

1 Dick Schneider #1 7/9/97

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6 Q: You received a PhD from the University of Wisconsin in 1966. When you were doing

7 that, did you have in mind a typical academic career path, or did you intend to something

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No, I was clearly on an academic path. I did my PhD in about three years, with no stop,

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even for a master's, and I did a postdoc at Wisconsin for a couple of months after I

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finished, because it was early, and then I accepted a postdoctoral position at MIT. So, I

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was clearly going in that direction, going the academic route.

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Q: And then what made you veer toward industry? Did some kind of opportunity pop

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More than I could ever imagine. This was in 1966, probably before you were born, but

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certainly a long time ago. At that time, there was a tremendous shortage of academic and

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PhD level trained scientists in the United States, and the number of jobs in industry was

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just overwhelming. I thought it would be kind of fun to just cast my net, to put my hand

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up and talk to some people, and I talked to ten companies and got ten offers. The other

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reason I changed my mind was that academic research was beginning to undergo a lot of

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difficulty getting adequate funding. We could just begin to see the tip of the berg, the

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size of the berg, however, wasn't known, but it's turned out to be monstrous. And as a

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result, some really high quality potential academic guys were turning toward industry that

33 overall brought the level of industrial science up to a very high level. Industry was then
34 allowing people to publish, allowing people to travel and do good science, at the highest
35 level. And I could see that, with financing being difficult and with the high quality of
36 research being done in industry that the number of opportunities was far greater, and the
37 last was that I was extremely and was always very interested in the application of science
38 to business. I didn't realize what that meant at the time, but when I started interviewing
39 for some of these industrial positions, it didn't take me long to figure out that we were in
40 harmony, more so than I was with academic colleagues. So, much to the chagrin of a
41 number of people at that time...

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45 Q: At Wisconsin?

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50 Both at Wisconsin and MIT. I decided to take a position. Now, in addition to that, I
51 would tell you that I had a very unusual circumstance. I did finally accept a position at a
52 large pharmaceutical company called Sandoz, New Jersey. Then I read an article that
53 appeared in Chemical & Engineering News, that we all got at that time, and they were
54 talking about a new company, a new group of people, starting a company in California.
55 I'm from California. And it was in an area that I was interested in. And even though I'd
56 already accepted this job, I hadn't reported to the job, but I'd accepted it, I decided to
57 write a letter to the people that had started it. And they invited me to come to California
58 and visit them on my next trip, and I did. To make a long story short, I ended up
59 accepting their offer to start a company from scratch with two other people. I was the
60 third employee in this company. And I had to go back to Sandoz, this big, famous, strong

61 company, and tell them that I wasn't coming. So, you can imagine the consternation.
62 First, I wasn't going to be an academic, and second, I wasn't going to go the company
63 about which they finally said, 'Yeah, that would be a good one to go to.' I was going to
64 go start one. So, to make a long story short, I didn't do what they thought I was going to
65 do.

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69 Q: This was Syva?
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79 Q: Did you perceive that as a risky move at the time, to go from Sandoz to a start-up?
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84 Well, remember that I never went to Sandoz. Even though I'd accepted the position, I
85 never reported to work. Did I consider that risky? Knowing what I know now, I consider
86 that insane, but knowing what I knew then, it seemed like an opportunity. I also felt that
87 once I got going, I remember that Syntex and Varian were the two financing founders of
88 this company, that was before there was any venture capital, and I figured, 'Man, if I do a
89 really good job at Syva, somebody at Syntex is going to see that.' I'm a chemist and they
90 were a chemistry company, and I was only twenty-six years old, twenty seven. I thought,
91 'Man, if I was ever going to take that kind of a risk,' of course, I didn't realize the
92 magnitude of the risk at the time, but that was the time to do it. And I never looked back.
93 It was the best thing I ever did.

94 Q: And when you arrived there, what kind of work did start doing?
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99 Well, that's a long, long time ago, over thirty years now. I was a lab scientist. I mean,
100 there were only three of us, that's pretty incredible. We were managed by the senior
101 managers of Syntex and Varian, the chairmen of their boards, and the Presidents of their
102 operating divisions were on our board at Syva, and I just had opportunities to interact
103 with Nobel-quality people all the time, at Stanford, at Syntex, and at Varian, and I started
104 working on some pretty esoteric projects. The money that was promised us from the two
105 companies was designed to last us about four years, but as things would have it, young
106 scientists being somewhat aggressive, trying to do too many things, we used the money
107 up in three. At the end of three years, we didn't have a product and the economy had
108 changed dramatically by 1970 and neither Syntex nor Varian had the extra cash to
109 support us, so it looked like the lights were going out. They didn't. Something happened
110 that caused us to keep them on.

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114 Q: And that was?

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119 Well, at that time, unfortunately the United States had a very massive involvement in
120 Southeast Asia. 500,000 men and women were over there for reasons that we don't have
121 to discuss, because everybody knows the history, but while they were there, they were
122 being exposed to some pretty noxious agents, namely drugs of all kinds, and there was
123 almost an hysteria in this country about bringing drug addicts back to the United States.
124 They, meaning the government, said, 'Look, we've just got to test all of these people.

125 We've got to know what we're going to get into when we bring them back.' One of our
126 scientific advisory board members, actually two of them, were involved in drugs of abuse
127 and were very concerned about this issue, and made a suggestion to us as we were about
128 running out money. They said, 'Look, you guys are so bright, you're working in these
129 very esoteric areas, maybe you could figure out a way to determine whether there's an
130 abused drug, any of twenty, in somebody's urine, and do it quickly.' Because the only
131 way that had been available to science in general at that time was a very labor intensive,
132 very costly method of either thin-layer chromatography or high-pressure liquid
133 chromatography, and imagine extracting 500,000 urine samples, shipping all that
134 chloroform, it weighs a ton as it is, over there, it was just totally impractical. And to
135 make a long story short, we came up with a method that would take one drop of urine,
136 could test for twelve different drugs, took a minute to do it, and require almost nothing,
137 just mix it with a reagent that we had developed and put it in a special instrument that we
138 had developed. And almost overnight, Syva went from as close to the brink of extinction
139 as you could get, to an operating company with sales and shipments, and people in Asia,
140 and airplanes, and we had a massive issue. And then, when these guys came home, we
141 developed some more assays that became useful, and were very generally useful, in
142 prison systems and all hospital emergency rooms. And the Syva broadened into
143 therapeutic assays in blood, serum, and others, for drugs that were being used
144 therapeutically to treat epilepsy, asthma, cardiac disease, what have you. And those
145 assays were extremely precise, very quantitative, and are used today to help physicians
146 determine the correct dosage of drug that an individual should be taking, a child or an
147 adult. And again, to make a very long story short, you know, the company became a

148 \$250 million a year, very profitable, wholly owned subsidiary of Syntex. By 1977, it was
149 already well on its way. I left in 1983, and I've been gone a long time, but that was a
150 very, very successful enterprise.

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154 Q: And when you had the first product, there was an immediately an explosion of growth,
155 you had to scale up to produce this, right? Was it at that point that you sort of
156 transitioned into management, away from the lab bench to other sorts of functions?

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161 Well, it was probably happening during all of that time. I was the guy who was leading
162 the group that was developing these products, and we had more to do than we could do,
163 and none of us knew anything about product development, and nothing about medicine,
164 at that time. You know, we were just scratching it out. We were young kids, basically.
165 And talk about opportunity, it was overwhelming. We had to learn quality assurance, we
166 had to learn manufacturing, we had to build a plant, we had to build instruments for these
167 products, we had to build a sale force, and eventually, we had 1,100 people in that
168 company. It became a very, very major enterprise. And being in the right place at the
169 right time, you know, good luck is being prepared for an opportunity, but nonetheless,
170 you have to have your eyes and ears open.

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175 Q: When did David Kabakoff come to Syva?

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180 Well, David, sure I hired him. I remember very well, I wish I could tell you the year. I
181 think it was around 1979. I may be off by a little bit, maybe '78. He was at Baxter down
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183 here in Southern California. I hired him and he became the assistant director of
184 development, and was just invaluable to us. We became very, very good friends. So,
185 anyway, he played an important role in it.

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189 Q: OK, let's see. You stayed at Syva until 1983, and then went to Liposome? What
190 made you decide then to leave Syva and do this other thing?

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194 You know, that's kind of a complicated story. It probably actually begins in 1979, when
195 Syntex sent me to the Advanced Management Program at Stanford Business School for
196 the summer. I left the company and lived at Stanford and went to business school, full-
197 time, seven days a week. I loved it. I was learning formally what I should have been
198 doing, you know, the years before. When I came back, I assumed my old responsibilities,
199 plus I became general manager of a new instrument company that we were starting. So, I
200 really had an opportunity, again, to start something new. It was a wholly owned
201 subsidiary of the company, we were at a \$20 million sales rate, with one customer,
202 internal. Just overnight, we were building instruments of all kind. During the next year
203 or so, they asked me to help start three other divisions, which we did, all of which
204 became reasonably successful, and I realized that what I really liked to do more than
205 anything was to start new things. I was not a very good long-distance runner, but I was a
206 pretty good sprinter. Running large organizations just didn't give me much of a thrill.
207 Sitting in meetings slows me down. I didn't care for that. So, that's really where a lot of
208 the thinking started about leaving the company, because it was just very big. I was just
209 feeling that there were other ways that I could leverage my time. There were other

210 complications at that time, 1981-82. Genentech had just appeared on the scene, and went
211 public in one of the most successful public offerings ever. In 1981, it opened at twenty-
212 five dollars a share and closed at eighty-one. Something clearly was happening in the
213 biology area, and I wanted to be part of it. You know, Cetus had started and then Chiron
214 and Biogen. In 1981-82, Ted Greene, who as you know, is a very prominent member of
215 the San Diego community, and Brook Byers came to see me and asked me to become the
216 VP of R&D at Hybritech, and I said no. I told them that I was perfectly happy at Syntex
217 and Syva, that this was my whole life, that I really loved doing it, and who are you guys
218 anyway? What kind of a crazy, wild-ass idea is that? And I suggested another guy who
219 we all all knew, Tom Adams, who at that time was at DuPont. And I said, 'Tom's
220 exactly the guy you need for that job,' and Tom did become the first VP of R&D for
221 Hybritech, and of course, David Kabakoff, who we mentioned before, was the second, an
222 interesting coincidence. One of the poorer mistakes I've made, one of the bigger
223 mistakes of my life, was not to take that one. Obviously, I left a lot on the table. But it
224 began to infect me with the idea that there was a huge amount of opportunity for people
225 who had the ability to implement new ideas and manage and lead people. So, I went to
226 Syntex and I was resigned. I wasn't quite sure what I was going to do. I did that three
227 times. On the third time, I really left. The first two, I was just kidding. On the third, I
228 really did leave and I became president of a company called Liposome Technology, now
229 known as Sequus. It's in the Bay Area. And to tell you the truth, I hated it, absolutely
230 hated it. After nineteen years of one success after another at Syntex, or Syva, whatever, I
231 really hit the mountain on that one. I didn't do my due diligence carefully. I did not fit
232 with the people and the culture. They hired me because somebody was making them

233 seek an outside guy, and the insiders really resented having anybody come in. I was the
234 wrong guy in the wrong place at the wrong time. And nine months later, I left the
235 company, practically shattered, I must add, I mean, I was just disillusioned completely. I
236 didn't do anything for a couple of months. The phone was ringing constantly with people
237 who said, 'Look, why don't you help us this, help us do that,' and I started a company
238 called Biomedical Consulting Associates, which is Dick Schneider. There isn't anybody
239 else. I did that for a number of years and basically, people would come with an idea, and
240 I would help them with a business plan, if I liked the idea, and I would try to get it
241 financed. Trying to get them financed provided the entree to venture capital, which I
242 knew nothing about, but I learned fast. During the years that I had Biomedical
243 Consulting Associates going, I involved in starting five companies.

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Q: Which were those?

252 The one that's best known today is one called Molecular Devices. That was also started
253 by another guy who started Syva years before. And there were some others, and at this
254 point it's irrelevant, but the fact is that I could see what I did well. I would go in as the
255 president and CEO, run that company, hire somebody who could run it long term, stay on
256 the Board, work with them, develop the strategy, recruit people, help develop the science
257 wherever I could, and then move on to the next. And of course, that provided the entree
258 to venture capital. I got an opportunity to join Sequoia, a very well-known venture
259 capital firm. I'm getting pretty long in the tooth by now, I'm an old guy, and they
260 suggested that I come in and help them with some business plans, and that they would

261 help me look at some things. Anyway, one thing led to another, and they suggested, and
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263 I concurred, that I really wanted, that a reasonable career path for me was to become a
264 professional venture capitalist full-time, and they made the suggestion that I join a firm,
265 and they made some introductions, a number of offers were made, and I ultimately
266 accepted one from a group called 3i Ventures, a very large source of money that came
267 from the UK, here in Orange County.

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271 Q: And was Gensia one of the first companies that you got involved with at 3i?

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276 Yes, the first. It was the first investment that I made. I was involved with other
277 companies, but the first one that I recommended that they invest in was Gensia. And the
278 venture capitalists that I met, I met a lot of them during the due diligence process, but the
279 most relevant and important one was a guy named Jim Blair. Jim Blair, of course, was
280 just starting Domain at that time, and Blair said look, 'If you do Gensia, with us, you can
281 become president. You become president, I'll become chairman, and we'll go find a
282 president.' Of course, we found David Hale. But that's where Gensia came from. We
283 met Harry and Paul at the lab at UC-San Diego, financed that company.

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287 Q: How did you evaluate, and maybe I could make this a general question, how do you

288 go about evaluating people and technologies?

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292 You don't have enough time to listen to that. I mean, that's what I do for a living. If you
293 can be more specific, I'll be happy to answer your question?

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296 Q: Why did you invest in Gensia? Why did you think this would work? What was it

297 about those guys and what they were doing?

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301 You could turn that around and say, 'Look, that was your first investment. How could

302 you have possibly known?' I probably didn't. Sometimes you just get a feeling about

303 things. That's not as quantitative as you'd like, but Harry and Paul, two bright, very

304 articulate, and real sincere young scientists. I liked what they were doing. I understood

305 them perfectly. I mean, on a technical basis, I understood them one for one. I thought I

306 could add a lot of value. We were on the same wavelength in many respects. They had

307 something that looked like it was a proprietary program in an area that was very

308 interesting, in very large markets, a hundred million plus dollars a year markets. As I

309 said, they had good technology, good people. It looked like it was in the realm of the

310 doable, meaning with the resources that one could actually obtain. We thought we could

311 attract good management around them, they were in San Diego, they were very highly

312 regarded. You know, you put all of that together and you say, 'Well, gee, this is what

313 you do for a living,' so you give them a hand and get it started.

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317 Q: You were involved in bringing David Hale in?

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Yes.

Q: What were the circumstances surrounding that?

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330 Man, all of this stuff is so interrelated. Remember, I told you that I left Syntex in '83 for
331 a short stint at LTI, and then did Biomedical Consulting Associates. The operative word
332 there is consulting. Hybritech hired me as a consultant, and I worked for David
333 Kabakoff, exactly the guy that used to work for me. So now we'd turned the tables.
334 Now while I was there, I got to meet and know personally Cam Garner, David Hale, Tim
335 Wollaeger, Tom Adams, and a list of guys, Kim Blickenstaff, Gunirs Valkirs, I mean, this
336 whole group, many of the people who are on your list here. I can tell you other stories.
337 You know, when you're as old as I am, sooner or later, you know almost everybody.
338 Cam Garner, for instance, the fellow, who as you know, is the very successful, wonderful
339 guy at Dura, was a sales rep when I first met him. When I was at Syva, I was a customer
340 of his, one of his best customers, but nonetheless, that's where I first met him. He was
341 working for a company out of Oberlin, Ohio called Guilford Instruments that sold
342 spectrophotometers. Imagine now, the circumstances, here we are, investors in his
343 company, Jim's on the Board, and we're investors in Spiros, and here's David Kabakoff
344 running Spiros. I mean, you talk about a spaghetti factory here, we're all connected.
345 And it happens because, it's going to bring you right back to this concept, it's the people.
346 These people. It's the people and their connections and knowing them and trusting them
347 and being friends with them, and having a lot of respect for them that you develop over
348 years and years. It's not a mistake, it's not a surprise, it's not an accident. I don't believe
349 that at all. There's a very good reason why all these people are where they are. Anyway,
350 just to finish that up, that's how I met David Hale. He was the CEO of the company that
351 was employing me as a consultant. When I got into the venture capital business, the

352 second deal I did was one, no actually, excuse me, I'll back up on that, was one called
353 Immunetech. Immunetech is the predecessor company of Dura, which is a whole story in
354 itself, in fact, there's a business school case written on Immunetech and Dura.

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358 Q: Whose case?

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362 I think it's at Darden. I have a copy of it if you'd like it. It's fascinating. I think it's
363 fascinating. It's really neat. We were trying to recruit David Hale to become president at
364 Dura while he was at Hybritech, because, you see, Hybritech was just sold at that time to
365 Lilly, so David was potentially hireable. Well, we never really convinced him to come to
366 Immunetech, but he did agree to join the board of Immunetech, nee Dura, and he still is on
367 the board. Well, we got to know him even better, Blair and I, two different firms, I at 3i,
368 Jim at Domain, and when Harry and Paul were rocking and rolling to get Gensia started,
369 we went to him again, and they had gone to him independently, so he knew them, and
370 again, to make a long story short, we convinced him to become the president, so we could
371 get a real president and get me out of there.

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376 Q: A couple of people have told me that when putting together Gensia, there were some
377 problems between Kleiner-Perkins and Domain about who would lead the deal, and that
378 you acted as a sort of intermediary in those negotiations.

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382 Well, I wouldn't exaggerate my role, but I would tell that it seems so incongruous today
383 that Jim Blair, one of the paradigms of virtue of the biotech industry, I mean, he had

384 done Amgen and Genzyme and Repligen and Immunex and Genetics, I mean, just a
385 million, and Brook Byers, who had done Hybritech and Genentech, etc., etc., huge things,
386 and the two of them had never met. They had never met.

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390 Q: Were you there when they met?

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395 I introduced them. Sure, in order to resolve this issue that you're now referring to. I
396 remember very well setting up that meeting. I think it was at the Hyatt here in town, or
397 maybe it was in San Diego, I don't remember anymore, exactly, but I mean, I remember
398 watching these two guys come together, and they became fast friends, and that was
399 resolved that afternoon. There was never another wrinkle in that.

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403 Q: Brook Byers didn't go on the board of Gensia. He put on Howard Birndorf as his
404 surrogate, is that how it worked out?

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408 Yes, for a very, very short time. Howard was really not on the board, he was not on that
409 board for very long. I don't remember how long, you can verify that, but he didn't stay
410 on that board too long. I just don't remember.

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414 Q: Did Kleiner-Perkins have a representative on the board?

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419 No.

420 Q: It was basically a Domain company?
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424 Well, no. I mean, there were other very significant people who played a role in that. 3i
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426 had a representative, that was myself. Oxford Bioscience, which at that time was called
427 Fairfield, Ned Olivier, I don't think he was on the board, but he was there, Jerry
428 Benjamin from Advent in the U.K. I'm sure I'm forgetting somebody in the early days,
429 and maybe Paul or Harry, I mean, I don't remember now, but there were other venture
430 groups involved.

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433 Q: What's your view on what happened with the clinical trials on the adenosine
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435 compound? I talked to Harry Gruber, and he blames David Hale for the problems that
436 cropped up.

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439 Well, you're jumping ahead. I don't mind doing it, but there was a whole lot of stuff
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441 happening in the meantime. My view is that you have to look at things in perspective.
442 You can't take things out of context, it's very dangerous to do. In addition, you have to
443 have a certain belief that the system works. If you don't believe in that, then we're all
444 doing the wrong thing. And what I mean by that is that the regulatory system works. My
445 belief is net, net, net, the compound didn't work, OK? You don't blame David Hale, you
446 don't blame Harry Gruber. It's Harry's child, so in a way, he's going to strike out and try
447 to protect it. And I'm not being critical of that, but it went through a very exhausting
448 trial, and net, net, a number of people much smarter than me looked at that data and
449 concluded that it was not statistically significantly better than the placebo. Sorry, bell

450 rings, bong! Now, maybe they picked the wrong indication, maybe they adjust the trila
451 properly, maybe they didn't administer it properly, maybe they didn't present it to the
452 FDA properly, maybe, maybe, maybe. Monday morning quarterbacks, irrelevant. Net,
453 net, whatever they tried to prove, they were unable to?

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457 Q: And in the years that we just skipped over, what, in your opinion, were the really
458 significant events that stand out?

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462 Well, look, I would say that David Hale was recognized as a very successful leader at
463 Hybritech and he brought an aura of a winner, of a leader, to Gensia, and he raised a lot
464 of money, a huge amount of money over multiple times. The stock, as you know, went
465 from four or five dollars to sixty- some odd dollars. They had a full portfolio of very
466 interesting compounds and products. They built other instruments, as well, the Gen-Esa
467 system came out of there. It's an absolutely clever scheme, originally proposed by Ron
468 Tuttle, a very, very clever guy. He recruited a superb board, guys like John Wilkerson
469 joined that board, from the Wilkerson Group, Steve Mandell, the ex-CEO of XOMA, and
470 currently the president of Prizm. These are, you know, wonderful, high-quality people.
471 He recruited a management team that was great, really wonderful people. Another one
472 was the acquisition of McGaw, which has now become Gensia Laboratories. There's a
473 whole lot to talk to you about that, and why that was done, and how it was done, and
474 what the scheme was, and what they were thinking about, all of that. This company was
475 clearly on a rocket ship. I mean, it had over a billion dollar market cap.. It was held up
476 as a paragon to other companies in San Diego and all over California, and all over the

477 U.S., so there were a lot of positive things going on there, but they took a couple of pretty
478 serious torpedoes.

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482 Q: One being the adenosine compound, the other?

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487 The Gen-Esa system was not approved, either, until just recently, but meanwhile, a lot of
488 damage was done when that did not get approved. It did get approved in Europe, as you
489 know, and it's being sold in Europe, but it had a huge impact, those two turn-downs in
490 the U.S. They went back and re-submitted and argued the point and negotiated their way,
491 and now they've got the approval for Gen-Esa, and they're going to be able to market it,
492 but meanwhile, a lot of water had run out of the dam. They lost the patina of a winner,
493 they lost people, they lost time, they lost a lot. And you know, they had to basically sell
494 the company and refinance it, and now it's Gensia-Sicor, but it's a credit to Hale to stay
495 in there and fight the fight, and he's going to win, but that was a tough, tough, tough
496 time.

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500 Q: Were you involved with Viagene, too? You were on the board there?

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505 Sure, certainly.

Q: Do you remember the discussions about spinning it off?

506 Sure, I remember it perfectly. This is the creativity of Harry Gruber. Harry had
507 developed some of these ideas, the use of retroviral delivery vehicles for gene therapy,
508 and it was clearly not within the original scope of Gensia, when we put it together, which
509 was principally a cardiovascular company, working in adenosine metabolism. But the
510 early founders of Gensia, the venture founders, said, 'Look, let's take a small amount of
511 money, and you guys putter around in the back room,' I think it was Brad Gordon and
512 Doug Jolly, 'and see if you guys can get a proof of principle, and then, if you can, we'll
513 talk about spinning this out separately.' It was not within Gensia's purview or business
514 plan, but if it's a good idea and it can stand on its own, if the technology is robust
515 enough, it ought to be financeable, and indeed it was financeable. And Gensia retained
516 an ownership of 20%, originally, of Viagene.

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521 Q: At certain points, Viagene had problems raising money?

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526 Sure, what company doesn't?

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531 Q: Well, in particular, Series D wouldn't close.

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536 I don't think I can tell you for certain it was the Series D.

Q: Doug Jolly said that this is one that seemed to go on forever.

537 Well, that may be. I don't remember. I would say that it was probably an earlier round
538 where they began to run into problems. One of the issues was that we had to make a
539 change in the president, and that was a pretty uncomfortable time. Anytime you have to
540 go in and change the senior management of a company, you run the risk of losing the
541 support of your existing investors, and you clearly damage the possibility of getting any
542 new ones until you get things settled down. We went without a CEO for some period of
543 time. That was a very precarious time for Viagene.

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547 Q: How important, then, is the role of the CEO for those kinds of things? I mean David
548 Hale was invaluable for Gensia. The problems that Viagene had, can you attribute them
549 to, you know, who the CEO was, or who wasn't the CEO?

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553 I think you have to, to some extent, to recognize that if there is a failure of strategy or im
554 plementation of strategy, it probably falls at the feet of the CEO. If there's a failure of
555 science, we can't manage biology. But a really hot management team would recognize
556 that the science isn't working and change course before they ride the horse over the cliff.

557 Well, in the early case of Viagene, the science was slow to develop and the science was
558 not being implemented properly, and so it was necessary to change the management.

559 How important is management in any of these companies? It's probably even more
560 important than the technology. It's probably the single most important element.

561 Q: When you started working with Hybritech as a consultant what precisely were you
562 working on?

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566 When they were first starting the company, Ted Greene and Brook Byers will tell you
567 that they asked me if I would join their management team and start the company with
568 them. I said no. Years later, many years later like three, four, five, by then Tom Adams
569 had brought in David Kabakoff, Kabakoff was running R&D, and I was out of Syntex
570 and Syva, running Biomedical consulting as a free agent, and David Kabakoff called me
571 up and asked if I would have some time available to act as a consultant on certain
572 elements of their business strategy, and I said yes. Well, remember, at the time, and even
573 now, I'm principally a scientist and my area of expertise was in diagnostics, and
574 particularly in immunodiagnostics, using antibodies to detect the presence of certain
575 antigens in small molecules, and Hybritech was a diagnostics company at that time. The
576 part that was being run by Dennis Carlo, in therapeutics, I had nothing to do with, but the
577 part of it that was diagnostics, which was Tom Adams and David Kabakoff, was right
578 down the throat of what I did, and I ran these groups for years at Syva, so I had some
579 contacts and expertise, and a small amount of knowledge, so David said, 'Hey, look, it
580 can't hurt, you know. If you don't screw anything up, come on in here and give me a
581 hand.' So, I was in there helping them with assays and automated assays, machines,
582 instrumentation.

583 Q: Let me jump ahead now, to Biosite. It sound like what you were doing at Syva is very
584 similar to what they've done at Biosite. You were familiar with the problems, told them
585 you didn't think it would work, and declined to invest?

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589 Yes, I declined twice. I was wrong twice. That's my second mistake. My first one was
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591 Hybritech.

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596 Q: What were the problems that you saw?

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600 I'm sure they told you that, because they love telling that story that Dick screwed up
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602 again, and he did. Interestingly enough, my partner, Jesse Treu, at Domain, did make the
603 investment, and I'm glad we did, because we made money on it. But 3i, the company
604 that I represented and turned them down twice, did not make the investment and it didn't
605 make any money. The reason I turned them down was that I felt, and I thought, that the
606 magnitude of the task was very large. It was larger than they had estimated. They
607 underestimated how hard it would be to mix all those antibodies at one time, to get them
608 all balanced and to behave properly. But what I underestimated was the ability of Gunars
609 and Kim and the other people that they had with them, one in particular, I can't think of
610 his name right now, but I will in a minute, I underestimated how smart they were and
611 how dedicated they were to getting it done, and it really taught me a lot about people and
612 their will. They literally made it happen. They are really good people. I don't mind,
613 they can tell that story all they want.

