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COURT OF COMMON PLEAS

CUYAHOGA COUNTY, OHIO

ROSE BASTIAN, et al.,)

Plaintiff,)

vs.)

Case No. 202353

KEITH R. KOEPKE,)
M.D.,)

Judge John L. Angelotta

Defendants.)

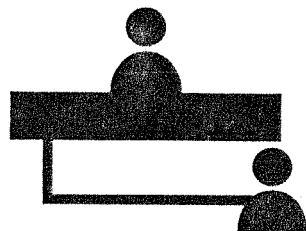
DEPOSITION UPON ORAL EXAMINATION OF:

BENJAMIN KIM, M.D.

TAKEN AT: 50 North Medical Drive, Salt Lake City, Utah

DATE: April 30, 1992

REPORTED BY: Rockie Dustin, CSR



**CAPITOL
REPORTERS**

175 South Main, #510
Salt Lake City, Utah 84111

(801) 363-7939

File No,

11879

A P P E A R A N C E S

For the Plaintiff: Clark A. Harms
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Salt Lake City, Utah 84111

For the Defendant: Susan M. Reinker
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* * *

I N D E XWITNESSPAGEBENJAMIN KIM, M.D.

Examination by Ms. Reinker 5

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* * *

E X H I B I T SEXHIBITSPAGE

No. 1 10-13-89 Letter to Fayiz A. Salwan 25
from Benjamin Kim

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1 THE DEPOSITION OF BENJAMIN KIM, M.D. was
2 taken on April 30, 1992, commencing at 9:30 a.m., at
3 the medical offices of Dr. Kim, 50 North Medical
4 Drive, Salt Lake City, Utah, before Rockie Dustin, a
5 Notary Public in and for the County of Salt Lake,
6 State of Utah.

7
8 BENJAMIN KIM, M.D.,
9 having been first duly sworn to tell the truth, the
10 whole truth and nothing but the truth, was examined
11 and testified as follows:

12
13 EXAMINATION

14 BY MS. REINKER:

15 Q. Dr, Kim, we met earlier today. My name is
16 Susan Reinker and, as I explained to you, I'm the
17 attorney who is representing Dr. Koepke, who is an
18 internist in the Cleveland area who has been sued by
19 Ms. Bastion, That's why we are here to take your
20 deposition, because you were the one who treated
21 Mrs. Bastian for her breast cancer, at least at
22 first.

23 I think the record should show that this
24 is the discovery deposition taken by the defendant,
25 as on cross-examination of Dr. Kim by agreement of

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(By Ms. Reinker)

1 counsel.

2 MS. REINKER: Is that correct?

3 MR. HARMS: That's correct.

4 MS. REINKER: And Mr. Harms is here today
5 on behalf of the plaintiff?

6 MR. HARMS: That's correct.

7 Q. Doctor, if you don't understand any of my
8 questions today, I want you to tell me that before
9 you try to answer the question, because we are going
10 to be relying on the testimony you give later on
11 when this case goes to trial.

12 A. All right.

13 Q. Would you state your name, please, for the
14 record?

15 A. Benjamin Kim.

16 Q. And your current business address?

17 A. Business address is Department of Surgery,
18 University of Utah, 50 North Medical Drive, Salt
19 Lake City.

20 Q. And your profession?

21 A. My profession is surgeon.

22 Q. How long have you been here in Utah?

23 A. I have been here two years.

24 Q. Do you know the approximate date you came
25 out here?

(By Ms. Reinker)

1 A. I came out here in the end of March of
2 1990.

3 Q. So your care, then, of Mrs. Bastian would
4 have ended in March of 1990 or February of 1990?

5 A. Yes.

6 Q. Would you tell us a little bit -- before
7 we get to that, have you had occasion prior to today
8 to discuss this lawsuit with anyone?

9 A. I was contacted by Mr. Harms' firm, and
10 from your firm as well, indicating that there may be
11 a suit.

12 Q. I know as far as my firm goes, my
13 secretary and one of my other associate's
14 secretaries called you to set up the deposition. I
15 don't think you had any conversations with either --
16 I know you didn't with me, or any other lawyer in
17 our office, did you, about the case?

18 A. Not specifically, no.

19 Q. Did you discuss the actual care rendered
20 to Mrs. Bastian with Mr. Harms or anyone else on
21 behalf of the plaintiff?

22 A. No.

23 Q. So you talked about setting up the
24 deposition?

25 A. Basically I was informed of the pending

(By Ms. Reinker)

1 lawsuits, at least potential lawsuits.

2 Q. How about the patient, did you ever talk
3 to her about the lawsuit?

4 A. I have not spoken with her about her
5 lawsuit, though prior to my moving out here she had
6 indicated some dissatisfaction, And I think the
7 potential might have been there.

8 Q. At that time did you give her any opinions
9 one way or another about the lawsuit?

10 A. Not to her.

11 Q. To whom?

12 A. I was contacted by -- and I don't recall
13 which firm it was -- I think it may have been
14 Mr. Harms' firm, about any evaluation assessments
15 for a potential lawsuit.

16 Q. And that would probably be one of the
17 lawyers back in Cleveland, Mr. Blakely or
18 Mr. Newman?

19 A. I don't recall.

20 Q. Do you have any notes from that
21 conference?

22 A. I do not have that currently. I have to
23 tell you that for some reason my notes on her did
24 not arrive with me here. I can gather that perhaps
25 it was left back in Cleveland, yes.

(By Ms. Reinker)

1 Q. Did you have more than one conversation
2 with Mr. Blakely's office about her case?

3 THE WITNESS: Let me clarify. Mr. Blakely
4 and you are the same firm?

5 MR. HARMS: Sure. Let me explain that.
6 It would be good to have this in the record, too.
7 Mr. Blakely is the plaintiff's attorney in this
8 matter. Mr. Blakely and his firm practice law in
9 Ohio. The case is pending now in Ohio. Yesterday
10 Mr. Blakely contacted me and asked if I would sit in
11 on this deposition for him. I'm just an attorney in
12 Salt Lake City. Our firm only practices in Salt
13 Lake City. And I have no other connection with the
14 case other than being here today.

15 THE WITNESS: Okay. So he has not said
16 anything to you, then, about anything?

17 MR. HARMS: He gave me a background of
18 what the case was and why we are going to have this
19 deposition.

20 Q. I believe my question was whether you had
21 more than one conversation with Mr. Blakely or the
22 plaintiff's lawyers about this case.

23 A. I have had more than one conversation with
24 Mr. Blakely and his law firm with regard to
25 Mrs. Bastian. I don't know whether those

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(By Ms. Reinker)

1 conversations -- how each of those directly related
2 to this case. At least one of them did.

3 Q. Were the other conversations just
4 basically about the patient's condition, how she was
5 doing?

6 A. About her -- yes, about her condition.

7 Q. Did you ever prepare any kind of a written
8 report for Mr. Blakely or his firm?

9 A. I prepared a report where I was asked to
10 give an assessment of potential lawsuit.

11 Q. And do you have a copy of that report?

12 A. I believe that should be back in
13 Cleveland, or it should be in Mr. Blakely's files.

14 Q. What do you recall of the substance of
15 that report?

16 A. The substance of the report was to say
17 that I did not feel she was materially -- let me
18 backtrack and say that I felt, you know, she had
19 optimal care rendered.

20 Q. You are referring to by Dr. Koepke?

21 A. No. To my care of her. And, you know,
22 the referral from Dr. Salwan, is it?

23 Q. So you did not make any comments about Dr.
24 Koepke's care?

25 A. Dr. Koepke's name, I believe, did not come

(By Ms. Reinker)

1 up until, yes, your office contacted me.

2 Q. Did you render -- well, are you aware that
3 the claim in this case is that there was a delay in
4 diagnosing Mrs. Bastian's breast cancer of about a
5 year?

6 A. That was not made clear to me.

7 Q. Were you ever asked to render any opinions
8 as to whether earlier diagnosis would have made any
9 difference in your management of this patient?

10 A. No.

11 Q. Doctor, have you ever had any
12 conversations, since coming out here, with
13 Dr. Silverman, Dr. Paula Silverman, about this case'?

14 A. No.

15 Q. Do you know Dr. Silverman?

16 A. Yes.

17 Q. How about Dr. Jean Stevenson, have you
18 ever discussed the case with her?

19 A. No.

20 Q. Do you know Dr. Larry Levy in Cleveland?

21 A. No.

22 Q. Have you ever discussed the case with Dr.
23 Levy?

24 A. No, I have not.

25 Q. Would you tell us just a little bit about

(By Ms. Reinker)

1 your background and training very briefly, where you
2 went to medical school, your residency, that kind of
3 thing?

4 A. I received my medical degree at Columbia
5 University in New York College of Physicians and
6 Surgeons. That was in 1978. I did my surgical
7 residency at Yale Newhaven Hospital. I spent two
8 years doing a surgical oncology fellowship at the
9 National Cancer Institute,

10 Q. What year did you conclude that?

11 A. That was between 1980 and 1982. And then
12 I joined the faculty at Case Western Reserve in
13 1985, and I was assistant professor there and also
14 both in general surgery and specifically surgical
15 oncology.

16 Q. What was your position there in 1989 when
17 you cared for Mrs. Bastian?

18 A. I was still on the faculty. I was
19 assistant professor in the department.

20 Q. In the department of surgery?

21 A. In the department of surgery. I had other
22 cross appointments in general medicine. I was also
23 in surgical oncology there, and I also was director
24 of the breast clinic there.

25 Q. Is your -- do you have a subspecialty

(By Ms. Reinker)

1 field in surgical oncology?

2 A. We consider surgical oncology sufficiently

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(By Ms. Reinker)

1 Q. In general surgery?

2 A. In general surgery.

3 Q. When you left in 1990, left Metro, did
4 Dr. Jean Stevenson take over your practice?

5 A. I wouldn't phrase it quite that strongly.
6 Many of my patients were referred over to Dr. Jean
7 Stevenson, and others were sent to doctor -- another
8 physician.

9 Q. Is Dr. Stevenson also a surgical
10 oncologist?

11 A. She is a general surgeon. I am not aware
12 of specific training that she received outside the
13 training that was given to her during her residency
14 in surgical oncology.

15 Q. Who was the chief of surgery at Metro at
16 the time you were there?

17 A. Dr. Anthony Imbembow was chief of surgery,
18 but he had left and I don't know the exact
19 transition dates. We **did** have an acting chief of
20 surgery at Metro, which was Dr. Monseur.

21 Q. Dr. Monseur?

22 A. Yes. And I don't know what year that was,
23 but I believe Dr. Monseur may have been the acting
24 chairman.

25 Q. Was he technically your boss, the head of

(By Ms. Reinker)

1 your department?

2 A. If he is acting chairman, yes, he would
3 be.

4 Q. Now, Doctor, you have had a chance a
5 little bit ago to look through the outpatient notes
6 and the records that we give you on Mrs. Bastian?

7 A. Right.

8 Q. And from what I can tell, your first
9 contact with this patient occurred on October 10th
10 of 1989; is that correct?

11 A. That is correct.

12 Q. And I believe she was sent down to you or
13 came to see you from Dr. Salwan at Parma?

14 A. Yes.

15 Q. I'm going to ask you some questions about
16 parts of your consult note, of your patient note,
17 and I want you to feel free to look at that note as
18 I ask you questions. The note is three pages long,
19 I think. Apparently on that date, Mrs. Bastian
20 brought with her the mammogram film, because you
21 refer to them in your note.

22 A. Yes.

23 Q. Do you have any recollection at all
24 sitting here today of what those mammogram films
25 looked like?

(By Ms. Reinker)

1 A. Very vaguely.

2 Q. Now, I think in your note you refer to the
3 first mammogram as being in February of 1988. It
4 was actually March 14th of 1988. You saw that film,
5 the first film?

6 A. (Pause.)

7 Q. It's in the middle of the second page.

8 A. Right, Yes.

9 Q. Did you ever see the x-ray interpretation
10 on that film?

11 A. I don't note that, so I have to assume
12 that I did not see that, to the best of my
13 recollection.

14 Q. Now, I have got that here and I would like
15 to show that to you. It was interpreted on March
16 15th of 1988. And, basically, the report that was
17 sent to Dr. Koepke finds no evidence of a malignancy
18 and describes a dense area of the parenchyma. I
19 think it's in the upper right quadrant?

20 A. Yes.

21 Q. Were you aware that on that date, March
22 15th of 1988, certainly around that point in time,
23 no one felt any breast lump?

24 A. I did not have notes from there, so I
25 can't comment on that.

(By Ms. Reinker)

1 Q. Now, the second mammogram that you looked
2 at that day was from September 14th of 1989, I
3 think it's probably the -- right there.

4 A. All right.

5 Q. And, again, had you ever seen that
6 interpretation before, the one you are now looking
7 at?

8 A. I don't recollect.

9 Q. Now, that interpretation describes an
10 ill-defined lesion; correct?

11 A. Yes.

12 Q. And I think they give a dimension on it
13 the of 1.8 by 1 centimeter?

14 A. Yes.

15 Q. Now, there was no description of any mass
16 lesion in the first interpretation you looked at:
17 correct?

18 A. Please ask me that again.

19 Q. If you want to just flip back, there was
20 no description or documented size of any lump or
21 mass in the first mammogram report in 1988, was
22 there? You don't give any dimensions?

23 A. The report just notes an area of
24 asymmetric dense mammary parenchyma.

25 Q. And they don't give any size,

(By Ms. Reinker)

1 measurements, or anything like that of size or
2 measure?

3 A. I can't read a size measurement, looking
4 at this note.

5 Q. Now, Doctor, **do** you have any knowledge
6 from your experience working with breast cancer
7 patients, of what size lesion -- what size does the
8 lump have to be before it's first visible on
9 mammography?

10 A. The technical resolution of the
11 mammography is extremely fine, meaning we could pick
12 up less than a 1 millimeter size lesion, such as
13 microcalcifications. The interpretation, however,
14 is quite different. So there may be variances
15 between physicians as far as what they call a mass.

16 Q. Generally speaking, is it about, what,
17 half a centimeter before they call it a defined
18 mass?

19 A. That totally depends, so you can't make
20 hard and fast rules.

21 Q. So, then, I gather you would have no
22 opinion as to what would have been seen on a
23 mammogram at any point in time prior to September
24 14th of '89? Let me try that one over again.

25 Do you have any opinion as to at what

(By Ms. Reinker)

1 point in time a specific mass lesion became visible
2 on a mammogram -- could have become a visible
3 mammogram for this lady?

4 A. Sometime between these two studies.

5 Q. But you have no opinion as to when
6 specifically?

7 A. Not exact times, no.

8 Q. Have you ever seen any notes from
9 Dr. Grant Libe? He was the first surgeon that
10 Mrs. Bastian saw after September of 1989.

11 A. Was that at the time?

12 Q. I haven't provided those to you.

13 A. As I recall, and looking over my notes, I
14 must have been privy to an operative note. But I
15 don't have a copy of that here with me.

16 Q. And you don't recall seeing his office
17 note?

18 A, No, I do not see an office note.

19 Q. Were you aware that when Dr. Koepke, my
20 client, examined Mrs. Bastian in September of '89,
21 he could not palpate a breast lump at that point in
22 time?

23 A. I was not aware.

24 Q. And were you aware that when Dr. Grant
25 Libe first saw her after the mammogram report came

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(By Ms. Reinker)

1 back, Dr. Libe felt he could only barely palpate a
2 lump?

3 A. I was -- I did not have those notes.

4 Q. In fact, he used needle localization to do
5 the biopsy because of the difficulty he had in
6 palpating the lump. Were you aware of that?

7 A. I knew he did a needle localization
8 biopsy.

9 Q. And that's why needle localizations are
10 done, aren't they, it's a lesion that's kind of hard
11 to find?

12 A. We do needle localization biopsies for
13 lesions that are either nonpalpable or difficult,
14 and when the abnormality that we are going after is
15 a mammographic abnormality.

16 Q. Meaning it's a mammographic abnormality,
17 not a palpable abnormality?

18 A. Or some in some breasts it's difficult to
19 examine. If we rely upon the abnormalities, the
20 mammographic ones, then we have to rely upon some
21 way of confirming. And that's done by needle
22 localization.

23 Q. By the way, do normal doubling time
24 theories, do they apply to infiltrating lobular
25 carcinoma as well as intraductal carcinoma?

(By Ms. Reinker)

1 **a.** Tell me what your doubling theories are,

2 Q. Well, do infiltrating lobular carcinomas
3 double as they grow from one cell to two to **four** to
4 eight? Do they develop the same way an intraductal
5 carcinoma develops?

6 A. You are asking an extremely complex
7 question here, so I think I would have to ask you to
8 be more specific. Cancer **cells**, in general,. unless
9 either treated in some way, or unless the body
10 defenses can handle it, grow- And growth means cell
11 divisions.

12 Q. Let's just leave it at that.

13 A. Okay.

14 Q. Now, when you saw Mrs. Bastian, she did
15 bring with her the pathology report on the biopsy
16 that Dr. Libe had done; correct?

17 A. Correct.

18 Q. And you mentioned that also in your note?

19 A. Yes.

20 Q. That's on the second page again. And I
21 think you say here that there was a '3.5 centimeter
22 well-demarcated fibrous area.

23 Do you see that?

24 A. Right.

25 Q. Is that the size that we are considering

(By Ms. Reinker)

1 to be the size of her original tumor, the 3.5
2 centimeter well-demarcated fibrous area?

3 A. This was -- I was quoting from the
4 pathology report. I was not assuming necessarily
5 that that was the tumor. It's just an area there
6 where they were suspicious, presumably, for the
7 tumor.

8 Q. Do you have any opinion as to approximate
9 size of her breast tumor as of September 1989?

10 A. I think you can probably say it's likely
11 that the minimal dimensions were the ones that
12 corresponded to the mammography.

13 Q. The 1.8 by 1 centimeter?

14 A. Yes.

15 Q. What is the relevance of this 3.5
16 centimeter area described in the pathology report
17 from Parma Hospital?

18 A. It suggests that perhaps the tumor may
19 have been 3.5 centimeters.

20 Q. So is it fair to say, in your opinion,
21 that the size of the tumor in September of '89 was
22 somewhere in between 1 by 1.8 centimeters and 3.5
23 centimeters?

24 A. Yes, grossly.

25 Q. Grossly? That means by --

(By Ms. Reinker)

1 A. That means by just view of the naked eye
2 and by palpation. That's different than
3 microscopically.

4 Q. Do we know what the size was
5 microscopically?

6 A. I do not know.

7 Q. Were you able to find that anywhere in the
8 records that I gave you to look at?

9 A. I'm not trying to trick you, Doctor,
10 because, basically, I couldn't find it or figure it
11 out. I could not. We make best guesses.

12 Q. And our guess is somewhere between 1.8 by
13 1 and 3.5, grossly?

14 A. Right. When we are asked for size, the
15 best estimate will probably be the 3.5.

16 Q. Now, your conclusion when you saw the
17 patient on October 10th was that, first of **all**, you
18 wanted to review some more things. That's down at
19 the bottom paragraph of that page. You wanted to
20 look at the pathology slides that she brought with
21 her; correct?

22 A. Yes.

23 Q. And you want to know what the ERPR
24 receptors were, the operative margins, and then you
25 were going to talk to the patient again?

(By Ms. Reinker)

1 A. Right.

2 Q. Now, in that last paragraph, which
3 continues on the last page, you basically said that
4 after you had done all these things you were going
5 to consider either reexcision or a mastectomy or
6 surgical treatment of this patient?

7 A. Yes.

8 Q. Now, a reexcision is really a lumpectomy?
9 It's taking out the local area: correct?

10 A. Reexcision is a local resection of that
11 area.

12 Q. Which the layperson calls a "lumpectomy"?
13 Is it fair to use that term?

14 A. I would be careful about that. A
15 lumpectomy is a little too loose.

16 Q. So let's call it a reexcision, which would
17 not involve taking off the whole breast, just that
18 local area?

19 A. **Yes.**

20 Q. Or a mastectomy, which involves taking off
21 the whole breast?

22 A. Yes.

23 Q. So of your knowledge, as of October 10th,
24 you felt that a reexcision might still be a
25 possibility?

(By Ms. Reinker)

1 A. It's a possibility, though not necessarily
2 a specific recommendation.

3 Q. But it was a possibility in that the 3.5
4 centimeter size alone did not rule out the
5 possibility of a reexcision?

6 A. That's correct.

7 Q. Now, after that visit with the patient,
8 you did look through the additional materials, some
9 additional materials. I could not find another note
10 where you saw her in her office again until after
11 her surgery. Would you agree with that?

12 A. That appears to be correct.

13 Q. And that's why I brought with me that copy
14 of a letter that you wrote to Dr. Salwan.

15 A. Yes.

16 Q. Do you have that there in front of you?

17 A. I do.

18 Q. I think I would like to have that marked.

19 (Exhibit 1 marked.)

20 Q. Now, this letter was written to
21 Dr. Salwan, who was the referring doctor, dated
22 October 13th, 1989?

23 A. Yes.

24 Q. And this was written, apparently, after
25 you had had an opportunity, especially, to look at

(By Ms. Reinker)

1 the pathology slides. And you looked at those
2 slides with a pathologist at Metro, Dr. Park?

3 A. Yes.

4 Q. And you came to the conclusion -- you and
5 Dr. Park together, I presume?

6 A. Yes.

7 Q. Did you, by the way, actually look at the
8 slides yourself? Do you remember? Or would you
9 normally do that?

10 A. I usually do look at the slides.

11 Q. And that diagnosis was a diffuse
12 infiltrating lobular carcinoma?

13 A. Yes.

14 Q. That's the type of cancer she had?

15 A. Yes.

16 Q. Now, it's my understanding that that type
17 of cancer is relatively rare in the overall picture
18 of breast cancers. I think it's about six to eight
19 percent of all cancers are infiltrating lobular?

20 A. Yes. You could argue over a few
21 percentage points, but that's the 10 percent.

22 Q. Clearly, this is not the most common type
23 of breast cancer that you see?

24 A. That is correct.

25 Q. Infiltrating lobular cancer, carcinoma,

(By Ms. Reinker)

1 has some characteristics that are different from
2 other kinds of breast cancer, does it not?

3 A. Yes.

4 Q. What is significant about this kind of
5 cancer to you?

6 A. The infiltrating lobular carcinoma has
7 basically two important things. One is that they do
8 have, relative to the other, the common scirrhous
9 type of breast cancer, a slightly more favorable
10 prognosis. But it carries an increased risk for a
11 bilateral/contralateral tumor.

12 Q. For the tumor appearing in the other
13 breast?

14 A. Yes.

15 Q. Are infiltrating lobular carcinomas also
16 thought to be, by some people at least, multicentric
17 and multifocal in the breast?

18 A. I think we have a higher suspicion for
19 multifocality, yes.

20 Q. Meaning that they can be present in more
21 than one spot in the breast?

22 A. That can be true for any tumor.

23 Q. Is there a higher incidence or suspicion
24 of that with infiltrating lobular carcinoma?

25 A. Usually infiltrating lobulars, when they

(By Ms. Reinker)

1 are infiltrating are one lesion. And because the
2 standard of treatment used to be mastectomy, that
3 sufficed the question of multifocality is a much
4 more difficult one because that depends very much
5 about how carefully the studies are done.

6 Q. When you wrote your letter to Dr. Salwan,
7 what did you mean by the use of the word "diffuse
8 infiltrating"?

9 A. I meant that there were areas that were
10 involved with the carcinoma throughout, that this is
11 not a well circumscribed lesion.

12 Q. There were areas involved throughout what,
13 throughout the breast?

14 A. The slides. The pattern of the cancer
15 cells were, one, suggestive of a more widespread
16 type of involvement.

17 Q. Now, you sent to Dr. Salwan a -- in your
18 letter you sent to him a copy of an article from
19 Johns Hopkins University, a Johns Hopkins study from
20 the Annals of Surgery?

21 A. Yes.

22 Q. Now, I think I got out the correct
23 article. Is that the one you are referring to?

24 A. Yes.

25 Q. Now, that article was a study of

(By Ms. Reinker)

1 infiltrating lobular cancer patients; correct?

2 A. Yes.

3 Q. I think there were 99 patients with this
4 type of cancer who were studied in that article.

5 A. Correct.

6 Q. **Is** it fair to say that the main focus of
7 that article was to give physicians advice as to how
8 to handle the other breast?

9 A. Yes.

10 Q. And then in that article, all 99 of those
11 patients had radical or modified radical
12 mastectomies?

13 A. Yes.

14 Q. The majority school of thought for
15 treatment of infiltrating lobular carcinoma, at
16 least in 1989, was that you do a mastectomy:
17 correct?

18 A. In the country at large, yes.

19 Q. If Mrs. Bastian had been diagnosed with
20 breast cancer sooner than she was at some point in
21 time, it would have still been infiltrating lobular
22 carcinoma: correct?

23 A. I don't know. How much sooner?

24 Q. Well, does ductal cancer become
25 infiltrating lobular? Can we assume that since her

(By Ms. Reinker)

1 diagnosis in September of '89 was infiltrating
2 lobular carcinoma, that if somehow somebody could
3 have made that diagnosis a few months earlier, or a
4 year earlier, it would have been the same type of
5 cancer?

6 A. Sure. Early diagnosis of the same cancer
7 would have given you the same cancer.

8 Q. That's my question. So if she were
9 diagnosed even a year earlier, it still would have
10 been infiltrating lobular carcinoma?

11 A. This is a little tricky. At some point a
12 process started that became infiltrating lobular
13 carcinoma, and it's one possibility that that's the
14 diagnosis you would have.

15 Q. I guess just to -- once this process
16 started, at whatever point in time it was biopsied,
17 it would have been the same histological type?

18 A. Carcinomas don't change type.

19 Q. In your letter to Dr. Salwan, the second
20 paragraph of that letter deals primarily with your
21 thought as to how to handle her other breast;
22 correct?

23 A. Yes.

24 Q. And that was because some physicians, or
25 some researchers, have recommended going ahead and

(By Ms. Reinker)

1 removing the opposite breast to avoid the risk of

2
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11 Q. But there are schools of thought that go
12 both ways?

13 A, Yes.

14 Q. And the Johns Hopkins article basically
15 arrives at the conclusion that you don't need to
16 remove the other breast?

17 A. That is correct.

18 Q. Which is the treatment that you
19 recommended for this lady?

20 A. That is correct.

21 Q. I should say that's the course you chose
22 to follow with her, is to not remove the other
23 breast?

24 A. That is correct.

25 Q. By the way, does anyone know why this

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(By Ms. Reinker)

1 particular type of cancer has a higher incidence of
2 developing in both breasts?

3 A. Not to my knowledge.

4 Q. If a patient gets this kind of cancer in
5 both breasts, can it be at the same point in time?

6 A. Sure.

7 Q. Can it be at different points in time?

8 A. Yes.

9 Q. In either of those two cases, is -- are
10 both breast lesions thought to be primary lesions?

11 A. That depends upon the pathology, but yes,
12 we believe that a contralateral type of cancer in
13 someone with a lobular debrisus or a second delobal
14 primary.

15 Q. It's not considered to be a metastasis
16 from the first breast?

17 A. No.

18 Q. Now, you recommended that Mrs. Bastian
19 have a mastectomy of her right breast?

20 A. Yes.

21 MS. REINKER: Off the record.

22 (Discussion off the record.)

23 Q. And was that based on the type of cancer
24 she had, the diffuse nature of her infiltrating
25 lobular cancer?

(By Ms. Reinker)

1 A. Yes. It was -- that recommendation was
2 based upon several factors. One was the type of
3 cancer that it was.

4 Q. You say type of cancer?

5 A. Lobular.

6 Q. The infiltrating lobular.

7 A. And the diffuse nature of it, the
8 uncertainty as far as the margins of resection and
9 the size of the lesion.

10 Q. Again, the majority school of thought back
11 in '89 was that a patient with this type of cancer
12 needed to have a mastectomy; correct?

13 A. I would back off from "majority." I would
14 say the most conservative, meaning the safest and
15 best curative option, would have been that.

16 Q. In your opinion, was that the strongest
17 indication for her to have a mastectomy, the
18 histological type of cancer that she had?

19 A. No.

20 Q. Did you discuss with her the possibility
21 of having a lumpectomy even at that point in time?

22 A. I had discussed with her various options
23 but left the specific recommendation open until I
24 had a chance to review the pathology and to review
25 all of the other studies and to order up further

(By Ms. Reinker)

1 clinical studies.

2 Q. Was it your decision or hers to have a
3 mastectomy as opposed to a **local** reexcision?

4 A. The decision is always the patient's.

5 Q. Did you discuss with Mrs. Bastian before
6 her surgery the option for reconstruction of her
7 breast?

8 A. I'm sure I mentioned that to her.

9 Q. Back in 1989, what sort of reconstruction
10 would you have discussed with her or offered to her?

11 A. I offered to refer her to a plastic
12 surgeon. I myself do not do the reconstructions. I
13 refer that specific discussion to a plastic
14 surgeon. I offer the patients the opportunity to
15 speak with a plastic surgeon if they so desire.

16 Q. The type of reconstruction that would have
17 been offered to this lady back in 1989, would that
18 have been the type that would have been started at
19 the time of her mastectomy?

20 A. There are many ways to do this, so I don't
21 make a specific recommendation.

22 Q. Do you know whether she could have had a
23 reconstruction that was done at the time of her
24 mastectomy?

25 A. Yes, she could have.

(By Ms. Reinker)

1 Q. Do you know whether she was aware of
2 that? Would you have told her that that was a
3 possibility, if she so chose?

4 A. I'm sure I would have if she chose to go
5 the reconstruction route. I would have mentioned
6 that.

7 Q. Do you have any recollection of her
8 response when you offered reconstruction?

9 A. Not specifically.

10 Q. Now, I would like you to take a look at
11 the surgery note. Do you see that there? Primarily
12 the pathology report from the surgery. Now, that
13 procedure was done on October 23, I believe, 1989?

14 A. Yes.

15 Q. Did you do the surgery yourself?

16 A. Yes.

17 Q. Do you recall anything specific about that
18 surgery?

19 A. Not extraordinary. It was a very smooth
20 operation.

21 Q. Now, I would like you to take a look at
22 the pathology report.

23 A. All right,

24 Q. My copy of that report is two pages long.

25 A. I have a copy here.

(By Ms. Reinker)

1 Q. Now, the entire breast that had been
2 removed was sent down to the pathology department;
3 correct?

4 A. Yes.

5 Q. I would like you to take a look at the
6 first page of the pathology report.

7 A. All right.

8 Q. There is a longish paragraph?

9 A. Right.

10 Q. Now, the final diagnosis was "residual
11 infiltrating lobular carcinoma of breast with
12 multifocal lobular carcinoma in situ and focal
13 lobular carcinoma in situ with apocrine features."

14 That was their final diagnosis; correct?

15 A. Right.

16 Q. Reading through the gross description
17 about six or seven lines down, there is a sentence
18 that begins, "On sectioning the specimen through the
19 nipple... ." Do you see that sentence?

20 A. Yes.

21 Q. It says, "An irregular area of fibrosis
22 deep to the nipple is noted, and that area measured
23 approximately 2 by 1.8 centimeters"?

24 A. Yes.

25 Q. That's one area. Then going on further

(By Ms. Reinker)

1 down there is another sentence, just four or five up
2 from the bottom of that paragraph. It starts, "On
3 serial sectioning of the breast..." Do you see that
4 sentence?

5 A. Yes.

6 Q. It says, and I'm quoting, "An irregular
7 area of fibrosis with fine nodularity is felt in the
8 inferior medial quadrant of the breast. That area
9 measured 3.5 by 2 centimeters"?

10 A. Yes.

11 Q. And then the report goes on to say, "The
12 rest of the breast shows irregular areas of
13 fibrosis"?

14 A. Yes.

15 Q. Now, there is no specific microscopic
16 description that I could find of those three areas.
17 It seems to me that -- did you ever see a
18 microscopic report on this pathology report that you
19 remember?

20 A. Yes, there should be.

21 Q. There is something missing here, isn't
22 there?

23 A. There may be something missing because you
24 have here labeled eight through W, all of the
25 sections that were done. So, two things; one is

(By Ms. Reinker)

1 that those would have been read out, and they have
2 also -- they should have the slides on file.

3 Q. We haven't been able to find any
4 microscopic report yet at Metro on that. But those
5 areas that I had you look at, the one deep to the
6 nipple, the one in the inferior medial quadrant, and
7 then the irregular areas of fibrosis in the rest of
8 the breast, could those areas have also been
9 infiltrating lobular cancer?

10 A. From what I see in the report, I would not
11 make that assumption, but it doesn't specify.

12 Q. I was just wondering if those were the
13 areas that the pathologist referred to in the final
14 diagnosis when they talked about multifocal lobular
15 carcinoma in situ.

16 A. You have to make a very clear distinction
17 here between infiltrating lobular carcinoma and in
18 situ lesion. They are very radically different
19 entities.

20 Q. But it's my understanding that there still
21 is -- people aren't certain what the relationship is
22 between the two; isn't that correct?

23 A. Right. We don't understand the full
24 pathogenesis and the relation between the two.

25 Q. And carcinoma, lobular carcinoma in situ,

(By Ms. Reinker)

1 is still thought to be precancerous, isn't it?

2 A. It is felt to be a marker of high risk for
3 infiltrating lobular carcinoma,

4 Q. So we just don't know the relevance of
5 those areas that you and I looked at before as to
6 what they were because we don't know the microscopic
7 report on them; is that correct, the area deep to
8 the nipple, the area in the inferior medial
9 quadrant, and the other areas of the breast?

10 A. Not from reading this. Not until you have
11 -- since they took the representative sections and
12 reviewed each of those.

13 Q. Not until we see the microscopic report on
14 those, we can't really tell what --

15 A. Right. The summary report does not
16 specify specifically what each of those sections
17 corresponded to.

18 Q. Is it even possible that those areas were
19 either infiltrating lobular or lobular carcinoma in
20 situ? That's a possibility, isn't it, from what we
21 see here?

22 A. My interpretation of this is that they saw
23 residual tumor, I presume, around the area of the
24 prior biopsy. But, as you point out, you wouldn't
25 want to make that assumption until you went back and

(By Ms. Reinker)

1 made correlations between the sections and the **areas**
2 where it was taken from.

3 Q. Women who have breast cancer generally
4 have had it, at least in microscopic form, for a
5 very long time before it reaches the size that it
6 can be diagnosable; correct?

7 A. You have to be more precise. What is "a
8 very long time"?

9 Q. Sometimes years.

10 A. It **could** be years.

11 Q. Isn't there a general feeling that to get
12 from that very first abnormal cell to a lesion that
13 can be either seen on mammogram or palpated, it can
14 be many years, sometimes?

15 A. It's a hypothesis.

16 Q. I mean, people don't just --

17 A. You don't go from one cell to a five
18 centimeter lesion in one week.

19 Q. It takes some period of time for it to
20 grow to the point it can be detected?

21 A. Yes.

22 Q. Now, in my copy of the Metro chart, I
23 could not find ERPR studies, estrogen receptor/
24 progesterone receptor studies done. Were they done
25 at Metro?

(By Ms. Reinker)

1 A. Yes. I have those reports here.

2 Q. What were they?

3 A. The estrogen receptor was 51 fentimals
4 (phonetic) per milligram of protein.

5 Q. That would make her ER positive; correct?

6 A. Yes.

7 Q. And the progesterone?

8 A. That level was five fentimals per
9 milligram of protein.

10 Q. Is that what you would consider
11 borderline? It's positive but it's a low positive?

12 A. The Nichols Institute considers five
13 through 100 positive.

14 Q. So she was high, but she was on the very
15 low end of it?

16 A. So she was positive, and she was at the
17 low end of positive.

18 Q. Now, you excised 17 lymph nodes at the
19 time of her surgery?

20 A. Yes.

21 Q. And they were all negative; correct?

22 A. Yes.

23 Q. And Dr. Stevenson, in her notes later on,
24 makes reference to DNA flow cytology studies?

25 A. Yes.

(By Ms. Reinker)

1 Q. I didn't see those either in my copy of
2 Metro charts.

3 A. I have those available,

4 Q. I'm going to ask you for copies of those
5 today before I leave.

6 A. Sure.

7 Q. She says, basically, that the cytometry
8 showed the tumor was diploid (phonetic)?

9 A. Yes.

10 Q. And there was a low S phase?

11 A. Yes.

12 Q. And all of those things are good
13 prognostic factors, are they not?

14 A. Yes.

15 Q. The absence of positive nodes, the
16 positive ERPR, the diploid nature of the tumor, and
17 the low S phase, those are all favorable
18 prognostically?

19 A. Yes, all favorable prognostic indicators.

20 Q. In addition to that, you did all the
21 series studies for metastatic diseases, looking for
22 tumors spread throughout her body?

23 A. Yes.

24 Q. And that was all negative as well?

25 A, Yes.

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(By Ms. Reinker)

1 Q. And that was also a positive finding?

2 A. It meant she had no metastasis detected.

3 Q. Good prognosis at that point?

4 A. It meant that, for the tumor, she had the
5 best prognosis.

6 Q. Dr. Stevenson, in some of her notes later
7 on, referred to this patient as a stage one. Would
8 you agree with that?

9 A. No.

10 Q. And why would you disagree with that?

11 A. Because a 3.5 centimeter lesion is
12 catagorically a stage two.

13 Q. Would that alone make her a stage two?

14 A. Yes.

15 Q. Dr. Stevenson referred to her as a low-
16 risk patient. Would you disagree with that?

17 A. She was biologically low risk.

18 Q. All those positive prognostic factors that
19 we talked about before, ERPR, **DNA** flow studies,
20 let's see here, the absence of nodal involvement,
the low S phase, the diploid nature of the tumor, we
22 can assume that those results would all have been
23 the same if this tumor would somehow have been
24 removed at some point in time, can we not?

25 A. It's reasonable.

(By Ms. Reinker)

1 Q. Now, your plan of treatment for this
2 patient after this surgery was what?

3 A. She was placed on adjuvant therapy,

4 Q. Now, my understanding is the only
5 treatment she got was tamoxifen?

6 A. Yes

7 Q. And that is a medication given to suppress
8 ovarian activity?

9 A. Tamoxifen is an antiestrogen.

10 Q. And the reason she was given tamoxifen was
11 because she had the ERPR positive findings; is that
12 correct?

13 A. She was given tamoxifen for four reasons.
14 The first was that she was estrogen receptor
15 positive. Her tumor was, I presume, at least 3.5
16 centimeters. She was postmenopausal. And tamoxifen
17 has relatively few side effects.

18 Q. Would she have gotten the tamoxifen if
19 diagnosis had been made at some point earlier,
20 assuming she was postmenopausal and ERPR positive
21 the point of diagnosis?

22 A. She may not have gotten it.

23 Q. At what point in time would she not have
24 needed it?

25 A. She may not have needed it, and you have

(By Ms. Reinker)

1 to remember that we are now in an area where there
2 is a tremendous amount of controversy. But if she
3 were a small lesion, one could make the argument
4 that she may not have required it, or that one might
5 not have recommended tamoxifen,

6 Q. How small?

7 A. Less than three centimeters, at least
8 according to the studies that have been done.

9 Q. Now, you told me earlier that it's
10 possible in this case that the tumor size described
11 -- that -- in this case we really don't know for
12 certain her exact tumor size; correct?

13 A. Exactly, we may not.

14 Q. So is it fair to say that in the height of
15 caution you decided to give her the tamoxifen?

16 A. I made the best reconstruction from the
17 data available of what I thought her tumor size, et
18 cetera, were.

19 Q. Does tamoxifen play any impact on the
20 development of cancer in the other breast?

21 A. It may.

22 Q. What is novadex?

23 A. Tamoxifen.

24 Q. Is that the brand name for tamoxifen?

25 A. Yes.

(By Ms. Reinker)

1 Q. Is there a reason, if these patients are
2 at higher risk of developing -- patients with
3 infiltrating lobular carcinoma can develop it
4 bilaterally, is there any reason that they are not
5 put on full chemotherapy?

6 A. We usually reserve chemotherapy for
7 treatment. So standard cytotoxic chemotherapy, I
8 don't know of anyone who would treat with that for
9 the other breast. Some people might argue that she
10 should be put on cytotoxin chemotherapy as an
11 adjuvant therapy. This business of tamoxifen, you
12 are going to see, as an adjuvant therapy, as quite
13 controversial right now.

14 Q. Now, this lady did not get routine
15 cytotoxic chemotherapy, did she?

16 A. No.

17 Q. Tamoxifen is not considered that kind of
18 chemotherapy?

19 A. It's a hormonal agent. It's called a
20 what?

21 A. A hormonal agent.

22 Q. What was your plan for follow-up with this
23 patient?

24 A. Follow-up would be examination every three
25 months, yearly contralateral mammography, blood work

(By Ms. Reinker)

1 when she was seen every three months, chest X rays
2 every six months, and bone scan yearly.

3 Q. There is some indication that this patient
4 may now have an abnormality in her other breast. If
5 that in fact is occurring, would that surprise you?

6 A. It wouldn't surprise me, no.

7 Q. That's part the bilateral nature of
8 infiltrating lobular carcinoma; correct?

9 A. It depends on what it is. Our index of
10 suspicion and care with which you follow the other
11 breast is heightened by lobular carcinoma.

12 Q. Assuming that she still does not have any
13 evidence of metastases, would the development of an
14 infiltrating lobular carcinoma in her other breast
15 change your prognosis?

16 A. Prognosis with respect to?

17 Q. For long-term survival.

18 A. It depends on the other breast lesion,
19 when that is detected, at what stage.

20 Q. This lady now apparently has developed
21 some form of leukemia. Are you aware of that?

22 A. I was aware of that in, I think, my last
23 conversation with the law firm.

24 Q. What did they tell you about that?

25 A. They said, I believe, chronic lymphocytic

(By Ms. Reinker)

leukemia.

Q. Does that have any relationship to her breast cancer?

A. Not that I am aware of.

Q. Do you have any -- have you been given any idea what her prognosis is from her leukemia?

A. No.

Q. Do you treat patients with that kind of leukemia?

A. I refer them to a medical oncologist.

Q. Based on your general knowledge, what is your prognosis for that kind of leukemia?

A. It varies.

Q. You have no opinion, in her case, what her prognosis is?

A. Not without looking at her smears, bone marrow, a whole series of evaluations.

Q. What is your opinion of what -- or what was your opinion back in 1989, when you were treating this lady as to her prognosis from her infiltrating lobular breast cancer, based on everything you knew after her surgery and all the prognostic studies that have been done?

A. I thought that she would fall into our best prognostic group, stage two carcinoma.

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(By Ms. Reinker)

1 Q. Which was what percentage?

2 A. Better than 70 percent survival in five
3 years.

4 Q. And you did not feel at that point that
5 she needed radiation treatment of any kind?

6 A. **Such** as?

7 Q. I don't know. Did you ever discuss
8 radiation with her?

9 A. When I discussed options for treatment, I
10 would have said to her that if she had -- or if she
11 decided, regardless of recommendations, that she
12 wanted a local recission, then I would have
13 suggested radiation. But not if she went total
14 mastectomy and resection.

15 Q. So if she would have chosen to have a
16 local recission, she would have gotten the radiation
17 treatment?

18 A. I would have recommended that.

19 Q. But since she did not choose to go that
20 route, she did not receive the radiation?

21 A. That is correct.

22 Q. You also did not feel she needed a full
23 course of adjuvant cytotoxic chemotherapy?

24 A. I did not feel that would be in her best
25 interest.

(By Ms. Reinker)

1 Q. Do you know what Dr. Stevenson's criteria
2 was for calling this lady a stage one?

3 A. No, I don't.

4 Q. Basically, you considered her to be a
5 stage two, based on an assumption as to what her
6 tumor size was; correct?

7 A. Yes.

8 Q. If, in fact, her tumor size was 1.8 by 1
9 centimeter, as the mammogram report shows, then she
10 would have been a stage one; correct?

11 A. Yes.

12 Q. And if that were in fact the true
13 dimension of the tumor, where would that put her,
14 prognostically, as a stage one?

15 A. Better than a stage two.

16 Q. And that's the best possible prognostic
17 stage to be, is a stage one, is it not?

18 A. That's the earliest, yes.

19 Q. Do you know the difference between -- you
20 said a better than 70 percent survival if she was a
21 stage two, based on tumor size alone. But if her
22 tumor size was under 2 centimeters, if in fact the
23 mammogram was right and it was 1.8 by 1, what
24 percentage rate of survival does she then have?

25 A. It would be better, but how much better

(By Ms. Reinker)

1 you can't say. These numbers are statistical
2 numbers. So for any given patient, you know, they
3 either get disease or they die from disease or they
4 don't. But these are -- if you take a thousand
5 women, they apply. So it gives you some sense of
6 how well they might do. But it doesn't predict the
7 outcome for each patient.

8 Q. Would her new development, the development
9 of the leukemia, shorten her life expectancy?

10 A. Without knowing the current status of her
11 leukemia, I can't answer that.

12 Q. How about it developing -- if indeed she
13 was developing cancer in the other breast, would
14 that shorten her life expectancy?

15 A. Not necessarily, if it's detected early
16 and treated.

17 Q. And again, if she does develop cancer in
18 the other breast, that's part of the course for some
19 women who have infiltrating lobular carcinoma;
20 correct?

21 A. And also for women who have in situ
22 lobular carcinoma,

23 Q. And that is not considered to be a
24 metastatic lesion from the first cancer?

25 A. Usually not, but you need to depend upon

(By Ms. Reinker)

1 the pathology. If, in fact, she develops it in the
2 other breast, the same factor would apply that we
3 talked earlier, that she has had it for a
4 considerable period of time for it to become
5 diagnosable. And, again, I think you have to be
6 careful about what the considerable time factor is.

7 Q. What does that mean to you? How long
8 would you say a woman has cancer cells in her breast
9 before it can be become diagnosable?

10 A. That's a tough question now, because of
11 mammography and all -- but occasionally we will pick
12 up carcinoma incidentally. So very often we may be
13 not looking at the cancer cells per se but a
14 secondary response to it. So that's -- you know, I
15 think you have to specify each situation.

16 Q. So you really have no opinion as to how
17 long a woman has had cancer cells in her breast
18 before they become diagnosable by mammography or any
19 other way?

20 A. I have opinions, but I think in this case,
21 especially with the in situ lesions, that would be
22 extremely difficult to know. Because remember that
23 there is a distinction between cancer cells per se
24 and cancer cells that invade outside. And so you
25 can have cancer cells, or what we call, you know, in

(By Ms. Reinker)

1 situlogical cancer cells, or has not yet developed
2 the characteristics of invasion.

3 Q. So are you saying that if the pathology
4 report on the breast that you removed found in situ
5 cancer cells, you are assuming she also had those in
6 situ cells in her other breast?

7 A. No, though she may, and I won't be
8 surprised if she could have those.

9 Q. Doctor, is there anything that I haven't
10 asked you about this case which you feel you would
11 like to render some opinions about or talk about?

12 A. Specifically?

13 Q. I'm just asking if there is anything in
14 your mind that you are burning to say, to tell me,
15 about this patient or this case?

16 A. No, other than I think that this -- we are
17 dealing with a histology that is infrequent, and we
18 are dealing with a fairly diffuse process. And that
19 makes decision making more complex than your
20 standard run-of-the-mill breast cancer.

21 Q. This kind *of* cancer is somewhat difficult
22 to diagnose, is it not?

23 A. I don't know how you mean that.

24 Q. I have read some articles that say that
25 sometimes this is a hard cancer to diagnose because

(By Ms. Reinker)

1 of the way it grows, or something like that.

2 A. You would have to be more specific.

3 Q. Do you have any opinion as to whether --
4 well, I think I asked you this earlier, whether
5 earlier -- at what point in time this breast cancer
6 could have been diagnosed?

7 A. You can always diagnose it earlier, but
8 the practical reality is, I don't know how much
9 earlier you can do that. You have to have
10 convergence *of* several things. One is what
11 modalities you use to study and your level of
12 suspicion.

13 Q. Well, I'm asking you to base that on the
14 assumption that nobody was ever really able to
15 palpate a breast lump. Dr. Libe thought maybe he
16 could, but there was never really any palpable
17 breast lump. We know the mammogram in September
18 reported only a 1 by 1.8 centimeter lesion.

19 I'm just wondering if you had any opinion,
20 with those assumptions, how much earlier it would
21 have even been possible to diagnose this lesion.

22 A. I mean, you could -- you know, we have
23 pathologic data showing that people sometimes will
24 find carcinomas that were never detected. So it's a
25 theoretical question which I think is, you know, not

(By Ms. Reinker)

1 helpful.

2 Q. So you really, then, don't have any
3 opinion as to when this became diagnosable?

4 A. Not from what I can gather back here.
5 Because you have an initial mammogram that
6 apparently was read as being low suspicion, And
7 then you have a second mammogram with a suspicious
8 lesion. And if it wasn't palpable, then we really
9 don't have other modalities as yet that would help
10 us.

11 MS. REINKER: I think that's all I have.

12 MR. HARMS: I have a few, if I could
13 follow up on a couple of things.

14

15 EXAMINATION

16 BY MR. HARMS:

17 Q. The first item is, when you were talking
18 earlier about the actual decision of what type of
19 surgical procedure to use, you indicated that you
20 had advised Mrs. Bastian that there were a wide
21 range of curative options. Did you recommend a
22 single one of those to her as probably her best bet
23 for a least intrusive yet best result, or did you
24 just say, "Here is what we can do"?

25 A. I made a very strong recommendation for

(By Mr. Harms)

1 her to undergo a total mastectomy and mass
2 redissection.

3 Q. And that is in fact what she underwent?

4 A. Yes.

5 Q. What were the considerations that you made
6 that strong recommendation upon? Why did you choose
7 that route rather than something less invasive or a
8 reexcision?

9 A. Based upon the pathologic review.

10 Q. So, in your opinion, at least, at the time
11 of the surgery you performed, a less -- a breast
12 conserving procedure, or less intrusive procedure,
13 other than the radical modified mastectomy that was
14 performed was not in her best interest?

15 A. That is correct.

16 Q. Dealing with this histological type of
17 carcinoma, just generally with an infiltrating
18 lobular carcinoma, is that a higher risk of
19 recurrence than a normal other type of breast cancer
20 or, you know, what goes into determining what the
21 risk of recurrence is?

22 A. The risk of recurrence is dependent upon
23 histology and then all the other process prognostic
24 factors that we talked about, like size of tumor,
25 the differentiation of the tumor, the ERPR status,

(By Mr. Harms)

1 and the S phase, and the lymph nodes.

2 Q. So the size of the tumor at excision
3 relates directly to the risk of recurrence?

4 A. It's one of the factors that weighs into
5 recurrence risk.

6 MR. HARMS: Other than getting back into
7 the doubling-rate-growth-chart area, I think
8 everything that I had has been covered. *So* I don't
9 think I have anything else.

10

11 FURTHER EXAMINATION

12 BY MS. REINKER:

13 Q. Doctor, are you aware of the current
14 recommendation as to the cutoff point at which
15 lumpectomies should no longer be performed?

16 A. There are different recommendations. In
17 general, it has to do with the size of the lesion
18 versus the size of the breast. And factored into it
19 are also the histologic features of the particular
20 tumor.

21 Q. Isn't it a fact that current thinking is
22 that any tumor less than four centimeters in size
23 can be safely handled with a lumpectomy?

24 A. Not necessarily.

25 Q. So you would disagree with any reports on

(By Ms. Reinker)

1 role in resection.

2 Q. And the fact that this lady had
3 infiltrating lobular carcinoma played a role in your
4 recommendation for a mastectomy?

5 A. The findings of positive margins from the
6 primary biopsy and the diffuse nature made me make
7 that recommendation, that she not undergo simple
8 lumpectomy.

9 Q. Doctor, how did you happen to leave Case
10 and come out here? Just out of curiosity,

11 A. I came up to head surgical oncology here.

12 Q. Just a different position you were
13 offered?

A. It's a bigger scope.

15 Q. What is your position here?

16 A. I am associate professor here, and chief
17 of surgical oncology for the University of Utah.

18 MS. REINKER: Okay. I have nothing else.

19 MR. HARMS: I have nothing else.

20 (Concluded at 11:00 a.m.)

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REPORTER'S CERTIFICATE

STATE OF UTAH)
COUNTY OF SALT LAKE) ss.

I, ROCKIE E. DUSTIN, Certified Shorthand Reporter and Notary Public for the State of Utah, certify:

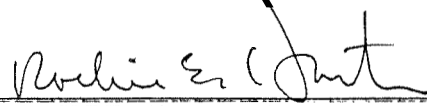
That the foregoing deposition of BENJAMIN KIM, M.D. was taken before me pursuant to notice at the time and place therein set forth, at which time the witness was put under oath by me;

That the testimony of the witness and all objections made at the time of the examination were recorded stenographically by me and were thereafter transcribed;

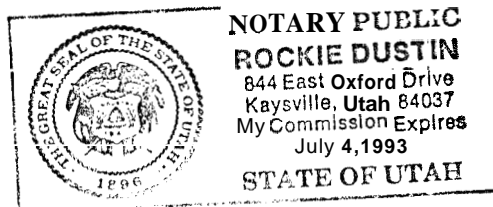
That the foregoing deposition is a true record of the testimony given by the witness and of all objections made at the time of the examination.

I FURTHER CERTIFY that I am neither counsel for nor related to any party to said action nor in anywise interested in the outcome thereof.

IN WITNESS WHEREOF, I have subscribed my name and affixed my seal this 5th day of May, 1992.


ROCKIE E. DUSTIN, CSR
Notary Public in and for the
County of Salt Lake, State of Utah

My Commission Expires:
July 4, 1993



Rockie Dustin * Capitol Reporters

WITNESS SIGNATURE CERTIFICATION

STATE OF UTAH)
COUNTY OF _____) ss.

BENJAMIN KIM, M.D. deposes and says: That he is the witness referred to in the foregoing deposition; that he has read the same and knows the contents thereof; that the same are true of his own knowledge,

BENJAMIN KIM, M.D.

SUBSCRIBED and SWORN to before me this ____ day of _____, 19____.

Notary Public

Residing at _____

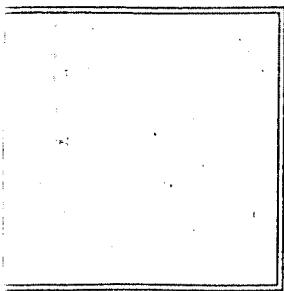
My commission expires:

Rockie Dustin * Capitol Reporters

C O R R E C T I O N S

Deposition of: BENJAMIN KIM, M.D.
 Taken: 4-30-92

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October 13, 1989

Fayiz A. Salwan, M.D.
6789 Ridge Road
Parma, Ohio 44129

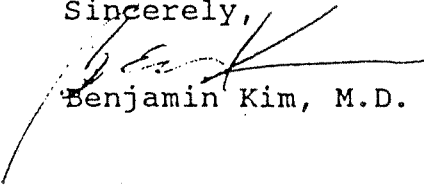
EXHIBIT 1
DATE 4-30-92
WITNESS Kim
ROCKIE DUSTIN, REPORTER/NOTARY

Dear Doctor Salwan:

Thank you for referring Roseaques Bastian. Her operative notes, pathology report, mammograms and pathology slides were all reviewed in conjunction with Dr. Yaegen. Radiologist, and Dr. Chung-Moon Park, of Pathology. The patient has a diffuse infiltrating lobular carcinoma and from Dr. Lieby's operative report and from the submitted slides it would appear reasonable to assume that there may be some residual tumor following biopsy. The mammogram shows bilateral calcifications which appear benign and the enlarging density for which the patient underwent biopsy. ~~Because the lesion is described as being 3 cm and because of the diffuse nature of its histology, I would recommend that the patient undergo a total mastectomy with axillary dissection.~~

The management of the contralateral breast in lobular carcinoma has been controversial, but because the patient's left mammogram appears normal and relatively easy to interpret, both on physical exam as well as mammographically, close follow-up with physical exam q.3 months and yearly mammograms should suffice without necessitating prophylactic biopsy or mastectomy. A review of the Johns Hopkins experience just published in Annals Of Surgery is enclosed for your perusal. I have conveyed these recommendations to the patient and her husband together with recommendation for a preoperative bone scan, CXR, and liver function test to exclude possible metastatic disease. Again, thank you for the referral. Please contact me if I can further elaborate.

Sincerely,


Benjamin Kim, M.D.

Benjamin Kim, M.D.
Assistant Professor of Surgery
Cleveland Metropolitan General Hospital
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Cleveland, Ohio 44109 (216)459-5358