

John Crawford

*Interview conducted by
Matthew Shindell, Historian
July 8, 2008*

SAN DIEGO TECHNOLOGY ARCHIVE



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John Crawford



Mr. John E. Crawford serves as Financial Advisor of Cebix Incorporated. Mr. Crawford serves as Principal Accounting Officer of NovaCardia, Inc. Mr. Crawford co-founded Cabrellis Pharmaceuticals Corporation in 2006 and served as its Chief Financial and Administrative Officer. Mr. Crawford served as Business Development Consultant at Lithera, Inc. He served as the Chief Financial Officer of Phenomix Corporation since October 2007. Mr. Crawford served as Chief Financial Officer of Conforma Therapeutics Corporation since February 16, 2006. He served as Chief Financial Officer of NovaCardia, Inc. from January 2007 to September 2007. From 1997 to December 2005, Mr. Crawford was a Consultant to life sciences and technology firms. During this time, Mr. Crawford served on the Board of Directors of several private companies and was a Consultant to Archimedes Technology Group, LLC. He served as Founding President of San Diego-based Corvas International, Inc. and held various positions within it from 1987 to 1999, including that of Chief Financial Officer and served as the first full-time executive of International Genetic Engineering, Inc. He was an Executive Officer of early-stage life science and technology firms since 1981. Since 1998, he has been a Consultant, Executive Officer and Board Member for several venture-backed and privately funded technology firms. He has 25 years of extensive experience in the areas of biopharmaceuticals and medical devices. He served as a Director of Lipid Sciences Inc. since May 17, 2006. Mr. Crawford serves on the Board of Directors of the Clarity Foundation for ovarian cancer. Mr. Crawford holds a BS degree in Mathematical Sciences from Stanford University and an MBA in Finance from the University of Chicago Graduate School of Business.

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THE SAN DIEGO TECHNOLOGY ARCHIVE

INTERVIEWEE: John Crawford

INTERVIEWER: Matthew Shindell, Historian

DATE: July 8, 2008

LOCATION: San Diego, CA

1 **SHINDELL:** So, this is July 8, 2008. This is an interview with John Crawford. The
2 interviewer is Matthew Shindell. This is the San Diego Technology History Project.
3 So, you can start pretty much wherever you want, but why don't you, if you'd like,
4 start by telling us basically who you are, and what you do, in your own words. And,
5 and then you can go from there to how you got involved in this sort of thing.

6 **CRAWFORD:** Okay. John Crawford. I live here in San Diego and have been here for
7 twenty-one years. From an academic background, I majored in math and economics
8 at Stanford, and took an MBA in finance from the University of Chicago. I spent
9 about five years in a large commercial bank lending to high-leverage firms, mostly in
10 the media industry.

11 **SHINDELL:** Can I interrupt you for just one second? Could you also give us say the
12 years, say, the year that you finished your degrees?

13 **CRAWFORD:** I graduated 1976. Took the MBA in 1979. I was in the night program,
14 so I worked at the First National Bank of Chicago from 1976 to 1981. My work there,
15 as I said, was with high-leveraged lending to media firms, and that prompted my
16 interest in venture capital and owner-manager situations, as I saw some very
17 interesting jobs where managers had a lot of control over what they were doing, and
18 but had to live sometimes month to month, but certainly year to year, not knowing
19 whether they'd be financed and whether their companies would survive and so forth,
20 which appealed to me. And, of course, the rewards of doing that appealed to me as
21 well. So, I moved from there into venture capital. I came back to California at that
22 point, having grown up in California in 1981. I went to work for the venture capital

unit of Atlantic Richfield for just a short time. I thought I'd be there for a while and maybe get some board experience and then eventually transition to a small company. But, one of my first opportunities there was to get Atlantic Richfield involved in this new science they were calling "genetic engineering," both offensively and defensively. Atlantic Richfield, as a fine chemicals producer, thought there might be some opportunities for producing these fine chemicals by other means than just refining them from petroleum, and also was concerned that this new technology might take markets away from them, including the energy market. And so, I got them involved through some UCLA professors. We assembled a group of professors from the different aspects of genetic engineering and got to know these guys pretty well, and we started talking about maybe the thing to do was for them to form a firm, and maybe ARCO could invest in it, or sponsor research with it. I soon went to ARCO and said, "Gee, I'd kind of like to do that. I know it's a conflict but, you know, do you mind?" And, they were very gracious about it and so I became the first employee of that firm called International Genetic Engineering. That was in 1981. We went public in 1986. That firm just stuck with the strategy of doing contract research and didn't retain proprietary rights in its products. It was the first profitable firm in the industry, a very small industry, and one of the first ten firms in the industry. That was to its credit, but the Street was looking for firms that had – the "Street" meaning Wall Street – was looking for firms that had proprietary interests and could really hit a homerun. If their product succeeded, that they would own it and all of the returns that could be gained from it, not just taking say a royalty stream from sponsored research. So, about a year after we went public I left and formed a firm down here with a group of Scripps investigators, this was in 1987. It was called Corvas. I was there for eleven years, full-time, was founding president. I assembled the financing and the strategy. I got some partners and so forth. I was with the firm a total of eleven years. The first two years I was CEO, taking it through two rounds of financing. And then, as it grew pretty fast and furiously, and I was really more interested in being part of something big and successful, and necessarily leading it, and I was finding I was spending an awful lot of time learning on the job. So, I worked out an arrangement with the board where we could hire someone else in and then I continued as CFO. That firm's focus was cardiovascular disease, strokes, heart attacks, and so forth. It went public in 1992 and had an up and down history as so many of these firms had, but in particular had a very promising stroke drug that was sponsored by Pfizer. That didn't, in the end, work out. The data were not strong enough to take it into Phase III, which was very difficult for the firm. It

59 had a couple of other anticoagulant or antithrombotic agents that also struggled in
60 the clinic that were sponsored by Schering Plough. I left just for personal reasons in
61 1999, having worked very hard and wanting to spend more time with my three
62 young sons. So, I consulted after that for about seven years. I worked for a number of
63 firms, usually in an advisory capacity and usually as an acting CFO or actual CFO in
64 helping them raise funds. During that time, that consulting period, I also worked on
65 my own firm for a while, whose technology didn't pan out, as is often the case. There
66 are a lot of nonsuccesses. I won't quite call them failures. For every . . .

67 **SHINDELL:** What was the technology that you were working on there?

68 **CRAWFORD:** In this case it was stereo-specific phosphorothioate oligonucleotides.
69 They have to do with antisense technology. Antisense molecules have a stereo
70 feature to them so when they're synthesized they aren't exactly as the natural ones
71 would be, which do not have this stereo feature at each stage in their synthesis. The
72 idea was that these would be more accurate and I think it's probably still a promising
73 idea, just the chemistry behind it, which theoretically looked correct, did not pan
74 out in the lab. We were not able to reproduce that adequately in the lab. So, without
75 that there wasn't much to go on. You know, good concept, good idea, but didn't
76 execute. So, in 2006 I went to work for Conforma, which had a technology platform
77 on heat shock protein 90, or HSP90, which had some applications in cancer and
78 other areas. I joined to take them public or help them with what we call the
79 "liquidity event," or getting the initial investors to the point where they can sell their
80 stock, whether that be by selling the whole company or by going public and then
81 being able to eventually sell their stock into the public market. Fairly quickly we
82 developed a couple of acquisition proposals, and one of those panned out and we
83 sold the firm to Biogen Idec, which is another local firm, or it's actually based in
84 Boston but has a very large presence here in San Diego due to its own merger. There
85 was a piece of that technology the firm didn't want. It didn't fit their profile. Quite
86 sensibly they didn't want it. We placed that in a subsidiary and divvied it out to
87 our shareholders and then our same venture capital investors invested in us. We had
88 had some interest by other parties in that product and so we were actually able to
89 sell that company as well about six months later. The Biogen Idec transaction was
90 about a \$250 million transaction. The second firm called Cabrellis had about \$10
91 million invested in it, or exactly \$10 million invested in it and we sold it to Pharmion,
92 a much larger biotech firm, for \$104 million, about six months later. From there I

went to NovaCardia, which was headed by Randy Woods, who had also been the third CEO at Corvas, so someone that I had worked with before. And, same objective, to take the company public or otherwise find an exit for it. We filed for a public offering, cleared the SEC, and were ready to go, but had also been talking about acquisitions, and at that point Merck made a strong indication of wanting to acquire us. So, we sold that firm for \$350 million in Merck stock in 2007. Next, I joined this firm, Phenomix in 2008 and, or actually late 2007, October of 2007, and actually we've gone through two rounds with the SEC and would be prepared to go public but the public markets are closed to IPOs right now. So, we won't have that opportunity with this firm. So, we're looking at alternatives, as well, for exit. So, the career has been primarily CFO and financial roles, although I have been CEO a couple of times, and been acting CEO in a situation where I was on the board and there was difficulty with the management team. But, it has been very much from a financial perspective that I followed things.

SHINDELL: Okay. Let's go back for a second to your educational background. Your background is in mathematics and economics. Were you at all interested in the biosciences prior to discovering biotech and genetic engineering?

CRAWFORD: Actually not. My last class in biology was as a sophomore in high school. And, as I think so often happens your interest in a field is very much biased by your first experiences with it and how you learn about it. Teacher was poor, and my math teacher in high school was superb. And, I think that, as I look back, had a lot to do with the fields that I went into. Also at Stanford, the biology was the premed field. I wasn't interested in becoming a doctor, and you had to be a little wary about taking premed classes because the competition was just incredibly intense. So, not having any particular interest in biology, not knowing any better, it made more sense to take physics and, you know, geology and things like that.

SHINDELL: Uhm-hmm. So, you went into mathematics. Did you have a notion that you would be going on to economics when you first chose mathematics, or were you just sort of interested in math?

CRAWFORD: My interest was in business. And I figured I probably would go the MBA route. I was ambitious with respect to making money, not having grown up with very much. And, I was comfortable with math and figured, "Well, it doesn't hurt, you know. It's something I was relatively good at and . . ."

126 **SHINDELL:** So, you wouldn't say it was a love of pure mathematics? You were more
127 interested in applied mathematics from the beginning?

128 **CRAWFORD:** Yes. In fact, I kind of skipped over something. I say I was a math
129 major. It was actually called the Program in Mathematical Sciences. It was a four-
130 discipline field with theoretical math, statistics, operations research . . . and there
131 was another one in there. [Laugh] I forget. I guess computers, there were some
132 computer science elements to it, and it also required the core physics program. So, it
133 was a, like you say, an applied math program.

134 **SHINDELL:** And then with the first, the first group that you formed with the UCLA
135 bioscientists, they had presumably never had a background in math or economics?
136 Or, well, they had some math, obviously. But, you know, they had not done any
137 biotech ventures prior to that? They had not tried to create any products?

138 **CRAWFORD:** None of them had any experience. Yeah.

139 **SHINDELL:** And, you didn't have experience with biology. So, what was that like,
140 sort of crossing worlds with your background and theirs?

141 **CRAWFORD:** Well, first of all nobody had any experience. You know, Genentech
142 had only been formed a couple of years earlier. Amgen was in formation at the time.
143 So, nobody had experience. So, we weren't at a disadvantage in that regard. I like
144 science, in general, and this applied math thing had given me exposure to several
145 sciences. So, the biology appealed to me. I mentioned there were four, or six rather,
146 UCLA professors. They were willing teachers and they even had some video tapes of,
147 you know, basic gene splicing and stuff. So, I watched those, read a couple of simple
148 books. I attended our weekly science meetings where we all got together and talked
149 about different products they might attempt to develop. And, so I started to pick up
150 the science. I love science. I respect it. I'm not afraid of it. And, I'm conceptual
151 enough that I didn't really need to spend, you know, five years getting a PhD in the
152 field to at least understand the basics of it and what made sense. And oddly enough,
153 that turned out to be quite an advantage with respect to getting the initial clients
154 that we had in that firm. Because, the heads of research in large pharmaceutical
155 firms at that time were chemists. Chemistry was basically what the drug industry
156 was based on. And, they knew about this genetic engineering. They were skeptical of
157 it. They were afraid of it. They were somewhat, some of them I would have to say

were pretty cynical about it. That this was just pie-in-the-sky boloney. So, when I went out initially to speak with these guys about explaining our firm and that we were seeking research contracts, or could give advisory service for helping them get into the industry. These were some of the large pharmaceutical firms that we were approaching. I would go out alone, partly just because the other guys were teaching and busy, and kind of do the first cold call, and I could say to them, "Look, I'm in the same boat you are. I'm not a biologist or anything, but let me kind of explain from my perspective how I've come to see the science and where it can apply," and would go through just some of the basic aspects of, you know, the function of hormones, which they were familiar with, but how, how these biological actors might come into play on it and how we might derive chemically-based drugs from that eventually, and so forth. So, it was very non-threatening. Whereas, when they got in the room later with the scientists and MDs and the jargon started bouncing around, and, and they started getting hit with words that they didn't know, then you could just kind of see the tension levels kind of rising. But, I got the first level of comfort going and then by coaching my team at InGene to be aware of that and avoid the jargon, to understand that these guys were top notch. I mean, you know, the head of research at Merck is a pretty sharp guy. He just didn't know our field. Our objective was to get them comfortable with it and to see us as a resource for that comfort, and it worked. We got a lot good contracts doing that. This, the whole strategy was still a bit flawed in the sense that we didn't keep anything for ourselves, to speak of. But they, my lack of knowledge in the biology area and learning it from an outsider's perspective turned out to be an advantage.

SHINDELL: Did you – it sounds like you got a sense that there were two really distinct cultures there, the culture of say the academic biologist, the UCLA guys that you were working with, versus the culture of Merck heads of research and development, who also presumably have a background in science as well but are doing science in a much more, I don't know, corporate or industrial environment. Do you think that's the case, that there were two distinct cultures or . . .

CRAWFORD: Oh yes. And they, and they have changed quite a bit over the years out of necessity.

SHINDELL: How would you characterize that?

CRAWFORD: In a large company like that you'd have a chemistry division that

191 might be located in one town or one state, and they would go through and develop a
192 whole bunch of these compounds that met criteria that had been set, and then they
193 would be signed off and shipped to another group that might screen or further
194 characterize, or try to develop animal models and so forth on them. And, the
195 feedback and communication between those two groups was often minimal and was
196 astounding to us coming from biotech. And not so much at InGene, but at Corvas,
197 the second firm that I was involved in. We had, on the same floor, running down one
198 side the chemistry group and running down the other side the biology and in vivo
199 testing group. And, you'd have compounds synthesized with expected properties and
200 then run through various quality control screens to make sure we had what we
201 thought they were. Next the compounds went to in vivo pharmacology and to be
202 tested in cell cultures and very often in mice, very rapidly. In a period of a week or
203 two feedback coming back as to a compound was reacting and whether or not it was
204 inhibiting the clotting function, for example. And then the chemist would take, you
205 know, there might be ten compounds that had been sent over and two might have
206 interesting properties, and they'd study what were the properties about that that
207 might be conferring this activity. And then, they'd go through a cycle in chemistry
208 where they would synthesize more of that class and then they'd cycle back through
209 in vivo pharmacology. Well, that was not possible, not done, in the large companies
210 at that time. They were distinct entities and the people hardly knew each other and
211 they weren't communicating. And, that's one thing that I think had a lot to do with
212 the rise of biotechnology was that chemistry and biology were eating lunch together
213 in the lunch room, or running into other in the restroom, or were face to face with
214 each other. And while sometimes we had some pretty significant cultural disputes
215 and differences as to, to how things were done and who was really doing the
216 important work, [Laugh] and so forth and so on, as scientists, as they got working
217 together and began to realize that by working together they got some pretty
218 interesting results, that encouraged everybody and built some momentum. This was
219 taking place across the industry. It was this kind of marriage between sciences. In
220 other words, chemistry, to my mind, had run the gauntlet in refinements.
221 Improvements in the field were pretty marginal, you know, getting down to really
222 fine points. And biology, while it was an exploding field, wasn't interacting with
223 chemistry. What we began to call biotechnology, to my mind, was in large part a
224 merger of two disciplines that hadn't talked much before. And, if you look, at DNA
225 and the very basis for life, it's chemical molecules. It can be synthesized. So, that
226 was, to me, the revolution in this thing. I mean, yes, just the biology aspect was a

227 revolution, but the merging of those two fields had a lot to do with it.

228 **SHINDELL:** Uhm-hmm. Now having, you know, multiple disciplines under one roof
229 in a new biotech company or venture, it seems like that is sort of the key to this, this
230 sort of new innovative process, but what is that the product of? Is it just the product
231 of the fact that these are startup companies, and they're small, and so people are
232 necessarily close together and working towards a common goal of making a thing
233 profitable, or is it more a result of sort of a new philosophy of how to do things, of
234 how science and technology . . .

235 **CRAWFORD:** Well, I think it's a mix of all of those. There is, in small companies,
236 less focus on self-interest and more focus on team interest simply because there's a
237 good chance the company won't make it. But, if it does make it, you make out very
238 well. When you're working in a large firm you don't have impact as an individual,
239 unless you're CEO or something, on the outcome of the firm. So, there tends to be a
240 lot more of, "What about me?" in the big company process. In small companies,
241 people are willing to make remarkable work commitments, and work long hours,
242 with the sense that their work could really matter, that it's one of the only things
243 that the company is doing so that there's a lot of attention to it. It's not one of fifty
244 projects that might get scratched because of budget constraints. I mean, a couple of
245 projects are the company. Scientists had the sense that they were going to be able to
246 go forward with it, and own it, including presenting their work to the Board of
247 Directors and to Scientific Advisory Councils and so forth. You could give them an
248 awful lot of ownership in the work. And, you know, I think that's a lot of what all of
249 us want is a sense of ownership in what we're doing. The small company does that in
250 a much better way than, than a large company. And, that's not only the scientist but
251 the financial guys, and human resources people, and everything. You have impact.

252 **SHINDELL:** Uhm-hmm. It's interesting, yeah. In academia, also, I think you
253 wouldn't see, well you definitely wouldn't see financial people working side by side
254 with scientists and, you know, chemists working as closely with biologists. Was it
255 difficult for academic scientists, from your observations, to work in such close
256 quarters or was it something that they adapted to fairly well?

257 **CRAWFORD:** It didn't seem – there was just so much enthusiasm, the sense of
258 starting something that could be big that those other things kind of fell to the
259 wayside. And I think, too, just the individual attention that you got, your importance

260 to the firm outweighed any, "Oh, this is, you know, biology versus chemistry." There
261 was rivalry among the fields, rivalry among individuals and so forth, but to me not
262 the politicking to the extent that you have in the large firms, where the best interests
263 of the company weren't necessarily panning out.

264 **SHINDELL:** Uhm-hmm. Okay. So, your second company was Corvas, and that was
265 your first San Diego company?

266 **CRAWFORD:** That's right.

267 **SHINDELL:** What year was that again?

268 **CRAWFORD:** We started in 1981.

269 **SHINDELL:** Nineteen eighty-one? So, biotech was sort of just getting off the ground
270 here at that point, is that right? Hybritech was in 19 . . .

271 **CRAWFORD:** Nineteen eighty-seven. I'm sorry.

272 **SHINDELL:** Oh, sorry. So, that was a little bit later?

273 **CRAWFORD:** I get my, I – yeah.

274 **SHINDELL:** So, biotech was at least maybe . . .

275 **CRAWFORD:** I get my dates – I've been through almost a dozen firms. I guess it is a
276 dozen firms now.

277 **SHINDELL:** I guess if you take Hybritech . . .

278 **CRAWFORD:** That was 1987.

279 **SHINDELL:** If you take Hybritech as the beginning of biotech here, then you were
280 entering maybe within the first decade of biotech here? What, what was the biotech
281 scene like here or would you say there even was much of a biotech scene?

282 **CRAWFORD:** Well, Hybritech largely was the scene. There were others. But, in '87
283 there were about twenty firms that were fully funded venture capital firms. It was a
284 banner year for biotech down here. And, there were some good reasons for it,

including Hybritech. Hybritech was acquired by Eli Lilly in 1986 and Lilly began to make changes there. And, it was a very entrepreneurial culture that was suddenly in a big corporate environment. So, there was an awful lot of talent. I believe there was nearly a thousand people there, and there were several hundred who were college educated biologists, and chemists, and so forth. Mostly biologists. Hybritech didn't have much chemistry. Immunologists, that kind of stuff. And, Hybritech was sometimes cutting back or these people were just interested in doing it again with a younger firm. So, there was an enormous pool of talent down here. So, we had the academic institutions, UCSD, Scripps, and some of the smaller places. I don't know that the Burnham Institute had started at that point, the La Jolla Cancer Foundation was active. So, you had the academic component. You had a large labor force, especially at the sub PhD level. PhDs, experts in the field, you had to hire from outside, or might be available in one of the local institutions. But, for every one of those you hired you needed five or six other people, and that other talent was local. It was available. So, in terms of forming a firm down here it, it was a lot easier. There was a huge labor pool and a lot of academic talent to draw upon as advisors and so forth. There's also just an attitude or a culture and I think it's long been the case in California of going for it, of doing something. And finally, you had a quality environment here. San Diego's an attractive place to live. Housing prices are a little higher, but especially at that time all you had to do was go a little ways east and a family with three kids and so forth could get, you know, a decent single-family house and then have, you know, a mother or father, or maybe both, could have a very exciting career in the biotech industry with good compensation and that stock option, which just might be their key to riches. So, Hybritech had everything to do with the process down here. And, in fact, there was an investigator, I think at SDSU who did something called a begetting pattern, tying all of the early firms back to Hybritech one way or another. And, Corvas was also tie-able back to that because the second CEO, David Kabakoff had been vice president for R&D at Hybritech. And, when we looked to finding another CEO for Corvas, the first place you'd look. And, in fact, I spoke with David and worked with David for four days short of a year before the Board and I convinced him to come over to the company. And, of course, he's still a big figure in the industry here.

SHINDELL: So, it does, it does seem like the local talent is always moving from company to company and sort of, you know, splitting up in one place, meeting up in another. And, it also, in some ways the way that you've talked about it in this

interview, it seems as though the companies are sort of being built not necessarily to produce a product but in a way to be a product themselves to be sold. Is that unique or is that a fair characterization even?

CRAWFORD: I think it's, I think it's a trend. You can't deny that that's a trend, that nowadays, even earlier on in the '80s, the model measure of success was going public and being a successful company. And sometimes it was just going public. And, you know, then quite a number of those firms started doing poorly when they went public, so then you had to add on going public and being a successful public company. At first, when there was a buyout of a firm, even if it was at a good return to the investors, there was a bit of a stigma attached to that. Management hadn't guided the firm all the way, didn't have all the prestige or the cache, and that has slowly shifted to being a very respectable exit for companies. In many ways for investors it has become a preferred exit, because it's immediate in terms of cash return rather than having the additional risk of being a public company and getting a following, and product failure in the late stage, especially with the FDA's tightening of restrictions. And, even when a, when a company's gone public it may be two or three years, or even longer, before an institutional investor, a venture capitalist, can exit, can actually sell their shares. So, in more recent years acquisition has become a more acceptable alternative, and I would have to say in the last several years it's probably become a preferred alternative. Sarbanes-Oxley and the additional costs and burdens of being a public company have been a deterrent as well to that as a liquidity vehicle, or financial exit.

SHINDELL: And based on your own observations, do you think that this trend helps or hinders the actual eventual product development once these smaller companies are bought by bigger companies and maybe their research becomes a division of a larger company?

CRAWFORD: Oh, you know, if I really get kind of conceptual or theoretical about this, it's a shame that the bias for investors, and I've been a part of this too, is for later-stage programs that already in the clinic. They are "derisked" from the R&D standpoint, the research standpoint in particular. Products are advanced in the clinic from, say, early Phase I or pre-clinical, through maybe Phase II, which still is fairly modest in cost and then sold to large pharma. So rather than having a ten year development time frame, which makes it very hard for early investors to get a good return, you've got a three year development time frame. The victim in that is the

early-stage, creative, scientific developments, "platform technologies" as they call them. They're still there and some firms still are able to get funded with a basic concept, but there's fewer of them and the system, the FDA process, the demands of the investors behind the venture capital firms and so forth have shifted to more short-term thinking from long-term investment in a program.

SHINDELL: Do you think . . .

CRAWFORD: That's too bad. I mean, you know, the innovation isn't what it used to be.

SHINDELL: Uhm-hmm. And, do you think that that is a product of the trends of the industry itself, what sort of has set the precedent for being a profitable exit strategy, or is this a product of the market today, that the market maybe is different than, than earlier, people were more willing to take a more long-term risk back in the '70s and '80s? What would your take on that be?

CRAWFORD: Sure, I think the market is more short-term, and I think the perspective of the institutional investors, who are the ones who drive the behavior of biotech firms as stocks, have the perspective that rather than jump in an IPO of something conceptual and you don't see many of those anymore, "Why not wait? Fine, I may have to pay twice as much but let's wait until the technology has proven itself out better. I'll get in a little later, but if it's really promising I'll still get a good play on it, and I'll see that return in a year or two instead of sitting on it for five." As a consequence of that it's been harder and harder to start conceptual firms, platform-technology based firms because everybody's attitude is, "Well, let somebody else do that and I'll come in later when it, when it's been 'derisked,'" as they like to say.

SHINDELL: Uhm-hmm. Do you think that over time and maybe as a product of these shifts that we were just talking about that the relationship between the university and local biotech companies has changed at all? It seems like (Crawford: Yeah.) a lot of the early development . . .

CRAWFORD: There used to be quite a stigma for academics to be associated with a commercial firm. They were somehow compromising themselves or tainting themselves. They were money grubbers. The fellas that I was involved with in the

385 initial two firms suffered some by that, and especially if they were successful. I think
386 there was a little envy on the part of their colleagues when they saw these firms like
387 Corvas take off and get funded, and have pretty generous budgets, and be, you know,
388 getting a lot of attention. Somebody who hadn't had the courage and the foresight to
389 get involved with a firm and hadn't put the time and effort into it could very easily
390 say, "Ah, well, I'm a pure academic, you know. He's compromising himself. He's
391 more interested in his activity at his company than he is in being, you know, a true
392 leading scientist." All that kind of stuff came into the picture. And, egos involved
393 with these scientists are a very big part of it. I mean their intellect is on the line all
394 the time and it's very competitive, very cut-throat competitive in many cases in these
395 fields where things are happening fast. So yeah, there were some where there was
396 some true friction there. Some left academia all together, others bridged, others
397 backed away from the commercial firms and stuck with academia.

398 **SHINDELL:** Uhm-hmm. It seems like maybe because of the influx of biotech money
399 and other sectors of industry as well into the biosciences at the university, maybe the
400 structure of the biosciences in academia has changed as a result. From the biotech
401 perspective, from outside of the university, does that seem to be the case?

402 **CRAWFORD:** Yes. I would think so, and in fact there's even been legislative biases
403 toward that: That work sponsored by the NIH has to have vehicles for getting into
404 the commercial environment now, and academic institutions like Scripps have to be
405 able to show the NIH that they've got a licensing group and they do license things
406 out. Part of the purpose of the NIH in fostering innovation and research is to
407 develop our economy and so forth. If academia and the places where that money is
408 going don't encourage it or at least don't fight it, [Laugh] then the will of the tax
409 payers is thwarted. So, there's been pressure to do that. There's been this SBIR
410 program, Small Business Innovation Research Grant Program, that's been very well
411 funded by NIH and the National Science Foundation, and others, which provides for
412 kind of a bridge between academic work and the other. Why don't we turn this off
413 for a minute. I'll see if they need to use this room.

414 **SHINDELL:** Sure.

415 **CRAWFORD:** Go ahead and then I can interrupt it again.

416 **SHINDELL:** I see. All right. Okay. Were you in the middle of . . . [Laughter] I sort of

417 lost track of where we were. Sorry.

418 **CRAWFORD:** I think I had finished that point.

419 **SHINDELL:** Okay. Well, let me move on to a different question then. How about the
420 sort of legal side of biotech? That must have been new to you, the whole patenting of
421 biotechnologies, which was new to the sciences as well? What role do you think
422 patents have played in biotech or how important have they been?

423 **CRAWFORD:** Well, they've been essential to the industry. I mean, the core value of
424 what we have is the ability to have exclusive rights to it for a period, especially in
425 pharmaceuticals. So, without that there would have been nothing. If it was all public
426 domain, such as the academic work was, nobody would have funded it.

427 **SHINDELL:** And there were points at which it seemed like it might not be possible
428 to patent some of the, you know, biological products that were being produced. Was
429 the industry affected by those, that, maybe you'd say the fears that, that these things
430 would not be patentable, or did they sort of take steps to try to ensure that they
431 would be patentable?

432 **CRAWFORD:** Well, both. Especially with respect to biologicals and things like cell
433 lines and so forth. There were questions about what was patentable and what wasn't.
434 I think there were some technologies that weren't developed as much because of
435 patenting concerns. But, we seemed to have worked through that. There was enough
436 that came out of it that was patentable and we were able to do it.

437 **SHINDELL:** Uhm-hmm. And, let's see, where should we go from there? How about
438 geography, if you have sort of a sense of how the geography of San Diego, Biotech
439 Beach, what a lot of people called Biotech Beach, how has that affected the rise of
440 biotech here? Some say that part of the reason that biotech has been so successful
441 here is because of this sort of geographical clustering phenomenon, which is
442 probably different from what you experienced when you were further north here in
443 California? So, did that . . .

444 **CRAWFORD:** Yeah, we do have a critical mass of the academic institutions, the
445 labor supply that was created by Hybritech, and the winding down of all the
446 Hybritech activity so that we knew that there was a lot of labor down here. An
447 attractive place to live. That had a lot to do with it. You bet.

SHINDELL: When you founded Corvas though, did you locate it to be close to Hybritech.

CRAWFORD: Yes. I was living in L.A., InGene was based in Santa Monica. And, I had three young children myself and wanted to raise them in an attractive place. I wanted to stay in California, because I had other family here. I was committed at that point to the biotech industry and San Diego made a lot of sense. It was an attractive place to live. There was a lot more activity that was going to happen down here because of all of these elements. And so, it was my place of choice. I said, you know, I'm going to form a firm and it's going to be in San Diego. And, in fact, before Corvas was funded we had started to form the firm, but before it was funded my wife and I moved down here with our kids, bought a house, because I was going to stay here and do biotech here. I don't think I would have done that in any other community. Housing would have been too expensive, and I don't think there were many other communities where I could have that much confidence that I would be able to find employment in biotech.

SHINDELL: Uhm-hmm. And, were there any key individuals down here that maybe drew you down here or made you feel that this would be a good place for biotech?

CRAWFORD: There was a, there was a community here and organizations that were fostering it. The CONNECT Program had a lot to do with that. And . . .

SHINDELL: Several people have brought up Bill Otterson with that group. Yeah.

CRAWFORD: Obviously, yeah. He was just starting that program as well and I was in touch with him, told him what I had in mind, what I had done, and he did provide some contacts and provided, or offered, I don't think I actually used it, but offered space at those trailers on Torrey Pines Road that they'd given him [Laugh] some space in and all. And he had a lot to do with fostering the early stage of "Hey, let's all work together," and created social events where you met each other. I started a group with another CFO called the Association of Bioscience Financial Officers here. We just happened to meet each other at the swimming pool. We did swim workouts together for a while and then discovered we were both CFOs and both had the problem that there wasn't a financial environment here. So, for ourselves and for the controllers and all that worked for us there wasn't enough of a forum to kind of educate and learn about these things. And, we talked about getting that together.

We learned that the Bay Area was already starting a group. So, we contacted them and said, "Hey, let's go national and we'll be another chapter." And so, we formed the local Association of Bioscience Financial Officers on that basis, which today is a national group and has about ten chapters and membership of maybe a thousand people and has all the CFOs of the industry, and meets annually at national conventions. In fact, they're every June and this last one was here in San Diego. So, the community was very, very open to that and has continued to be through Biocom and other organizations.

SHINDELL: Now, you've mentioned that, in, that in 1987 when you started Corvas was, you said, "a banner year in biotech here in San Diego for several reasons." Were there any other sort of "banner years" or turning points in biotech here, or maybe moments that sort of indicated the coming of age of a strong biotech sector?

CRAWFORD: Well, the group, the '87 group that formed, in large measure, a large percentage of those, you know, fully-funded venture-capital-backed firms went public in '91 and early '92. So then, you had firms that in their first round of financing got two and three million and then done a Series B and gotten five million or more, then suddenly they were doing an IPO. And, back in that day a thirty or forty million dollar IPO was a pretty good sum. It was a fair amount of money then. So now you had, I don't know, somebody will have statistics on this, but maybe a dozen firms in San Diego that year that went public that had, you know, a mandate from their investors to grow substantially. So, of course, I'm going to see things from a financial perspective, but I think that was probably a banner year. After that, it was more just, you know, regular course of business and more and more firms coming out, and one firm or another having a spectacular growth profile. Invitrogen's been an interesting firm to watch, and in fact with that industry the supply industry to biotech wound up growing here. I think those folks came out, the initial ones came out of Stratagene, which was a very bright play by an individual who saw that, "Hey, there's going to be a need for product, and specialized product, custom products, in this industry," and created that firm.

SHINDELL: And who was that?

CRAWFORD: Joe Sorge. Ultimately, they like to say that spawned Invitrogen and ultimately it was more successful. But, both were very successful, although I don't think that particularly contributed to the growth here of the industry in general. It

513 was . . .

514 **SHINDELL:** It was more of a response to the growth, right?

515 **CRAWFORD:** Yeah. It was response to the growth, and maybe a bit of a response to
516 the nature of this community that you could do stuff like that. He was an
517 investigator at, at Scripps I believe, he's an MD. I don't recall exactly.

518 **SHINDELL:** So, could you give me a brief rundown, maybe, of what all is involved
519 in, in starting a biotech company, [Laugh] from your own experience here? What are
520 the most important steps to this process?

521 **CRAWFORD:** The people. With respect to the scientific side, the people with the
522 ideas and the caliber of research and respect of the community, and so forth, that
523 they can draw interest from the investor community. But then also, a pragmatic and,
524 a word that comes to mind, "mature" perspective on what can be achieved and how
525 to go about it. People who can do that are pretty rare. Next you need the
526 complementary people to them, the CEOs and other officers like CFOs, who can
527 meet the scientists' need and understand them at the same time that you meet needs
528 of investor and the venture capitalists and so forth. And then, have the vision thing.
529 That's very, very big in the industry as well. And, the vision isn't one that is set in
530 stone when the firm starts with the academic, but someone or some people, and this
531 can be on the Board, it can be the CEO, it can be another person inside the
532 company, or a mix of those who can track a changing landscape, can adjust the
533 firm's goals, and see well into the future. I was very fortunate at Corvas that the
534 founding scientist was a great visionary with respect to science and medicine.

535 **SHINDELL:** And who was that?

536 **CRAWFORD:** Tom Edgington.

537 **SHINDELL:** Tom Edgington.

538 **CRAWFORD:** He was willing to trust me to do what I could do. I had experience,
539 but not a lot, maybe about five years when we started it. That was kind of risky to, at
540 the age of thirty-two, trust me to being CEO of it, but nobody else had any more
541 experience because of the nascent industry. But still, to trust and take chances on
542 that and, you know, he was fairly assertive on opinions and so forth, and also

543 someone that you'd work with and discuss these things with, and all. There aren't
544 many people that have started a firm and taken it all the way through to commercial
545 realization, and that's true in the biotech industry as well as in other industries.
546 These firms usually go through transitions. You know, if you look at something like
547 Hewlett Packard, where it was literally started by Hewlett and Packard, and who
548 were with it for the entire forty years of their career, and remained chairman and
549 president during the whole thing. That's pretty rare in any industry. There have been
550 examples of it, and I think it's even rarer today because we're much more specialized
551 as a society and as a marketplace and the numbers of experts, the types of expertises
552 that you need in these firms today is, is overwhelming, and the leadership need
553 shifts as a company matures from research to clinical to marketing. And, one of the
554 reasons I think Corvas was successful and had, had a good run for so many years and
555 eventually reached a market cap of \$700 million was that we did transition through
556 three CEOs but I was there for the entire period of those transitions. So, there was a
557 certain amount of stability as well, and I was willing to, was able somehow to
558 transition these things, set myself aside a little bit and keep the organization stable
559 but allow it to go through these transitions. And, it worked. Ultimately the science
560 determines the product and, you know, the drugs that we put the most money in
561 just didn't quite perform in the clinic. Actually, the trials in that case were
562 successful, but Pfizer, the sponsor, and I didn't agree with them at the time but as
563 I've gotten away from it I agree with them, that they weren't compelling enough to
564 take on the multi-hundred million dollar Phase III trial. But, yeah, the transitions are
565 a big thing and a lot of companies are destroyed in the process of needing to make a
566 transition or failing to make a good transition.

567 **SHINDELL:** Overall do you think that this is a healthy environment for biotech?

568 **CRAWFORD:** San Diego?

569 **SHINDELL:** Yeah.

570 **CRAWFORD:** Oh yeah.

571 **SHINDELL:** Still today is healthy?

572 **CRAWFORD:** We've never had a large investment community down here. We do
573 have a decent presence, you know, Domain, and Enterprise, and ProQuest, and some

of the venture firms are down here, Sofinnova and Sanderling. But, the money is really in the Bay Area and in Boston, and there tend to be satellite offices here. They tend to be very good ones. I think the investors down here have done very well within those firms, but it's kind of sad that we've never kind of developed a strong local presence. You know, Sand Hill Road in Menlo Park is where most of the money comes from.

SHINDELL: Uhm-hmm. So, we're in our last ten minutes of the hour now. So, I'd like to ask you these sorts of questions that we save for the end that are sort of your own evaluations of your career and your time in biotech. So, some of these you may have already touched on, but maybe you can elaborate on a little bit more. So, first of all, what do you think was the most important change in biotech, in San Diego biotech during your time here?

CRAWFORD: Well, I don't know about change. I would say what distinguished us from others, and of course I've only been in this community, but one of the advantages we had was a strong sense of community and cooperation within the community. While we may have been competitive with each other, sometimes in the marketplace say two firms that were both addressing cardiovascular products, or we may have been competing for funds with each other, there was always a spirit of cooperation and not one of rivalry and competition. There's some of that, and I think it was healthy, but within the community there was always a sense of working together and making it better in the community. I have always been comfortable in calling people, fellow CFOs, and CEOs, and so forth, getting advice on things, and I see people still doing that regularly.

SHINDELL: And how, how, if that distinguishes San Diego, how has San Diego fostered that? How is it that that sort of atmosphere . . .

CRAWFORD: Come back to Bill Otterson, and he's been well recognized for that. He was a great advocate for it. He was taking Interferon, which was one of the early promising drugs, and, you know, he could stand up in front of an audience. I think he even used to carry around an empty bottle sometimes and said, "You know, I take this drug and I would have been dead, you know, six or eight years ago if it hadn't been for this drug," and he'd disappear off the scene once in a while when his cancer flared back up. And just say, "You know, this is what it is," and then he'd come back and say, you know, "Well, biotech saved me again." I mean, and he was a very, very

effective guy. He knew because of his illness that he couldn't be a hard hitting CEO again and it had to be something where he could come and go a little bit, depending on how he was doing, and I think he always, at least the first year or two, figured that he only had a year or two left and was going to make the most of it. But, golly, how can you, and he was a charming guy, how can you turn down a, you know, a personality like that? How can you be selfish when you see someone doing that? And, there were other, a lot of other leaders in the community. I mean, when Ted Greene left Hybritech he formed a VC firm and fostered six or eight firms, including Amylin. And, another one, Pixis that was very successful and bought out very early on. So, there were, you know several leaders that had the same perspective. Ted Greene, while he had no interest in any of the firms that I was involved in, financial interest, was a resource to me. I could give him a call and say, "Hey, you know, I've got this issue. What would you do?" It might only be ten minutes but, you know, I'd get a response on that. Jim Bergman, one of the early venture capitalists in the industry at Enterprise Partners decided not to invest in Corvas, but told me he'd be happy to talk with me and, counsel me if I thought I would need it. And, I said, "You know, I'd like that," and I did call him a few times and he did respond, just because he's a nice guy who wanted to see the industry here succeed. And, there are lots of examples of that in the community. I do it too, for that matter.

SHINDELL: Oh. Let's see. I think that you've just told me then what made Biotech Beach successful. Is that what you would . . .

CRAWFORD: Yeah. I would say. Yeah.

SHINDELL: Yeah. And, have there been any, any major events here or, you know, any forces here that have threatened the success of this sector? What would be the major threats that you've witnessed?

CRAWFORD: I think the FDA has been a challenge and continues to be a challenge. Their rules aren't clear a lot of the time. For Big Pharma that's a problem but it's not nearly the problem it is for us because we bet on one product and if we don't have a clear path as to where we're going to go with it, if you go through a process of a year or two of clinical trials, you do your best to construct them as you think they should be, then you discuss them with the FDA and then they come back and say, "Oh gee, you know, we really wish you'd done this instead." That's pretty tough and that happens with a lot of firms. It's a political process and I can be too harsh. I mean

640 you're messing with peoples' lives with pharmaceutical development so you do have
641 to be careful and all. But, I feel a lot of decisions that are taken by the FDA have to
642 do with trends at the time, or the biases of the individual serving on advisory
643 committees, and so forth, and it makes it pretty tough. Tougher than you'd like. The
644 marketplace is also fickle, but somehow it makes it tougher when it's individuals,
645 influential individuals on the panels that seem inconsistent. It's different than when
646 a product is introduced and the market doesn't accept it. You know, like pulmonary
647 delivery of insulin. Everybody thought that was going to be great and folks got it out
648 there and spent a lot of money on the products and all and it just didn't work. It
649 surprised the heck out of me, a lot of people, and a lot of big firms. But, that's
650 something that is difficult to accept but easier to accept than a regulatory or
651 administrative process that could have told you that yesterday, kind of thing.

652 **SHINDELL:** Has that been a problem in your career? Has this happened to you?

653 **CRAWFORD:** In smaller doses. Yeah. But, I've had enough colleagues in companies
654 where it's destroyed the company. It may not be readily identifiable as that, but as
655 you look at it, you know, what turned the company was an additional trial or
656 additional steps that had to be taken that were not expected. It wasn't a failure to
657 foresee. It was, you know, some external factor that, that emerged, that delayed
658 something and then caused them to get dilutive financing and maybe caused
659 management changes and began an unwinding process.

660 **SHINDELL:** Uhm-hmm. Okay. I, is there anything about San Diego biotech, the
661 sector here, or the cluster, that you would have liked to have changed, that you
662 thought wasn't working properly but could have been better?

663 **CRAWFORD:** Gosh, nothing comes to mind, at this point. There was a period when
664 the press was very negative about the industry, and that did change eventually but
665 there were a couple of reporters at the major newspaper that were very cynical.
666 Reporting in generals tends to be negative. You focus on the negative. But, gee,
667 there's been a wonderful development at a firm that, you know, was very
668 newsworthy and very positive and there would be a "however" and that "however" -
669 the negative little thing about it - would be drummed up and that would become the
670 title. That occurred in the '87 to '89 and '90 period where it was very, very hard to
671 see that kind of publicity for an industry that, I felt, deep in my heart was going to be
672 very important to San Diego and not being embraced. I think, if anything, that may

673 have unified the industry a little bit more and helped these organizations like BIO
674 and so forth, and Biocom, and CONNECT, get a stronger foothold here, and
675 eventually the press coverage changed. That came around and . . .

676 **SHINDELL:** Was that just the idea that, that they had to form a group to sort of
677 promote their own image, their own interests? Or . . .

678 **CRAWFORD:** Well, it was amazing to me how much influence a couple of writers
679 had. And, when they were no longer writing on the industry and there was more
680 positive stuff coming out then it, that felt better. But there was a particular period
681 there of a couple of years where I think it was discouraging for all of us to see, to feel
682 like we were being blasted in the papers, when in fact we were making a potentially
683 great contribution to the economy here.

684 **SHINDELL:** Uhm-hmm. Did you experience any tension, yourself, between pursuing
685 your own intellectual interests and the goals of say the, the people who were funding
686 the companies or the projects that you were helping to develop?

687 **CRAWFORD:** Uhm . . .

688 **SHINDELL:** Or, were you always at a level sort of high enough where you could sort
689 of escape?

690 **CRAWFORD:** Well, I was always at an officer level here. I kind of started the career,
691 my career at an officer level, because, you know, like I said nobody knew what they
692 were doing anyway. So, I might have been a little insulated from that. I mean, I did
693 experience those kinds of things, but I always had access to the Board room, direct
694 access to the Board. And so, I could at least express my opinion, eye-to-eye, with
695 either the management team I was working with or the Board of Directors. I might
696 not get my way, but there's value to knowing you've been heard, and having the
697 opportunity to speak your mind.

698 **SHINDELL:** All right. And how, personally, did your experience with Biotech and
699 with San Diego biotech in particular, change the course of your life, or affect the
700 course of your life, I guess? It's hard to know how it might change it.

701 **CRAWFORD:** Oh, well, it's been my career and you know a very fulfilling one for
702 me. I liked very much being in small environments. I grew up in a small town, and I

knew everybody, and worked in a large company where it didn't feel very personal. These firms are personal, for better and for worse, because sometimes you don't get along with people, or don't agree with them. But you're in there close with them and I think that was very, very key to me, this personal feel, this family feel. I don't know whether I answered the question well, or stayed on topic with it. But . . .

SHINDELL: That seems like a good answer.

CRAWFORD: That was, that was really key to me. Long hours, a lot of travel sometimes, a lot of pressure on my family. My wife has stood by me for twenty-eight years, who did not grow up in this kind of environment. When I was going to quit working for a major oil firm to go to work for what she called a "stack of resumes," [Laugh] namely the UCLA professors, it was, "I'll support you, but I'm skeptical about this." [Laughter] And, you know, especially with kids, and all it's a tough career choice. My wife's attitude was, "Well, if I have to I'll go back and get a job. I don't want to do that." And, one of the reasons that I did the consulting for many years had to do with wanting to spend time with my own kids and be away from the pressures of it as well. While my three sons were in high school I only worked part-time as a consultant. So to make these things work it takes an awful lot of commitment, and I will say sometimes people, you know, naturally they resent when a firm is successful and you make a lot of money on it and all of that, [Laugh] and sometimes they don't appreciate how much you've been through, how much risk there's been, how many times you nearly failed. You know how long and how many hours you've worked. Weekends, and you're kind of always on-call. And, these darn BlackBerries have made that [Laugh] even more the case.

SHINDELL: Uhm-hmm. Well, since it is, as you say, so, so personal compared to other industries, I would imagine that it has the potential to affect you personally a lot more than another career might have, that a lot of personal development occurs because of your work?

CRAWFORD: Oh, absolutely. Yeah.

SHINDELL: That, you know, you might not be getting otherwise? Or, you know, do you, do you find it – maybe this is too personal of a question. It's not really on my list. But, because of that personal nature do you find it difficult to not take your work home with you or to not make a division between say your, your home life and

735 your work life?

736 **CRAWFORD:** Well, I do take my work home with me. Although, I tend, especially
737 when I had kids, to come back in and work at the office and it was often just ten
738 minutes from home, rather than take it home. Because, at least when I was home I
739 wanted really be there. And, I wasn't good at, you know, "Go away. I'm working." I
740 just couldn't do that. I would, instead, spend some time home. And, my wife always
741 insisted that I come home for dinner and have some time with the kids in the
742 evening, even if it meant I needed to go back. I'm also a morning person, so it's not
743 uncommon to get in at six or so to kind of make up that difference. And, the same
744 thing Saturday morning, you know. I might come in for a few hours, if I really need
745 to catch up on stuff. But, it is a part of my life and when the kids had moved on and
746 we'd been successful, my wife and I sat down and said, "Okay, what are we going to
747 do here? We like to work. This is a fun time in our careers." She's also in the
748 industry, in finance. She works with very early stage firms, developing their
749 infrastructure, and doing all the non-science, nonclinical stuff in a firm, from
750 facilities to accounting to HR and so forth. And, we both decided, "Hey, we want to
751 do this." We take the weekends off and do a lot of stuff. But, it's very much a part of
752 our lives and we like it. You know, it is fun. It's very fulfilling and it's nice, now that
753 our kids are grown up that we don't have that competition, that kind of angst of
754 moments spent at work are moments not spent with the children. And, it's a whole
755 lot easier when it's just a married couple [Laughter] as opposed to a family of five
756 that you're kind of juggling.

757 **SHINDELL:** Uhm-hmm. So, this is sort of in the way of wrapping things up, but is
758 there any question that I should have asked you, or anything, you know, you really
759 would like to talk about before we really wrap things up?

760 **CRAWFORD:** Well I think, you know, for me I'm proud to have been a part of the
761 community and build things, and I'm just proud that the spirit of this community
762 has been so cooperative in the process. That fit very much with my personality and
763 desires, and it's a neat thing when it happens. We're competitive with each other in a
764 lot of ways, but we're so supportive, and this is across all of these organizations. The
765 networking that takes place and the spirit of helping is terrific. And that, to me, is
766 one of the pleasures of being in the community. It's a small community in the sense
767 of number of people and you can be here for a couple of years and know most of the
768 key people in the biotech community. Very open.

769 **SHINDELL:** All right. And then, one last question. Is there anyone you would
770 recommend that we interview for this project that might not be on our list?

771
772 **CRAWFORD:** Well, I don't know who's on your list. I mean, the natural thing to do,
773 I think, is to hit the highlights and the guys that have been prominent in the
774 community. I mean, I'm sure you've talked with Ted Greene. He's kind of one of the
775 granddaddies, if you will. And then, you know, some people like David Hale have
776 been very influential. I think a lot of Randy Woods. I think he'd be a good guy to
777 speak to. He's CEO of Sequel Pharmaceuticals now, which is a spin-out from
778 NovaCardia. I was surprised a bit that I was selected. I mean, I've been in the
779 community for a long time, but I am not a CEO and haven't been one for quite a
780 while. And I, you know, I commend you for not necessarily for talking to me but for
781 guys like me, getting perspectives of other than CEOs. I think some Scripps
782 investigators might be interesting, or UCSD investigators that have been involved in
783 the process and getting that academic perspective would be an interesting exercise
784 to do. You know, some of the lawyers in town have been very influential in the
785 process. Wain Fishburn at Cooley Godward comes to mind as a guy who's been
786 involved with many, many firms through the years. But gee, Rich Mejia, who just
787 retired as head of the local office at Ernst & Young for accounting, will have a lot of
788 perspective on the industry. Some of those folks really, really key to the industry,
789 were able to communicate with each other sometimes through attorneys and
790 accountants, you know. "Who else has seen this situation?" And, they'll know and
791 they may have to call them first and say, "You know, I want to tell this guy this, or he
792 wants to call you," and they'll say, "Fine, have him call." In all they've, they've been
793 good communication devices for us. I don't know what your list is, but I think some
794 of the service providers would be important.

795 **SHINDELL:** Okay. All right. Well, if you don't have any, any other last statements --

796 **CRAWFORD:** No. That's fine.

797 **SHINDELL:** -- then we can conclude it. Thank you very much.

798 **CRAWFORD:** You're welcome.

END INTERVIEW

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The San Diego Technology Archive (SDTA), an initiative of the UC San Diego Library, documents the history, formation, and evolution of the companies that formed the San Diego region's high-tech cluster, beginning in 1965. The SDTA captures the vision, strategic thinking, and recollections of key technology and business founders, entrepreneurs, academics, venture capitalists, early employees, and service providers, many of whom figured prominently in the development of San Diego's dynamic technology cluster. As these individuals articulate and comment on their contributions, innovations, and entrepreneurial trajectories, a rich living history emerges about the extraordinarily synergistic academic and commercial collaborations that distinguish the San Diego technology community.