNOSTICS OR CURES WOULD BE SOLD AT THE LOWEST PRICE
23 TO PUBLICLY FUNDED HEALTHPLANS. THAT OFTEN IS THE
24 MEDICAID PRICE, BUT I THINK SPECIFICALLY THE LOWEST IS
25 A BETTER WAY TO FRAME IT.

1 AND THE NOTION OF ACCESSIBILITY, WE REALLY
2 BELIEVE THAT THERE SHOULD BE A PATENT POOL, THAT THIS
3 WOULD MAXIMIZE ACCESSIBILITY FOR EVERYBODY. IT'S TRUE
4 THAT, YOU KNOW, IT TOOK A WORLD WAR I TO FORCE THE
5 PATENT POOL THAT MADE AIRPLANES FLY AND REALLY COULDN'T
6 HAVE DEVELOPED AIRPLANES WITHOUT GETTING ALL THE
7 PATENTS TOGETHER IN ONE PLACE. AND I GUESS IT WAS
8 FRANKLIN ROOSEVELT THAT WAS GIVEN THE CHARGE TO
9 ESSENTIALLY ORDER A PATENT POOL, AND THAT'S WHAT LED TO
10 FIGHTER PLANES BEING BUILT FOR WORLD WAR I.

11 IN SOME WAYS I THINK WE'RE IN A SIMILAR
12 SITUATION WITH STEM CELL RESEARCH, AND SOMEBODY HAS GOT
13 TO STEP UP TO THE PLATE AND SAY THERE ARE TOO MANY
14 PATENTS. THEY NEED TO GET POOLED, SO WE CAN GO AHEAD
15 ON THIS.

16 OTHER POINTS THERE ARE SOME OF THE SIMILAR
17 ONES THAT WE'VE HAD IN THE NONPROFIT RULES. AND YOU
18 CAN ALL READ THEM. WELL, I THINK THE NOTION THAT A
19 BUSINESS WOULD HAVE TO PROVIDE A PLAN FOR ACCESS TO
20 UNINSURED PATIENTS IS IMPORTANT. AND AGAIN, I CAN
21 FORESEE A SITUATION WHERE EXCLUSIVITY SOMETIMES WORKS
22 AGAINST DISSEMINATION, AND CIRM OUGHT TO BE ABLE TO BAR
23 AN EXCLUSIVE LICENSE IN THAT SORT OF SITUATION.

24 THE SAME RULES IN THE NEXT SLIDE AS ARE IN
25 THE FOUR -- EXCUSE ME -- THE NONPROFIT RULES. I THINK

1 THE ACCESS TO RESEARCH FOR RESEARCH PURPOSE IS
2 CRITICAL. THAT NEEDS TO BE MAINTAINED. AND THIS LAST
3 ITEM UNDER ACCESSIBILITY, I THINK THAT IN THOSE
4 CIRCUMSTANCES WHERE AN EXCLUSIVE LICENSE IS GRANTED,
5 IT'S NECESSARY TO GRANT IT ON A DISEASE-SPECIFIC BASIS.
6 THIS PRECLUDES ONE COMPANY FROM ESSENTIALLY GETTING A
7 LICENSE FOR A DRUG AND THEN ONLY GOING DOWN ONE ROUTE,
8 BUT BLOCKING EVERYONE ELSE FROM DOING IT.

9 SO I THINK THAT IF YOU WANT TO LICENSE
10 SOMETHING EXCLUSIVELY, IT SHOULD BE, OKAY, YOU'VE GOT
11 THIS PARTICULAR THERAPY. WE'RE LICENSING IT
12 EXCLUSIVELY FOR YOU TO DO SOMETHING WITH DIABETES. AND
13 THEN ANOTHER EXCLUSIVE LICENSE, IF NECESSARY, TO GO
14 INTO A DIFFERENT DISEASE. I THINK THAT WOULD KEEP
15 PROGRESS GOING AHEAD FASTER.

16 AS I SAID BEFORE, ALL THE GREAT RULES IN THE
17 WORLD REALLY ARE ONLY SO MUCH WORDS AND PAPER UNLESS
18 THERE IS A MEANS OF ENFORCING ACCOUNTABILITY. WE
19 BELIEVE THAT THE ATTORNEY GENERAL EXPLICITLY NEEDS TO
20 HAVE INVOLVEMENT IN SOME OF THIS. IT IS TRUE THAT
21 THROUGH THE OFFICE OF ADMINISTRATIVE LAW THAT
22 ULTIMATELY YOUR REGULATIONS ARE LAWS, BUT I THINK IN
23 SOME AREAS THERE NEEDS TO BE MORE EXPLICIT INVOLVEMENT
24 OF THE ATTORNEY GENERAL AS THE ENFORCER, AND I THINK
25 THAT COMES SPECIFICALLY ALONG THE NOTION OF

1 UNREASONABLE PRICING.

2 AND I THINK THERE THAT IF DRUGS, CURES,
3 THERAPIES ARE FOUND TO BE PRICED UNREASONABLY, THEN THE
4 ATTORNEY GENERAL NEEDS TO HAVE SO-CALLED MARCH-IN
5 RIGHTS, THE RIGHTS TO INTERVENE. AND AGAIN, TO
6 REITERATE, REASONABLE PRICING IS PRICING THAT TAKES
7 INTO ACCOUNT THE ACTUAL DEVELOPMENT COST OF THE DRUG
8 AND THE PUBLIC'S INVESTMENT IN IT. AG NEEDS MARCH-IN
9 RIGHTS IF OTHER PUBLIC BENEFIT REQUIREMENTS AREN'T MET.
10 SHOULD BE MARCH-IN RIGHTS FOR CIRM IF BUSINESS FAILS TO
11 DEVELOP SOMETHING THAT THEY'VE HAD A LICENSE FOR. THE
12 NEXT SLIDE. MARCH-IN RIGHTS FOR PUBLIC HEALTH AND
13 SAFETY REASONS.

14 AND THEN FINALLY, WE WOULD HOPE THAT THERE'S
15 GOING TO BE A NUMBER OF START-UP COMPANIES THAT COME
16 OUT OF THIS WHOLE PROCESS, BUT WE THINK THAT BECAUSE
17 IT'S PUBLIC MONEY THAT'S GOING INTO SOME OF THESE KINDS
18 OF THINGS, THERE DOES NEED TO BE COMPLETE TRANSPARENCY.
19 AND, THEREFORE, INVESTORS AND MAJOR SHAREHOLDERS WHO
20 ARE INVOLVED IN START-UP COMPANIES THAT COME ABOUT
21 THROUGH PROPOSITION 71-FUNDED RESEARCH NEED TO FILE
22 DISCLOSURE FORMS THAT WOULD BECOME PUBLIC RECORDS.

23 SO IN CONCLUSION, THE SUMMARY, THE HIGH
24 POINTS OF WHAT WE THINK ARE IMPORTANT IS, ONE, IP
25 POLICY IS THE MEANS TO ASSURE THAT THE PROMISES OF

1 PUBLIC BENEFIT THAT WERE INCORPORATED IN PROPOSITION 71
2 ARE KEPT. NO. 2, THAT IF A PATENT COMES ABOUT
3 BECAUSE -- DIRECTLY BECAUSE OF PROP 71 MONEY, THEN THE
4 STATE SHOULD HAVE THAT PATENT. WE THINK THAT
5 ABSOLUTELY ESSENTIAL IN BOTH THE NONPROFIT RULES, AND
6 WE'LL BE ADDING THOSE COMMENTS IN THE APPROPRIATE
7 PUBLIC COMMENT TIME, AND IN THE FOR-PROFIT RULES IS
8 REQUIREMENT FOR REASONABLE PRICING.

9 NOW, WE DON'T SAY THAT COMPANIES SHOULDN'T
10 MAKE A PROFIT. WE DON'T THINK THAT. WE THINK THAT THE
11 PROFIT NEEDS TO BE REASONABLE. I MENTION THE NONPROFIT
12 IP RULES NEED IMPROVEMENT, AND WE'LL BE CONTINUING TO
13 WORK THROUGH ALL OF THE PROCESSES THAT HAVE BEEN MADE
14 AVAILABLE TO US TO TRY AND BRING ABOUT THAT
15 IMPROVEMENT, BUT THEY ARE A MINIMUM STARTING POINT.
16 AND THERE'S SOME IMPORTANT PRINCIPLES THAT ARE IN THOSE
17 THAT NEED TO BE CARRIED OVER FOR SURE INTO THE
18 FOR-PROFIT RULES.

19 FINALLY, YOU KNOW, THE VENTURE CAPITALISTS
20 WHO HAVE BEEN TALKING ABOUT CIRM MONEY AS, QUOTE, FREE
21 MONEY OR ALMOST LIKE FREE MONEY AND INDEED THEY HAVE --
22 I'VE BEEN AT CONFERENCES WHERE I'VE HEARD IT CALLED
23 THAT -- NEED TO UNDERSTAND, AND I THINK THAT MANY
24 RESPONSIBLE COMPANIES DO UNDERSTAND THIS, IS THAT WITH
25 PUBLIC MONEY COMES THE REQUIREMENT OF MEETING SOME

1 FAIR, EQUITABLE PUBLIC BENEFIT REQUIREMENTS.

2 THANK YOU VERY MUCH. ANY QUESTIONS, I'D BE
3 GLAD TO ANSWER THEM. AND I GUESS IF WE HAVE ONE MORE
4 SLIDE, IF YOU WANT TO SEND YOUR SLINGS AND ARROWS,
5 THERE'S A PLACE THERE AGAIN WHERE YOU CAN REACH ME ANY
6 TIME.

7 CHAIRMAN PENHOET: THANK YOU, JOHN. SO FIRST
8 OF ALL, ANY QUESTIONS FROM THE TASK FORCE HERE IN SAN
9 DIEGO? FRANCISCO.

10 DR. PRIETO: YES. FRANCISCO PRIETO. ONE
11 COULD SAY ABOUT THE 120 MILLION FOR STEM CELL RESEARCH,
12 IS THAT IN THE U.S. OR IS THAT WORLDWIDE?

13 MR. SIMPSON: MY UNDERSTANDING WAS THAT WAS
14 IN THE U.S. THAT WAS VENTURE CAPITAL INVESTMENT IN THE
15 UNITED STATES. I WOULD HAVE TO DOUBLE-CHECK THAT, BUT
16 MY RECOLLECTION OF THE FIGURE IS THAT THAT WAS IN THE
17 UNITED STATES.

18 DR. PRIETO: I THINK ONE OF THE QUESTIONS FOR
19 A LOT OF US IS, YOU KNOW, WHETHER, INDEED, WE WILL BE
20 THE WORLD'S BIGGEST, OR ARE WE REALLY FAR BEHIND THE UK
21 AND SINGAPORE AND ISRAEL. AND I DON'T KNOW WHAT THE
22 DOLLAR ARE IN THOSE PLACES AND WHAT THE FUNDING SOURCES
23 ARE, BUT I'D BE VERY INTERESTED TO LEARN.

24 CHAIRMAN PENHOET: WE'LL DO SOME HOMEWORK AND
25 CAN GET BACK TO YOU ON IT. JOHN.

1 DR. REED: JOHN REED. YEAH, I ALSO HAVE SOME
2 QUESTIONS ABOUT THOSE NUMBERS. THERE WAS A 5.9 BILLION
3 MENTIONED, AND I UNDERSTAND THAT, BUT WE CAN DECLARE
4 LATER.

5 I WAS CURIOUS ABOUT THE GROUP'S CALCULATION
6 OF THE 25-PERCENT ROYALTY. CAN YOU EXPLAIN HOW YOU
7 TARGETED THAT AMOUNT, WHAT WAS YOUR THINKING THERE?
8 YOU'RE SUGGESTING A 25-PERCENT ROYALTY ON ANY PRODUCTS.

9 CHAIRMAN PENHOET: I THINK THAT'S NOT -- WHAT
10 THE NONPROFIT POLICY SAYS IS 25 PERCENT OF THE ROYALTY,
11 NOT A 25-PERCENT ROYALTY. AND THERE'S BEEN SOME
12 MISUNDERSTANDING ABOUT THAT.

13 MR. SIMPSON: IN OTHER WORDS, WHAT WE WERE
14 SAYING IS IF THERE IS A REVENUE STREAM. AND I CAN
15 ENVISION -- EVEN IF THE STATE HELD SOME PATENTS, I CAN
16 ENVISION A SITUATION WHERE THERE WOULD BE ROYALTIES
17 COMING BACK BECAUSE THERE MIGHT BE A CASE WHERE OTHER
18 PATENTS WERE INVOLVED OR WHATEVER. SO THE COMPANY IS
19 GETTING ROYALTIES. AND IF PROPOSITION 71 MONEY HAS
20 GONE INTO THAT ROYALTY, THEN WE THINK IT SHOULD BE 25
21 PERCENT OF THE ROYALTY. AND THAT'S COMPATIBLE WITH
22 WHAT WAS DONE WITH THE NONPROFITS.

23 THE SAME SITUATION THERE, ALTHOUGH THE
24 TROUBLESOME THING WITH THE NONPROFITS IS IF THE ROYALTY
25 KICKS IN ONLY AFTER A \$500,000 THRESHOLD, WHICH SHOULD

1 BE LOWER BECAUSE IT'S NOT GROSS, IT'S NET, AND WE THINK
2 IT NEEDS TO BE A LOWER THRESHOLD BEFORE. BUT IT'S THE
3 SAME MODEL.

4 AND IF YOU'RE TALKING ABOUT ROYALTY RATES OR
5 COMMISSION RATES IN THE SENSE OF WHAT THEY WOULD BE,
6 THAT'S WHERE I COMPARE IT TO THE UNIVERSITY OF
7 CALIFORNIA WHERE USUALLY YOU NEGOTIATE RATES MAYBE 2 TO
8 6 PERCENT OF THE GROSS. SO THE 25 PERCENT WOULD BE OF
9 A -- OF THAT KIND OF THING.

10 DR. REED: THANK YOU FOR CLARIFYING.

11 MS. KING: THIS IS MELISSA AT UC SAN DIEGO.
12 I JUST WANTED TO LET EVERYBODY KNOW ON THE PHONE AND
13 HERE AS WELL IF EVEN BOARD MEMBERS COULD, BEFORE
14 SPEAKING, ADDRESS YOURSELF BY YOUR OWN NAME, STATE WHO
15 YOUR FOR THE BENEFIT OF THE TRANSCRIBER, WHO'S ON THE
16 PHONE, THAT WOULD BE GREAT. SAME WITH MEMBERS OF
17 PUBLIC. YOU DON'T HAVE TO IDENTIFY YOURSELVES, BUT IF
18 YOU WOULD, THAT'S VERY HELPFUL TO US. THANK YOU.

19 DR. PRIETO: FRANCISCO PRIETO AGAIN IN SAN
20 DIEGO. TWO QUESTIONS AND ONE IS SORT OF A COMMENT.
21 REGARDING PATENT POOLS, I DON'T KNOW VERY MUCH ABOUT
22 THAT AND WOULD LIKE TO LEARN MORE AND WHAT MODELS THERE
23 ARE FOR PATENT POOLS AND WHO MANAGES THEM AND WHAT
24 COSTS ARE ASSOCIATED WITH MANAGING THAT.

25 AND THE OTHER IS A COMMENT REGARDING THE

1 POINT YOU MADE ABOUT DISEASE-SPECIFIC LICENSES. ONE OF
2 THE PROBLEMS I HAVE WITH THAT JUST CONCEPTUALLY IS THAT
3 I CAN IMAGINE A TREATMENT OR A THERAPY THAT CAME OUT OF
4 THIS RESEARCH FOR DEMYELINATING DISEASES SUCH AS
5 MULTIPLE SCLEROSIS, BUT THE INFORMATION GLEANED FROM
6 THAT COULD HAVE VERY DIRECT APPLICATIONS TO PARKINSON'S
7 DISEASE, TO DIABETIC NEUROPATHY, TO SPINAL CORD INJURY.
8 SO WHAT IS THE DISEASE? YOU KNOW, YOU HAVE RESEARCH
9 LEADING IN PERHAPS SEVERAL DIFFERENT AREAS, AND I DON'T
10 KNOW THAT YOU COULD LIMIT THAT.

11 MR. SIMPSON: PATENT POOLS FIRST. THEY'VE
12 BEEN SUCCESSFUL IN A NUMBER OF INDUSTRIAL SORTS OF
13 SITUATIONS. I ALLUDED TO THE AIRCRAFT INDUSTRY. AND
14 WHEN THE AIRCRAFT INDUSTRY GOT STARTED, THE WRIGHTS HAD
15 PATENTS, THE CURTISES HAD PATENTS, AND THEY ESSENTIALLY
16 BLOCKED EACH OTHER FROM DEVELOPING SUCCESSFULLY
17 COMMERCIALY VIABLE AIRPLANES, AS I UNDERSTAND IT. AND
18 THEY WERE REQUIRED TO POOL THEM. THEY PUT THEM IN A
19 POOL. THEN EVERYBODY GOT A ROYALTY BACK THAT WAS
20 PRORATED.

21 I BELIEVE THEY HAVE BEEN VERY SUCCESSFUL IN
22 THE SEWING MACHINE BUSINESS AT ONE POINT OR ANOTHER.
23 AND AS I UNDERSTAND IT, THERE ARE A NUMBER OF POOLS NOW
24 THAT WORK IN ELECTRONICS. AND THE NOTION ESSENTIALLY
25 IS THAT IF YOU HAVE A LOT OF PATENTS THAT ARE REQUIRED

1 TO DEVELOP SOMETHING, IF A COMPANY WANTS TO DO IT, THEY
2 HAVE TO GO OUT AND NEGOTIATE WITH 50 DIFFERENT PATENT
3 HOLDERS, AND THAT BECOMES TIME-CONSUMING AND A
4 DISINCENTIVE. AND BY HAVING A POOL, PARTICULARLY IF
5 YOU'RE IN A SITUATION WHERE A LOT OF THE PATENTS ARE IN
6 UPSTREAM RESEARCH, IT GIVES YOU A PLACE WHERE ONE SORT
7 OF ONE-STOP SHOPPING. YOU GO IN, YOU GET EVERYTHING
8 YOU NEED, THERE'S A ROYALTY RATE THAT IS FAIR TO
9 EVERYBODY, AND IT HOLDS THE PATENT, EVERYBODY GETS SOME
10 OF THE REVENUE BACK, AND GENERALLY THE POOL IS GOVERNED
11 BY SOME REPRESENTATION OF THE PEOPLE WHOSE PATENTS ARE
12 IN IT.

13 ALSO, I THINK IF YOU HAD A STRONG ENOUGH POOL
14 WITH ENOUGH IMPORTANT PATENTS IN IT, IT WOULD GIVE YOU
15 IN THE WORST-CASE SCENARIO, IF THE OVERREACHING,
16 OUTRAGEOUS PATENTS THAT WARF HOLDS ON HUMAN EMBRYONIC
17 STEM CELLS AREN'T BROKEN BY SOMEBODY, AND THEY DO NEED
18 TO BE BROKEN AND THEY DO NEED TO BE CHALLENGED, BUT IF
19 FOR SOME REASON THEY'RE NOT, A PATENT POOL THEN GIVES
20 YOU SOMETHING THAT'S IN LEVERAGE TO BE ABLE TO PERHAPS
21 WORK AND NEGOTIATE WITH THEM, WHICH IS ANOTHER REASON
22 THAT IT WORKS.

23 THERE IS A LEGISLATIVE BILL RIGHT NOW, I
24 THINK, THAT MAY SET UP AN OFFICE OF STATE INTELLECTUAL
25 PROPERTY, WHICH I BELIEVE CIRM IS SPECIFICALLY EXCLUDED

1 FROM AT THE MOMENT, BUT I MEAN CONCEIVABLY AN OFFICE
2 LIKE THAT COULD MANAGE THE POOL. I'M NOT SURE THAT
3 CIRM HAS THE RESOURCES. THE THING TO ME WITH THE
4 PATENT POOLS IS THAT EVERYONE I'VE TALKED IN THE FIELD
5 SAYS THAT IT'S TREMENDOUSLY INTERESTING AND VALUABLE,
6 BUT WE'RE NOT THERE YET. AND I GUESS MY POINT IS IF WE
7 DON'T STEP UP AND SAY WE'VE GOT TO GET THESE THINGS
8 INTO A POOL SO EVERYONE CAN USE THEM, WE'RE GOING TO
9 HAVE, AS WE DO NOW, 1400 PATENTS AND PATENT THICKETS
10 THAT ARE REALLY GETTING IN THE WAY OF DEVELOPMENT IN
11 RESEARCH.

12 AND THE SECOND QUESTION HAD TO DO -- I'M
13 SORRY.

14 DR. PRIETO: REGARDING DISEASE SPECIFIC.

15 MR. SIMPSON: RIGHT. THERE MAY BE SEVERAL
16 DIFFERENT WAYS OF GETTING AROUND WHAT I'M CONCERNED
17 ABOUT, BUT WHAT I'M CONCERNED WITH IS SOMEONE GETTING
18 AN EXCLUSIVE LICENSE AND ONLY GO IN ONE WAY WHEN THERE
19 ARE OTHER WAYS THAT THAT TREATMENT COULD BE USED, AND
20 THAT FOR COMMERCIAL REASONS, THEY JUST DON'T GO THERE.
21 SO IT SEEMS TO ME THAT THERE NEEDS TO BE SOME WAY OF
22 DEALING WITH THAT POSSIBILITY. I MEAN I SUPPOSE SOME
23 KIND OF MARCH-IN RIGHTS THAT SAID, LOOK, WE KNOW THAT
24 IF YOU DEVELOP -- IF YOU ALLOW IT TO BE LICENSED IN THE
25 PUBLIC INTEREST BECAUSE THIS COULD BE TREATED THAT WAY

1 AS WELL, THAT WOULD WORK TOO. BUT THE CONCERN
2 ESSENTIALLY IS BLOCKING FRUITFUL AVENUES OF TREATMENT
3 WITH A SINGLE LICENSE. I THINK THAT WOULD BE A BAD
4 THING.

5 CHAIRMAN PENHOET: ANY COMMENTS FROM ANY OF
6 OUR OFF-SITE SITES? IRVINE, QUESTIONS THERE? ANY
7 QUESTIONS FROM NUVELO? FROM LOS ANGELES?

8 UNIDENTIFIED SPEAKER: NO.

9 CHAIRMAN PENHOET: FROM CHICO?

10 DR. WRIGHT: NO.

11 CHAIRMAN PENHOET: ANY COMMENTS FROM THE
12 PUBLIC IN SAN DIEGO? YES.

13 MR. ROBINS: ALLAN ROBINS FROM NOVOCELL. I
14 JUST WANTED TO MAKE A COUPLE OF COMMENTS, JOHN. FIRST
15 OF ALL, IN TERMS OF THE FUNDING. I THINK THERE'S A LOT
16 OF CONFUSION WHEN PEOPLE TALK ABOUT STEM CELL RESEARCH
17 GENERICALLY, AND THEY TALK ABOUT HUMAN EMBRYONIC STEM
18 CELL RESEARCH. SO I THINK THE NUMBERS, AT LEAST THE
19 NIH NUMBERS YOU QUOTED, WAS \$30 MILLION (INAUDIBLE).
20 THE NIH PUT 200 MILLION.

21 MR. SIMPSON: I'M SORRY. I DIDN'T THINK MAKE
22 THAT CLEAR.

23 MR. ROBINS: RIGHT. BUT I MEAN I THINK
24 THEY'RE IMPORTANT POINTS. WHEN STEM CELL RESEARCH
25 INVESTMENT IS REPORTED, YOU HAVE TO MAKE SURE YOU'RE

1 COMPARING APPLES TO APPLES. I THINK THERE'S A LOT MORE
2 EFFORT GOING INTO STEM CELL THERAPY THAN JUST --

3 THE SECOND THING I WANTED TO DO IS MAYBE
4 POINT OUT SOME DIFFERENCES BETWEEN YOUR THEORY OF
5 BUYING A HOUSE MODEL AND WHAT WE'RE REALLY DOING IN A
6 COMPANY. AND I THINK THERE ARE TWO MAJOR POINTS HERE.
7 ONE IS RISK PROFILE. WHEN YOU'RE GOING TO BUY A HOUSE
8 AND YOU CONTRACT A BUILDER AND YOU'RE GOING TO THE BANK
9 AND YOU BORROW THE MONEY, IT'S A PRETTY SURE THING WHAT
10 YOU ARE GOING TO END UP WITH AT THE END OF THE DAY IS A
11 HOUSE. AND MOSTLY THEY APPRECIATE IN VALUE MOST OF THE
12 TIME. SO I THINK THERE'S A SORT OF A FLAW IN YOUR
13 MODEL WHEN YOU'RE USING THAT TO COMPARE IT TO SORTS OF
14 THINGS (INAUDIBLE).

15 AND THE SECOND THING THAT I WANTED TO POINT
16 OUT IS THAT BUILDING A HOUSE IS REALLY A DISCRETE
17 THING. YOU GO, YOU BY A BLOCK OF LAND, AND YOU BUILD
18 THE HOUSE BASICALLY. RESEARCH IS A CONTINUUM, SO IF
19 CIRM OR ANY OTHER ORGANIZATION IS GOING TO FUND A
20 COMPANY WHO HAS BACKGROUND RESEARCH THAT WE LOOK AT AND
21 CIRM FUNDING WILL NOT TAKE IT THROUGH TO A PRODUCT.
22 THERE WILL BE OTHER INVESTMENT THAT'S NEEDED. AND
23 ALONGSIDE THAT, OUR INVESTMENT OF MONEY IN AT THE SAME
24 TIME. SO I DON'T THINK IT'S DISCRETE, AND I THINK
25 DIFFICULT TO MAKE THAT COMPARISON IN THAT WAY. I MEAN

1 THERE ARE OTHER REASONS, BUT THEY'RE THE TWO
2 FUNDAMENTAL ONES.

3 AND THE THIRD QUESTION/COMMENT THAT I WANTED
4 TO MAKE IS THAT YOU'VE ALLUDED TO THE FACT THAT THERE'S
5 ALREADY SORT OF WIDE-REACHING INTELLECTUAL PROPERTY,
6 AND YOU'VE ALLUDED TO ONE SORT OF INTELLECTUAL
7 PROPERTY, BUT THERE ARE, IN FACT, OTHER ISSUED PATENTS
8 OR PATENTS THAT ARE WORKING THEIR WAY THROUGH THE USPTO
9 WHICH ARE ALSO VERY BROAD AND JUST AS TROUBLESOME.

10 SO I'M WONDERING IN TERMS OF YOUR PATENT
11 POOL, THESE PATENTS CLEARLY SIT OUTSIDE OF THAT POOL.
12 AND ANY COMPANY, LIKE OURSELVES, THAT WOULD BE LOOKING
13 AT CONTRIBUTING SOMETHING TO A PATENT POOL, CLEARLY ARE
14 AT A DISADVANTAGE. SO I'M WONDERING IF YOU'RE SORT OF
15 CLOSING THE GATE AFTER THE HORSE HAS BOLTED, OR WHAT
16 YOU WOULD PROPOSE TO DO ABOUT THOSE BROAD-RANGING
17 PATENTS THAT ARE ALREADY IN EXISTENCE OR WILL BE
18 DRAFTED.

19 MR. SIMPSON: TO THE HOUSE MODEL, NO ANALOGY
20 WORKS PERFECTLY, I GUESS, AND I TAKE SOME OF YOUR
21 POINTS. BUT WHAT I ESSENTIALLY WAS TRYING TO SAY IS
22 THAT IT'S -- AT SOME POINT IF PUBLIC MONEY HAS GONE
23 INTO IT AND, YES, THE HOUSE -- IF YOU'RE BUILDING THE
24 HOUSE IN A RISKY PLACE, MAYBE IT GETS BLOWN DOWN BY
25 HURRICANES AND SO ON AND SO ON AND SO FORTH AND YOU

1 NEVER CAN GET YOUR MONEY BACK, BUT WITH RESEARCH IT'S
2 EVEN RISKIER. BUT IF YOU DO GET SOMETHING THAT
3 GENERATES REVENUE, THAT'S WHERE I SAY, BECAUSE THE
4 STATE HAS PUT THE MONEY IN, TAXPAYERS DESERVE TO GET
5 SOMETHING BACK. THAT'S THE KIND OF THING, AND I WAS
6 TRYING TO DO IT IN A WAY THAT WAS AT LEAST SOMETHING
7 THAT A POOR OLD NEWSPAPERMAN'S BRAIN COULD GET HIS ARMS
8 AROUND, WHICH WAS THE HOUSE MODEL.

9 AS FAR AS THE PATENTS GO, I THINK THE WARF
10 ONES ARE JUST INCREDIBLY OVERREACHING. I MEAN THE
11 NOTION OF -- IT'S NOT EVEN A MYTHOLOGY. IT'S
12 COMPOSITION OF MATTER. IT'S ALL HUMAN EMBRYONIC STEM
13 CELLS, AND RECOGNIZED NOWHERE ELSE IN THE WORLD. AND I
14 THINK THAT THERE ARE A NUMBER OF THINGS THAT NEED TO BE
15 DONE. I THINK THOSE -- I THINK THEY HAVE TO BE
16 CHALLENGED, AND I THINK THAT THEY LIKELY WILL BE BROKEN
17 IF THEY ARE.

18 MR. ROBINS: -- PROVIDING THE DATA.

19 MR. SIMPSON: I'M NOT SURE WHO WOULD DO THAT
20 AT THIS POINT. I MEAN THERE ARE A NUMBER OF DIFFERENT
21 WAYS THAT IT COULD BE DONE. I MEAN A COMPANY THAT
22 WANTED TO JUST GO AHEAD AND DO WHAT THEY WANT TO DO AND
23 LET WARF COME SUE THEM WOULD BE IN A POSITION TO
24 CHALLENGE THEM IN COURT. I MEAN IF WARF SAYS YOU'RE
25 INFRINGING, IT WOULD SEEM TO ME THAT IT WOULD BE AN

1 APPROPRIATE THING FOR THE BIOTECH INDUSTRY OR AN
2 INDUSTRY ASSOCIATION TO TAKE ON AND TO HELP FUND.
3 THERE ARE OTHER WAYS THAT YOU CAN DO IT. YOU CAN GO TO
4 THE PTO AND YOU CAN ASK FOR A REEXAMINATION. SOMEBODY
5 COULD DO THAT.

6 I THINK THAT CLEARLY WARF PATENTS ARE RULING
7 OVER THIS. I'VE BEEN TOLD SPECIFICALLY THAT A NUMBER
8 OF VENTURE CAPITALISTS HAVE NOT PUT MONEY INTO HUMAN
9 EMBRYONIC STEM CELL RESEARCH PRECISELY BECAUSE OF THOSE
10 PATENTS.

11 CHAIRMAN PENHOET: IF I COULD, I THINK THAT
12 WILL LOCK THE SUBJECT UP. IT'S AN IMPORTANT POINT, BUT
13 NOT DIRECTLY RELATED TO WHAT WE'RE TRYING TO GRAPPLE
14 WITH HERE.

15 ARE THERE COMMENTS FROM THE PUBLIC IN IRVINE?

16 DR. BRYANT: NO.

17 CHAIRMAN PENHOET: SAN CARLOS?

18 UNIDENTIFIED SPEAKER: YES.

19 MR. REED: THIS IS DON REED. THE UNMENTIONED
20 SO FAR GORILLA IN THE ROOM IS THAT WHILE WHAT WE'RE
21 DOING NOW IS VERY HELPFUL, WORKING OUT PROBLEMS IN THE
22 PUBLIC, I DON'T THINK MOST OF US WANT THE IP POLICIES
23 TO BE LEGISLATIVELY IMPOSED. AND THAT'S HAPPENING VERY
24 FAST RIGHT NOW UP IN SACRAMENTO. TO THE BEST OF MY
25 KNOWLEDGE, THERE'S ONLY ONE MORE COMMITTEE MEETING ON

1 SB 401, THE ORTIZ-RUNNER LEGISLATION, AND THAT'S THE
2 APPROPRIATIONS MEETING. THEY'RE TYPICALLY HELD ON
3 WEDNESDAYS. IT PROBABLY IS NEXT WEDNESDAY, AND THAT'S
4 THE LAST ONE BEFORE IT GOES TO THE FLOOR. IT'S ALREADY
5 GONE THROUGH THE SENATE BECAUSE IT WAS DONE WITH A
6 GUT-AND-AMEND DEAL WHERE A BILL ALREADY HALF PASSED WAS
7 HOLLOWED OUT AND SB 401 WAS PUT INSIDE OF IT. AND IT
8 WENT THROUGH VERY FAST.

9 I WENT UP AND DID SOME DOOR TO DOOR, TALKED
10 TO ALL 18 MEMBERS, LEGISLATIVE AIDES OF THE 18 MEMBERS
11 OF THE APPROPRIATIONS COMMITTEE YESTERDAY. AND IT WAS
12 SHOCKING THAT DEMOCRATS DID NOT KNOW ABOUT IT, AND THEY
13 SAID, "OH, WELL, IF ORTIZ WANTS IT, IT MUST BE GOOD FOR
14 STEM CELL RESEARCH." SO IF ANYBODY WANTS TO HAVE AN
15 IMPACT ON THIS, I STRONGLY SUGGEST THAT YOU CONTACT THE
16 18 MEMBERS OF THE -- OR LEAST THE HEAD, JUDY CHU, OF
17 THE APPROPRIATIONS COMMITTEE, BECAUSE THE SITUATION MAY
18 BE TAKEN OUT OF YOUR HANDS. THANK YOU.

19 CHAIRMAN PENHOET: ANY COMMENTS FROM LOS
20 ANGELES? FROM CHICO?

21 DR. WRIGHT: NO.

22 CHAIRMAN PENHOET: OKAY. THANK YOU VERY
23 MUCH, JOHN. WE'LL MOVE ON TO OUR SECOND SPEAKER, WHO
24 IS DAVID GOLLAHER, PRESIDENT AND CEO OF THE CALIFORNIA
25 HEALTH CARE INSTITUTE HERE IN LA JOLLA. DAVID.

1 MAYBE WHILE THEY'RE CHANGING THE LOGISTICS, I
2 WOULD REMIND EVERYONE THAT THE IP POLICY WE GENERATE
3 FOR BUSINESSES WILL HAVE TO BE MANAGED BY THE CIRM
4 ITSELF. SO WE WILL HAVE TO HAVE STAFF THAT ADMINISTERS
5 THIS POLICY WITHIN CIRM. AND IT'S BECOMING ABUNDANTLY
6 CLEAR, I THINK, THROUGH ALL OF THE CONVERSATIONS WE'VE
7 HAD, INCLUDING JOHN'S PRESENTATION THIS MORNING, THE
8 CONTRAST TO THE SITUATION WITH THE NONPROFITS WHERE
9 IT'S LIKELY THAT THE VAST MAJORITY OF WHAT WE DO WILL
10 BE IN THE FORM OF GRANTS. WITH COMPANIES THERE ARE
11 POSSIBILITIES TO MAKE LOANS, TO MAKE GRANTS, TO BUY
12 SERVICES. SO EACH ONE OF THESE WILL HAVE TO BE DEALT
13 IN SOME DEGREE IN A SEPARATE WAY.

14 WE'VE ALSO HEARD THAT WHILE FROM JDRC, FOR
15 EXAMPLE, WHO HAS POINTED OUT THAT THEY ESSENTIALLY HAVE
16 TO NEGOTIATE DIFFERENT CONTRACTS EVERY TIME THEY MAKE
17 AN ARRANGEMENT WITH A FOR-PROFIT ENTITY. SO TO SOME
18 DEGREE THAT IS A LAYER OF COMPLEXITY. I THINK IN THIS
19 SENSE WHAT WE'RE DOING HERE WILL OVERLAP WITH THE
20 STRATEGIC PLANNING PROCESS, WHICH WILL DEFINE SOME OF
21 THESE CHARACTERISTICS. BUT WE'RE LIKELY TO END UP WITH
22 AT LEAST THREE WAYS, DIFFERENT WAYS, IN WHICH WE COULD
23 CONCEIVABLY PROVIDE FUNDING.

24 DR. REED: JOHN REED HERE. LAST TIME WE
25 DISCUSSED THIS I BELIEVE WE DECIDED THAT THE INVENTORS

1 OF THE TECHNOLOGY WOULD BE RESPONSIBLE FOR COVERING THE
2 PATENT COSTS, THAT THE STATE WOULD NOT DO THAT; IS THAT
3 CORRECT?

4 CHAIRMAN PENHOET: WE DON'T HAVE ANY FUNDS TO
5 DO THAT.

6 DR. REED: OKAY. I THINK THAT'S AN IMPORTANT
7 THING TO ALSO KEEP IN MIND WITH RESPECT TO WHATEVER
8 POLICY WE ARRIVE AT INASMUCH AS IT'S NOT UNCOMMON TO
9 SPEND ABOUT A QUARTER MILLION DOLLARS PER PATENT FOR
10 WORLDWIDE PROTECTIONS AS YOU GO THROUGH THE WHOLE
11 PROCESS WITH EACH OF THE TERRITORIES. AND WE DON'T
12 WANT TO DO THINGS THAT WOULD DISINCENTIVIZE COMPANIES
13 TO PROTECT THEIR TECHNOLOGY AND TO, THEREFORE, PROTECT
14 THE INVESTMENT THAT THE TAXPAYERS ARE MAKING IN THAT
15 TECHNOLOGY. IF IT'S NOT PROTECTED BY PATENTS, THEN
16 IT'S NOT GOING TO HAVE THE OPPORTUNITY TO REALLY RETURN
17 FUNDS HERE, REVENUE HERE TO THE STATE. SO I THINK THAT
18 HAS TO BE AN IMPORTANT ELEMENT OF WHATEVER WE --
19 WHATEVER POLICY WE ARRIVE AT.

20 CHAIRMAN PENHOET: OKAY. WE'RE NOW READY TO
21 GO WITH DAVID GOLLAHER'S PRESENTATION. DAVID.

22 MR. GOLLAHER: THANK YOU, ED. I'M DAVID
23 GOLLAHER. I'M PRESIDENT AND CEO OF THE CALIFORNIA
24 HEALTH CARE INSTITUTE. CHI'S BACKGROUND, WE'RE ABOUT
25 260 MEMBERS STATEWIDE, AN ORGANIZATION THAT INCLUDES

1 ACADEMIC RESEARCH CENTERS, UNIVERSITY OF CALIFORNIA,
2 CSU, SALK, IF NOT ALL OF THE ACADEMIC INSTITUTIONS THAT
3 WILL BE CANDIDATES FOR CIRM FUNDING FOR GRANTING STEM
4 CELL RESEARCH ALONG WITH COMMERCIALIZED SCIENCES. SO
5 MOST MAJOR BIOTECH COMPANIES, I THINK ALL OF THEM, IN
6 FACT, IN CALIFORNIA ARE MEMBERS OF OURS ALONG WITH
7 MEDICAL DEVICES, DIAGNOSTICS COMPANIES LIKE GENPRO, AS
8 WELL AS BIOINFORMATICS, RESEARCH TOOLS COMPANIES LIKE
9 INVITROGEN. SO BROADLY SPEAKING, WE REPRESENT THE LIFE
10 SCIENCES IN CALIFORNIA, AND I THINK IT'S REFLECTIVE OF
11 OUR MEMBERS ON OUR BOARD.

12 JUST TO FRAME THE DISCUSSION, AND I THINK
13 IT'S NOT INCIDENTAL THAT CALIFORNIA IS BOTH THE STATE
14 THAT PASSED PROP 71, BUT ALSO THE ROLE OF
15 BIOTECHNOLOGY. THE INDUSTRY ORIGINATED HERE SOME 30
16 ODD YEARS AGO. PERHAPS 40 PERCENT OF ALL TOTAL U.S.
17 BIOTECH JOBS ARE HERE. AND, OF COURSE, WE HAVE THE
18 WORLD'S PREMIERE ACADEMIC INSTITUTIONS, PUBLIC AND
19 PRIVATE. AND IMPORTANTLY, A HIGHLY EVOLVED
20 INFRASTRUCTURE.

21 WHAT'S IMPORTANT ABOUT THAT IS THAT THE
22 FUNDING FOR EMBRYONIC STEM CELL RESEARCH, THE PERHAPS
23 \$300 MILLION A YEAR OVER TEN YEARS, WOULD BE HIGHLY
24 LEVERAGED. BEING ABLE TO COME INTO A MARKET IN WHICH
25 IT DOESN'T HAVE TO CREATE INFRASTRUCTURE FROM THE

1 GROUND UP. IT CAN TAKE ADVANTAGE OF THE SCIENCE, THE
2 INFRASTRUCTURE, AND COMMERCIAL NETWORK THAT'S ALREADY
3 HERE.

4 THIS IS JUST A CHART YOU CAN'T READ VERY
5 WELL, BUT IT SHOWS ON THE LEFT HAND CALIFORNIA'S
6 BIOTECH COMPANIES COMPARED TO EVERYONE ELSE. THE NEXT
7 CLOSEST IS MASSACHUSETTS. AS YOU CAN SEE, IT'S A VERY
8 DISTANT SECOND.

9 NOW, SOME OF YOU KNOW MICHAEL PORTER'S WORK
10 ABOUT CLUSTERS, AND MICHAEL PORTER IS A PROFESSOR AT
11 HARVARD BUSINESS SCHOOL AND HAS SPENT YEARS STUDYING
12 THE DEVELOPMENT OF CLUSTERS OF INNOVATION, HE CALLS
13 THEM. AND I JUST WANTED TO RUN THROUGH THIS BECAUSE IT
14 WILL COME BOOK TO A POINT THAT I THINK IS IMPORTANT IN
15 THINKING ABOUT INTELLECTUAL PROPERTY. IN PARTICULAR,
16 HOW COMPANIES, COMMERCIAL COMPANIES, WORK ALONGSIDE
17 ACADEMIC AND PUBLICLY FUNDED INSTITUTIONS TO PRODUCE
18 PUBLIC BENEFIT.

19 I'LL RUN THROUGH THE LIST, AND THEY'RE ONES
20 THAT YOU KNOW. YOU CAN LOOK AT A CLUSTER LIKE SAN
21 DIEGO HERE IN THE TORREY PINES AREA WHERE WE HAVE A
22 STRONG SCIENCE BASE, ENTREPRENEURIAL CULTURE, A GROWING
23 COMPANY BASE. IT'S IMPORTANT TO HAVE COMMERCIAL
24 COMPANIES THAT ARE THRIVING, ABSORBING, SPENDING IT ON
25 TALENT, WORKING WITH INSTITUTIONS, THE ABILITY TO

1 ATTRACT TALENT, THE INFRASTRUCTURE, ACCOUNTING FIRMS,
2 LAW FIRMS, REAL ESTATE COMPANIES THAT ARE EXPERTS AT
3 BUILDING FACILITIES. SOURCES OF RISK CAPITAL. YOU CAN
4 HAVE ALL OF THESE ELEMENTS, BUT IF YOU DON'T HAVE
5 SOURCES OF RISK CAPITAL, YOU DON'T HAVE THE CATALYST,
6 AND YOU DON'T HAVE THE INITIATIVE TO MAKE IT WORK.
7 LARGER COMPANIES IN RELATED INDUSTRIES, VERY IMPORTANT,
8 SO THAT THE PRESENCE OF AN INVITROGEN OR A BIOGEN IDEC
9 IS AN IMPORTANT ELEMENT IN CREATING A CLUSTER.
10 OBVIOUSLY A SKILLED WORKFORCE, EFFECTIVE NETWORKING,
11 ORGANIZATIONS LIKE CONNECT, EDC, WHO ARE IN THE ROOM
12 HERE, THAT PROVIDE CHANNELS OF COMMUNICATION AND
13 NETWORKING FOR EXECUTIVE SCIENTISTS AND SO FORTH TO
14 SHARE INFORMATION AND ACCELERATE THIS KNOWLEDGE
15 INDUSTRY. AND FINALLY, SUPPORTIVE PUBLIC POLICY. I
16 WILL COME BACK TO THAT IN A SECOND.

17 SO IN PORTER'S VIEW, WHAT MAKES CLUSTERS
18 PRODUCTIVE, AND YOU CAN LOOK AROUND THE WORLD AND SEE
19 ONES THAT ARE HIGHLY PRODUCTIVE, ONES THAT HAVE MANY OF
20 THE ELEMENTS THAT I JUST TALKED ABOUT THAT DON'T WORK
21 VERY WELL, DON'T WORK VERY WELL IN TERMS OF PRODUCING
22 INNOVATION AND PUSHING THAT INNOVATION OUT TO THE
23 MARKETPLACE WHERE IT CAN BE USED BY CONSUMERS, BY
24 PATIENTS, AND HELP PEOPLE.

25 SO THE INTERACTIVITY WITHIN A NETWORK, VERY

1 IMPORTANT HOW DYNAMIC IT IS, AND COMPETITION WITHIN THE
2 NETWORKS. PORTER MAKES A VERY GOOD AND, I THINK,
3 DEEPLY IMPORTANT POINT IN THAT IF YOU ARE IN A BUSINESS
4 AND YOU'RE A SINGLE BUSINESS PERSON, AND YOU LOOK
5 ACROSS THE STREET AND YOU SEE YOUR COMPETITOR RISING UP
6 AND DOING WELL, YOU TEND TO RESIST THAT AND THINK
7 COMPETITION IS A BAD THING. BUT WHAT HE'S FOUND IN THE
8 STUDY OF CLUSTERS, IN FACT, THAT THE MORE COMPETITIVE
9 THEY ARE, THE MORE BUSINESSES THAT ENTER THE SPACE TO
10 COMPETE TO PROVIDE SUPERIOR PRODUCTS, THE MORE
11 INNOVATION HAPPENS AND THE FASTER IT'S PUSHED OUT IN
12 THE MARKET.

13 FINALLY, THE VELOCITY OF TECHNOLOGY TRANSFER,
14 CRITICAL POINT. AND I'LL DRILL DOWN INTO THIS A LITTLE
15 BIT MORE IN A MINUTE. HOW QUICKLY TECHNOLOGY CAN BE
16 TRANSFERRED FROM BASIC SCIENCE INSTITUTIONS FROM
17 PERHAPS GOVERNMENT-SPONSORED UNIVERSITIES OUT INTO
18 COMMERCIAL COMPANIES. SO THE COMMERCIALIZATION OF
19 ACADEMIC SCIENCE IS ONE OF THE HALLMARKS OF SUCCESSFUL
20 CLUSTERS.

21 SOME PEOPLE ASK AND THERE'S ACTUALLY A CHORUS
22 OF CRITICS, PARTICULARLY IN THE PHARMACEUTICAL
23 INDUSTRY, RIGHT NOW WHO ASK WHY COMMERCIALIZATION IS
24 NECESSARY IN THE FIRST PLACE. THERE'S WHAT I WOULD
25 THINK OF AS THE MARCIA ANGEL THEORY. MARCIA ANGEL WAS

1 THE EDITOR OF THE *NEW ENGLAND JOURNAL* AND HAS WRITTEN A
2 BOOK ABOUT THE DRUG INDUSTRY THAT BASICALLY PROPOUNDS
3 THE THEORY THAT THE GOVERNMENT, THROUGH NIH, CREATES
4 BASIC RESEARCH AND COMPANIES COME ALONG AND
5 COMMERCIALIZE IT AND BASICALLY TAKE THE INTELLECTUAL
6 PROPERTY THAT WILL BE FUNDED FOR THEIR OWN PURPOSES AND
7 THEIR SHAREHOLDERS' PURPOSES. AND THERE'S NO REAL
8 RETURN TO SOCIETY FOR THAT. I THINK THAT'S WRONG. I
9 THINK IT COMPLETELY MISCHARACTERIZES AND MISUNDERSTANDS
10 THE ROLE OF COMMERCIALIZATION, THE ROLE OF THE MARKET,
11 AND THE ROLE OF PRIVATE COMPANIES.

12 FIRST THING IS THAT BASIC RESEARCH PRODUCES
13 UNDERSTANDING, BUT IT DOESN'T PRODUCE PRODUCTS. SECOND
14 THING IS THAT THE INSTITUTIONS THAT ARE GOOD AT
15 PRODUCING BASIC RESEARCH LACK THE SKILLS AND RESOURCES
16 TO PRODUCE PRODUCTS. THEY LACK THE CAPITAL, AND IT'S
17 LARGELY AN EXPENSIVE PROCESS. THE LACK THE APPLIED
18 RESEARCH AND REGULATORY EXPERTISE THAT IS A HUGE PART
19 OF PRODUCT DEVELOPMENT. AND OBVIOUSLY THEY LACK BOTH
20 THE MANUFACTURING CAPACITY, NOT TO MENTION THE
21 MARKETING AND DISTRIBUTION. THE MARKETING AND
22 DISTRIBUTION IS AN IMPORTANT POINT BECAUSE
23 UNDERSTANDING MARKETS, PARTICULARLY DISEASE MARKETS,
24 UNDERSTANDING HOW PHYSICIANS AND HOSPITALS AND THE
25 WHOLE COMPLICATED HEALTHCARE SYSTEM WORKS AND ABILITY

1 TO MOVE A PRODUCT THROUGH THAT IS NOT TRIVIAL.

2 SO WHAT DOES IT TAKE TO TRANSFORM A RESEARCH
3 DISCOVERY INTO A SUCCESSFUL DRUG? WELL, INDUSTRY
4 CRITICS CLAIM THAT A RESEARCH INPUT CREATES MOST OF THE
5 PRODUCTS' END VALUE. BUT THERE'S NO GOOD MEASURE OF
6 THE REAL PERCENTAGE OF WHAT IT TAKES TO GET A PRODUCT
7 FROM THE LABORATORY INTO THE HANDS OF THE PATIENT
8 THROUGH A PHYSICIAN. SOME INDUSTRY STUDIES HAVE
9 SUGGESTED THAT 90 PERCENT OF THE VALUE OF AN END
10 PRODUCT IS DOWNSTREAM. BUT I THINK THE TRUTH IS THAT
11 NO ONE REALLY KNOWS. WE DO KNOW THAT IT'S ENORMOUSLY
12 TIME-CONSUMING AND EXPENSIVE, AND HERE'S ANOTHER SLIDE
13 THAT YOU CAN'T READ VERY WELL, BUT AT THE BOTTOM IS
14 YEARS, AND IT GOES FROM ZERO TO 16 YEARS. AND THE
15 FIRST BLOCK UP ON THE LEFT-HAND CORNER IS THE DISCOVERY
16 PROCESS, AND THEN WE MOVE FROM THAT PERIOD THROUGH A
17 WHOLE SET OF PRODUCT DEVELOPMENT AND REGULATORY PHASES
18 UNTIL WE FINALLY GET A PRODUCT OUT INTO THE PIPELINE.

19 AND IF YOU LOOK AROUND AT COMPANIES THAT WE
20 KNOW OF, THERE'S A GREAT EXAMPLE IN SAN DIEGO, BIOGEN,
21 IS NOW -- SORRY -- IDEC, WHICH IS NOW BIOGEN IDEC AFTER
22 A MERGER, TOOK ABOUT 13 YEARS TO MOVE FROM ITS ORIGINAL
23 ANTIBODY SCIENCE TO PRODUCING THE FIRST MONOCLONAL
24 ANTIBODY FOR CANCER FOR NON-HODGKIN'S. AND DURING THAT
25 PERIOD CAME WITHIN THREE OR FOUR WEEKS OF RUNNING

1 ENTIRELY OUT OF MONEY AND TURNING OUT THE LIGHTS. HAD
2 THAT HAPPENED, THOUSANDS UPON HUNDREDS OF THOUSANDS OF
3 PATIENTS WOULD HAVE BEEN DENIED AN EFFECTIVE DRUG.

4 THE END RESULT OF BIOGEN IDEC OR IDEC'S
5 RESEARCH AND DEVELOPMENT WAS TO PRODUCE A DRUG THAT IS
6 OF ENORMOUS BENEFIT TO PATIENTS WHO HAVE TERRIBLE
7 DEBILITATING AND OFTEN FATAL DISEASE.

8 SO IN A SENSE, IF WE LOOK AROUND GLOBALLY,
9 WE'VE BEEN DOING AN INTERESTING EXPERIMENT WITH RESPECT
10 TO INTELLECTUAL PROPERTY AND HOW IT'S TRANSFERRED INTO
11 COMMERCIAL INSTITUTIONS OR COMMERCIAL COMPANIES. AND
12 RECENTLY I HAD THE OPPORTUNITY TO BE IN THE UK AND TO
13 TALK TO SOME PEOPLE ABOUT THE ATTEMPT IN THE UNITED
14 KINGDOM TO ACCELERATE THE TECHNOLOGY TRANSFER OF
15 SCIENCE INTO COMPANIES MORE OR LESS ALONG THE U.S.
16 MODEL. AND THIS WAS SPURRED IN PART BY THE REALIZATION
17 THAT WHILE UK UNIVERSITIES, LIKE OXFORD AND CAMBRIDGE
18 AND THE UNIVERSITY OF THE PLYMOUTH, WERE QUITE GOOD AT
19 BASIC RESEARCH AND TEACHING. THEY HAVE BEEN EXTREMELY
20 BACKWARD WITH RESPECT TO MOVING ANY OF THAT SCIENCE
21 INTO THE COMMERCIAL SPHERE.

22 THEY HAD ALSO EXPERIENCED A BRAIN DRAIN, SO
23 THAT IF YOU WALKED AROUND CALIFORNIA OR ROUTE 128
24 CORRIDOR IN BOSTON, YOU WOULD BUMP INTO A BUNCH OF
25 EX-PAT BRITS WHO HAVE MOVED TO THE UNITED STATES IN

1 ORDER TO MOVE THEIR TECHNOLOGY, THEIR EXPERTISE INTO
2 COMMERCIAL SPACE.

3 WHAT THE UK ALSO DISCOVERED IS THAT THEY HAVE
4 A HUGE RESERVOIR OF WHAT MIGHT BE CALLED ORPHAN IP;
5 THAT IS, INTELLECTUAL PROPERTY TO WHICH A UNIVERSITY
6 HAD RIGHTS, BUT THERE WAS NO ABILITY NOR INTEREST IN
7 DEVELOPING THAT AND MOVING IT FORWARD INTO THE MARKET.

8 SO IN 1999 THE DEPARTMENT OF TRADE AND
9 INDUSTRY IN THE UK PROPOSED A THIRD MISSION, WHICH IS
10 SUPPORT INDUSTRY. AND SINCE THEN GRADUALLY, AND WITH
11 CLINIC OF THE UK, IS BEGINNING TO DEVELOP ITS FIRST
12 LIFE SCIENCES CLUSTERS, I BELIEVE ONE AROUND CAMBRIDGE
13 UNIVERSITY, TO ACCELERATE SCIENCE MOVING FORWARD.

14 I WANT TO SHIFT FOCUS TO TALK JUST IN A
15 GENERAL WAY ABOUT TWO MODELS THAT ARE WORTH THINKING
16 ABOUT IN THE WAYS THAT GOVERNMENT FUNDS SCIENCE WITHIN
17 THE CONTEXT OF COMMERCIAL COMPANIES. AND I WANT TO
18 START WITH DARPA. I JUST PUT DARPA'S MISSION STATEMENT
19 DOWN HERE. I THINK MOST OF US WOULD SEE DARPA AS A
20 REASONABLY SUCCESSFUL LONG-TERM INITIATIVE IN WHICH
21 PUBLIC MONEY, I.E., DEPARTMENT OF DEFENSE FUNDS, ARE
22 SPECIFICALLY TARGETED IN QUITE CAREFUL WAYS CHanneled
23 INTO COMMERCIAL OPERATIONS TO DEVELOP BASIC TECHNOLOGY
24 THAT THEN MAY BE APPLIED TO DEFENSE APPLICATIONS.

25 THE PART THAT I UNDERLINED HERE IS FROM

1 DARPA'S MISSION STATEMENT IN WHICH IT SAYS THAT IT
2 MANAGES AND DIRECTS SELECTED BASIC AND APPLIED RESEARCH
3 AND DEVELOPMENT PROJECTS FOR DOD, TO PURSUE THIS
4 RESEARCH AND TECHNOLOGY WHERE RISK AND PAYOFF ARE BOTH
5 VERY HIGH AND WHERE SUCCESS MAY PROVIDE DRAMATIC
6 ADVANCES FOR TRADITIONAL MILITARY ROLES AND MISSIONS.

7 IT STRIKES ME, IN THINKING ABOUT CIRM, THAT
8 THIS IS A GOOD ANALOGY TO WHAT WE'RE LOOKING AT IN THE
9 EARLY STAGE OF EMBRYONIC STEM CELL RESEARCH. HIGH RISK
10 AND A POTENTIALLY VERY HIGH PAYOFF, GREAT UNCERTAINTY,
11 AND A PLACE WHERE PERHAPS PUBLIC MONEY CAN BE A
12 CATALYST.

13 NOW, DARPA'S PRINCIPLES ARE TO PROMOTE
14 PRIVATE SECTOR COMPETITION. THEY MAKE GRANTS TO LOTS
15 OF DIFFERENT COMPANIES AND OFTEN TO COMPETITORS WITHIN
16 THE SAME FIELD. THE THOUGHT IS THAT HERE GOVERNMENT IS
17 A FUNDING CATALYST IN AREAS OF PARTICULARLY HIGH
18 IMPORTANCE, HIGH RISK WORK. ULTIMATELY THERE ALSO IS
19 THE BENEFITS TO TAXPAYERS, BENEFITS TO THE PUBLIC DON'T
20 ACCRUE FROM A DIRECT RETURN IN THE FORM OF A ROYALTY OR
21 MONEY PAYMENTS BACK TO GOVERNMENT, BUT RATHER FROM THE
22 PROMULGATION OF INVENTIONS INTO A SPECIFIC SECTOR, IN
23 THIS CASE THE NATIONAL DEFENSE.

24 AND THIS IS AN IMPORTANT THEORETICAL, IF YOU
25 WILL, ISSUE FOR CIRM AND, IN FACT, FOR THE LEGISLATURE

1 AND FOR THE PUBLIC TO CONSIDER. THAT IS, WHETHER THE
2 MAXIMUM PUBLIC BENEFIT FROM STEM CELL RESEARCH WILL
3 COME FROM THE FASTEST POSSIBLE DEVELOPMENT OF THE BEST
4 SCIENCE AND THE FASTEST ACCELERATION OF THAT SCIENCE
5 INTO COMMERCIAL PRODUCTS OR FROM ROYALTIES, FROM MONEY
6 PAYMENTS FROM THESE INVENTIONS BACK TO THE PUBLIC
7 COFFERS.

8 SO MY VIEW IS THAT CIRM SHOULD DISCOUNT
9 NAYSAYERS WHO MISREPRESENT -- WE'VE SEEN THIS FROM THE
10 VERY BEGINNING OF PROP 71 -- THE INTERRELATIONSHIP ON
11 THE ACADEMIC AND COMMERCIAL SCIENCE. IT'S MORE
12 COMPLICATED THAN MOST PEOPLE APPRECIATE. IT'S FAR FROM
13 THE CASE THAT ACADEMIC SCIENCE CREATES DRUGS AND THAT
14 COMMERCIAL DEVELOPMENT OF THOSE DRUGS IS INCIDENTAL OR
15 AN ADD-ON.

16 CIRM SHOULD DISCOUNT ANY CLAIM THAT STEM CELL
17 SCIENCE CAN BENEFIT PATIENT GROUPS WITHOUT AGGRESSIVE
18 PARTICIPATION OF COMMERCIAL COMPANIES. THE FACT IS
19 THAT WE LIVE IN A MARKET-BASED, MARKET DRIVEN ECONOMY.
20 AND THE THEORY THAT MOST COMMERCIAL COMPANIES OPERATE
21 ON IS THE THEORY OF CAPITALISM, AND THE BEST
22 DISTRIBUTION OF GOODS AND SERVICES HAPPENS THROUGH THE
23 ACTIVE PARTICIPATION OF THE MARKET AND OF COMMERCIAL
24 ENTERPRISE. AND, AGAIN, ANY OF US WHO SPENT
25 SIGNIFICANT TIME IN ACADEMIC INSTITUTIONS OR WITHIN

1 GOVERNMENT INSTITUTIONS, I THINK, HAVE A LOW LEVEL OF
2 CONFIDENCE IN THE ABILITY OF THOSE ORGANIZATIONS TO
3 QUICKLY MOVE TECHNOLOGY THROUGH THE WIDEST POSSIBLE
4 ALLOCATIONS FOR THE GREATEST PUBLIC BENEFIT.

5 FINALLY, CIRM SHOULD DISCOUNT ATTEMPTS TO
6 REGULATE COMMERCIAL TRANSACTIONS IN WAYS THAT
7 DISCOURAGE PARTICIPATION OF THE BEST COMPANIES AND
8 ENTREPRENEURS. I THINK THE CONCERN THAT WE'VE HEARD
9 MORE THAN ONCE IS THAT THE MORE STRENUOUS CONDITIONS,
10 ROYALTIES, CAVEATS THAT ARE ATTACHED TO TECHNOLOGY
11 TRANSFER RELATIONSHIPS, THE MORE RELUCTANT COMMERCIAL
12 PARTICIPANTS ARE TO CAPITALIZE ON THOSE RELATIONSHIPS
13 AND THOSE TECHNOLOGIES.

14 SO I THINK FROM CHI'S POINT OF VIEW, GUIDING
15 PRINCIPLES FOR CIRM WITH RESPECT TO COMMERCIAL IP
16 POLICY SHOULD BE, FIRST, MAXIMUM ACCELERATION OF THE
17 BEST SCIENCE. AND WHETHER THAT SCIENCE AS TECHNOLOGY
18 EXISTS IN A COMMERCIAL COMPANY OR IN AN ACADEMIC
19 INSTITUTION, CIRM SHOULD BE RIGOROUS AND THOUGHTFUL IN
20 MAKING INVESTMENTS BEHIND THE BEST SCIENCE. CIRM HAS
21 COLLECTED AN ASTONISHINGLY GOOD SET OF EXPERT
22 SCIENTIFIC REVIEWERS, AND THE ABILITY OF CIRM TO MAKE
23 INTELLIGENT JUDGMENTS WITH RESPECT TO WHERE THE BEST
24 SCIENCE IS AND WHERE FUNDING OPPORTUNITIES LIE, I
25 THINK, IS SUPERB AND SHOULD BE APPLAUDED.

1 SECOND, INVESTMENT IN IMPORTANT AREAS OUTSIDE
2 ACADEMIC EXPERTISE ARE IMPORTANT FOR ANY COMMERCIAL
3 PRODUCT OR WIDE-SCALE APPLICATION OF STEM CELL
4 RESEARCH. SO, FOR EXAMPLE, SCALED PRODUCTION IS AN
5 AREA THAT'S UNLIKELY TO BE FOUND IN A UNIVERSITY AND
6 MAY WELL EXIST IN A COMPANY THAT MIGHT BE A TARGET FOR
7 CIRM FUNDING. AND I'M SURE THERE ARE MANY, MANY OTHER
8 AREAS.

9 FINALLY, THAT CIRM SHOULD FOSTER THE
10 DEVELOPMENT OF BUSINESS MODELS FOR COMMERCIAL STEM
11 CELL-BASED PRODUCTS. PART OF THE REASON, IN FACT,
12 PERHAPS THE MAIN REASON, THAT VENTURE CAPITALISTS HAVE
13 NOT BEEN FUNDING STEM CELL RESEARCH IN A LARGE WAY IS
14 THAT WE DON'T HAVE BUSINESS MODELS TODAY THAT SUPPORT
15 THAT. AND WHEN WE DO AND IF THERE'S A CLEAR PATH TO
16 PROFITABILITY, THE LANDSCAPE WILL CHANGE DRAMATICALLY.

17 SO I HAVE THREE EXAMPLES FOR YOU. THIS IS
18 THE FUN PART OF THE PROGRAM. SO HERE'S A
19 GOVERNMENT-FUNDED CAR. COST ABOUT \$22 MILLION, AND IT
20 WORKED VERY, VERY WELL FOR ITS PURPOSE. THIS IS A
21 GOVERNMENT-BUILT TRUCK. THIS WAS BUILT IN 1956 IN THE
22 SOVIET UNION. THIS IS A PUBLIC-PRIVATE PARTNERSHIP
23 CAR, AND IT'S INTERESTING. IT REPRESENTS AN
24 EXTRAORDINARY CONTRIBUTION OF, IN THIS CASE ORIGINALLY
25 IN JAPAN, OF PUBLIC MONEY CHanneled INTO TOYOTA OF A

1 PRIVATE COMPANY ON THE THEORY THAT IF TOYOTA BECAME
2 MORE COMPETITIVE AND SOLD MORE CARS, IT WOULD BENEFIT
3 THE JAPANESE ECONOMY. SO FAR SO GOOD.

4 SO THANK YOU FOR YOUR ATTENTION. AND I'D BE
5 HAPPY TO ENTERTAIN ANY QUESTIONS OR COMMENTS.

6 CHAIRMAN PENHOET: THANK YOU VERY MUCH,
7 DAVID. SO STARTING HERE IN SAN DIEGO WITH THE MEMBERS
8 OF THE TASK FORCE.

9 DR. PRIETO: FRANCISCO PRIETO. I GUESS MY
10 QUESTION UNDER THE DARPA MODEL THAT YOU CITED IS WHAT
11 IS THE BENEFIT TO TAXPAYERS BECAUSE THAT SEEMS PERHAPS
12 NEBULOUS?

13 AND ALSO, JUST A COMMENT THAT I WOULD SEE
14 SHARING IN ROYALTIES AS NOT A TAX, BUT MORE AN
15 EXPECTATION THAT AN INVESTOR MIGHT PUT IN. AND, OF
16 COURSE, HERE THE INVESTORS ARE THE STATE OF CALIFORNIA.

17 MR. GOLLAHER: SURE. WHAT'S THE PUBLIC
18 BENEFIT FROM DARPA? THE THEORY, AND YOU CAN ACCEPT IT
19 OR NOT, IS THAT PUBLIC SAFETY THROUGH THE MILITARY
20 BEING EQUIPPED WITH THE MOST ADVANCED TECHNOLOGY IS THE
21 PUBLIC BENEFIT. DOES THAT HAVE AN ECONOMIC
22 IMPLICATION? AND I THINK THE ANSWER WOULD BE YES. SO
23 I GUESS THAT'S AS FAR AS I CAN TAKE THAT.

24 IN TERMS OF A ROYALTY PERCENTAGE RETURN BEING
25 A TAX, I DON'T KNOW HOW ELSE I WOULD CONSTRUE THAT. I

1 MEAN IT IS A TAX. ON THE TRANSACTION, TAXES ARE NOT
2 NECESSARILY BAD THINGS, SO I'M NOT SUGGESTING THAT IT'S
3 WRONG OR BAD. I'M ONLY SUGGESTING THAT IT BECOMES PART
4 OF THE EQUATION IN THE COMMERCIAL TRANSACTION THAT
5 SOMETHING THAT HAS TO BE TAKEN INTO ACCOUNT BECAUSE IT
6 HAS ECONOMIC VALUE. I'M NOT AGAINST TAXES.

7 DR. PRIETO: I JUST, YOU KNOW, WONDER HOW
8 ELSE WOULD AN INVESTOR EXPECT TO GET A RETURN?

9 MR. GOLLAHER: WELL, I THINK THIS CUTS TO THE
10 VERY HEART OF THE MATTER, AND ACTUALLY CUTS TO THE
11 HEART OF JOHN SIMPSON'S PRESENTATION TOO. AND THAT HAS
12 TO DO WITH WHAT IS THE RETURN. AND I THINK THAT TO
13 MAKE A VERY EXTREME -- TO PRESENT AN EXTREME DICHOTOMY,
14 ON THE ONE HAND, YOU COULD ARGUE THAT THE GREATEST
15 RETURN TO PUBLIC HEALTH WOULD BE THE GREATEST
16 DISSEMINATION OF THESE TECHNOLOGIES INTO PRODUCTS FOR
17 PATIENTS. THAT PRODUCES A PUBLIC HEALTH RETURN.

18 IF, IN ADDITION TO THAT, YOU LOOKED FOR A
19 FINANCIAL MONETARY RETURN, I THINK THAT'S AN ADDITION
20 TOO. BUT THE ARGUMENT THAT WE HAD DURING THE
21 REASONABLE PRICING ATTACHMENT TO CRATA'S AT NIH AND THE
22 CONGRESS HAD OVER MONTHS AND MONTHS REVOLVED EXACTLY
23 AROUND THIS ISSUE ABOUT WHETHER NIH IN ITS INTELLECTUAL
24 PROPERTY TRANSFER COOPERATIVE RESEARCH AND DEVELOPMENT
25 AGREEMENTS WITH ACADEMIC INSTITUTIONS SHOULD EXPECT A

1 FINANCIAL RETURN OR WHETHER THE PUBLIC WOULD BENEFIT
2 MOST FROM FEWER STRINGS AND THE FASTEST POSSIBLE
3 ACCELERATION OF THE SCIENCE.

4 CHAIRMAN PENHOET: ED PENHOET. IF I MIGHT, I
5 THINK THE ISSUES OF DIRECT FINANCIAL RETURN VERSUS
6 INDIRECT FINANCIAL RETURN. THE OTHER ARGUMENT THAT'S
7 MADE, DARPA, FOR EXAMPLE, WAS THE ONLY FUNDER OF THE
8 EARLY DAYS OF THE TECHNOLOGY REVOLUTION. AN EXAMPLE IS
9 THE COMPUTER INDUSTRY PROVIDED GOVERNMENT FUNDING. IT
10 HAS ALREADY BEEN RETURNED, IN ADDITION TO DIRECT
11 RETURN, WHICH IS WHAT YOU'D BE TALKING ABOUT IF THERE
12 WAS A DIRECT RELATIONSHIP. THE INDIRECT RETURNS ARE A
13 MORE ROBUST ECONOMY IN FINANCIAL TERMS, MORE ROBUST
14 BUSINESS SECTOR PAYING TAXES.

15 MR. GOLLAHER: THAT'S WELL SAID.

16 DR. REED: JOHN REED. I WAS WONDERING IF YOU
17 COULD MAKE SOME GENERAL COMMENTS. I'M SURE YOUR
18 ORGANIZATION MUST LOOK AT THIS. AND THAT IS, ON THE
19 TOPIC OF THE STATE OF CALIFORNIA AND ITS RELATIVE
20 COMPETITIVENESS IN THE BIOTECHNOLOGY AND LIFE SCIENCE
21 INDUSTRY, AND SPEAK TO THE ISSUE ABOUT OUR STATE AND
22 ITS INCENTIVES AND BUSINESS FRIENDLY PRACTICES RELATIVE
23 TO OUR STATE AND OTHER STATES IN TERMS OF DISINCENTIVES
24 AND BUSINESS UNFRIENDLY PRACTICES AND HOW THAT WILL
25 WEIGH IN, DO YOU THINK, ON THE ULTIMATE BENEFIT TO

1 PATIENTS AND RETURNED ECONOMICALLY TO THE STATE.

2 MR. GOLLAHER: THAT'S A GREAT QUESTION, AND
3 THE TENTATIVE LANDSCAPE IN BIOTECH AND THE LIFE
4 SCIENCES IS CHANGING VERY QUICKLY RIGHT NOW, CERTAINLY
5 OVER THE PAST FEW YEARS. I THINK THE FIRST COMMENT IS
6 THAT OUR GLOBAL LEADERSHIP IN LIFE SCIENCES DIDN'T
7 HAPPEN AS THE RESULT OF ANY STATE POLICY EXCEPT FOR
8 ONE, AND THAT WAS THE FUNDING AND BUILDING OF RESEARCH
9 EXCELLENCE AT THE UNIVERSITY OF CALIFORNIA. AND THAT
10 BECAME AN EXTRAORDINARILY IMPORTANT ENGINE,
11 INTELLECTUAL ENGINE, BEHIND THE GROWTH OF THE LIFE
12 SCIENCES INDUSTRY.

13 NONETHELESS, OVER THE PAST 20 OR 30 YEARS,
14 CALIFORNIA HAS TURNED INTO AN INCREASINGLY DIFFICULT
15 PLACE TO DO BUSINESS, AND MUCH OF IT IS OUTSIDE THE
16 CONTROL OF GOVERNMENT, HIGH HOUSING PRICES, LOTS OF
17 INFRASTRUCTURE PROBLEMS, AND SO FORTH. AND YET WE HAVE
18 SO MANY OF THOSE ELEMENTS THAT I ALLUDED TO WITH
19 RESPECT TO CLUSTER THAT WE'RE STILL A MAGNET.

20 NOW, AT THE MARGIN OTHER COMPANIES AND
21 OTHER -- I'M SORRY -- OTHER COUNTRIES AND STATES HAVE
22 UNDERSTOOD THAT THESE ARE TERRIFIC CLUSTERS TO DEVELOP,
23 THAT THEY'RE HIGH PAYING JOBS, KNOWLEDGE WORKERS, THE
24 KIND OF COMPANIES AND UNIVERSITIES THAT YOU WOULD WANT
25 YOUR CHILDREN TO WORK FOR, AND THE KINDS OF THINGS THAT

1 YOU WANT IN YOUR COMMUNITY THAT HAVE BECOME
2 INCREASINGLY AGGRESSIVE. SO WE'RE SEEING OHIO,
3 FLORIDA, MARYLAND, SINGAPORE WITH EXTRAORDINARY
4 ECONOMIC PACKAGES TO ENTICE COMPANIES OR ENTICE PARTS
5 OF THEIR OPERATIONS OR, INDEED, MANUFACTURING TO MOVE
6 OFFSHORE.

7 WE ARE PERHAPS AT AN INFLECTION TIME NOW IN
8 WHICH THE NEXT STAGE OF DEVELOPMENT IN OUR INDUSTRY
9 WILL HAPPEN IN SOUTHERN CALIFORNIA, SOME OUTSIDE, AND I
10 THINK YOU AT BURNHAM ARE SEEING THE DESKTOP EDITION, IN
11 THIS CASE WITH RESPECT TO FLORIDA, IN WHICH THE
12 GOVERNOR, THE LEGISLATURE PERSONALLY BECOME INVOLVED IN
13 ATTEMPTING TO ATTRACT THE INDUSTRY.

14 NONETHELESS, THE STEM CELL INITIATIVE IN PROP
15 71 AND CIRM, AS IT MOVES FORWARD, IS TREMENDOUSLY
16 ATTRACTIVE IN CALIFORNIA, PARTICULARLY BECAUSE IT DOES
17 LEVERAGE INSTITUTES LIKE BURNHAM AND UCSD'S LIVE AND SO
18 FORTH.

19 CHAIRMAN PENHOET: THANK YOU. ANY COMMENTS
20 OR QUESTIONS FROM IRVINE?

21 DR. STEWARD: NO.

22 CHAIRMAN PENHOET: SAN CARLOS?

23 UNIDENTIFIED SPEAKER: NONE HERE.

24 CHAIRMAN PENHOET: LOS ANGELES?

25 UNIDENTIFIED SPEAKER: NONE.

1 CHAIRMAN PENHOET: CHICO.

2 DR. WRIGHT: NO.

3 CHAIRMAN PENHOET: OKAY. ANY COMMENTS FROM
4 THE AUDIENCE IN SAN DIEGO? AUDIENCE IN IRVINE? SAN
5 CARLOS? LOS ANGELES?

6 UNIDENTIFIED SPEAKER: NO.

7 CHAIRMAN PENHOET: OKAY. THANK YOU VERY
8 MUCH, DAVID. I THINK JULIE MEIER WRIGHT WILL PROBABLY
9 TALK MORE ABOUT THE COMPETITIVE INITIATIVE IN
10 CALIFORNIA. AND LIKE I SAID, REMIND PEOPLE ONCE AGAIN
11 ONE OF THE GOALS OF PROP 71 WAS THE ENHANCEMENT OF
12 CALIFORNIA'S POSITION IN THE BIOTECH INDUSTRY
13 GENERALLY. WE WILL HEAR MORE ABOUT THAT FROM JULIE,
14 FOR SURE, LATER ON.

15 OUR NEXT SPEAKER, AND I SAY IT'S VERY NICE TO
16 HAVE SOMEBODY ACTUALLY FROM A STEM CELL COMPANY FOR A
17 CHANGE TO GIVE US THE BENEFITS OF THEIR FACTS ON THIS
18 ISSUE. ALLAN ROBINS IS THE VICE PRESIDENT AND CHIEF
19 TECHNICAL OFFICER OF NOVOCELL, WHICH IS A STEM CELL
20 COMPANY, AND LOOK FORWARD TO YOUR COMMENTS.

21 MR. ROBINS: THANK YOU. FIRST OF ALL, I JUST
22 WANT TO THANK ED AND MARY FOR THE OPPORTUNITY TO TO
23 SPEAK TO YOU TODAY ABOUT THE IMPORTANT POLICY THAT
24 YOU'RE DEVELOPING. IT'S A PIONEER EFFORT.

25 NOVOCELL, I JUST WANT TO MAKE A FEW OPENING

1 COMMENTS THAT --

2 UNIDENTIFIED SPEAKER: CHICO CAN'T HEAR YOU.

3 MR. ROBINS: IS THAT BETTER? NOVOCELL IS A
4 SMALL BIOTECHNOLOGY COMPANY WITH THREE SUBS, ONE HERE
5 IN SAN DIEGO, ONE IN ORANGE COUNTY, AND WHERE I'M FROM
6 IN ATHENS, GEORGIA. AND WE'RE FOCUSED ON THE
7 DEVELOPMENT OF HUMAN ES CELLS TO PRODUCE A PRODUCT TO
8 CURE TYPE 1 DIABETES INITIALLY. THE COMPANY ALSO HAS
9 ENCAPSULATION TECHNOLOGY WHICH IS IN CLINICAL TRIALS AT
10 THIS POINT IN TIME.

11 SO I WANTED TO START THE TALK BY MAKING A
12 COUPLE OF GENERAL POINTS AND THEN GO ON TO A NUMBER OF
13 QUESTIONS THAT MARY POSED OF INTEREST. I'M SURE WE ALL
14 KNOW THIS, BUT I WANTED TO USE IT TO FRAME MY TALK.
15 STEM CELL BUSINESS, IT'S A TRUE BIOTECH OPPORTUNITY IN
16 MUCH THE SAME WAY AS RECOMBINANT DNA WAS A TRUE BIOTECH
17 OPPORTUNITY HERE IN CALIFORNIA IN THE '70S. AND I'M
18 OLD ENOUGH TO HAVE SORT OF LIVED THROUGH THAT AND
19 WORKED BEFORE RECOMBINANT DNA TECHNOLOGY, AND I KNOW
20 THERE WAS A LOT OF EXCITEMENT AT THE TIME, AND THERE
21 WAS A LOT OF CONCERNS, SAFETY ISSUES, AND THERE WAS A
22 LOT OF SPECULATION WHETHER WE WERE ACTUALLY GOING TO
23 GET ANYTHING COMMERCIAL THAT CAME OUT OF THAT. AND
24 THAT'S ALL HISTORY NOW WITH GENENTECH.

25 OBVIOUSLY AT THIS POINT IN TIME, IT'S A VERY

1 HIGH RISK, BUT THERE'S HIGH REWARD. AND IT'S DRIVEN BY
2 INNOVATION OF TECHNOLOGIES TO DISCOVER NOVEL PRODUCTS
3 AND SERVICES BASED ON THESE TECHNOLOGIES. BASICALLY IT
4 ENCAPSULATES THE STEM CELL OPPORTUNITIES BEFORE US
5 TODAY.

6 THE THING THAT REALLY EXCITES US AS A COMPANY
7 IS THE POTENTIAL TO PRODUCE TRUE DISEASE MODIFYING
8 TREATMENTS AND CURES FOR MANY DISEASES WHICH ARE
9 DEGENERATIVE DISEASES. TODAY WE TREAT THOSE DISEASES.
10 WE TREAT THE SYMPTOMS FOR THE DISEASES, BUT WE DON'T
11 TREAT THE CAUSES, WHICH IS DEGENERATION OF CELLULAR
12 SYSTEMS, AND SO WE HAVE NO CURES FOR THAT. WE'RE
13 SAYING THIS IS A NEW ERA IN MEDICINE, AND THAT THE
14 2000S WILL BE THE CENTURY OF THE CELL. SO THIS MAY BE
15 ONE OF THE LAST FRONTIERS IN HUMAN MEDICINE THAT WE ARE
16 VERY EXCITED TO BE INVOLVED IN.

17 I THOUGHT I'D GIVE THE PITCH FOR YOUR HOME
18 STATE. WHY CALIFORNIA? IT'S -- THIS POINT HAS BEEN
19 MADE, SO I WON'T BELABOR IT, BUT IT'S THE HUB OF
20 BIOTECHNOLOGY INNOVATION. YOU'VE GOT A LOT OF STEM
21 CELL TALENT; YOU'VE GOT A LOT OF BIOTECH TALENT.
22 YOU'VE GOT GREAT RESEARCH INSTITUTIONS. YOU'VE GOT
23 GREAT INFRASTRUCTURE, GOT THINGS LIKE SAN DIEGO
24 CONSORTIUM. YOU'VE GOT A CONGREGATION OF VERY
25 EXPERIENCED VENTURE CAPITALISTS. WE FEEL GOOD ABOUT

1 THIS BECAUSE THEY'VE MADE MONEY IN AREAS THAT THEY'VE
2 INVESTED IN BEFORE THAT WOULD SEEM TO BE HIGHLY RISKY.
3 AND, OF COURSE, PROPOSITION 71 AND THE INSTITUTE FOR
4 REGENERATIVE MEDICINE.

5 WHAT ARE THE CONCERNS FROM A COMPANY POINT OF
6 VIEW AND FROM OUR INVESTORS' POINT OF VIEW? WHAT ARE
7 THE CONCERNS THAT FACE US? THERE'S A FINANCIAL
8 UNCERTAINTY. WE CAN'T GO OUT AND RAISE FIVE YEARS
9 MONEY AND SIT ON THAT AND SAY TO OUR INVESTORS COME
10 BACK IN FIVE YEARS AND WE'LL SHOW YOU WHAT WE'VE
11 DEVELOPED. SO IT'S PRETTY MUCH HAND TO MOUTH. POINTS
12 WERE MADE EARLIER. MANY VENTURE CAPITALISTS ARE
13 SITTING ON THE SIDELINES. JOHN MADE THE POINT ABOUT
14 INTELLECTUAL PROPERTY. IT'S REALLY EXTREMELY
15 COMPLICATED. THERE'S ABOUT 1500 PATENTS THAT HAVE BEEN
16 ISSUED WITH THE WORD "INFILL" IN THE CLAIM. AND
17 THERE'S ABOUT ANOTHER 2,500 PATENTS THAT HAVE BEEN
18 PUBLISHED AND ARE WORKING THEIR WAY -- SORRY -- PATENT
19 APPLICATIONS THAT HAVE BEEN PUBLISHED AND ARE WORKING
20 THEIR WAY THROUGH THE USPTO. SO IT'S AN EXTREMELY
21 COMPLICATED AREA.

22 THE TIMELINES FOR CELL PRODUCTS ARE VERY
23 UNCERTAIN, AND (INAUDIBLE) BE FOR A WAY WHEN WE'VE
24 PRODUCED TIMELINES. BUT THE FACT IS THIS IS A
25 FIRST-IN-CLASS DEVELOPMENT OF NEW PRODUCTS, AND NOBODY

1 KNOWS THE (INAUDIBLE). AND WE ALL KNOW ABOUT THE
2 POLITICAL/ETHICAL TENSIONS THAT FACE THIS AREA,
3 PARTICULARLY WITH HUMAN EMBRYONIC STEM CELLS.

4 SO TO MARY'S SPECIFIC QUESTIONS THAT SHE
5 POSED FOR ME. ONE IS THE VALUE OF RESEARCH GRANTS TO
6 COMPANIES. WELL, THE FIRST ONE REALLY IS THAT IT
7 DECREASES OUR INVESTORS' RISK A LITTLE BIT. WE SEE
8 THIS AS A PARTNERSHIP WHERE WE TRY TO DEVELOP CURES FOR
9 DISEASES AND WHERE ALL PARTIES SHOULD BENEFIT. AS I
10 SAID, A LOT OF INVESTORS ARE SITTING ON THE SIDELINE,
11 AND THAT'S BECAUSE OVER THE LAST 10 OR 15 YEARS, WE'VE
12 SEEN TECHNOLOGY SUCH AS ANTI-SENSE, CANCER VACCINES,
13 GENE THERAPIES, GENOMICS, XENOTRANSPLANTATION IS
14 ANOTHER ONE. WE'VE SEEN A LOT OF MONEY SPENT, AND SO
15 FAR WE HAVEN'T SEEN ANY PRODUCTS.

16 BUT WE CAN ALSO LOOK AT TECHNOLOGIES LIKE THE
17 DEVELOPMENT OF MONOCLONAL ANTIBODIES, WHICH FOR A LONG
18 TIME DIDN'T WORK, A LOT OF MONEY WAS SUNK INTO IT, AND
19 THEN WHEN PEOPLE REALIZED THAT WITH MONOCLONAL
20 ANTIBODIES, YOU HAD SOMETHING THAT WAS VALUABLE. AND
21 THAT SECTOR OF THE INDUSTRY IS GROWING AT A VERY RAPID
22 RATE AT THE MOMENT. OBVIOUSLY ANY INVESTMENT IN
23 RESEARCH IS NONDILUTIVE FOR THE COMPANY. AND IT CAN
24 POTENTIALLY DECREASE TIMELINES BECAUSE, AGAIN, JOHN
25 POINTED OUT THAT THE BASIC INVESTMENT IN THIS AREA OVER

1 THE LAST YEAR WAS MINIMAL, AND SO WE CAN'T DO ALL THE
2 THINGS WE'D LIKE TO DO. WE CAN'T PARALLEL TRACK ALL
3 THE THINGS WE'D LIKE TO DO. AND I THINK HAVING
4 ADDITIONAL INVESTMENT THROUGH THE INSTITUTE WOULD MAKE
5 THOSE THINGS POSSIBLE.

6 IT WILL ALLOW GREATER INNOVATION. DAVID MADE
7 THE POINT THERE'S A LOT OF INNOVATION GOES ON IN
8 UNIVERSITIES AND RESEARCH INSTITUTIONS, BUT INNOVATION
9 ALSO GOES ON IN COMPANIES. AND I THINK IN TERMS OF THE
10 SORT OF RESEARCH COMPONENT OF IT, SOME OF OUR NOVOCELL
11 WORK IS THE ETHICAL PART. ONE OF MY COLLEAGUES
12 PUBLISHED A PRESTIGIOUS PAPER IN *NATURE BIOTECHNOLOGY*
13 AT THE END OF LAST YEAR. SO WE DO DO GOOD RESEARCH IN
14 COMPANIES TOO. IT'S NOT ONLY THE TRANSLATIONAL
15 RESEARCH.

16 THE LAST POINT IS THAT YOU CAN PROVIDE PEER
17 REVIEWED EVALUATION. SO OUR INVESTORS, TO A LARGE
18 EXTENT, DON'T HAVE ACCESS TO THE SORT OF SOPHISTICATED
19 PEER REVIEW EVALUATION THAT THE INSTITUTE WILL HAVE
20 ACCESS TO. AND I THINK IT WOULD GIVE OUR INVESTORS
21 SOME CONFIDENCE IF THERE WAS A THIRD-PARTY EXTERNAL
22 REVIEW THAT LOOKED AT THE SCIENCE IN A VERY HARD WAY.

23 SECOND QUESTION MARY ASKED US ABOUT OUTSIDE
24 GRANT EXPERIENCE THAT CALIFORNIA HAS HAD. I'M USING
25 TODAY THE EXAMPLE OF JDRF BECAUSE WE HAVE HAD A COUPLE

1 OF GRANTS FROM JDRF AND WE JUST OUTLINED TO YOU WHAT WE
2 FUND WITH THEM. WE'VE ALSO HAD NIH GRANTS THROUGH SBIR
3 AND THROUGH INFRASTRUCTURE GRANTS, BUT I THINK THEY'RE
4 LESS RELEVANT TO WHAT WE'RE TALKING ABOUT TODAY.

5 AND LEVEL WITH THIS IS ABSOLUTELY APPLICABLE
6 TO IP, BUT I JUST WANTED TO RUN THROUGH THE MAJOR
7 POINTS OF A GRANT THAT WE HAVE AT THE MOMENT. FIRST OF
8 ALL, THERE'S A STEERING COMMITTEE, SO IT'S A
9 PARTNERSHIP BETWEEN JDRF AND NOVOCELL, TWO FROM
10 NOVOCELL AND TWO FROM JDRF. WE HAVE TO PROVIDE
11 PROGRESS REPORTS, BOTH TECHNICAL AND FINANCIAL, AND
12 THEY'RE SIX MONTHLY. WE HAVE TO PROVIDE AT LEAST THE
13 TECHNICAL REPORTS FOR UP TO FIVE YEARS AFTER THE
14 COMPLETION OF THE GRANT. WE HAVE TO MAKE REASONABLE
15 EFFORTS TO EITHER PUBLISH OR DISSEMINATE THE RESULTS,
16 WHICH MEANS TALK ABOUT THEM AT CONFERENCES OR WHATEVER,
17 WE DON'T NECESSARILY HAVE TO PUBLISH IT. AND IT IS
18 REASONABLE EFFORT, WHICH IS A DIFFERENT LEGAL MEANING
19 FROM BEST EFFORT. BUT NEVERTHELESS, THERE'S A
20 GOOD-FAITH EXPECTATION.

21 THE INTELLECTUAL PROPERTY, IF THERE IS ANY,
22 THAT'S DEVELOPED IS OWNED BY NOVOCELL. BUT IF WE WANT
23 TO ABANDON ANY INTELLECTUAL PROPERTY DURING
24 PROSECUTION, JDRF HAS A RIGHT OF NEGOTIATION TO THAT
25 INTELLECTUAL PROPERTY. WE HAVE TO REPORT ANY

1 THIRD-PARTY AGREEMENT FOR INTELLECTUAL PROPERTY THAT'S
2 DEVELOPED UNDER THE AGREEMENT OR MIGHT BE COVERED UNDER
3 THE AGREEMENT FOR UP TO TEN YEARS.

4 I GUESS THE MODEL WITH JDRF, AND IT'S TRUE
5 WE'VE HAD A COUPLE OF CONTRACTS AND THEY'RE NEGOTIATED
6 ON AN INDIVIDUAL BASIS, BUT THIS PARTICULAR ONE ASKED
7 FOR REPAYMENT OF THREE TIMES OF THE TOTAL FUNDING, AND
8 IT'S BASED ON PRODUCT SALES IN WHOLE OR IN PART ON
9 RESULTS -- SORRY -- WHERE THE PRODUCT IS IN WHOLE OR IN
10 PART FROM RESULTS OF THE RESEARCH PROGRAM. AND SO IT'S
11 UNLIKELY THAT SOME FUNDED PIECE OF RESEARCH, AND IT
12 COMES BACK TO THE CONTINUUM IDEA THAT I WAS TALKING
13 ABOUT THE OTHER DAY -- I'M SORRY -- A LITTLE EARLIER IN
14 YOUR TALK, IT'S UNLIKELY THAT YOU'RE GOING TO FUND
15 SOMETHING THAT'S GOING TO TAKE, AT LEAST FROM A CONCEPT
16 TO A PRODUCT, BUT IF THERE'S SOME INTELLECTUAL PROPERTY
17 IN THERE, THEN THIS IS THE WAY THAT JDRF PROPOSES TO
18 SHARE IT.

19 OR 5 PERCENT OF THIRD-PARTY PAYMENT. WHAT
20 THAT REALLY COMES DOWN TO, AND JOHN USED A NUMBER OF 25
21 PERCENT, WHICH IN THE INDUSTRY WOULD BE HIGH. I MEAN
22 IT'S REALLY A SUBLICENSING FEE. SO IF WE HAD
23 SUBLICENSING TECHNOLOGY TO A THIRD PARTY, SAY A BIG
24 PHARMA COMPANY, TO COMMERCIALIZE IT, THEN SOME PORTION
25 OF THAT SUBLICENSING FEE CAN GO BACK TO JDRF, GO BACK

1 TO THE INSTITUTION. AND THE DEALS THAT I'VE DONE WITH
2 UNIVERSITIES OR WITH JDRF OR WHATEVER, THIS NUMBER IS
3 NORMALLY SORT OF 5 TO 10 PERCENT.

4 THERE'S NO REPAYMENT IF COMMERCIALIZATION
5 DOESN'T OCCUR. WE ALL KNOW THAT THERE'S HIGH RISK,
6 HIGH REWARD. OUR INVESTORS ARE IN THERE ALONGSIDE
7 THOSE OF JDRF, AND SO IT SEEMS FAIR, A FAIR PRINCIPLE
8 THAT IF NO COMMERCIALIZATION OCCURS, THERE WILL BE NO
9 REPAYMENT.

10 WE CANNOT ASSIGN OR SUBCONTRACT THE GRANT OR
11 INTELLECTUAL PROPERTY WITHOUT PRIOR CONSENT. THAT'S
12 EXCEPT IN THE CASE OF A MERGER OR ACQUISITION.

13 ONE THING THAT MARY ASKED ME ABOUT IS
14 POTENTIAL INTELLECTUAL PROPERTY AND REVENUE SHARING
15 MODELS. AND I PUT UP THREE OR FOUR DIFFERENT MODELS,
16 AND TO ME THE JDRF MODEL IS VERY ACCEPTABLE BECAUSE ON
17 DAY ONE THE COMPANY KNOWS WHAT THEY'RE SIGNING UP FOR.
18 WE'RE GOING TO BE GIVING, IF THERE'S A PRODUCT, THAT
19 ENSURES HERE'S THE RETURN. NOW, IT'S A 3 X RETURN, SO
20 THAT SEEMS REASONABLE. AND I KNOW JDRF NEGOTIATES ON
21 THAT. I'VE SEEN CONTRACTS OF 2 X, AND I'VE SEEN
22 CONTRACTS OF 4 X.

23 THE OTHER THING THAT I THINK IS GOOD IS
24 THERE'S BEEN A LOT OF FOCUS ON INTELLECTUAL PROPERTY,
25 BUT THE MODEL IS NOT BASED ON INTELLECTUAL PROPERTY,

1 BUT PRODUCT SALES. SO DAVID MADE THE POINT, AND IT'S A
2 VERY GOOD ONE, A LOT OF THE THINGS THAT COMPANIES WILL
3 DO IS TRANSLATING BASIC RESEARCH INTO A PRODUCT. I'LL
4 MAKE THIS UNDER MY NEXT POINT, THE ROYALTY MODEL, WHERE
5 I'LL TELL YOU IT MAY WORK IF IT'S LIMITED TO NEW IP,
6 BUT MY VISION OF WHAT CIRM WOULD LIKELY FUND IN A
7 COMPANY SCENARIO, I WOULD LOOK AT ACTIVITIES SUCH AS
8 CLINICAL TRIALS, SUCH AS SCALE-UP THAT DAVID MENTIONED,
9 SUCH AS MANUFACTURING PROCESS TOGETHER. AND MOST
10 IMPORTANT, NECESSARILY LEADS TO INTELLECTUAL PROPERTY.
11 SO A ROYALTY MODEL BASED ON INTELLECTUAL PROPERTY, I
12 THINK, MIGHT HAVE THE INSTITUTE MISSING OUT ON REVENUES
13 THAT IT WOULDN'T OTHERWISE.

14 THE OTHER THING WITH A ROYALTY MODEL IS THAT
15 THIS IS A VERY COMPLEX FIELD, AND WHOEVER
16 COMMERCIALIZES A PRODUCT IS GOING TO REQUIRE MULTIPLE
17 LICENSES. YOU ARE GOING TO WANT TO HAVE ANTISTACKING
18 PROVISIONS IN THOSE LICENSES, AND THESE ROYALTIES GET
19 OUT OF HAND VERY, VERY QUICKLY. WE KNOW BECAUSE WE'RE
20 DEALING WITH A FEW PARTNERS AT THE MOMENT, AND WE KNOW
21 THAT ADDING ROYALTY AFTER ROYALTY IS VERY UNATTRACTIVE
22 TO FUTURE PARTNERS AS OPPOSED TO A MODEL WHERE THEY
23 KNOW WHAT THEY'RE SIGNING UP FOR.

24 OTHER POTENTIAL MODELS, WE TALKED ABOUT
25 LOANS, AND I THINK A LOAN COULD WORK FINE, AN INTEREST

1 BEARING LOAN, WHICH IS FORGIVABLE IF NO PRODUCT WITHIN
2 X YEARS. SO UNLIKE JOHN'S MODEL, THIS IS NOT LIKE
3 BUILDING A HOUSE BECAUSE THERE'S NOT A SURETY THAT
4 WE'RE GOING TO HAVE A PRODUCT AT THE END OF THE DAY.
5 OUR INVESTMENT SCHEME AND MONEY IS ON THE LINE. CIRM'S
6 MONEY IS ON THE LINE. IT BEING FAIR THAT THAT WOULD BE
7 FORGIVABLE IF NO PRODUCT WAS DEVELOPED.

8 IT CAN BE MESSY ACCOUNTINGWISE. IT WOULD SIT
9 ON OUR BALANCE SHEET, AND SO I KNOW THAT OUR FINANCIAL
10 FOLKS WOULDN'T LIKE THAT; BUT NEVERTHELESS, IT'S A
11 MODEL THAT COULD WORK.

12 AND I DON'T KNOW WHAT THE RULES ARE FOR CIRM
13 IN TERMS OF HOLDING EQUITY IN COMPANIES, BUT ANOTHER
14 THING THAT I THOUGHT OF, AND THIS IS JUST IDEAS I'M
15 THROWING OUT THERE, IS CONVERTIBLE DEBT. AND SO IT'S
16 BASICALLY A LOAN AND IT BEARS INTEREST, BUT IT CAN BE
17 CONVERTED AT THE OPTION OF THE COMPANY OR YOU COULD
18 ARGUE AT THE OPTION OF CIRM. AND, OF COURSE, THE
19 COMPANY WOULD RATHER AT THE OPTION OF THE COMPANY AT
20 SOME POINT IN TIME OR WHEN SOME EVENT HAPPENS. THIS IS
21 A MODEL THAT COULD WORK.

22 GETTING BACK TO THE BENEFITS TO CALIFORNIA,
23 THE LAST MODEL I THOUGHT ABOUT WAS A SORT OF AN
24 EXCLUSIVE MARKETING TOOL. PATIENTS CAN COME FROM ALL
25 OVER THE WORLD, AND THEY DO GO TO VARIOUS PLACES FOR

1 VARIOUS TREATMENTS. SO WHAT ABOUT IF FOR A PERIOD OF
2 TIME, YOUR PRODUCTS WERE OFFERED ONLY IN CALIFORNIA SO
3 THAT THE BUSINESS IN CALIFORNIA IS GROWING. AND THAT
4 STIMULATES THE STATE ECONOMY. IT'S AN INDIRECT WAY OF
5 GETTING MONEY BACK INTO THE STATE. BUT, AGAIN,
6 PARTNERS WON'T LIKE THAT IF YOU'RE GOING WITH A BIG
7 BIOTECH COMPANY THAT WORKS INTERNATIONALLY, BUT IT'S
8 SOMETHING THAT I, AT LEAST, THINK YOU SHOULD DO.

9 I THINK THIS HAS ALREADY BEEN ADDRESSED BY
10 ED. CAN CIRM MAKE UNIFORM POLICIES FOR COMMERCIAL
11 ENTITIES, UNIFORM POLICIES? I THINK IT'S DIFFICULT,
12 BUT BASICALLY (INAUDIBLE) JDRF MODEL WHERE THEY'RE
13 INDIVIDUALLY NEGOTIATED. WHILE THEY'RE MUCH MORE
14 COMPLICATED FOR CIRM, I THINK IT'S THE ONLY WAY TO DO
15 IT. IT'S NOT REALLY POSSIBLE TO DO IT ANY OTHER WAY.

16 I'LL JUST PUT UP A COUPLE OF POINTS FOR
17 COMPANIES AT VARIOUS STAGES. I WOULD NOTE WHAT SORT OF
18 A DEAL YOU CAN CUT WITH A COMPANY WILL -- A VIRTUAL
19 COMPANY, YOU'VE PROBABLY GOT A LOT MORE LEVERAGE OVER
20 THAN A PRIVATE COMPANY WITH VC INVESTORS. AND, OF
21 COURSE, A PUBLIC COMPANY HAS ITS SHAREHOLDERS TO DEAL
22 WITH. AND SO ALL THESE GROUPS ARE VERY DIFFERENT WHEN
23 YOU LOOK AT THEM. AND THERE ARE STEM CELL COMPANIES
24 THAT FIT INTO ALL THESE CATEGORIES. AND THEN, OF
25 COURSE, THERE ARE BUSINESSES. WE WOULD SEE OURSELVES

1 AS A THERAPEUTICS BUSINESS. OUR INVITROGEN FRIENDS
2 WOULD SEE THEMSELVES AS BEING IN THE REAGENTS BUSINESS,
3 AND, OF COURSE, ONE CAN CONCLUDE DIAGNOSTICS IS A
4 BUSINESS. AND THE SORTS OF RETURNS ON THOSE BUSINESSES
5 ARE DIFFERENT, AND SO ARE THE SORT OF DEALS YOU HAVE
6 ARE DIFFERENT TOO.

7 COLLABORATIVE CONSORTIUMS ARE SORT OF LIKE
8 THE LAST QUESTION, AND I REALLY STRUGGLED WITH THIS.
9 COLLABORATIVE CONSORTIUMS BETWEEN PUBLIC COMPANIES ARE
10 BASICALLY DRIVEN BY TRYING TO INCREASE SHAREHOLDER
11 VALUES. IT IS VERY DIFFICULT. AND SO I THINK IF SUCH
12 PROGRAMS ARE SET UP, THEY NEED TO BE DRIVEN BY
13 COMMERCIAL CONSIDERATIONS. COMPANIES ARE ONLY GOING TO
14 WORK TOGETHER WHEN THEY BOTH HAVE SOMETHING TO GAIN
15 FROM IT. AND I THINK CORPORATE CULTURE IS VERY
16 IMPORTANT.

17 WE MET WITH A COMPANY RECENTLY, AND WHETHER
18 THEIR TECHNOLOGY GOOD OR BAD, ALAN IS LAUGHING BECAUSE
19 HE WAS THERE WITH ME, I CAN TELL YOU AFTER AN HOUR WITH
20 THIS GUY, THERE'S JUST NO WAY WE CAN EVER WORK WITH
21 THEM.

22 ANOTHER IMPORTANT POINT IS THAT COMPANIES
23 NEED TO BE ABLE TO MAINTAIN CONTROL OF CONFIDENTIAL
24 INFORMATION AND INTELLECTUAL PROPERTY. AND I'VE HEARD
25 A LOT OF TALK TODAY ABOUT THE NEED TO SHARE

1 CONFIDENTIAL INFORMATION AND INTELLECTUAL PROPERTY, AND
2 MAYBE IN THE PUBLIC SECTOR THAT CAN WORK, BUT IN THE
3 PRIVATE SECTOR IT DOESN'T WORK. I SUSPECT EVEN IN THE
4 PUBLIC SECTOR WHERE PEOPLE ARE APPLYING FOR NIH GRANTS
5 AND THEY NEED LEVERAGE, IT'S VERY DIFFICULT TO MAKE
6 THIS WORK. SO I DO NOT THINK CONSORTIUMS SHOULD BE
7 SOME SORT OF A REQUIREMENT OF CIRM, BUT IT COULD HAVE
8 SPECIFIC GRANTS, MUCH LIKE THE NIH HAS CONSORTIUMS LIKE
9 GRANTS, THEIR PROGRAM GRANTS, AND I THINK IT'S MORE
10 LIKELY TO BE ACADEMIC-ACADEMIC OR COMPANY TO ACADEMIC
11 COLLABORATION THAN IT IS COMPANY TO COMPANY.

12 I'LL JUST FINISH WITH THINGS THAT ARE
13 SLIGHTLY OFF TOPIC, BUT OCCUR TO ME TO BE IMPORTANT
14 THINGS TO CONSIDER. ONE IS HOW WILL CIRM PROTECT
15 CONFIDENTIAL INFORMATION DIVULGED IN GRANT
16 APPLICATIONS? AND HERE I THINK THE NIH MODEL PROBABLY
17 DOESN'T WORK. COMPANIES GET VERY PROTECTIVE, A LITTLE
18 BIT PARANOID ABOUT PROTECTING THEIR INTELLECTUAL
19 PROPERTY AND THINGS THAT THEY KNOW. AND SO THE NIH
20 MODEL WHERE YOU HAVE TO SUBMIT A GRANT IN GREAT DETAIL
21 AND GIVE ALL THESE BACKGROUND RESEARCH RESULTS, AND I
22 KNOW YOU WILL HAVE YOUR REVIEWERS SIGN CONFIDENTIALITY
23 AGREEMENTS, BUT A WELL-KNOWN FRIEND OF MINE ONCE TOLD
24 ME THE SECRET IS SOMETHING THAT TWO PEOPLE KNOW, AND
25 ONE OF THEM IS DEAD.

1 IT'S AMAZING BECAUSE WHEN ED SUBMITTED HIS
2 PAPER TO *NATURE BIOTECHNOLOGY*, LITERALLY I DIDN'T EVEN
3 KNOW HE HAD SUBMITTED IT, BUT A FRIEND CALLED ME FROM
4 SINGAPORE WHO HAD BEEN TOLD BY ONE OF THE REVIEWERS
5 THAT THIS PAPER HAS BEEN SUBMITTED. THAT'S HOW I FOUND
6 OUT ED HAD SUBMITTED THE PAPER, AND SO I'VE NO FAITH IN
7 THE CONFIDENTIALITY AGREEMENT.

8 SO I THINK IF YOU CAN PRESENT A GRANT WHERE
9 YOU'RE PRESENTING RESULTS, BUT YOU'RE NOT GIVING
10 DETAILS ABOUT HOW YOU MAY HAVE GOTTEN TO A PARTICULAR
11 POINT, OF COURSE, YOU HAVE TO PROVE WHAT YOU'VE GOT IS
12 WHAT YOU'VE GOT. THAT'S A BETTER MODEL THAN WHERE YOU
13 HAVE TO HAVE GREAT DETAIL.

14 THE OTHER THING IS MOST NIH GRANTS ARE
15 HYPOTHESIS DRIVEN. AND WHEN WE'RE TALKING ABOUT
16 COMPANIES DOING TRANSLATIONAL RESEARCH, CLINICAL
17 RESEARCH, DEVELOPING MANUFACTURING PROCESSES DEVELOPED
18 IN DARPA, THIS IS REALLY HYPOTHESIS DRIVEN. AND SO I
19 THINK THAT NEEDS TO BE TAKEN INTO ACCOUNT.

20 THE OTHER THING IS WHEN I WENT BACK AND READ
21 THE PROP 71 LEGISLATION, THE PRIORITY FOR STEM CELL
22 RESEARCH, UNLIKELY TO RECEIVE FEDERAL FUNDING, TO ME
23 THAT MEANS THERE'S REALLY GOING TO BE A FOCUS ON HUMAN
24 ES CELL RESEARCH, ON THERAPEUTIC CLONING, AND THINGS OF
25 THAT NATURE, AND I'M JUST WONDERING HOW THAT WILL BE

1 IMMERSSED WITHIN THE ORGANIZATION. I REALIZE THAT'S NOT
2 PURELY AN IP TASK FORCE ISSUE, BUT IT IS AN ISSUE.

3 AND THEN WE'VE ALREADY DISCUSSED THE POINT I
4 RAISED WITH JOHN. HOW WILL CIRM HANDLE ALREADY
5 EXISTING INTELLECTUAL PROPERTY WHICH IS OVERARCHING?
6 AND A LOT WAS SAID ABOUT THE WARF INTELLECTUAL
7 PROPERTY, THAT GERON HAS SOME VERY BROAD PATENTS IN THE
8 AREA, WHICH COULD BE VERY TROUBLESOME, AND THERE ARE
9 OTHER GROUPS. IT'S NOT AS CUT AND DRY AS (INAUDIBLE)
10 HAVING COMPETITION OF (INAUDIBLE). THERE ARE OTHER
11 PATENTS OUT THERE THAT ARE VERY BROAD AND NEED TO BE
12 CONSIDERED. AND I THINK IF YOU'RE GOING TO SPEND A LOT
13 OF MONEY TRYING TO DEVELOP PRODUCT, YOU NEED TO KNOW
14 HOW YOU ARE GOING TO HANDLE IT. THAT'S THE END OF MY
15 TALK. THANK YOU.

16 CHAIRMAN PENHOET: THANK YOU VERY MUCH. VERY
17 THOUGHTFUL PRESENTATION. DO WE HAVE QUESTIONS FROM OUR
18 TASK FORCE WITH US HERE IN SAN DIEGO? IN IRVINE?

19 DR. STEWARD: NO QUESTIONS.

20 CHAIRMAN PENHOET: SAN CARLOS?

21 UNIDENTIFIED SPEAKER: NO QUESTIONS.

22 CHAIRMAN PENHOET: LOS ANGELES?

23 UNIDENTIFIED SPEAKER: NO.

24 CHAIRMAN PENHOET: CHICO?

25 DR. WRIGHT: NO.

1 CHAIRMAN PENHOET: UNUSUALLY QUIET GROUP OUT
2 THERE THIS MORNING. HOW ABOUT FROM THE AUDIENCE IN SAN
3 DIEGO? JOHN SIMPSON.

4 MR. SIMPSON: THAT WAS A FANTASTIC
5 PRESENTATION, VERY INTERESTING. I HAVE A QUESTION. I
6 SENSE THAT YOU, ALTHOUGH YOU MIGHT NOT COME DOWN AT THE
7 SAME SPECIFICS, BUT YOU DO BUY INTO THE NOTION THAT IF
8 THERE IS PUBLIC MONEY GOING INTO THIS, THAT THERE IS
9 SOME SORT OF REQUIREMENT FOR PUBLIC BENEFIT TO COME
10 BACK.

11 MR. ROBINS: THE LEGISLATION APPROVED THAT,
12 SO THAT'S VERY CLEAR TO US. WE'RE NOT GOING TO CHANGE
13 THE LEGISLATION, SO I AGREE.

14 MR. SIMPSON: THAT'S DELIGHTFUL TO HEAR. AND
15 THEN THE SECOND QUESTION, I GUESS, WOULD BE YOU
16 ALLUDED --

17 MR. ROBINS: I DON'T SAY I AGREE. OUR
18 COMPANY AGREES.

19 MR. SIMPSON: YEAH. I UNDERSTAND. THE
20 NOTION OF THE POOLING OF PATENTS, DO YOU SEE A
21 POTENTIAL SOLUTION FOR THAT AND SOME KIND OF PATENT
22 POOL ON THE UPSTREAM RESEARCH PROCESS? WOULD THAT BE
23 USEFUL IF YOU COULD GO TO ONE PLACE AND ONE-STOP
24 SHOPPING, SO TO SPEAK?

25 MR. ROBINS: IT'S A BIT LIKE SOCIALISM OR

1 COMMUNISM. THEY HAVE GREAT IDEALS, BUT THE DEVIL IS IN
2 THE DETAIL OF HOW YOU DO THAT. I THINK IN A COMMERCIAL
3 WORLD, WHICH IS, YOU KNOW, LET'S FACE IT AND GET DOWN
4 TO THE (INAUDIBLE). IT'S VERY DIFFICULT TO DO THAT.

5 CHAIRMAN PENHOET: ANY COMMENTS FROM THE
6 PUBLIC IN ANY OF THE OTHER SITES? IF NOT, WE'VE NOW
7 HAD THREE STIMULATING PRESENTATIONS. MAYBE WE'LL TAKE
8 A TEN-MINUTE BREAK, BIOBREAK OR WHATEVER OTHER KIND OF
9 BREAK. THERE'S FOOD OUT HERE ON THE VERANDA IN SAN
10 DIEGO. WE'LL RECONVENE AT 10:15.

11 (A RECESS WAS TAKEN.)

12 MS. KING: SUSAN BRYANT. MICHAEL GOLDBERG.
13 SHERRY LANSING. IS EVERYBODY'S PHONE ON MUTE POSSIBLY?
14 TED LOVE.

15 DR. LOVE: YES, WE'RE HERE.

16 MS. KING: ED PENHOET.

17 CHAIRMAN PENHOET: HERE.

18 MS. KING: PHIL PIZZO. FRANCISCO PRIETO.
19 JOHN REED.

20 DR. REED: HERE.

21 MS. KING: JEFF SHEEHY. OSWALD STEWARD.

22 DR. STEWARD: HERE.

23 MS. KING: JANET WRIGHT.

24 DR. WRIGHT: HERE.

25 MS. KING: I HEARD SOMEONE RING IN, SO I'M

1 GOING TO GO BACK THROUGH THE FIRST FEW NAMES AGAIN.
2 SUSAN BRYANT.

3 DR. BRYANT: HERE.

4 MS. KING: MICHAEL GOLDBERG.

5 MR. GOLDBERG: HERE.

6 MS. KING: SHERRY LANSING.

7 CHAIRMAN PENHOET: THERE'S REPRESENTATION.
8 THANK YOU FOR HANGING IN WITH US.

9 NEXT SPEAKER IS JULIE MEIER WRIGHT. JULIE IS
10 CURRENTLY THE SAN DIEGO REGIONAL ECONOMIC DEVELOPMENT
11 CORPORATION'S CEO. IN A PAST LIFE JULIE WAS SECRETARY
12 OF TRADE AND COMMERCE IN THE STATE OF CALIFORNIA AND
13 DID A LOT TO RAISE THE VISIBILITY OF THE BIOTECH
14 INDUSTRY IN THE STATE GOVERNMENT AND TO PROMOTE
15 CALIFORNIA'S BIOTECH INDUSTRY BROADLY. SO, JULIE,
16 THANK YOU VERY MUCH FOR JOINING US TODAY.

17 AND WE ASKED JULIE TO GIVE US A PERSPECTIVE
18 ON THE IMPORTANCE OF CALIFORNIA'S BIOTECH INDUSTRY TO
19 THE ECONOMY OF THE STATE AND PARTICULARLY OBVIOUSLY SAN
20 DIEGO. SO IT'S THE COMPETITIVE POSITION THAT JULIE IS
21 GOING TO ADDRESS HERE THIS MORNING.

22 MS. WRIGHT: EXACTLY. THANK YOU VERY MUCH,
23 ED. IT'S A PLEASURE TO BE HERE THIS MORNING. I THINK
24 YOU WILL FIND THAT MANY -- FOR THOSE OF YOU AT REMOTE
25 SITES, I HAVE A BRIEF POWERPOINT PRESENTATION THAT IS

1 KIND OF IN THE MIDDLE, SO I'LL TELL YOU WHEN I'M GOING
2 TO CUE IT UP.

3 I WANTED TO TAKE A SLIGHTLY DIFFERENT TACK
4 AND TALK ABOUT HOW -- WHAT ARE THE ECONOMIC DEVELOPMENT
5 IMPLICATIONS. AND ALTHOUGH I DIDN'T TALK TO
6 DR. GOLLAHER AT ALL ABOUT HIS PRESENTATION, I THINK YOU
7 WILL FIND THAT THEY ARE VERY COMPLEMENTARY. I AM
8 SHARING MY PERSPECTIVE OF BEING SECRETARY OF TRADE AND
9 COMMERCE FROM '91 THROUGH '97 AND ALSO CHAIR OF THE
10 GOVERNOR'S ADVISORY COUNCIL ON BIOTECHNOLOGY AS WELL AS
11 ON THE NATIONAL STRENGTH COMMITTEE FOR THE STUDY OF
12 REGIONAL CLUSTERS AND INNOVATION THAT DR. GOLLAHER
13 REFERRED TO.

14 AND IN THE LAST FEW YEARS, OUR MESSAGES ABOUT
15 STATE ECONOMIC DEVELOPMENT HAVE BEEN PRETTY NEGATIVE.
16 WE'RE VIEWING WHAT THE GAIN ISSUE HAS BEEN SIMILAR, I
17 THINK. BUT OUR MESSAGES ABOUT ECONOMIC DEVELOPMENT IN
18 CALIFORNIA HAVE BEEN PRETTY NEGATIVE. SOME KEY TAX
19 CREDITS HAVE EXPIRED, MANY OF WHICH BENEFIT THE LIFE
20 SCIENCES INDUSTRY. AND GOVERNOR GRAY DAVIS DISMANTLED
21 MY AGENCY IN 2003. SO I THINK THAT IN STARK CONTRAST,
22 THE PASSAGE OF PROP 71 SENT ONE OF THE MOST POWERFUL
23 MESSAGES THAT CALIFORNIA HAS EVER SENT ABOUT BEING AT
24 THE FOREFRONT OF INNOVATION AND SPURRING ECONOMIC
25 DEVELOPMENT.

1 SO PREPARING FOR TODAY, I TALKED TO A NUMBER
2 OF PEOPLE, INCLUDING IVOR ROYSTON WHO WAS INVITED TO
3 PRESENT TODAY, BUT IS IN CHINA. HE'S ONE OF SAN
4 DIEGO'S BIOTECH PIONEERS AND AN ACTIVE LIFE SCIENCE
5 INVESTOR. ALSO LAST WEEK MY ORGANIZATION POSTED --

6 (INTERRUPTION IN PROCEEDINGS.)

7 DR. WRIGHT: ANYWAY, I DO WANT TO EMPHASIZE
8 THAT IT HAS BEEN IN STARK CONTRAST THAT THE PASSAGE OF
9 PROP 71 SENT A HUGELY POWERFUL MESSAGE, MAYBE ONE OF
10 THE MOST IMPORTANT THAT CALIFORNIA HAS EVER SENT, ABOUT
11 BEING AT THE FOREFRONT OF INNOVATION AND SPURRING
12 ECONOMIC DEVELOPMENT.

13 AND IN TALKING TO IVOR ROYSTON AND HOSTING
14 THIS SAN DIEGO CONFERENCE ON COMPETITIVENESS, I HAVE A
15 LOT OF FOOD FOR THOUGHT JUST IN THE LAST FEW DAYS. SO
16 I WANT TO TALK A LITTLE BIT ABOUT THE COMPETITIVE
17 CLIMATE IN WHICH INNOVATION IS TAKING PLACE BECAUSE IT
18 IS REALLY INTENSIFYING. THE PASSAGE OF PROP 71 SPURRED
19 MANY OTHER STATES TO ACTION. AND WHILE THEIR EFFORTS
20 ARE IN SOME CASES NOT AS BOLD AS CALIFORNIA'S, NEITHER
21 ARE THEY TIED UP IN LITIGATION, WHICH HAS PREVENTED THE
22 STATE FROM TRULY BUYING A COMPETITIVE EDGE. AND I DO
23 WANT TO CONGRATULATE CIRM ON A SOLID WIN IN THE
24 CALIFORNIA SUPERIOR COURT LAST WEEK, BUT THE FACT IS
25 THAT TIME WAS ON OUR SIDE AND THE LAWYERS HAVE

1 INTERFERED.

2 SO I'M GOING TO GIVE YOU A FEW EXAMPLES OF
3 THE COMPETITIVE LANDSCAPE IN THIS COUNTRY ALONE.
4 WISCONSIN, THE STATE, MENTIONED EARLIER, \$350 MILLION
5 LIFE SCIENCES DISCOVERY FUND THAT WILL OUT-TAKE \$35
6 MILLION A YEAR. PENNSYLVANIA IS TALKING ABOUT CREATING
7 A \$500 MILLION JONAS SALK LEGACY FUND FOR BIOSCIENCES
8 FACULTY AND RECRUITMENT AND FACILITIES. THAT ONE KIND
9 OF BUGS ME SINCE WE HAVE THE SALK INSTITUTE RIGHT UP
10 THE ROAD HERE. MISSOURI IS PROPOSING A \$450 MILLION
11 LEWIS AND CLARK DISCOVERY INITIATIVE. OHIO HAS ALREADY
12 ALLOCATED 300 MILLION, 60 PERCENT TO BIOSCIENCE-RELATED
13 INITIATIVES OUT OF 1.6 BILLION. CONNECTICUT, A HUNDRED
14 MILLION OVER TEN YEARS USING SOME TOBACCO SETTLEMENT
15 MONEY. FLORIDA, 30 MILLION LIFETIME GRANT THAT COULD
16 BE EXPANDED.

17 I THINK WHAT WE ALL KNOW ABOUT THEIR
18 AGGRESSIVE EFFORTS TO WIN THE SCRIPPS RESEARCH
19 INSTITUTE AND ARE NOW WORKING ON AN EXPANSION FROM
20 BURNHAM. THESE INSTITUTES WILL MAINTAIN A STRONG
21 PRESENCE IN SAN DIEGO, BUT THESE OPPORTUNITIES ARE TOO
22 ATTRACTIVE FOR THEM TO IGNORE AS THEY SEEK TO EXPAND
23 THEIR RESEARCH BASE.

24 NEW JERSEY, 11 AND A HALF MILLION
25 APPROPRIATED, \$150 MILLION IN CAPITAL SPENDING AND

1 250 -- 30 MILLION, SORRY, FOR RESEARCH PROPOSED.
2 MARYLAND, 12 MILLION, AND ALSO IN THE 5 TO 10 MILLION
3 RANGE, ILLINOIS AND SOUTH CAROLINA. I WAS GOING TO ADD
4 WISCONSIN UNTIL I LEARNED IN AN E-MAIL FROM ALAN LEWIS
5 OF NOVOCELL THAT GOVERNOR DOYLE WAS AT BIO WITH A
6 STRONG PITCH FOR STEM CELL RESEARCH, AND HE HIGHLIGHTED
7 50 MILLION IN DONATIONS BY THE MORGRIDGE FAMILY, JOHN
8 MORGRIDGE, CALIFORNIA CEO OF CISCO. 50 MILLION FROM
9 THE WISCONSIN ALUMNI RESEARCH FOUNDATION THAT'S BEEN
10 TALKED ABOUT THIS MORNING, AND ANOTHER 50 MILLION FROM
11 THE STATE TO CREATE THE WISCONSIN (INAUDIBLE) OF
12 DISCOVERY.

13 TIME DOESN'T PERMIT TO ME TO FOCUS ON THE
14 INTERNATIONAL COMPETITION, BUT I JUST WANT TO USE
15 SINGAPORE AS ONE EXAMPLE OF MANY. I HAVE A COPY OF AN
16 ARTICLE. I APOLOGIZE FOR NOT GETTING IT TO THE REMOTE
17 SITES, BUT I KNOW THAT MELISSA WILL DO SO AS A PDF
18 FILE. IT IS AN ARTICLE FROM *WIRED MAGAZINE* CALLED
19 "SINGAPORE WANTS YOU." AND LET ME QUOTE, "THE
20 FUTURE-FRIENDLY CITY-STATE HAS AN OFFER BIOSCIENTISTS
21 CAN'T REFUSE: UNRESTRICTED RESEARCH, TOPNOTCH TALENT,
22 AND LIMITLESS FUNDS. (JUST LEAVE OF CHEWING GUM AT
23 HOME)." IT ALSO GOES ON TO SAY, "SINGAPORE IS TREATING
24 HUNDREDS OF SCIENTISTS LIKE FREE AGENTS, PROMISING
25 FIRST CLASS LABORATORIES, TOPNOTCH EQUIPMENT, AND MORE

1 THAN ENOUGH MONEY TO PURSUE WORK THAT'S NOT FUNDABLE,
2 OR IS TOO CONTROVERSIAL, BACK HOME. THE GOVERNMENT IS
3 INVESTING MORE THAN \$2 BILLION INTO RESEARCH OF ALL
4 STRIPES HOPING TO ATTRACT LEADERS IN THERAPEUTIC
5 CLONING, DRUG DISCOVERY, CANCER RESEARCH, AND OTHER
6 AREAS. BIOSCIENCE ALL-STARS WHO WILL IN TURN HELP
7 BUILD A LOCAL COMMUNITY THAT WILL BOLSTER THE ECONOMY."

8 I FRANKLY THINK THAT'S WHAT WE OUGHT TO BE
9 TALKING ABOUT DOING IN CALIFORNIA WITH RESPECT TO STEM
10 CELL RESEARCH. OUR OWN ED HOLMES, WHOM YOU ALL KNOW,
11 WHO'S THE DEAN OF UCSD'S MEDICAL SCHOOL, THIS FALL IS
12 GOING TO SPEND 20 WEEKS A YEAR IN SINGAPORE RUNNING
13 THEIR EQUIVALENT TO NIH. AND HIS WIFE IS GOING TO
14 BECOME THE HEAD OF A NEW INSTITUTE FOR TRANSLATIONAL
15 MEDICINE. IN A CONVERSATION WITH ED THIS WEEK, HE
16 SAID, "THEY ARE HARD CHARGING, FOCUSED. WE CANNOT
17 COAST."

18 THE LIFE SCIENCES SECTOR IS UNIQUELY TARGETED
19 BY PEOPLE LIKE ME ALL OVER THE WORLD AND GOVERNMENTS
20 FROM ALL OVER THE WORLD WHO ARE PASSIONATE ABOUT
21 EARNING THEIR SHARE OF THIS EXCITING INDUSTRY.

22 AND NOW IF YOU WANT TO PULL UP THE SLIDES,
23 I'M GOING TO USE A FEW QUICK SLIDES FROM THE LIFE
24 SCIENCE COMMUNITY'S MAJOR INTERNATIONAL CONFERENCE BIO,
25 WHICH THIS YEAR ATTRACTED MORE THAN 20,000 ATTENDEES

1 FROM AROUND THE WORLD. THIS CONFERENCE OF 2006 IN
2 CHICAGO ATTRACTED MORE THAN 20,000 PEOPLE GLOBALLY.
3 AND THE IRONY IS IF YOU WALK THE FLOORS OF BIO, IT'S AS
4 MUCH ABOUT ECONOMIC DEVELOPMENT AS IT IS ABOUT SCIENCE
5 ANYMORE.

6 I'M GOING TO GO THROUGH SOME SLIDES VERY
7 QUICKLY. YOU CAN SEE, CALIFORNIA THROW EVERYTHING WE
8 HAVE THE SEVENTH LARGEST ECONOMY IN THE WORLD, HAS A
9 RELATIVELY MODEST PRESENCE AT BIO. LOOK AT OTHER
10 SMALLER STATES THAT IN MANY CASES HAVE NOT VERY MUCH IN
11 THE LIFE SCIENCES: GEORGIA, ILLINOIS, IOWA. DUANE
12 ROTH WILL TELL YOU THAT THE GOVERNOR OF IOWA OUT AND IS
13 ACTIVE TO RECRUIT COMPANIES FROM CALIFORNIA. MICHIGAN,
14 MISSOURI, NEBRASKA, NEW YORK SITE IS BLURRY. THESE ARE
15 AMATEUR PHOTOGRAPHS OBVIOUSLY.

16 IT'S AN INTERNATIONAL COMPETITION. SOME OF
17 THE INTERNATIONAL COMPETITORS DO MAJOR THINGS. AND
18 INCIDENTALLY, I WANT TO SAY THAT LOOKING AT THESE
19 PHOTOGRAPHS OF EXHIBITS IS BUT ONE COMPONENT OF THEIR
20 OVERALL EFFORT, WHICH INCLUDES INCENTIVES, INVESTMENT
21 IN RESEARCH, AND A VARIETY OF OTHER THINGS.

22 THIS FROM BIO 2003. SINGAPORE IS BACK THERE.
23 THIS BIG UPSIDE DOWN UMBRELLA, I GUESS, IS GERMANY.
24 THAT'S MARYLAND. THERE'S THE GERMAN BOOTH. THESE ARE
25 PAVILIONS, NOT JUST BOOTHS, I SHOULD SAY. WASHINGTON

1 STATE. IRELAND. CONNECTICUT. KENTUCKY. PLACES WE
2 DON'T EVEN THINK ABOUT DAY-TO-DAY. OF COURSE, ISRAEL.
3 THESE GO BACK TO 2001. THESE ARE SUSTAINED PRESENCE.
4 THESE COMPANIES AND COUNTRIES AND GOVERNMENTS AND
5 REGIONS ARE THERE YEAR AFTER YEAR. PHOENIX. BERLIN.
6 THE WALLONA REGION OF BELGIUM. MARYLAND AGAIN. AND
7 THIS IS THE LAST SLIDE. THIS IS AUSTRALIA'S EXHIBIT IN
8 2001.

9 MANY OF THE EFFORTS OF THESE PLACES ARE GOING
10 TO FAIL, BUT THAT'S THE KIND OF PASSION THAT PURSUES
11 BIOTECHNOLOGY AND MORE RECENTLY STEM CELL RESEARCH.
12 SOME OF THESE, HOWEVER, WILL SUCCEED, AND THEY'LL MAKE
13 MAJOR INROADS IN BUILDING AND INDUSTRY. IT'S KIND OF
14 LIKE RESEARCH TRIANGLE PARK IN THE '70S AND FLORIDA'S
15 LIFE SCIENCES EFFORTS TODAY. A LOT OF VISION AND
16 ENOUGH MONEY CAN REALLY TRANSFORM A STATE.

17 THIS IS A GLOBAL COMPETITION. WE CAN'T PUT
18 UP WALLS EVEN IF WE WANTED TO. AND THAT'S WHY I THINK
19 THAT CALIFORNIA CAN'T TAKE ITS CURRENT LEADERSHIP FOR
20 GRANTED. AND IT'S WHY SETTING UP THE BEST FRAMEWORK
21 FOR INTELLECTUAL PROPERTY AND INVESTMENT IN FOR-PROFIT
22 COMPANIES IS SO IMPORTANT.

23 AND I VIEW THE ISSUES AROUND INTELLECTUAL
24 PROPERTY AND ROYALTIES TO HAVE SIMILAR IMPLICATIONS, SO
25 I JUST WANTED TO MAKE A FEW POINTS FROM MY PERSPECTIVE.

1 WHAT YOU ARE TRYING TO DO IS TO STIMULATE
2 CURES, WHICH WHETHER THEY ARE INITIALLY DEVELOPED IN A
3 RESEARCH LAB, A NOT-FOR-PROFIT ENVIRONMENT, OR A YOUNG
4 BIOTECH COMPANY ARE ULTIMATELY GOING TO REQUIRE
5 MILLIONS OF DOLLARS TO PROGRESS THROUGH CLINICAL TRIALS
6 TO THE MARKETPLACE. AND IRONICALLY THIS DISCUSSION IS
7 CENTERING AROUND FOR-PROFIT COMPANIES WHEN, IN FACT,
8 FOR MANY OF THE COMPANIES THAT YOU WILL FUND,
9 PROFITABILITY IS A DISTANT DREAM. IN ACTUALITY IF YOU
10 CAN FOCUS ON THE GAP WHERE CAPITAL ACCESS HAS BEEN
11 DIFFICULT, YOU MAY FILL SOME VOIDS THAT EXIST TODAY.

12 WE'VE SPENT A LOT OF TIME IN SAN DIEGO
13 LOCATING WHERE THOSE GAPS ARE, AND ALTHOUGH I'M NOT
14 GOING TO TALK ABOUT IT, YOUR NEWEST BOARD MEMBER, DUANE
15 ROTH, IS SETTING UP A LIFE SCIENCE ACCELERATOR PROGRAM
16 TO FOCUS ON PROMISING EARLY STAGE PROOF OF CONCEPTS.
17 AND I REALLY ENCOURAGE YOU TO HEAR FROM HIM. IT'S A
18 VERY INTERESTING BUSINESS MODEL.

19 \$3 BILLION IN CALIFORNIA SOUNDS LIKE A LOT OF
20 MONEY, BUT IT WILL BE SPREAD OVER TEN YEARS AND OVER A
21 BROAD ARRAY OF INVESTMENTS FOR MANY WORTHWHILE
22 ENDEAVORS. SO EVEN IF CIRM MAKES WHAT FOR IT WOULD BE
23 A SIGNIFICANT INVESTMENT IN A FOR-PROFIT COMPANY
24 DEVELOPING NEW THERAPIES, THIS INVESTMENT WILL LIKELY
25 BE A SMALL PERCENTAGE OF THE TOTAL INVESTMENT REQUIRED

1 TO BRING THESE THERAPIES TO COMMERCIAL SUCCESS.

2 SO IF YOU ASSUMED A \$10 MILLION INVESTMENT,
3 AND I DON'T KNOW WHAT YOUR THRESHOLD FOR INVESTMENT IS
4 GOING TO BE, BUT A \$10 MILLION INVESTMENT IN A YOUNG
5 COMPANY THAT ULTIMATELY REQUIRES \$500 MILLION BEFORE
6 COMMERCIAL SUCCESS, CIRM'S INVESTMENT IS 2 PERCENT. SO
7 MY QUESTION IS HOW DOES A 2-PERCENT INVESTMENT WARRANT
8 ONEROUS ROYALTIES OR IP REQUIREMENTS?

9 I THINK IT IS INSTRUCTIVE TO LOOK AT THE
10 FEDERAL GOVERNMENT AND PROGRAMS LIKE DARPA THAT
11 DR. GOLLAHER MENTIONED AND ALSO THE SCIR PROGRAM, BUT
12 PRETTY TRADITIONALLY THE FEDERAL GOVERNMENT DOES NOT
13 INSIST ON ROYALTIES OR INTELLECTUAL PROPERTY OWNERSHIP,
14 THOUGH IT WILL, RIGHTLY IN MY OPINION, USE MARCH-IN
15 RIGHTS TO ENSURE THE BROADEST APPLICATION AND THE BEST
16 OF THE RESEARCH IT FUNDS. AND MARCH-IN RIGHTS,
17 PARTICULARLY IF THEY'RE REASONABLE, WOULDN'T CONSTITUTE
18 A DISINCENTIVE TO PRIVATE INVESTMENT.

19 SO AS YOU LOOK AT VARIOUS MECHANISMS
20 AVAILABLE TO CIRM, LOANS, ALTHOUGH I UNDERSTAND, ALLAN,
21 THE CONCERN ABOUT ACCOUNTING COMPLEXITIES, THEY MIGHT
22 BE A MORE ATTRACTIVE VEHICLE BECAUSE FAVORABLE INTEREST
23 RATES WITH REPAYMENT THAT DOES NOT OCCUR UNTIL THE
24 PRODUCT IS COMMERCIALY VIABLE, AND I THINK ALSO THAT
25 THE LOAN IS FORGIVEN IF THE RESEARCH GOES SIDEWAYS, IT

1 MIGHT BE A CLEAN WAY TO GO. YOU COULD AVOID SOME OF
2 THE QUESTIONS ABOUT A RETURN ON INVESTMENT WITH A
3 HEALTHY REPAYMENT SCHEME BACK TO THE CIRM WHILE
4 POTENTIALLY PROVIDING A SMALL COMPANY NEEDED LEVERAGE.
5 I THINK POLITICALLY, WHICH IS UNFORTUNATELY THE CLIMATE
6 IN WHICH WE ARE OPERATING, IT MIGHT BE A MORE
7 ATTRACTIVE OPTION.

8 THE FRAMEWORK THAT YOU'VE DEVELOPED FOR
9 NOT-FOR-PROFIT INSTITUTIONS WAS THE 25-PERCENT ROYALTY
10 PAYMENT WOULD TRULY BE A DISINCENTIVE ON THE PRIVATE
11 SIDE TO ATTRACTING PRIVATE CAPITAL. AND I THINK MOST
12 PRIVATE INVESTORS WOULD TELL YOU THAT THEY DON'T EXPECT
13 MORE THAN 1- OR 2-PERCENT ROYALTIES ON THERAPEUTICS AND
14 MAYBE 5 PERCENT OR SO ON MEDICAL DEVICES. SO YOU COULD
15 HAVE THE UNINTENDED CONSEQUENCE OF DISCOURAGING PRIVATE
16 SECTOR INVESTMENT WITH SOME THINGS THAT WOULD BE VERY
17 GOOD, AND IT COULD PREVENT CIRM FROM REALLY BEING A
18 CATALYST FOR THE BEST IDEAS AND THE BEST SCIENCE AT A
19 REALLY CRITICAL TIME IN THE LIFE CYCLE OF A FOR-PROFIT
20 COMPANY.

21 SO MY CONCLUSION IS THAT THE STATE MIGHT BE
22 AN ABSOLUTELY VITAL INVESTOR AT A CRITICAL STAGE, BUT
23 IT'S NOT LIKELY TO BE THE MAJOR INVESTOR. SO YOU COULD
24 SAY THAT THE STATE'S INVESTMENT MIGHT BE
25 DISPROPORTIONATELY SIGNIFICANT FOR GETTING TO CIRM'S

1 GOAL, WHICH IS CURES, BUT FINANCIALLY INSIGNIFICANT IN
2 THAT IT WILL ULTIMATELY BE A SMALL PERCENTAGE OF THE
3 TOTAL INVESTMENT, ALBEIT INVESTMENT PROVIDED AT A
4 CRITICAL TIME.

5 I WANT TO GIVE YOU ONE OTHER MEASURE OF
6 LONG-TERM SUCCESS. I DON'T KNOW IF YOU HAVE THE
7 FLEXIBILITY TO FOCUS ON IT, BUT I THINK IT'S A VERY
8 IMPORTANT ONE, AS I LOOK AT CALIFORNIA AND I LOOK AT
9 CLUSTERS.

10 WHAT THE STATE CAN FOCUS ON IS THE REVENUES
11 THAT IT WILL ULTIMATELY GARNER BY REMAINING AT THE
12 FOREFRONT OF INNOVATION, INCLUDING STEM CELL RESEARCH.
13 AND THESE INCLUDE PERSONAL AND CORPORATE INCOME TAXES,
14 PROPERTY TAXES, SALES TAXES. THE LAST THING, REALLY
15 THE LAST THING, RETURN TO THE STATE OF CALIFORNIA FOR
16 TAKING THE BOLD STEPS TO MAINTAIN LEADERSHIP IN THIS
17 VERY IMPORTANT TECHNOLOGY.

18 HALF OF CALIFORNIA'S TAX REVENUES ARE BASED
19 ON PERSONAL INCOME TAXES TODAY, SLIGHTLY LESS THAN
20 THAT. SO THE WEALTH CREATION OF A VIBRANT AND
21 IMPORTANT INDUSTRY IS IMPORTANT. AND I'VE SPOKEN TO
22 GOVERNOR JEB BUSH IN FLORIDA ABOUT THIS, BUT THAT IS
23 BEHIND HIS THINKING IN HIS SINGLE-MINDED FOCUS TO
24 CREATE A STRONG RESEARCH BASE IN FLORIDA SO THAT
25 FLORIDA CAN EMULATE IN MANY WAYS WHAT CALIFORNIA HAS

1 ALREADY DONE.

2 INCIDENTALLY, I SHOULD TELL YOU THAT 95
3 PERCENT OF THE STATE'S GENERAL FUND REVENUES IN 2005
4 WERE ATTRIBUTABLE TO THREE TAXES THAT WEALTH CREATION
5 DRIVES: PERSONAL INCOME TAXES, CORPORATE TAXES, AND
6 SALES AND USE TAXES. SO I TRULY THINK WE NEED TO TAKE
7 THE LONG VIEW. WE NEED TO BEGIN TO THINK ABOUT
8 OURSELVES AS A STATE THAT INVESTS IN INNOVATION BECAUSE
9 INNOVATION HAS ALREADY MEANT SO MUCH TO CALIFORNIA'S
10 ECONOMIC PROSPERITY AND OUR ECONOMIC LEADERSHIP
11 GLOBALLY, AND IT'S AN ABSOLUTELY CRITICAL PART OF OUR
12 FUTURE. SO I REALLY APPRECIATE THE OPPORTUNITY TO
13 SPEAK TO YOU TODAY, NOT ONLY ABOUT THE ISSUES OF
14 INTELLECTUAL PROPERTY, BUT OF THE MUCH LARGER CONTEXT
15 IN WHICH YOUR WORK IS TAKING PLACE. HAPPY TO ANSWER
16 ANY QUESTIONS.

17 CHAIRMAN PENHOET: THANK YOU VERY MUCH.
18 FIRST OF ALL, QUESTIONS FROM OUR MEMBERS IN SAN DIEGO.
19 WE'LL GO THROUGH THE USUAL ROTATION. IRVINE?

20 DR. BRYANT: NO.

21 CHAIRMAN PENHOET: SAN CARLOS.

22 UNIDENTIFIED SPEAKER: NO.

23 MR. GOLDBERG: YES. WE HAVE A SPLIT VOTE. I
24 HAVE A QUESTION FOR THE PRESENTER, PLEASE. HOW WOULD
25 YOU SUGGEST WE DEAL WITH THE ISSUES OF AFFORDABILITY

1 AND ACCESSIBILITY AS PROPOSED BY JOHN SIMPSON?

2 MS. WRIGHT: I GUESS I THINK THAT IF -- THE
3 MOST IMPORTANT THING WE CAN DO IS TO DEVELOP THE KIND
4 OF CURES THAT WILL BENEFIT EVERYONE. AND I THINK AT
5 SOME POINT THE GOVERNMENT HAS A ROLE IN PROVIDING
6 CURES, THERAPEUTICS, AND DIAGNOSTICS TO THE LEAST AMONG
7 US. BUT I HAVEN'T GONE DOWN THAT PATH IN TRYING TO
8 UNDERSTAND EXACTLY HOW THE FRAMEWORK SHOULD BE SET UP
9 FOR FOR-PROFIT COMPANIES, BUT I DO KNOW THAT ANYTHING
10 THAT IS MORE COSTLY THAN THE MODEL THAT IT TAKES TO
11 ENCOURAGE PRIVATE SECTOR INVESTMENT TODAY IS SOMETHING
12 THAT'S NOT GOING TO WORK, AND THEN WE WON'T GET TO THAT
13 DISCUSSION AT ALL.

14 CHAIRMAN PENHOET: LOS ANGELES? CHICO?

15 DR. WRIGHT: NO.

16 CHAIRMAN PENHOET: ANY COMMENTS BY THE
17 PUBLIC? FIRST IN SAN DIEGO? IN IRVINE?

18 DR. BRYANT: NO.

19 CHAIRMAN PENHOET: SAN CARLOS?

20 MR. REED: YES. I WOULD JUST WISH SO MUCH
21 THAT YOU COULD SQUEEZE INTO YOUR BUSY SCHEDULE TO
22 APPEAR AT THE APPROPRIATIONS COMMITTEE WHEN SUCH
23 INFORMATION AS YOU HAVE TO OFFER WOULD BE JUST HUGELY
24 BENEFICIAL. I DO NOT THINK THOSE ARGUMENTS ARE KNOWN
25 BY THE LEGISLATORS AT ALL. THANK YOU.

1 MS. WRIGHT: WELL, I WILL TELL YOU IN MY TIME
2 IN SACRAMENTO, I WAS APPROACHED BY A STATE SENATOR ONCE
3 THAT SAID, "WHY SHOULD THE STATE PAY MONEY IN THE FORM
4 OF A TAX CREDIT FOR SOMETHING COMPANIES MIGHT DO
5 ANYWAY?" ALTHOUGH THE OPERATIVE WORD IS MIGHT. MAYBE
6 MIGHT NOT. BUT THANK YOU. I WILL KEEP TRACK OF THE
7 SCHEDULE, AND IF YOU ALL WILL LET ME KNOW, I'LL TRY TO
8 BE THERE.

9 MR. REED: THANK YOU.

10 CHAIRMAN PENHOET: ANY COMMENTS IN CHICO?

11 DR. WRIGHT: NO.

12 CHAIRMAN PENHOET: OKAY. THANK YOU, JULIE,
13 FOR A VERY THOUGHTFUL PRESENTATION.

14 WE NOW COME TO OUR FINAL SPEAKER, A FREQUENT
15 VISITOR TO OUR MEETINGS. TODAY WE WILL GIVE JOYDEEP
16 GOSWAMI IS A FULL HALF HOUR INSTEAD OF HIS USUAL
17 COMMENTS FROM THE FLOOR. I WILL APPRECIATE YOUR
18 CONTINUING EFFORTS TO INFORM US AND FOR YOUR
19 PRESENTATION. THANK YOU.

20 MR. GOSWAMI: FIRST OF ALL, THANK YOU FOR
21 INVITING ME TO COMMENT IN THIS PRESENTATION AND FORMAT.
22 YOU KNOW, I THINK, FOR US IT'S -- I'M TRYING TO BRING
23 TO CIRM A VIEWPOINT THAT IS DIFFERENT FROM THE PEOPLE
24 CLOSER TO THE PATIENT IN SOME WAYS SUCH AS ALAN AND
25 NOVOCELL. FROM THE TOOLS COMPANIES AND THE KIND OF

1 PERSPECTIVE WE BRING MAY BE SLIGHTLY DIFFERENT AND I
2 THINK HOPEFULLY THIS WILL ENGENDER SOME QUESTIONS.

3 SO QUICK OVERVIEW OF INVITROGEN. WE'RE A
4 PART OF THE RESEARCH TOOLS MARKET. IT'S SUPPOSED TO BE
5 ABOUT \$17 BILLION OR SO. IT'S THE LINCHPIN OF MOST OF
6 LIFE SCIENCE INDUSTRIES; AND, OF COURSE, WITHOUT TOOLS,
7 THE BASIC RESEARCH THAT UNDERLIES A LOT OF THE
8 TREATMENTS THAT YOU SEE, I THINK, WILL NOT BE POSSIBLE.
9 WE'RE DEFINITELY HEADQUARTERED IN CALIFORNIA. OF
10 COURSE, AS YOU SEE THERE, MOST PEOPLE TALK ABOUT ONE
11 THERAPY GETTING TO MARKET. WE'RE A VERY FRACTURED
12 MARKET WITH ABOUT 15,000 DIFFERENT PRODUCTS SOLD
13 PREDOMINANTLY TO THE RESEARCH MARKET. AND, OF COURSE,
14 UNDERLYING MOST OF OUR PRODUCTS ARE PATENTS THAT ALLOW
15 INNOVATIVE SOLUTIONS TO SCIENTISTS' PROBLEMS, AND WE
16 ARE AN ACTIVE LICENSOR OF OR LICENSEE, I SHOULD SAY, OF
17 TECHNOLOGIES FROM UNIVERSITIES, INCLUDING UC, WITH OVER
18 40 LICENSES EXECUTED ANNUALLY THAT -- TECHNOLOGY AND
19 LICENSES EXECUTED ANNUALLY FOR OUR COMPANY.

20 WE ALSO HAVE MADE A MAJOR COMMITMENT TO STEM
21 CELL RESEARCH. WE'VE CREATED A BUSINESS VENTURE IN
22 THIS PARTICULAR AREA. HIRED MAHENDRA RAO FROM THE NIH
23 TO HEAD UP THE STEM CELL RESEARCH GROUP, AND CURRENTLY
24 DEDICATE OVER 20 RESEARCHERS TO THIS FIELD AND, OF
25 COURSE, LEVERAGE THAT WITH OUR OTHER RESEARCH

1 COMMITMENTS THROUGHOUT THE COMPANY OF OVER 600
2 RESEARCHERS. AND AS JEANNE AND OTHERS WILL ATTEST TO,
3 WE HAVE BEEN ABLE AN ACTIVE PARTICIPANT IN THE STEM
4 CELL COMMUNITY ALREADY. WE FUND A LARGE NUMBER OF STEM
5 CELL COURSES WORLDWIDE, NOT ONLY IN CALIFORNIA, OTHER
6 STATES IN THE UNITED STATES, AND, OF COURSE, NOW IN
7 OTHER COUNTRIES AS WELL, INCLUDING INDIA AND HOPEFULLY
8 SINGAPORE, KOREA.

9 MAHENDRA HAS TAUGHT MANY OF THESE COURSES,
10 AND, OF COURSE, OUR PRODUCTS GO INTO MOST OF THE
11 PROTOCOLS THAT ARE TAUGHT AT THESE COURSES.

12 I WANT TO SPEND SOME TIME ACTUALLY GIVING YOU
13 A BIRD'S-EYE VIEW OF HOW WE LOOK AT STEM CELL RESEARCH
14 AND THE BASIC ELEMENTS OF THE WORK FLOW OF STEM CELL
15 RESEARCH. IT WILL COME BACK BECAUSE THESE BASIC FOUR
16 ELEMENTS THAT I HAVE ON PAGE 4 REALLY THEN TRANSLATE
17 ACROSS FROM BENCH TO BEDSIDE WHETHER YOU'RE LOOKING AT
18 BEDSIDE RESEARCH OR BASIC RESEARCH. AND I THINK, IN
19 OUR OPINION, THE SAME FOUR STEPS WILL BE VERY IMPORTANT
20 IN ANY THERAPEUTIC THAT IS AFFORDABLE FOR THE PUBLIC.

21 SO THE FOUR STEPS ARE ESSENTIALLY ISOLATION
22 OF STEM CELLS AND THEIR PROGENY AND DIFFERENTIATED
23 PROGENY, A CHARACTERIZATION OF THESE CELLS, AND THIS IS
24 REALLY IMPORTANT BECAUSE THESE CELLS, I LIKE TO CALL
25 THEM, AS INHERENTLY UNSTABLE BECAUSE THEY WANT TO

1 BECOME OTHER TYPES OF CELLS; AND, THEREFORE, HAVING
2 THEM CHARACTERIZED AS BEING PLURIPOTENT OR HAVING THE
3 ABILITY TO FORM OTHER CELLS IS QUITE IMPORTANT. AND,
4 OF COURSE, ONCE YOU'VE GONE DOWN THE PATH OF
5 DIFFERENTIATION, AGAIN, IT'S IMPORTANT TO CHARACTERIZE
6 THE DIFFERENTIATED CELLS AND MAKE SURE THAT THEY ARE
7 WHAT YOU THINK THEY ARE.

8 EXPANSION OF CELLS, THIS IS ANOTHER KEY PART
9 OF HAVING THERAPIES THAT ARE AFFORDABLE BECAUSE YOU
10 NEED LOTS OF CELLS IN ANY KIND OF CELL THERAPY AND YOU
11 NEED THEM TO BE PRODUCED IN ANIMAL ORIGIN FREE
12 CONDITIONS HOPEFULLY AND, OF COURSE, IN A FASHION THAT
13 PROVIDES THAT EACH VIAL OF CELLS IS NOT A MILLION
14 DOLLARS IN PRICE. AND THEN THE LAST PART IS
15 DIFFERENTIATION. AGAIN, HERE THE CRITICAL CONDITIONS,
16 AGAIN, ARE ANIMAL ORIGIN FREE AND, OF COURSE, THE FACT
17 THAT YOU CAN DO THIS EFFICIENTLY IN AN ECONOMICAL
18 MANNER.

19 GIVEN THESE, HOW WE THINK ABOUT IT IS ALONG
20 TECHNOLOGY PLATFORMS THAT ALIGN ACROSS THESE FOUR
21 ELEMENTS OF THE WORK FLOW. AND THERE ARE SEVERAL.
22 THIS IS BY NO MEANS A COMPLETE LIST. BUT, YOU KNOW, IF
23 YOU LOOK AT THE ISOLATION AND CHARACTERIZATIONS PART OF
24 THINGS, THERE ARE THINGS LIKE PRIMARY ANTIBODIES, THERE
25 ARE THINGS LIKE BEAD-BASED CELL SEPARATION SYSTEMS

1 WHICH ARE ABSOLUTELY CRITICAL IN GETTING AND ISOLATING
2 THE RIGHT KIND OF CELL THAT YOU WANT. IN FACT, SOME OF
3 THESE ARE ALREADY IN THE CLINIC. SOME OF YOU MAY HAVE
4 HEARD BAXTER'S CD34 POSITIVE CELLS THAT ARE BEING USED
5 IN CARDIAC ISCHEMIA ACTUALLY USES MAGNETIC BEADS TO
6 ISOLATE THE CD34 POSITIVE CELLS. AND I'M SURE THIS
7 WILL BE A LINCHPIN OF OTHER THERAPIES THAT ARE COMING
8 TO THE MARKET AS WELL. BUT AGAIN, THE SAME
9 TECHNOLOGIES ARE USED BOTH IN BASIC RESEARCH AND THEN
10 HOPEFULLY WILL TRANSLATE, ARE SCALABLE WHEN YOU MOVE TO
11 THE CLINIC.

12 LABELING AND DETECTION SYSTEMS INCLUDES A LOT
13 OF FLUOROPHORES AND THINGS LIKE QUANTUM DOTS WHICH
14 ALLOW YOU TO GET A QUANTITATIVE ASSESSMENT OF YOUR
15 EXPERIMENTS AND, OF COURSE, DIFFERENT CELL TYPES. THEN
16 THERE ARE, OF COURSE, THE BASIC PROTEIN PURIFICATION
17 ANALYSIS TOOLS, INCLUDING QUANTITATIVE PROTEOMICS. A
18 LOT OF EFFORT ON OUR SIDE IS GOING INTO DNA/RNA-BASED
19 CHARACTERIZATION OF CELLS WHICH INCLUDES BASIC QPCR,
20 BUT ALSO THINGS LIKE MICRO-RNA ANALYSIS, METHYLATION
21 ANALYSIS OF CELLS. AND THEN WHAT WE DO OFFER TO THE
22 COMMUNITY IS SERVICES, AND THESE COULD TAKE MANY FORMS.
23 THESE COULD TAKE GENERATION OF CUSTOM ANTIBODIES. I
24 THINK SOMEONE MENTIONED GROWING A CELL. ALLAN, YOU
25 TALKED ABOUT GNP FACILITIES OR GNP TESTING OF CELLS

1 ETC.

2 AND, OF COURSE, ON THE EXPANSION
3 DIFFERENTIATION SIDE, THERE'S A HOST OF STEM CELL
4 CULTURE MEDIA, INCLUDING ANIMAL ORIGIN FREE, WHERE WE
5 WORK CLOSELY, NOT ONLY WITH ACADEMIA, BUT ALSO PEOPLE
6 SUCH AS NOVOCCELL IN DEVELOPING, MAKING THESE AVAILABLE
7 TO THE BROADER SCIENTIFIC COMMUNITY. GROWTH FACTORS
8 CYTOKINES, AGAIN, VERY KEY FOR SERUM FREE CULTURE OF
9 CELLS, GENE REGULATION AND TRACKING OF CELLS ONCE
10 THEY'RE INTO SMALL ANIMALS OR EVEN ACTUALLY HUMANS AT
11 SOME POINT. AND THEN THE LAST PART, AGAIN, SERVICES
12 ALONG THESE DIMENSIONS, WHICH I'VE TALKED ABOUT
13 EARLIER.

14 THIS IS HOW WE SEE THIS WORLD. THERE IS
15 OBVIOUSLY INVESTMENTS THAT WE AND OTHER COMPANIES HAVE
16 MADE. THE ISSUE IS THEN TAKING THESE TO THE NEXT
17 LEVEL, MAKING THOSE MUCH MORE RELEVANT TO STEM CELL
18 RESEARCHERS IN CALIFORNIA AND, OF COURSE, AROUND THE
19 GLOBE. AND I TALKED A LITTLE BIT ABOUT ENABLING
20 PLATFORMS FOR STEM CELL THERAPY. AGAIN, I'M SHOWING
21 YOU A VERY SIMPLIFIED VIEW OF THIS. BUT YOU NEED THE
22 FOUR ELEMENTS THERE TO MAKE THIS AFFORDABLE, YOU NEED
23 LARGE QUANTITIES OF ECONOMICAL CELLS, IF YOU WILL, AND,
24 OF COURSE, BASE CELLS. YOU NEED TO ISOLATE THE KIND OF
25 CELL THAT YOU REALLY WANT TO PUT INTO THE PATIENT. YOU

1 NEED TO CHARACTERIZE AND BACK THESE CELLS AND, OF
2 COURSE, BE ABLE TO SHIP THEM TO PLACES. AND THE LAST
3 THING IS YOU NEED TO MAKE SURE THAT THE CELLS ARE SAFE,
4 FREE FROM TOXINS, FREE FROM VIRUSES, ETC. ALL OF THOSE
5 REQUIRE INVESTMENT. ALL OF THESE REQUIRE INVESTMENT IN
6 GNP CONDITIONS, AND MOSTLY FOR ALL OF BIOTECH, THIS HAS
7 COME FROM THE PRIVATE SECTOR AND NOT FROM THE ACADEMIC
8 SECTOR. AND SOME OF THESE, OF COURSE, WILL REQUIRE NEW
9 TECHNOLOGIES TO BE DEVELOPED, AND I THINK SOME OF THIS
10 CAN BE SUPPORTED BY ORGANIZATIONS SUCH AS THE CIRM.

11 OF COURSE THE TALK WOULDN'T BE COMPLETE
12 WITHOUT A CURE GRAPHIC HERE. WHAT I WANTED TO SHOW YOU
13 HERE IS THE WAY WE HAVE SEEN OVER THE LIFE OF OUR LIFE
14 AS A COMPANY IS HOW PUBLIC RESEARCH FUNDING ACTUALLY
15 HELPS ULTIMATELY NEW TREATMENTS TO BE DERIVED. I
16 THINK, FIRST OF ALL, THE MAJOR PART OF WHERE PUBLIC
17 FUNDING GOES IS ACTUALLY NOT TO FOR-PROFIT COMPANIES
18 LIKE US, BUT TO ACADEMIC INSTITUTIONS. BUT THE
19 INTERPLAY BETWEEN COMPANIES LIKE US, THE TOOLS
20 COMPANIES, IS MAINLY THEN TAKING THE FUNDAMENTAL
21 RESEARCH THAT'S CARRIED OUT BY ACADEMIC RESEARCHERS AND
22 THE GOVERNMENT AND NONPROFITS AND THEN LICENSING SOME
23 KEY TECHNOLOGIES AND, OF COURSE, MAKE THESE
24 TECHNOLOGIES AVAILABLE TO THE COMMERCIAL RESEARCHERS
25 AND, OF COURSE, BACK TO THE ACADEMIC RESEARCHERS, GIVE

1 A PAYBACK IN TERMS OF LICENSING FEES AND, OF COURSE,
2 ROYALTY PAYMENTS, AND THEN, OF COURSE, THEN THE TOOLS
3 THEN HELP, HOPEFULLY, DEVELOP NEW TREATMENTS, WHETHER
4 IT'S DRUG SCREENING, WHETHER IT'S, YOU KNOW, GROWING
5 CELLS EFFICIENTLY IN ANIMAL ORIGIN FREE ENVIRONMENTS.

6 AND THIS HAS WORKED OUT, THIS HAS BEEN
7 WORKING VERY WELL FOR US AND FOR OUR COMPANIES IN OUR
8 FIELD, BUT I THINK THE KEY PART OF THIS IS REALLY
9 AROUND THE ABILITY TO LICENSE TECHNOLOGIES IN AND
10 KEEPING THE ABILITY TO LICENSE THINGS WITHOUT HAVING
11 ONEROUS TERMS, WITHOUT HAVING RIGHTS WHICH BASICALLY
12 PROVIDE DISINCENTIVES TO LICENSING OFF THESE
13 TECHNOLOGIES.

14 SO WHY DO COMPANIES SEEK GRANTS AND RESEARCH
15 GRANTS TO PURSUE RESEARCH OR OTHER THINGS? FIRST OF
16 ALL, I THINK ONE OF THE THINGS HERE IS RESEARCH THAT
17 THEY CANNOT FUND NOW OR IT DOESN'T MAKE ECONOMIC SENSE
18 TO FUND IMMEDIATELY. SO THAT ALLOWS AN AVENUE FOR US
19 TO FUND THESE KINDS OF RESEARCH. OF COURSE, BEING A
20 PUBLIC COMPANY, AND ALLAN ALLUDED TO THIS, THERE ARE
21 SOME VERY SERIOUS CONDITIONS OF THE KINDS OF RISK
22 PROFILE OF INVESTMENTS WE MAKE AND THE RESEARCH WE
23 CONDUCT. SO HAVING PUBLIC FUNDS OR GOVERNMENT FUNDS
24 SOMETIMES ALLOWS US TO TAKE ON RISK PROFILES AND
25 PROJECTS THAT THE MARKET WOULD NOT, FRANKLY, LIKE US TO

1 TAKE, RIGHT, BUT IT COULD HAVE REWARDS FOR THE RESEARCH
2 COMMUNITY AND OTHERS.

3 RESEARCH, OF COURSE, WHERE THE LEADING
4 PRODUCT OR YOU LEAVE YOUR PRODUCT FOR THE GOVERNMENT IS
5 A MAJOR CONSUMER. AND THEN THERE ARE RESEARCHERS WHERE
6 THERE ARE INTERESTING PARTNERS IN THE SENSE THAT,
7 RATHER I SHOULD SAY INTERESTED, NOT INTERESTING WAY.
8 THERE'S A PARTICULAR INTEREST OR CONSORTIUM OR
9 GOVERNMENT CONSORTIUM THAT HAS COME TOGETHER, FUNDED
10 MAYBE BY THE NIH OR OTHERS.

11 INVITROGEN TRADITIONALLY HAS NOT SOUGHT THESE
12 GRANTS. THERE'S A VERY CLEAR REASON. MOST OF THESE
13 GRANTS TEND TO GO TO SMALLER COMPANIES, AND WE CLEARLY
14 DON'T FIT THAT PROFILE ANY LONGER. BUT WE DO HAVE
15 CONTRACTS WITH GOVERNMENT AGENCIES, AND SEVERAL OF OUR
16 ACQUISITIONS HAVE BEEN PART OF THIS SBIR, U.S. COMMERCE
17 DEPARTMENT ADVANCED ATP GRANTS.

18 AND, OF COURSE, THE IP ARRANGEMENTS HERE ARE
19 VERY CLEAR. THE COMPANY OWNS THE IP, AND THEN THE
20 GOVERNMENT RETAINS THE GOVERNMENT USE LICENSE. AND
21 LASTLY, THERE IS NO CASE OF US HAVING A REVENUE SHARING
22 ARRANGEMENT FOR ANY OF THESE CASES.

23 MORE TO CIRM GRANTS AND WHY WOULD WE WANT TO
24 USE THESE GRANTS AND WHAT BENEFITS MIGHT RESULT TO THE
25 STEM CELL COMMUNITY. I THINK THE BIGGEST THING FOR US

1 IS TO MAKE ROBUST PLATFORMS, TECHNOLOGY PLATFORMS,
2 AVAILABLE FOR THE WORK FLOW THAT I SHOWED YOU EARLIER.
3 THERE IS CLEARLY A NEED FOR MUCH MORE STANDARDIZED
4 CHARACTERIZATION TOOLS FOR THIS INDUSTRY, AND BOTH FOR
5 THE STEM CELLS AND THEIR DIFFERENTIATED PROGENY,
6 WHETHER IT BE IDENTITY, STABILITY, QUALITY,
7 DIFFERENTIABILITY, ETC. THERE'S CLEARLY A NEED FOR THE
8 NEXT GENERATION OF ANIMAL ORIGIN FREE MEDIA AND
9 REAGENTS THAT ARE CRITICAL FOR ANY CELL THERAPY TO BE
10 APPROVED. CRYOPRESERVATION MEDIA AND REAGENTS TARGETED
11 TO STEM CELLS IS AN UNMET NEED AS FAR AS WE CAN TELL SO
12 FAR. AND TRANSFECTION REAGENTS, I THINK THIS IS MUCH
13 MORE FOR, SO FAR AT LEAST, FOR THE RESEARCH SIDE OF
14 THINGS, ARE CRITICAL. THESE CELLS, AS FAR AS WE CAN
15 TELL, ARE PRETTY HARD TO TRANSFECT. AND THE NORMAL
16 TRANSFECTION REAGENTS WE HAVE ON THE MARKETPLACE DO NOT
17 OFTEN SUFFICE.

18 THE BENEFITS THAT MIGHT RESULT TO THE STEM
19 CELL COMMUNITY, OF COURSE, IS INCREASED RANGE OF
20 PRODUCTS, SO WE CAN, WE AND OTHER COMPANIES, SUCH AS
21 US, WILL HAVE AN INCENTIVE TO INTRODUCE A WIDER RANGE
22 OF PRODUCTS ADDRESSING MANY DIFFERENT KINDS OF STEM
23 CELLS. WE COULD IMPROVE THE ROBUSTNESS OF THESE TOOLS.
24 AND, YOU KNOW, ONE OF THE MAJOR ISSUES NOW IS THAT STEM
25 CELLS, BECAUSE OF MAYBE THE WAY THE NIH HAS FUNDED

1 THEM, THEY DON'T TEND TO BE ALIVE. SO EVEN HUMAN
2 EMBRYONIC STEM CELLS FROM ONE LAB AND ANOTHER LAB
3 BEHAVE VERY DIFFERENTLY, AND TOOLS ARE OFTEN DEVELOPED
4 JUST FOR ONE TYPE OF STEM CELL OR ANOTHER. AND THEN
5 THE LAST THING IS SPEED, WHICH IS, AGAIN, CRITICAL,
6 ESPECIALLY WHEN YOU'RE COMPETING, NOT ONLY AGAINST
7 OTHER STATES, OTHER UNIVERSITIES, BUT AS SEVERAL OF THE
8 SPEAKERS HAVE POINTED OUT, THIS IS A VERY, VERY GLOBAL
9 ENVIRONMENT. PROBABLY FOR THE FIRST TIME IN THE
10 HISTORY OF BIOTECH THAT THE COMPETITION HAS BEEN SO
11 GLOBAL.

12 SO I WANTED A TAKE AWAY HERE IS REALLY WE
13 WOULD SEEK THESE GRANTS TO LEVERAGE INTERNAL
14 INVESTMENTS THAT WE ARE CONTINUING TO MAKE AND THEN
15 ACCELERATE NOVEL TOOL AVAILABILITY.

16 SO FAR SOME TERMS HERE AND, AGAIN, THIS IS
17 EARLY THINKING, BUT WE ARE COGNIZANT OF THE FACT THAT
18 THIS IS TAXPAYER MONEY, AND THERE IS A NEED FOR US TO
19 GIVE BACK TO THE STATE. SO YOU WILL SEE A LOT OF THAT,
20 AND I'M HOPING THERE ARE QUESTIONS TO DISCUSS SOME OF
21 THESE. BUT I THINK ONE OF THE THINGS WHICH HAS TO BE
22 THERE IS THAT THE GRANTEE MUST RETAIN THE IP. MOST
23 COMMERCIAL ORGANIZATIONS WOULD NOT ENGAGE IN THIS SORT
24 OF GRANT FUNDING UNLESS THERE IS CLARITY ON THE IP.

25 SECOND, I THINK THE FEDERAL POLICY'S EMPHASIS

1 ON STIMULATING RESEARCH ADVANCES AND ECONOMIC
2 DEVELOPMENT, I THINK, ED, YOU REFERRED TO THIS EARLIER,
3 THE DIRECT VERSUS THE INDIRECT BENEFITS BACK TO THE
4 STATE, I THINK THEY ARE CLEARLY THERE. WE DO INTEND TO
5 DO A LOT OF OUR RESEARCH IN CALIFORNIA. AND, YOU KNOW,
6 I THINK THE STATE WILL BENEFIT FROM THAT.

7 BUT THERE ARE OTHER MODELS TO PROVIDE MORE
8 DIRECT PAYBACK TO THE STATE AND TO CIRM. FIRST OF ALL,
9 IT COULD BE THROUGH ROYALTIES ON PRODUCTS
10 COMMERCIALIZED USING IP DEVELOPED WITH CIRM FUNDING.
11 THIS IS NOT UNCOMMON. WE'VE DONE THIS WITH OTHER
12 COMPANIES AND OBVIOUSLY INSTITUTIONS. I'M PUTTING
13 QUOTES AROUND THE ROYALTY BECAUSE IN SOME WAYS WHEN YOU
14 LICENSE THINGS, THE ROYALTY IS OWNED BY THE ACADEMIC
15 INSTITUTION. NOW, HERE YOU ARE -- YOU WOULD ENABLE US
16 TO DEVELOP IP, BUT THAT WOULDN'T BE OUR PROPERTY WHERE
17 WE WOULD BE OPEN TO ACTUALLY PROVIDING YOU BACK A
18 ROYALTY ON PRODUCTS DEVELOPED THROUGH THAT IP.

19 THE SECOND THING IS, AND I THINK CIRM SHOULD
20 REALLY CONSIDER THIS, IS THIS CONCEPT OF LEVERAGING
21 YOUR FUNDS, RIGHT. SO I THINK WE WOULD BE OPEN TO
22 INVESTING OURSELVES IN A MATCHING MANNER TO FUNDS THAT
23 YOU PROVIDE US TO, YOU KNOW, TO DOUBLE OR TRIPLE THE
24 AMOUNT OF INVESTMENT THAT IS PROVIDED BY CIRM.

25 AND THEN THE LAST PART IS REASONABLE

1 CONSIDERATION OF MANUFACTURING ACTIVITIES IN
2 CALIFORNIA. I THINK THAT'S AN IMPORTANT CONSIDERATION,
3 AND THAT GIVES BACK TO THE STATE IN GENERATING
4 EMPLOYMENT.

5 WE WOULD BE OPEN TO STANDARD THIRD-PARTY
6 AUDIT OF RESEARCH AND ROYALTIES. THIS IS SOMETHING WE
7 DO WITH EVERY INSTITUTION THAT WE HAVE. SOME OF THE
8 CONCERNS THAT ALLAN RAISED IN TERMS OF MAINTAINING THE
9 PROPRIETARY NATURE OF THE RESEARCH IN THIS CASE DO NEED
10 TO BE ADDRESSED, BUT I THINK -- WE HAVE ADDRESSED THAT
11 WITH PUBLIC INSTITUTIONS, SUCH AS THE UNIVERSITY OF
12 CALIFORNIA.

13 PROGRESS REPORTS ON OUR PROGRESS ON THE
14 RESEARCH, AGAIN, THAT'S SOMETHING WE'RE OPEN TO.

15 AND THEN THE LAST BIT IS RESEARCH USE AND
16 OTHER IP PROVISIONS NEED TO PRESERVE A COMMERCIAL
17 OPPORTUNITY. AND I THINK WE'VE DEBATED SOME OF THIS ON
18 THE NONPROFIT PART.

19 ONE OF THE THINGS THAT'S NOT HERE, AND I
20 THINK, AGAIN, MANY PEOPLE HAVE ALLUDED TO, IS WHAT IS
21 THE IMPLICATION OF HAVING OVERARCHING IP SUCH AS THE
22 WISCONSIN IP ON FUNDAMENTAL STEM CELL PATENTS. WHAT IS
23 THE IMPACT OF THAT ON COMMERCIAL ENTITIES, RIGHT, AND
24 OUR ABILITY TO PROVIDE TOOLS AND REAGENTS TO THIS
25 COMMUNITY? I KNOW CIRM IS CONSIDERING THAT. I WANT TO

1 TELL YOU FROM A BUSINESS POINT OF VIEW WHAT IS
2 HAPPENING IS THAT PEOPLE DEMAND A CERTAIN AMOUNT OF
3 MONEY FROM EVERY COMMERCIAL COMPANY TO DO EVEN VERY,
4 VERY SMALL, INCREMENTAL, I WOULD SAY, TOOLS TO THE
5 COMMUNITY. AND THE MONEY FAR OUTSTRIPS ANY COMMERCIAL
6 BENEFIT THAT WE COULD HAVE BY SELLING THESE TOOLS AND
7 REAGENTS.

8 THE INTERESTING SIDE OF THIS IS THESE TOOLS
9 AND REAGENTS, ALTHOUGH REVENUE MAY BE SMALL, ARE
10 EXTREMELY IMPORTANT TO THE COMMUNITY. SO THINK ABOUT
11 WHAT SOME ENGINEERED STEM CELL LINES COULD DO FOR THE
12 RESEARCH COMMUNITY IN BEING ABLE TO INSTANTLY TELL YOU
13 WHEN DIFFERENTIATION OCCURS UNDER WHAT CONDITIONS.
14 TODAY THAT IS AN ONEROUS PROCESS, BUT THERE ARE
15 MOLECULAR BIOLOGY TOOLS THAT WE ARE FAMILIAR WITH WHICH
16 COULD CHANGE THIS INDUSTRY. THE PROBLEM IS WE CANNOT
17 SELL THESE CELL LINES TO ANYONE IN THE UNITED STATES,
18 AND I WANT TO STRESS THAT, THIS IS ONLY A PROBLEM IN
19 THE UNITED STATES. OF COURSE, IN CALIFORNIA, PART OF
20 THE UNITED STATES, WE CANNOT MAKE THESE TOOLS AVAILABLE
21 BECAUSE THE -- IF YOU HAVE TO PAY THE LICENSE FEES THAT
22 PEOPLE ARE DEMANDING, IT'S A NONECONOMICAL VENTURE.

23 I ALSO WANT TO STRESS THIS IS WHEN YOU'RE
24 LOOKING AT COMPETITIVENESS OF CALIFORNIA AND, INDEED,
25 THE UNITED STATES AGAINST PLACES LIKE SINGAPORE, PLACES

1 LIKE KOREA, CHINA, EVEN MEXICO, BRAZIL, WE ARE AT A
2 COMPETITIVE DISADVANTAGE. AND I THINK CIRM NEEDS TO
3 CONSIDER SOME OF THAT WHEN IT LOOKS AT IT BROADLY ABOUT
4 COMPETITIVENESS OF THE STATE, COMPETITIVENESS OF THE
5 COMPANIES, AND COMPETITIVENESS OF RESEARCH THAT IS
6 CARRIED OUT HERE.

7 ANYWAY, I JUST WANTED TO MAKE THAT POINT AND
8 MOVE ON FROM HERE. HAPPY TO TAKE QUESTIONS ABOUT THAT.
9 WE'VE THOUGHT A LOT ABOUT THE IMPLICATIONS OF THIS
10 PARTICULAR PATENT, AND I KNOW OTHERS SUCH AS JEANNE IN
11 THE AUDIENCE WILL ALSO TALK ABOUT IT.

12 I WANT TO SPEND TWO MINUTES ON --

13 CHAIRMAN PENHOET: YOU'RE NOT PROPOSING WE
14 SECEDE FROM THE UNION?

15 MR. GOSWAMI: I THINK, MARY, I MAY HAVE
16 MISUNDERSTOOD, BUT YOU DID WANT US TO COMMENT ON BOTH
17 NONPROFIT. WE DO HAVE SOME CONCERNS ABOUT THE
18 NONPROFIT POLICY, AND I THINK IT GOES BACK TO THAT
19 DIAGRAM I SHOWED EARLIER. WE DON'T WANT TO BREAK THAT
20 LINK BETWEEN ABILITY TO LICENSE FROM NOT-FOR-PROFIT
21 ORGANIZATIONS. AND REALLY THE IMPORTANT PART THERE IS
22 THAT THERE SHOULDN'T BE CLAUSES IN CIRM'S FUNDING OF
23 NOT-FOR-PROFIT ORGANIZATIONS THAT PROVIDE DISINCENTIVES
24 FOR COMPANIES SUCH AS US TO LICENSE. AND I THINK THE
25 BIGGEST CONCERN WE HAVE RIGHT NOW IS AROUND THE

1 RESEARCHER'S EXEMPTION THAT REQUIRES CIRM-FUNDED
2 INVENTIONS TO PROVIDE ALL CALIFORNIA INSTITUTIONS FOR
3 RESEARCH PURPOSES AT NO COST. RIGHT. AND I THINK THIS
4 IS -- IF YOU LOOK AT IT, IF SOMEBODY ELSE NEEDS TO BUY
5 THIS REAGENT AT NO COST OR MAKE IT FOR FREE, WHY WOULD
6 SOMEONE WANT TO LICENSE IT AND SPEND THE MONEY AND
7 INVEST IN BRINGING IT TO THE MARKET IF THERE'S NO
8 GUARANTEE THAT YOU WON'T GET SIDESTEPED BY A PUBLIC
9 ENTITY IN THIS MANNER.

10 I'M GOING TO LEAVE THIS FOR -- THIS IS OUR
11 VIEWPOINT, AT LEAST, THAT SOME OF THE RESEARCH USE
12 POLICY FROM CIRM CONCERNS ELIMINATE THE POSSIBILITY
13 THAT THE COMMERCIAL SECTOR WILL HELP IN DISSEMINATING
14 SOME OF THESE NEW RESEARCH TOOLS. THE LIMIT, IN OUR
15 MIND AT LEAST, STEM CELL RESEARCH PROGRESS, WHICH IS A
16 LOT OF THIS, IS BASED ON HAVING GOOD, STANDARDIZED
17 TOOLS AVAILABLE EARLY TO THE MARKET. AND THEN, OF
18 COURSE, IT ALSO ELIMINATES POTENTIAL ROYALTY BACK TO
19 THE STATES. LOOK, I TOTALLY UNDERSTAND THE ROYALTY
20 AMOUNTS FOR RESEARCH TOOLS ARE SMALL, BUT STILL IT IS
21 SOMETHING THAT IS GIVEN BACK TO THE STATE EARLY.

22 AND THEN, OF COURSE, IF YOU TAKE THE FRUIT
23 SIDE OF IT, IF COMPANIES SUCH AS US DO NOT LICENSE IN
24 TECHNOLOGIES SUCH AS ANTIBODIES AND OTHERS, IT CREATES
25 AN UNDUE BURDEN ON THE STATE AND THE NONPROFIT

1 INSTITUTIONS TO PROVIDE THESE AND MAKE THESE AVAILABLE
2 TO RESEARCHERS.

3 AND ANYWAYS, I THINK I'VE BELABORED THIS
4 POINT. I THINK WE DON'T WANT THINGS THAT CROSS OUT AND
5 PREVENT LICENSING IN BOTH DIRECTIONS AND, OF COURSE,
6 PAYBACK TO THE STATE. THAT'S IT.

7 CHAIRMAN PENHOET: VERY GOOD. THANK YOU.
8 ONE QUESTION. THE 40 LICENSES THAT YOU NOW HAVE, OF
9 THE 40, HOW MANY ARE EXCLUSIVE LICENSES AND HOW MANY
10 ARE NONEXCLUSIVE?

11 MR. GOSWAMI: THAT'S 40 PER YEAR.

12 CHAIRMAN PENHOET: FORTY PER YEAR.

13 MR. GOSWAMI: I WOULD SAY ABOUT 20 PERCENT OR
14 SO TEND TO BE EXCLUSIVE, RIGHT. AND IT DEPENDS VERY
15 MUCH ON THE TECHNOLOGY THAT WE LICENSE. SO GENERALLY
16 THINGS LIKE ANTIBODY LICENSES ARE NONEXCLUSIVE JUST
17 BASED HOW THE COMPANIES HAVE GONE ABOUT DOING THIS.
18 WHERE IT MAKES SENSE TO HAVE EXCLUSIVE LICENSES IS
19 WHERE THE AMOUNT OF INVESTMENT THAT WE HAVE TO PUT IN
20 IS QUITE SIGNIFICANT AND, THEREFORE, WE WOULD WANT
21 THAT -- THE LICENSE TO BE EXCLUSIVE TO PROVIDE US A
22 RETURN ON OUR INVESTMENT BEFORE IT REACHES OR WHEN IT
23 REACHES THE MARKET.

24 CHAIRMAN PENHOET: OTHER QUESTIONS?

25 DR. PRIETO: I WAS WONDERING IF YOU COULD

1 SPEAK A LITTLE BIT ON WHAT YOU THINK THE IMPACT OF THE
2 WARF PATENTS IS? ARE THEY A SIGNIFICANT OBSTACLE TO
3 PRIVATE INVESTMENT?

4 MR. GOSWAMI: I THINK SO. I THINK THEY ARE.
5 YOU KNOW, I KNOW A LOT OF YOU KNOW ABOUT THE WARF
6 POLICY FOR NOT-FOR-PROFIT RESEARCH TENDS TO BE
7 SOMEWHAT, WELL, SOMEWHAT GENEROUS, ALTHOUGH MANY
8 SCIENTISTS THAT WE HAVE SPOKEN TO ARE QUITE TAKEN ABACK
9 AT SOME OF THE RESTRICTIONS THAT ARE PUT ON ACADEMIC
10 INSTITUTIONS IN TERMS OF COLLABORATING WITH COMPANIES
11 SUCH AS US. FOR PRIVATE INSTITUTIONS, I THINK IT IS A
12 BIG DISINCENTIVE BECAUSE WHAT WARF DOES IS IT LOOKS AT
13 THE SIZE OF THE INSTITUTION OR THE COMPANY THAT WANTS
14 TO LICENSE THIS PRODUCT AND WANTS TO CHARGE AN UPFRONT
15 PAYMENT, WHICH IS VERY STEEP IN ANY LICENSING TERMS,
16 AND THEN, OF COURSE, ROYALTIES OBLIGATIONS WHICH ARE
17 TWOFOLD. ONE, A DIRECT ROYALTY OBLIGATION WHICH
18 DEPENDS ON PRODUCTS DIRECTLY PRODUCED USING STEM CELLS,
19 HUMAN EMBRYONIC OR PRIMATE EMBRYONIC STEM CELLS, BUT
20 THEY ALSO HAVE A REACH-THROUGH ROYALTY CLAUSE WHERE
21 THEY SAY ANYTHING YOU'VE INVENTED REMOTELY BY USING
22 HUMAN EMBRYONIC STEM CELLS WILL NOW HAVE A ROYALTY
23 OBLIGATION BACK TO WARF. SO, FOR INSTANCE, EVEN IF YOU
24 HAD A TREATMENT THAT WAS SOMEHOW BROUGHT TO THE MARKET
25 THAT HAS TOUCHED AN EMBRYONIC STEM CELL IN THE UNITED

1 STATES, YOU WILL HAVE TO PAY A ROYALTY BACK TO WARF.

2 AND I THINK THIS IS PARTICULARLY ONEROUS. WE
3 ARE SOMEWHAT IN THAT CAMP, RIGHT, BECAUSE I THINK, EVEN
4 IF WE'RE NOT ON THE THERAPEUTIC SIDE, LET'S SAY WE MAKE
5 A DISCOVERY OF A TOOL USING AT SOME POINT A HUMAN
6 EMBRYONIC STEM CELL. THEY WANT A ROYALTY BACK ON THAT
7 PARTICULAR PRODUCT. AND NOT ONLY DO THEY WANT A
8 ROYALTY BACK ON SALES OF THAT PRODUCT TO THE EMBRYONIC
9 STEM CELL MARKET, BUT ANY MARKET THAT WE TOUCH, THEY
10 WANT A ROYALTY BACK ON THAT PARTICULAR TOOL. SO IT IS
11 QUITE ONEROUS.

12 DR. PRIETO: EVEN MARKETS OUTSIDE THE UNITED
13 STATES?

14 MR. GOSWAMI: YES. IN GENERAL THEY WOULD.
15 SEE, THE OTHER PART IS, YOU KNOW, IF YOU LOOK AT
16 COMPANIES IN THE UNITED STATES VERSUS OUTSIDE, RIGHT,
17 WHICH IS ACTUALLY EVEN MORE INTERESTING. SO LET'S SAY
18 THERE'S A COMPANY IN INDIA, RIGHT. FOR ANYTHING THEY
19 DO WITH RESPECT TO HUMAN EMBRYONIC STEM CELL RESEARCH,
20 THEY OWE NO ROYALTIES TO WARF. THEY DO NOT HAVE AN
21 OBLIGATION TO WARF IN ANY MANNER, WAY, SHAPE, OR FORM,
22 AND THEY CAN DEVELOP EXACTLY THE SAME THERAPIES
23 COMPANIES IN THE UNITED STATES ARE TRYING TO DEVELOP
24 WITHOUT ANY KIND OF RESTRICTIONS OR OBLIGATION TO WARF.
25 SO FROM THAT POINT OF VIEW, SETTING UP A

1 COMPANY TO DO STEM CELL THERAPY IN THE UNITED STATES, I
2 THINK, IS A MAJOR DISINCENTIVE, AND PEOPLE ARE NOT
3 REALIZING THIS BUSINESS ISSUE. I DON'T THINK, AT LEAST
4 FROM OUR POINT OF VIEW, THE ISSUE IS NOT WHETHER THE
5 PATENT IS LEGITIMATE OR NOT, BUT THERE ARE SERIOUS
6 BUSINESS IMPLICATIONS OF THE PATENT IN MAKING CURES
7 AVAILABLE TO THE PUBLIC HERE.

8 THE REPORTER: I'M SORRY. THIS IS THE
9 REPORTER. I DIDN'T CATCH THE NAME OF THE PERSON WHO
10 ASKED THE LAST TWO QUESTIONS.

11 DR. PRIETO: I'M SORRY. THIS IS FRANCISCO
12 PRIETO IN SAN DIEGO.

13 THE REPORTER: OKAY. THANK YOU.

14 DR. REED: JOHN REED HERE IN SAN DIEGO. JUST
15 A QUESTION ABOUT THIS CONCEPT OF INDIRECT RETURN TO THE
16 STATE. INVITROGEN IS CLEARLY ONE OF THE SUCCESS
17 STORIES OF THE CALIFORNIA BIOTECHNOLOGY INDUSTRY. ANY
18 IDEA WHAT YOU'RE PAYING IN CORPORATE INCOME TAXES EVERY
19 YEAR IN THE STATE OF CALIFORNIA?

20 MR. GOSWAMI: I DON'T KNOW OFF THE TOP OF MY
21 HEAD, BUT THAT'S PUBLIC INFORMATION. I'M SURE THAT WE
22 CAN GET THAT INFORMATION TO YOU. IT SHOULDN'T BE AN
23 ISSUE. WE HAVE A MAJORITY OF OUR BUSINESS, OR I SHOULD
24 SAY RESEARCH AND MANUFACTURING ACTIVITY, LOCATED IN THE
25 STATE. SO I'M SURE A MAJORITY OF THE TAX ACTUALLY IS

1 PAID IN THE STATE.

2 CHAIRMAN PENHOET: IT MUST BE \$150 MILLION A
3 YEAR.

4 DR. REED: SALES ARE ABOUT 1.5 BILLION?

5 MR. GOSWAMI: YEAH. 1.3 BILLION OR SO.

6 DR. REED: SO IT'S GOT TO BE SOMETHING IN
7 THAT BALLPARK?

8 MR. GOSWAMI: YEAH.

9 DR. REED: THAT'S ONE COMPANY. JUST
10 SOMETHING TO KEEP IN MIND.

11 MR. GOSWAMI: YEAH, I THINK WE ARE A LARGE
12 COMPANY. I THINK THE DECISIONS COMPANIES MAKE TO
13 INVEST IN THE FIELD, OF COURSE, CONSIDER THE SIZE OF
14 THE FIELD AND NOT -- YOU KNOW, IT IS SOMEWHAT OF A
15 ANIMAL WITHIN THE COMPANY AND WITH ITS OWN INVESTMENT
16 DECISIONS TO BE MADE.

17 CHAIRMAN PENHOET: OKAY. ANY COMMENTS FROM
18 IRVINE? QUESTIONS?

19 DR. BRYANT: NO.

20 CHAIRMAN PENHOET: FROM SAN CARLOS?

21 UNIDENTIFIED SPEAKER: NO.

22 CHAIRMAN PENHOET: LOS ANGELES? CHICO?

23 DR. WRIGHT: NO.

24 CHAIRMAN PENHOET: OKAY. NOW DO WE HAVE
25 COMMENTS FROM THE AUDIENCE IN SAN DIEGO? THIS IS ALLAN

1 ROBINS.

2 MR. ROBINS: ALLAN ROBINS IN SAN DIEGO. I
3 JUST -- I THOUGHT FRANCISCO BROUGHT UP A VERY GOOD
4 POINT. AND, JOYDEEP, I JUST WANTED TO ADD A LITTLE BIT
5 TO YOUR ANSWER. IS THAT IF A COMPANY IN THE FIELD OR
6 THE UNITED KINGDOM OR SINGAPORE DEVELOPS A PRODUCT OR
7 HUMAN EMBRYONIC STEM CELLS AND THAT PRODUCT IS
8 DIFFERENT FROM THE HUMAN EMBRYONIC STEM CELL, THAT
9 PRODUCT CAN BE IMPORTED INTO THE U.S.A. AND SOLD HERE,
10 AND THAT WOULD NOT INFRINGE THE LINE. AND SO YOU
11 REALLY ARE PUTTING COMPANIES THAT OPERATE IN THE U.S.
12 AT A DISADVANTAGE.

13 MR. GOSWAMI: THAT'S A GREAT POINT.

14 DR. PRIETO: FRANCISCO PRIETO AGAIN. THAT
15 WAS SORT OF WHAT I WAS STARTING TO GLEAN FROM THIS THAT
16 AS THEY CURRENTLY OPERATE, THAT THESE PATENTS ARE A
17 DISINCENTIVE TO THE RESEARCH IN THE UNITED STATES.

18 MR. GOSWAMI: IT IS ENORMOUS. AND THE
19 PATENTS ONLY, AS ALLAN RIGHTLY POINTS OUT, ARE TO THE
20 COMPOSITION OF MATTER OF THE PRIMATE EMBRYONIC STEM
21 CELL AND THE METHOD OF DERIVATION. SO ABSOLUTELY
22 RIGHT.

23 MR. ROTH: HI, IT'S DUANE ROTH IN SAN DIEGO.
24 I'M GOING TO -- I WOULD LIKE TO MAKE JUST A FEW
25 COMMENTS SPEAKING FROM MY POSITION AT CONNECT HERE IN

1 SAN DIEGO. OUR MISSION IS TO HELP TURN GOOD SCIENCE
2 AND GOOD IDEAS INTO BUSINESSES AND NEW COMPANIES, NEW
3 COMPANY CREATION.

4 I THOUGHT THE COMMENTS THAT WERE MADE BY
5 JOYDEEP AND ALLAN WERE APPROPRIATE FOR EXISTING
6 COMPANIES THAT USE THIS TECHNOLOGY, GET IT INTO THEIR
7 COMPANY TO DELIVER PRODUCTS TO THE MARKET. BUT FROM
8 THE START-UP STANDPOINT, THERE'S THREE COMMENTS I WOULD
9 MAKE ABOUT INTELLECTUAL PROPERTY.

10 FIRST, IT'S ABSOLUTELY NECESSARY, AND I'LL
11 COME BACK TO THAT IN A SECOND.

12 NO. 2 IS IT'S EXTRAORDINARILY COMPLICATED.
13 YOU HEARD THE NUMBER OF STEM CELL PATENTS THAT ARE
14 ALREADY OUT THERE, THE ONES THAT ARE ISSUED, AND THEN
15 YOU HAVE THOUSANDS THAT ARE IN THE PROCESS OF BEING
16 REVIEWED.

17 AND THE THIRD IS THAT INCREASINGLY IT IS
18 VERY, VERY EXPENSIVE. AND I CALL THEM THE SHUNS, BUT
19 YOU FIRST HAVE THE PREPARATION, WHICH IS A NECESSARY
20 AND EXPENSIVE PART OF IT, AND THERE'S THE PROSECUTION
21 OF THOSE PATENTS WORLDWIDE AND THE FEES YOU HAVE TO
22 PAY, ALL THE THINGS YOU HAVE ENTER INTO EVEN IN THE
23 APPLICATION PHASE AND THEN AT ISSUE.

24 THEN THE THIRD IS NEGOTIATION. YOU END UP
25 WITH ALL THIS INTELLECTUAL PROPERTY AND MILLIONS AND

1 MILLIONS OF DOLLARS BEING SPENT BY LAWYERS TRYING TO
2 NEGOTIATE THE RIGHT TO PRACTICE, WHICH IS INCREASINGLY
3 TIME AND MONEY THAT NOBODY ACCOUNTS FOR.

4 AND THEN THE FINAL ONE IS LITIGATION. AND
5 YOU END UP IN MANY, MANY CASES OVER MAJOR PATENTS,
6 THAT'S EXACTLY WHAT YOU'RE ALL TALKING ABOUT HERE,
7 LITIGATION. WHO'S GOING TO PAY FOR THAT? AND WHAT'S
8 THE ULTIMATE OUTCOME GOING TO BE? SO I THINK THOSE
9 THINGS NEED TO BE KEPT IN MIND.

10 BUT LET ME GO TO THE NECESSARY. WHAT I THINK
11 YOU'RE TRYING TO DO WITH INTELLECTUAL PROPERTY FOR
12 START-UP COMPANIES IS CREATE AN ENVIRONMENT WHERE YOU
13 CAN LEVERAGE FINANCING AND TRY TO GET PEOPLE TO INVEST
14 IN AN IDEA, A DREAM. AND THEY AREN'T GOING TO DO THAT
15 UNLESS THEY THINK, IF THEY GO THROUGH ALL THIS WORK AND
16 EXPENSE AND RISK, THAT THERE IN THE END IS RETURN.
17 THAT'S THE FIRST THING IP NEEDS TO DO.

18 THE SECOND IS YOU WANT THAT IP TO LEAD TO
19 PRODUCTS THAT ACTUALLY GET TO PEOPLE. AND FROM OUR
20 STANDPOINT, YOU START THESE THINGS, AND THAT'S WHAT
21 IT'S ALL ABOUT, GETTING PRODUCTS TO PATIENTS. TAKING
22 GOOD SCIENCE AND DEVELOP PRODUCTS AND GET THEM TO THE
23 PEOPLE THAT NEED THEM.

24 THE THIRD CONSIDERATION, AND AN IMPORTANT
25 ONE, IS ROYALTY. IF THERE IS A CONTRIBUTION TO THE

1 INTELLECTUAL PROPERTY THAT EVENTUALLY LEADS TO A
2 PRODUCT, WHICH IS A RARE OCCASION, THEN THERE SHOULD BE
3 A SHARING OF THE PROFITS THAT ARE DERIVED FROM THAT. I
4 DON'T THINK ANYBODY THAT'S SPOKEN TODAY WOULD DISCOUNT
5 THAT. BUT I WANT TO REMIND YOU, AND I USE THIS
6 CONSTANTLY, A HUNDRED PERCENT OF NOTHING IS NOTHING.
7 SO IT HAS TO BE VERY CAREFULLY USED, AND MANY THINGS
8 THAT ALLAN TALKED ABOUT EARLIER, STACKING ROYALTIES,
9 THESE THINGS COME UP CONSTANTLY.

10 THE CONCERN IS THE RISK OUTWEIGHS THE
11 OPPORTUNITIES. WHATEVER WE END UP WITH, I THINK IT HAS
12 TO FOLLOW THOSE GUIDELINES, PARTICULARLY FOR START-UP
13 COMPANIES.

14 CHAIRMAN PENHOET: THANK YOU. JOHN SIMPSON.

15 MR. SIMPSON: JUST ONE QUICK QUESTION. YOU
16 TALKED ABOUT 40 LICENSES A YEAR THAT YOU'RE DEVELOPING.
17 I HATE TO KEEP GOING BACK TO THE SAME QUESTION, BUT I
18 RARELY HAVE AN OPPORTUNITY TO TALK TO EXECUTIVES OF
19 COMPANIES WITH PRACTICE ON THIS. SO WOULD A PATENT
20 POOL WITH APPROPRIATE PATENTS PROVIDE YOU AN
21 OPPORTUNITY TO NEGOTIATE ONE-STOP SHOPPING TO GET
22 LICENSES THAT YOU NEED? WOULD THAT BE BENEFICIAL IF
23 THERE WERE SUCH A THING?

24 MR. GOSWAMI: YOU KNOW, AGAIN, IT'S A TOUGH
25 QUESTION TO ANSWER, RIGHT, BECAUSE I THINK IF YOU ASK

1 ME THAT QUESTION AND ALLAN THAT QUESTION, THE PORTFOLIO
2 OF PATENTS WOULD BE DIFFERENT.

3 NOW, THE WARF ISSUE THAT PEOPLE HAVE BROUGHT
4 UP EARLIER IS ONE SUCH PATENT THAT TRANSCENDS ANY WORK
5 THAT PEOPLE WOULD DO IN THE HUMAN EMBRYONIC STEM CELL
6 FIELD. SO THAT'S WHY YOU GET THIS PATENT COMING UP
7 AGAIN AND AGAIN. AND SOMEHOW AND SOMEBODY AT THE WORLD
8 STEM CELL CONFERENCE BROUGHT UP THIS THING. IF YOU
9 THINK BACK, ONE OF THE SIMILAR PATENTS WAS THE COVARO
10 PATENT WHICH INHIBITED MOST OF BIOTECHNOLOGY. AND THE
11 REASON THAT PATENT WAS SO SUCCESSFUL IN SOME WAYS IS
12 BECAUSE IT WAS LICENSED OUT TO PEOPLE ON WHAT THEY
13 CONSIDERED REASONABLE TERMS, RIGHT. THERE'S RISK THAT
14 ONE HAS TO TAKE UP FRONT. DOESN'T SEEM TO BE A
15 CONSIDERATION FOR HOW THE LICENSING MODEL IS WORKING
16 WITH WISCONSIN, WHICH IS HUGE UPFRONT PAYMENTS FOR A
17 FIELD THAT IS, YOU KNOW, PARDON THE PUN, BUT EMBRYONIC.
18 AND JUST FROM A BUSINESS POINT OF VIEW, IT MAKES VERY
19 LITTLE SENSE. AND THAT'S WHY MAYBE CIRM COULD HELP IN
20 ENABLING THAT PARTICULAR PATENT BECAUSE IT WILL BE USED
21 BY ALMOST EVERYBODY THAT IS ENGAGED IN THIS FORM OF
22 RESEARCH IN THE STATE OF CALIFORNIA.

23 SO THAT'S WHY I BROUGHT THAT UP. BUT I DON'T
24 THINK THERE WOULD BE ONE PORTFOLIO THAT WOULD MAKE
25 SENSE TO EVERYBODY. IT GETS TOO MESSY TO TRY AND DO

1 THAT.

2 CHAIRMAN PENHOET: ANY OTHER PUBLIC COMMENTS
3 IN SAN DIEGO? HOW ABOUT COMMENTS IN IRVINE?

4 DR. BRYANT: NO.

5 CHAIRMAN PENHOET: SAN CARLOS?

6 UNIDENTIFIED SPEAKER: NO.

7 CHAIRMAN PENHOET: LOS ANGELES?

8 UNIDENTIFIED SPEAKER: NO.

9 CHAIRMAN PENHOET: CHICO?

10 DR. WRIGHT: NO.

11 CHAIRMAN PENHOET: WE WILL THANK YOU, GUYS,
12 THANK YOU. VERY INFORMATIVE PRESENTATION. THAT BRINGS
13 OUR PRESENTATIONS TO A CLOSE, BUT I THINK IT WOULD BE
14 APPROPRIATE AT THIS TIME TO OPEN THE FLOOR TO ISSUES,
15 QUESTIONS, THAT MAY RELATE TO THESE, MAY NOT RELATE TO
16 THESE ON THE GENERAL SUBJECT OF TODAY'S MEETING, WHICH
17 IS AN INTELLECTUAL PROPERTY POLICY FOR, JUST CALL IT,
18 GRANTS. I THINK WE'VE HEARD THERE MAY BE GRANTS, THERE
19 MAY BE LOANS, THERE MAY BE CONTRACTS, A NUMBER OF
20 DIFFERENT VEHICLES FOR FUNDING THIS KIND OF WORK IN THE
21 PRIVATE SECTOR. SO NOW I'LL LOOK FORWARD TO ANY
22 COMMENTS ON TODAY'S PRESENTATIONS OR OTHERWISE.

23 BUT SPECIFICALLY FOR THOSE OF YOU ON THE TASK
24 FORCE, WE'D LIKE YOU ALSO TO BE THINKING ABOUT ANY GAPS
25 IN WHAT WE'VE HEARD SO FAR. WE HAVE HEARD FROM THE

1 FEDERAL GOVERNMENT, AT LEAST FROM THE NIH. WE HAVE
2 HEARD FROM INDUSTRY SOURCES. WE HAVE HEARD FROM
3 MR. SIMPSON. WE HAVE HEARD FROM A NUMBER OF DIFFERENT
4 GROUPS SO FAR. WE HAVE HEARD FROM THE JDRF. AND
5 PERHAPS MARY AND I, IF WE HAVE TIME, WILL SHARE A
6 CONVERSATION WE HAD WITH THE WELLCOME TRUST, WHICH HAS
7 ACTUALLY NOW BASED ITSELF IN FUNDING FOR-PROFIT
8 ENTITIES. IT'S THE SECOND LARGEST FOUNDATION IN THE
9 WORLD ACTUALLY AFTER GATES FOUNDATION. I THINK FOR
10 SIMILAR REASONS, THAT THEY ARTICULATED TO US AT LEAST,
11 WHAT THEIR VIEW IS IS BASICALLY THEY SHOULD FUND
12 ORGANIZATIONS WHICH ARE MOST LIKELY TO MOVE A FIELD
13 FORWARD. AND IF THAT HAPPENS TO BE A FOR-PROFIT
14 AGENCY, THAT'S WHAT THEY SHOULD FUND. SO THEY ACTUALLY
15 HAVE A MAJOR COMPONENT OF THEIR ACTIVITIES IN THAT
16 DIRECTION.

17 WE HOPE IN THE NEXT MEETING TO ACTUALLY HAVE
18 HERE SOMEONE FROM THE TRUST COME TO SPEAK TO US IN
19 PERSON, BUT ALSO BY PHONE. WOULD THAT -- LET'S OPEN
20 THE FLOOR TO THE TASK FORCE MEMBERS FIRST FOR COMMENTS
21 ON ANY OF THE THINGS WE'VE BEEN DISCUSSING HERE OR
22 QUESTIONS OF EACH OTHER, ETC. OPEN FORUM. JOHN REED.

23 DR. REED: I'LL START THE DIALOGUE OFF HERE.
24 JOHN REED. I WANTED TO THANK ALL THE SPEAKERS HERE FOR
25 THEIR PRESENTATIONS. I THOUGHT THEY WERE HIGHLY

1 INFORMATIVE AND GAVE US, I THINK, INSIGHTS TO SOME OF
2 THE SPECTRUM OF VIEWS THAT WE'RE LIKELY TO EXPERIENCE
3 AS WE TACKLE WITH THIS ISSUE OF INTELLECTUAL PROPERTY.

4 I WAS REALLY COMPELLED BY THE PRESENTATION
5 THAT REFERRED BACK TO WHAT WE EARLY LEARNED ABOUT THE
6 WAY JDRF FUNDS WORK IN FOR-PROFIT COMPANIES AND
7 PERSONALLY FIND THAT TO BE A VERY ATTRACTIVE MODEL, ONE
8 THAT SEEMS THAT MOST OF THE COMPANIES FIND ACCEPTABLE,
9 AND ONE THAT I THINK, JUST FROM MY OWN KIND OF GUT
10 REACTION TO, SEEMS QUITE FAIR IN TERMS OF THE ECONOMIC
11 BENEFIT THAT THE STATE COULD HOPE TO RECEIVE IN
12 EXCHANGE FOR PROVIDING SEED FUNDING TO COMPANIES FOR
13 THIS TYPE OF TECHNOLOGY.

14 SO I THOUGHT I MIGHT OPEN THE DIALOGUE BY
15 THROWING THAT OUT AS A SUGGESTION OF WHETHER THE JDRF'S
16 APPROACH TO THIS IS SOMETHING THAT WE MIGHT ADOPT. I
17 WOULD ALSO JUST ADD PARENTHETICALLY THAT I'M A BIG
18 BELIEVER OF NOT REINVENTING THE WHEEL IF YOU DON'T HAVE
19 TO. AND IF THERE'S ANOTHER MODEL WE CAN TURN TO THAT
20 SEEMS TO WORK, YOU KNOW, WHY NOT ADOPT IT?

21 CHAIRMAN PENHOET: WELL, FOR WHAT IT'S WORTH,
22 THE WELLCOME TRUST MODEL IS VERY SIMILAR TO THE JDRF
23 MODEL, AND THEY HAVE BEEN IN DISCUSSIONS WITH EACH
24 OTHER ABOUT THAT. SO THERE'S AT LEAST TWO AGENCIES
25 THAT HAVE PRIOR EXPERIENCE, HAVE VERY SIMILAR THINKING

1 ON THIS ISSUE.

2 I THINK THE CHARACTERISTICS THAT WE SET MIGHT
3 DEFINE THOSE, AS FAR AS WE CAN TELL, BY JUST GOING TO
4 THESE CONVERSATIONS. NO. 1, THAT THERE IS A RETURN IF
5 THE TECHNOLOGY IS SUCCESSFUL. NO. 2, THAT IT'S CAPPED,
6 WHICH I THINK DEALS WITH THIS CERTAINTY OR UNCERTAINTY
7 ISSUE THAT'S BEEN BROUGHT UP MANY TIMES. PEOPLE GO,
8 OKAY, WHEN THEY KNOW THE SIZE OF THE OBLIGATION, ETC.,
9 THAT IT HAS A FINITE LENGTH OF TIME ASSOCIATED WITH IT.
10 AND THAT THERE'S SOME KIND OF GOOD REPORTING THAT'S
11 ESSENTIAL TO UNDERSTANDING WHAT'S GOING ON, THAT THE
12 FUNDING AGENCY HAS THE ABILITY TO HAVE QUITE A BIT OF
13 TRANSPARENCY TO WHAT'S GOING ON. AND WE PROBABLY DON'T
14 HAVE THE STAFF TO DO A MANAGEMENT GROUP LIKE YOU
15 REFERRED TO WITH JDRF WITH TWO PEOPLE.

16 BY THE WAY, I WOULD MENTION THIS IS A
17 SIGNIFICANT ISSUE FOR US. CIRM AS AN AGENCY IS
18 DRAMATICALLY UNDERFUNDED GIVEN THE SCOPE OF ACTIVITIES
19 THAT IT WANTS TO UNDERTAKE IN THE FUTURE. JUST THE
20 LICENSING, WE'RE CAPPED AT 50 TOTAL PEOPLE. YOU KNOW,
21 A ROBUST LICENSING GROUP, A GRANT-MAKING GROUP, MAKING
22 A NUMBER OF DIFFERENT KINDS OF, LET'S CALL IT, THE
23 WHOLE FIELD OF GRANTS FOR THE MOMENT, THAT HAS TO BE
24 DONE DIRECTLY BY CIRM WOULD PROBABLY INVOLVE AT LEAST
25 10 PERCENT OF OUR TOTAL STAFF. IT'S A SIGNIFICANT

1 BURDEN, BUT I THINK ONE WE'VE ACCEPTED.

2 THOSE ARE SOME OF THE CHARACTERISTICS I
3 HEARD. MAYBE THE OTHERS, IF ANYBODY WANTS TO ADD TO
4 THAT LIST.

5 DR. REED: JOHN REED AGAIN. I ALSO NOTICED
6 THAT, I BELIEVE IN THAT JDRF CONFIGURATION, THE
7 COMPANIES THAT INVENT THE TECHNOLOGY OWN THE PATENTS,
8 BUT JDRF HAS CERTAIN RIGHTS TO ASSUME THE PATENTS IF
9 THE COMPANY ELECTS NOT TO PURSUE THEM AND PAY THE
10 ASSOCIATED COSTS. I FOUND THAT TO BE ATTRACTIVE AS
11 WELL BECAUSE THIS BUSINESS OF MANAGING PATENT STATES IS
12 VERY COMPLICATED AND COMES WITH SOME PRETTY ONEROUS
13 OBLIGATIONS TO DO IT WELL. AND I THINK, THEREFORE,
14 THERE'S A STRONG ARGUMENT TO BE MADE TO PUT THAT
15 MANAGEMENT OF PATENTING IN THE HANDS OF THE INVENTORS
16 AND ALLOW THEM TO OWN THE PATENTS, BUT OBVIOUSLY HAVE,
17 YOU KNOW, OTHER PROVISIONS THAT WOULD ALLOW US TO MARCH
18 IN IF THEY FAILED TO PURSUE THEM, OR POSSIBLY ALSO TO
19 MARCH IN IF THERE WERE ISSUES WHERE A PATENT HAD A
20 BLOCKING POSITION THAT WAS PREVENTING PROGRESS AND
21 WHERE ALL THE BEST EFFORTS TO GET THE PARTIES TO AGREE
22 TO MOVE THE FIELD FORWARD HAD FAILED, AND THEN WE MIGHT
23 HAVE CERTAIN RIGHTS TO ENFORCE CERTAIN USES OF
24 TECHNOLOGY ON A NONEXCLUSIVE BASIS.

25 BUT OVERALL I FOUND THE JDRF'S APPROACH TO

1 HOW THE PATENT WAS HANDLED ALSO TO BE QUITE ATTRACTIVE.

2 CHAIRMAN PENHOET: FRANCISCO, YOU HAVE ANY
3 GENERAL COMMENTS YOU'D LIKE TO SHARE AT THIS POINT?
4 YOU'VE ASKED A LOT OF QUESTIONS AS WE'VE GONE ALONG.

5 DR. PRIETO: NO. AND I'VE HEARD A LOT OF
6 INTERESTING ANSWERS. I THINK I WOULD LIKE TO HEAR A
7 LITTLE MORE FROM SOME OF THE OTHER GRANTOR
8 ORGANIZATIONS LIKE JDRF. I THINK I'M GOING TO TALK TO
9 THE PEOPLE AT MY OWN ASSOCIATION WHICH COOPERATES WITH
10 JDRF. I WONDER ABOUT THE GATES FOUNDATION, HOWARD
11 HUGHES, YOU KNOW, IF THEY'RE HANDLING THIS IN A SIMILAR
12 MANNER OR DIFFERENTLY.

13 CHAIRMAN PENHOET: MANY OF THE FOUNDATIONS IN
14 THE U.S. DON'T SUPPLY FUNDS TO FOR-PROFIT AGENCIES
15 BECAUSE OF THE TAX RULES RELATED TO HOW THEY DO THIS,
16 BUT WE DID STUDY A NUMBER OF THOSE MODELS WHEN WE WERE
17 DISCUSSING THE NOT-FOR-PROFIT PORTION.

18 UNIDENTIFIED SPEAKER: I'D LIKE TO MAKE ONE
19 COMMENT ABOUT THAT.

20 CHAIRMAN PENHOET: MAYBE WE'LL GET BACK TO
21 YOU AFTER WE'VE HAD THE TASK FORCE MEMBERS GIVEN AN
22 OPPORTUNITY. IN IRVINE, ANY GENERAL COMMENTS THAT
23 YOU'D LIKE TO MAKE FROM IRVINE?

24 DR. STEWARD: WELL, WE WERE WONDERING ABOUT
25 WHETHER IT WOULD BE WORTH HEARING FROM WARF.

1 MS. KING: DR. STEWARD, COULD YOU JUST STATE
2 YOUR NAME FOR THE TRANSCRIBER, PLEASE?

3 DR. STEWARD: YES. OS STEWARD.

4 CHAIRMAN PENHOET: WE'VE ASKED THEM TO COME
5 PRESENT TO OUR GROUP.

6 DR. STEWARD: WELL, IT WOULD BE INTERESTING
7 TO HEAR WHAT THEY HAD TO SAY ABOUT POSSIBLE
8 INTERACTIONS, LET'S CALL IT.

9 CHAIRMAN PENHOET: OKAY. WE'LL SEE WHETHER
10 THAT MAKES SENSE FROM SOME OTHER PERSPECTIVES, BUT IT
11 WOULD BE AN INTERESTING SESSION, FOR SURE. ANY OTHER
12 COMMENTS? SUSAN, DO YOU HAVE ANY?

13 DR. BRYANT: WELL, I AGREE WITH THAT. I
14 THINK IT WOULD BE TERRIFIC TO HEAR FROM WARF. BUT THE
15 OTHER THING MIGHT BE USEFUL TO HEAR FROM INTELLECTUAL
16 PROPERTY LAWYERS ON THIS ISSUE THAT SPECIALIZE IN STEM
17 CELLS MAYBE, SEE HOW THEY'RE DEALING WITH IT IN OTHER
18 PARTS OF THE COUNTRY.

19 CHAIRMAN PENHOET: IT'S A VERY DIFFICULT
20 ISSUE. AND I'M NOT HERE TO DEFEND WARF IN ANY WAY,
21 SHAPE, OR FORM; BUT, YOU KNOW, THERE IS A THREAD OF
22 THESE CONVERSATIONS. PATENTS ARE EXTREMELY IMPORTANT,
23 BUT EXCEPT THE ONES THEY DON'T LIKE. YOU KNOW, YOU
24 CAN'T HAVE IT BOTH WAYS. AND SO HOPEFULLY THE FIELD AS
25 A WHOLE CAN REACH SOME SENSIBLE ACCOMMODATION TO WARF

1 FOR THE PIONEERING WORK THAT THEY DID DO, BUT AT THE
2 SAME TIME NOT STOP THE ENTIRE FIELD FROM GOING FORWARD.
3 IT'S A HARD PROBLEM ACTUALLY.

4 FROM SAN CARLOS?

5 DR. LOVE: YEAH. I GUESS ON THIS END I JUST
6 WANTED TO REINFORCE --

7 CHAIRMAN PENHOET: WHO'S SPEAKING?

8 DR. LOVE: THIS IS TED LOVE.

9 CHAIRMAN PENHOET: THANK YOU.

10 DR. LOVE: -- TO REINFORCE WHAT JOHN SAID. I
11 THINK THE JDRF PROPOSAL REPRESENTS A SENSIBLE BALANCED
12 PROPOSAL THAT COULD WORK. AND I THINK MICHAEL GOLDBERG
13 HERE FROM THE D.C. SET HELPS SIMILARLY. SO I JUST
14 WANTED TO RE-ECHO THAT.

15 MR. GOLDBERG: YEAH. I AGREE WITH WHAT TED
16 JUST SAID. I ALSO WONDER WHETHER IN CONNECTION WITH
17 THE COMMENT THAT I THINK, MAYBE IT WAS SUE BRYANT MADE,
18 ABOUT INTELLECTUAL PROPERTY COUNSEL. I'D BE INTERESTED
19 PERHAPS IN SOMEBODY OR PERSONS WHO ARE EXPERTS, WHETHER
20 THEY'RE INTELLECTUAL PROPERTY LAWYERS, WHETHER THEY'RE
21 JUST LICENSING EXECUTIVES, TO KIND OF PUT INTO THE
22 RECORD THE RANGE OF ROYALTIES THAT ARE ORDINARY AND
23 REASONABLE AS PERCEIVED BY INDUSTRY FOR THE VARIOUS
24 SEGMENTS THAT WE MIGHT HAVE OCCASION TO APPLY THEM TO;
25 NAMELY, THERAPEUTICS, DIAGNOSTICS, AND REAGENTS.

1 BECAUSE I THINK THOSE THINGS ARE ALL VERY DIFFERENT,
2 AND I THINK THE WAY WE WERE ABLE TO DEAL WITH THE
3 ROYALTY ISSUE FOR THE NOT-PROFIT GRANTEES IS A LITTLE
4 MORE COMPLICATED THAN WE'RE GOING TO FIND IN DEALING
5 WITH THE DIFFERENT SETS OF POTENTIAL FOR-PROFIT
6 APPLICANTS THAT WE FACE. SO THAT'S FOR YOUR
7 CONSIDERATION.

8 CHAIRMAN PENHOET: AS A PERSON WHO'S ENTERED
9 INTO MANY DIFFERENT ROYALTY PAYING AGREEMENTS,
10 UNFORTUNATELY THEY SPAN A HUGE RANGE FROM THE LOWEST, I
11 THINK, I CAN REMEMBER EVER PAYING MYSELF AS AN
12 EXECUTIVE WAS ABOUT HALF A PERCENT, AND THE HIGHEST I
13 EVER GOT FROM A THIRD PARTY WAS CLOSE TO 40 PERCENT.
14 AND IT TOTALLY DEPENDS ON THE VALUE OF THE PATENT
15 YOU'RE TALKING ABOUT. IT'S VERY HARD TO HAVE A GENERIC
16 ANSWER TO THAT QUESTION BECAUSE SOME INVENTIONS ARE
17 EXTRAORDINARILY VALUABLE, COMPLETELY CHANGE THE FIELD,
18 ETC. AND OTHER ONES MAKE A TRIVIAL CONTRIBUTION TO
19 SOMETHING, NEVERTHELESS THEY'RE A PATENT. SO I THINK
20 IT'S VERY HARD TO FIND A GENERIC ANSWER TO THAT
21 QUESTION, BUT WE COULD AT LEAST EXPLORE WHAT THE RANGE
22 IS.

23 MR. GOLDBERG: I WOULD -- I AGREE WITH YOU,
24 ED, EXCEPT I THINK ONE OF THE THINGS THAT MAKES OUR
25 SITUATION A LITTLE EASIER, PERHAPS, IS THERE ISN'T

1 ANYTHING TO LICENSE AT THE TIME SOMEBODY IS ENTERING
2 INTO A GRANT RELATIONSHIP WITH CIRM. I MEAN THERE'S
3 JUST -- THEY'RE GOING TO DEVELOP -- WHAT COMES OUT OF
4 IT IS UNKNOWN. THEY DON'T KNOW WHETHER IT'S VALUABLE
5 OR NOT VALUABLE AS OPPOSED TO IF I KNOW THAT A
6 UNIVERSITY WITH A CIRM-DEVELOPED GRANT DEVELOPS
7 SOMETHING OF GREAT VALUE, IT'S VERY UNDERSTANDABLE THAT
8 ONE MIGHT BE WILLING TO GO TO THE HIGH END OF THE RANGE
9 BECAUSE THE INFORMATION IS KNOWN AS OPPOSED TO UNKNOWN.
10 SO...

11 THE REPORTER: I'M SORRY TO INTERRUPT. WHO
12 WAS JUST TALKING?

13 MR. GOLDBERG: TED LOVE.

14 MS. KING: MICHAEL GOLDBERG.

15 DR. LOVE: IT WAS A CHICO.

16 CHAIRMAN PENHOET: ANY MORE COMMENTS FROM SAN
17 CARLOS? FROM LOS ANGELES? FROM CHICO?

18 DR. WRIGHT: NO. I WOULD LIKE TO THANK THE
19 PRESENTERS AND THIS DISCUSSION BECAUSE IT REAFFIRMS HOW
20 COMPLEX THE ISSUE IS, BUT IT ALSO MAKES IT A LITTLE
21 MORE APPROACHABLE FOR THOSE OF US WHO HAVEN'T LIVED AND
22 BREATHED THIS.

23 MS. KING: THAT WAS JANET WRIGHT.

24 DR. WRIGHT: YES. SORRY.

25 MS. KING: WITH A W.

1 THE REPORTER: THANK YOU.

2 CHAIRMAN PENHOET: BUT SHE'S OFTEN RIGHT AS
3 WELL. YOU WERE UNUSUALLY QUIET TODAY, JANET.

4 DR. WRIGHT: I AM, AREN'T I?

5 CHAIRMAN PENHOET: WE DO HAVE SOME PUBLIC
6 COMMENTS NOW FROM SAN DIEGO.

7 MR. BAERGE: ED BAERGE FROM NOVOCELL. I WAS
8 JUST GOING TO MAKE A COMMENT WITH REGARD TO THE OTHER
9 INSTITUTIONS THAT SUPPORT BASIC RESEARCH AROUND THE
10 WORLD AND THE U.S. LIKE HOWARD HUGHES AND JDRF.
11 THERE'S ANOTHER INSTITUTION THAT SUPPORTS A LOT OF
12 RESEARCH HERE IN SAN DIEGO CALLED THE CLAGUE
13 FOUNDATION, AND THAT INSTITUTION ALSO HAS INVESTED IN
14 COMPANIES. THEY'VE INVESTED IN US TO DEVELOP STEM CELL
15 THERAPY FOR DIABETES. AND THEY TOOK AN EQUITY
16 INVESTMENT, SO THAT WAS JUST ONE THING THAT ALLAN
17 BROUGHT UP. I MEAN THAT'S ANOTHER WAY IN WHICH
18 TO -- THAT THE CIRM CAN THINK ABOUT THIS.

19 CHAIRMAN PENHOET: UNFORTUNATELY, I BELIEVE
20 THE STATE OF CALIFORNIA IS NOT ALLOWED TO HOLD EQUITY
21 IN COMPANIES UNFORTUNATELY.

22 MR. SIMPSON: I THINK YOU ARE CORRECT. WE'VE
23 BEEN LOOKING INTO THAT AS WELL. YOU MAY BE ABLE TO USE
24 WARRANTS. I'M NOT SURE.

25 CHAIRMAN PENHOET: AN EQUITY SURROGATE.

1 MR. SIMPSON: WE'RE LOOKING INTO THAT.

2 CHAIRMAN PENHOET: THANK YOU. ANOTHER
3 COMMENT FROM SAN DIEGO?

4 MS. LORING: THIS IS JEANNE LORING FROM THE
5 BURNHAM INSTITUTE. I JUST WANT TO PUT A PLUG IN FOR
6 SOME BACKGROUND INFORMATION. I RECENTLY HAD AN ARTICLE
7 PUBLISHED IN *SCIENCE*, THE MARCH 24TH ISSUE, I BELIEVE,
8 THAT IS EXACTLY TO THIS POINT. THAT IS, THE EFFECT OF
9 THE WARF PATENTS ON EMBRYONIC STEM CELL RESEARCH IN
10 BOTH THE ACADEMIC AND INDUSTRY AREA.

11 AND THE IDEA BEHIND RUNNING THIS ARTICLE WAS
12 TO EXPLAIN TO PEOPLE WHO WERE SCIENTISTS OR LAY PEOPLE
13 WHAT THE -- HOW THE PATENTS THAT WERE ISSUED, WHY IT
14 WAS ISSUED, WHAT THE OPTIONS ARE NOW. SO IT'S MARCH
15 24TH *SCIENCE*, THE POLICY FORUM.

16 DR. REED: JOHN REED. YOU MIGHT MENTION WHO
17 YOUR CO-AUTHOR WAS ON THAT JEANNE.

18 MS. LORING: OH, YES. I ALWAYS FORGET MY
19 CO-AUTHORS. MY CO-AUTHOR ON THAT WAS KATHERINE
20 CAMPBELL, WHO IS AN IP LAWYER, AND SO WE COLLABORATED
21 TO BE ABLE TO GET THE SCIENCE AND THE LAW STRAIGHT.

22 CHAIRMAN PENHOET: THANK YOU. SOME OF YOU
23 WHO DON'T SUBSCRIBE TO *SCIENCE*, WE WILL GET COPIES OF
24 THAT AND SEND IT TO ALL THE MEMBERS OF THE TASK FORCE.

25 MR. SIMPSON: JOHN SIMPSON, FOUNDATION FOR

1 TAXPAYER AND CONSUMERS RIGHTS. COUPLE OF QUICK THINGS.
2 FIRST OF ALL, I JUST WANT TO SAY HOW IMPRESSED I WAS
3 WITH THE DIVERSITY OF COMMENTS THAT YOU GOT FROM ACROSS
4 THE BOARD. AND I THINK THAT THE COMMITTEE SHOULD BE
5 COMMENDED FOR THAT APPROACH, AND IT WILL, I THINK,
6 INEVITABLY RESULT IN POLICY THAT IS FOR THE BEST PUBLIC
7 POLICY. I REALLY WANT TO COMMEND YOU FOR THAT.

8 THE ONE THING THAT I HEARD IN THE REQUESTS
9 FOR COMPARISON TO SOME OTHER FOUNDATIONS, I THINK
10 THAT'S USEFUL MODELS, BUT ONE OF THE THINGS THAT
11 STRIKES ME IS THAT THEY ARE ALL FOUNDATIONS. IT WOULD
12 BE INTERESTING TO TRY TO FIND OUT THE APPROACH OF
13 GOVERNMENT PUBLICLY FUNDED RESEARCH PERHAPS FROM OTHER
14 COUNTRIES. I'D BE INTERESTED IN LEARNING MORE ABOUT
15 HOW SINGAPORE HANDLES IT, HOW THE UK HANDLES IT. I'M
16 NOT SUGGESTING THAT YOU HAVE A FIELD TRIP NECESSARILY,
17 BUT THERE MIGHT BE USEFUL INFORMATION THERE.

18 AND FINALLY, I ALSO WANTED TO ADD MY TWO
19 CENTS WORTH ON THE ARTICLE THAT JEANNE LORING
20 CO-AUTHORED. IT'S A VERY HELPFUL, USEFUL ARTICLE.

21 THERE'S ANOTHER ONE THAT I WOULD ALSO COMMEND
22 TO EVERYONE'S ATTENTION, WHICH WAS WRITTEN BY JEAN
23 WASHBURN -- I'M SORRY -- JENNIFER WASHBURN. IT
24 APPEARED RECENTLY IN THE *LOS ANGELES TIMES*, ALSO TO THE
25 WARF PATENTS. THANK YOU VERY MUCH.

1 CHAIRMAN PENHOET: WE'VE BEEN EXPLORING
2 GETTING SOME FEEDBACK FROM OUR COUNTRIES. THE EU DOES
3 HAVE A PROGRAM OF FUNDING RESEARCH IN COMPANIES, AND
4 WE'VE BEEN TRYING TO TRACK DOWN WHO IT IS THAT'S
5 RESPONSIBLE FOR THAT PROGRAM NOW AND TRYING TO GET THEM
6 HERE OR AT LEAST ON THE PHONE FOR OUR NEXT MEETING.

7 MR. GOSWAMI: JOYDEEP GOSWAMI. JUST A QUICK
8 THOUGHT ON THE QUESTION OF ROYALTY THAT WAS BROUGHT UP.
9 YOU KNOW, I THINK THERE MIGHT BE SOME RANGE IN THERE.
10 I THINK YOU'RE ABSOLUTELY RIGHT. THE RANGES ARE PRETTY
11 BROAD. BUT ONE OF THE THINGS THAT SHOULD BE CONSIDERED
12 IN THIS PARTICULAR CASE IS HOW MUCH INVESTMENT OTHER
13 PARTIES MAKE, INCLUDING THE COMPANY THAT IS DEVELOPING
14 THIS, BECAUSE IT IS DIFFERENT FROM A LICENSE THAT
15 YOU'RE TALKING ABOUT. SO ANY POLICY THAT CIRM COMES UP
16 WITH SHOULD HAVE THAT CONSIDERATION, AND MAYBE THERE'S
17 SOME OPTIONS THAT THE COMPANY CAN REFER TO IN TERMS OF
18 THE ROYALTY RATE, WHICH IS DEPENDENT ON SOME SCALE TO
19 THE AMOUNT OF INVESTMENT THAT IS MADE BY OTHER PARTIES
20 OR BY THE COMPANY ITSELF.

21 MR. ROTH: DUANE ROTH AGAIN. ONE IMPORTANT
22 CONCEPT THAT I HEARD TODAY, AND I THINK THE COMMITTEE
23 SHOULD BEAR IN MIND WHEN THEY MAKE THEIR FINAL
24 RECOMMENDATION, IS FLEXIBILITY. I DON'T THINK YOU'RE
25 GOING TO FIND A POLICY THAT IS GOING TO BE A COOKIE

1 CUTTER THAT'S GOING TO WORK FOR YOU. I THINK IT'S
2 GOING TO HAVE TO BE VERY, VERY FLEXIBLE. AND,
3 THEREFORE, ANY CONSIDERATION SHOULD TAKE THAT INTO
4 CONSIDERATION.

5 THE SECOND THING IS REALLY A QUESTION. HAS
6 THE COMMITTEE HAD ANY ACCESS TO THE CONFERENCE
7 DISCUSSION THAT JUST TOOK PLACE RECENTLY?

8 CHAIRMAN PENHOET: ACCESS IN WHAT SENSE?

9 MR. ROTH: WELL, A COUPLE OF OUR PEOPLE --

10 CHAIRMAN PENHOET: PEOPLE PARTICIPATING IN
11 THAT MEETING --

12 MR. ROTH: BUT I BELIEVE THERE ARE
13 TRANSCRIPTS, AND I THINK THE WHOLE THING WAS
14 VIDEOTAPED. AND I WOULD HIGHLY RECOMMEND THAT GET PUT
15 INTO THIS RECORD SOMEHOW BECAUSE MANY OF THE THINGS I
16 HEARD DISCUSSED HERE WERE REALLY COVERED IN DETAIL,
17 INCLUDING THINGS THAT INVOLVE WARF AND OTHERS. AND IT
18 WOULD BE HELPFUL TO HAVE THAT. THANK YOU.

19 CHAIRMAN PENHOET: IS THERE PUBLIC COMMENT IN
20 IRVINE?

21 DR. BRYANT: NO.

22 CHAIRMAN PENHOET: SAN CARLOS?

23 UNIDENTIFIED SPEAKER: NO.

24 CHAIRMAN PENHOET: LOS ANGELES?

25 UNIDENTIFIED SPEAKER: NO.

1 CHAIRMAN PENHOET: JANET, YOU MUST BE BY
2 YOURSELF IN CHICO. WE HAVEN'T HEARD ANY PUBLIC COMMENT
3 FROM CHICO.

4 DR. WRIGHT: I DON'T HAVE TO REVEAL THAT
5 INFORMATION. NO PUBLIC COMMENTS.

6 CHAIRMAN PENHOET: OKAY. GOOD. WELL, JUST A
7 COUPLE OF REMINDERS. FIRST OF ALL, THE NONPROFIT --
8 OUR PROPOSAL FOR A NONPROFIT POLICY WAS FILED WITH THE
9 OFFICE OF ADMINISTRATIVE LAW EARLY THIS WEEK. AND IF
10 THEY MOVE IT ALONG, THE PROPOSED REGULATION SHOULD BE
11 PUBLICLY NOTICED ON THAT OAL WEBSITE, OAL IS THE OFFICE
12 OF ADMINISTRATIVE LAW, NEXT FRIDAY. THAT MEANS A WEEK
13 FROM TOMORROW, FORMALLY OPENING THE PUBLIC COMMENT
14 PERIOD FOR THE NONPROFIT POLICY. SO WE KNOW SOME OF
15 YOU WILL HAVE SOME COMMENTS TO MAKE.

16 MS. KING: AND IT WILL ALSO BE ON OUR WEBSITE
17 AT THAT TIME.

18 CHAIRMAN PENHOET: IT WILL ALSO BE ON THE
19 CIRM WEBSITE. BUT OUR PROPOSED POLICY HAS BEEN ON OUR
20 WEBSITE FOR SOME TIME.

21 MS. KING: CORRECT.

22 CHAIRMAN PENHOET: SO THIS IS A CHANCE FOR
23 ANOTHER BITE AT THE APPLE FOR ANYBODY WHO CHOOSES TO
24 PURSUE IT. FRIDAY A WEEK FROM OAL. AND I GUESS WITH
25 THAT, I THINK WE DON'T HAVE ANY MORE BUSINESS TO DO

1 TODAY. WE WILL SCHEDULE AT LEAST ONE MORE MEETING.
2 WE'VE GOT A LOT OF OTHER THINGS GOING ON IN THIS TIME,
3 SO IT MAY NOT BE FOR SEVERAL MONTHS, BUT WE WILL
4 CONTINUE TO DILIGENTLY PURSUE THIS POLICY.

5 MARY REMINDS ME THAT WE'RE TRYING FOR JULY.
6 WE'LL SEE. IT'S ALWAYS HARD TO PICK DURING VACATION
7 TIME. SO IF THERE ARE NO MORE COMMENTS, I'LL BRING
8 THIS MEETING TO A CLOSE. AND REALLY I THINK WE HAD
9 EXTRAORDINARILY THOUGHTFUL PRESENTATIONS ALL AROUND. I
10 THINK CERTAINLY THESE PRESENTATIONS IN ONE SENSE
11 EMPHASIZE THE DIFFERENCES IN VIEWS, BUT I THINK A LOT
12 OF COMMON ELEMENTS CAME OUT OF THIS DISCUSSION TODAY.
13 AND HOPEFULLY WE CAN GO FORWARD AND TRY TO DEVELOP A
14 CONSENSUS VIEW WHICH MAKES SENSE FOR EVERYONE. SO
15 THANK YOU ALL.

16 (APPLAUSE.)

17 (MEETING WAS THEN ADJOURNED AT LL:39
18 A.M.)

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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 IP TASK FORCE OF 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

UNIVERSITY OF CALIFORNIA SAN DIEGO
9500 GILMAN DRIVE
EUCALYPTUS POINT
LA JOLLA, CALIFORNIA
ON
APRIL 27, 2006

WAS HELD AS HEREIN APPEARS AND THAT THIS IS THE ORIGINAL TRANSCRIPT THEREOF AND THAT THE STATEMENTS THAT APPEAR IN THIS TRANSCRIPT WERE TRANSCRIBED BY ME FROM A DIGITAL RECORDING TO THE BEST OF MY ABILITY TO HEAR AND UNDERSTAND THE RECORDING. I ALSO CERTIFY THAT THIS TRANSCRIPT IS A TRUE AND ACCURATE RECORD OF THE PROCEEDING.

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