

STATE OF OHIO)
) ss:
 COUNTY OF LAKE)

COPY

IN THE COURT OF COMMON PLEAS

MARY ANN FEATHERS,)
)
 Plaintiff,) Case No. 01CV000824
) Martin Parks, J.
 vs.)
)
 ROBERTA BROWN, M.D., et al.,)
)
 Defendants.)

VIDEOTAPED DEPOSITION OF HOWARD OZER, M.D.

TAKEN ON BEHALF OF THE DEFENDANTS

IN OKLAHOMA CITY, OKLAHOMA

ON JULY 24, 2002

REPORTED BY: ELIZABETH CAUDILL, CSR, RMR, CRR



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S T I P U L A T I O N S

IT IS HEREBY STIPULATED AND AGREED by
and among the attorneys for the respective
parties hereto that the deposition of HOWARD
OZER, M.D. may be taken on behalf of the
Defendants on JULY 24, 2002 in Oklahoma City,
Oklahoma, by Elizabeth Caudill, Certified
Shorthand Reporter within and for the State of
Oklahoma, pursuant to agreement.

IT IS FURTHER STIPULATED AND AGREED by
and among the attorneys for the respective
parties hereto that all objections, except as to
the form of the question, are reserved until the
time of trial, at which time they may be made
with the same force and effect as if made at the
time of the taking of this deposition.

* * * * *

* * * * *

HOWARD OZER, M.D.,
after having been first duly sworn at 2:03 p.m.,
deposes and says in reply to the questions
propounded as follows, to wit:

DIRECT EXAMINATION

BY MS. REID:

Q I'm just going to assume you had your
right hand up there.

A I did. It's being videotaped so you
can confirm.

Q That's right. I forgot about that.

Doctor, my name is Christine Reid, and
I represent Dr. Roberta Brown in a case being
brought by Mary Ann Feathers, and we're here for
your deposition today.

I assume you've had a deposition taken
before?

A I have.

Q All right. If at any time throughout
this deposition, you, or the court reporter, for
that matter, cannot hear me or don't understand a
question, please let me know and I'll attempt to
rephrase.

A I will. Thank you.

Q You're welcome. Could you give me your
2 current business address, please?

3 A Sure. I'm at the -- I'm in the section
4 of hematology-oncology in the Cancer Center at
5 the University of Oklahoma Health Sciences
6 Center, which is in Oklahoma City, Oklahoma.

7 Q Okay. And how long have you been with
8 the University of Oklahoma?

9 A Little over two years.

10 Q And your specialty is oncology?

11 A It's hematology and oncology, yes.

12 Q All right. Is there any particular
13 type of cancer you subspecialize in?

14 A Well, I do my research in the area of
15 leukemia, lymphoma, and now in pancreatic
16 carcinoma, but I see patients with all
17 malignancies.

18 Q Okay. Can you quantify the percentage
19 of your practice that involves the treatment of
20 breast cancer patients?

21 A Certainly. It's about 30 percent.
22 Because of the demographics, breast cancer being
23 so common, I would say about a third of all my
24 patients are breast cancer.

25 Q Donna was kind enough to hand me a copy

1 of your CV. It's a 35-page document. I'm trying
2 to see if there's a date on it. My question is
3 how can I -- maybe you can help me with this --
4 how I can determine whether this is the most
5 current copy of your CV.

6 A It's accurate, I would estimate, within
7 six months. I have not updated it in the last
8 six months and have a few more publications, but
9 I've not remade my CV since that one.

10 a Okay. Off the top of your head without
11 me reading your entire CV, are there any
12 publications listed here that DEAL with the
13 diagnosis and treatment of breast cancer?

14 A There are a couple from the time when I
15 was at Chapel Hill where we were looking at sort
16 of physician's approaches to the diagnosis of
17 breast cancer relating mostly to surgeons. And
18 those are -- are listed in the CV. There's only
19 a few of those. They deal with populations
20 rather than the specifics of breast cancer.

21 And beyond that, there really isn't
22 anything.

23 Q Okay. I want to talk to you for a
24 minute about your practice, and specifically AS
25 it relates to the treatment of breast cancer

1 cancer patients.

2 I assume that you see most patients
3 after the diagnosis of cancer has been made?

4 A Yes and no. We -- in medical oncology,
5 we follow patients for life, so that means that
6 if we have patients who have either a hematologic
7 or malignant problem that is cured, we follow
8 them for 15 and 20 years. So we do do annual
9 mammography, and we do breast exams serially in
10 our patient population that is doing well.

11 Q Okay. But your first contact with a
12 patient, would it typically be after there's been
13 the diagnosis of some type of malignancy?

14 A That would be more typical, yes.

15 Q All right. So you're not typically
16 involved in the diag -- the initial diagnostic
17 process?

18 A Well, I'm not sure what you mean by
19 that. Generally, we're referred patients who
20 have a breast lump. The -- by -- if you mean by
21 initial diagnostic process, we're -- we see
22 patients, and I actually run the breast
23 conference, which is this afternoon, where
24 patients will come in with a lump and have an
25 abnormal mammogram. So the very first visit to

1 the mammographer I might miss, but from the
2 mammogram on, we're participants.

3 Q All right. So you at that point might
4 get involved in ordering surgical consultations,
5 biopsies, et cetera?

6 A That's correct. We have a
7 multi-disciplinary team approach at OU where,
8 once a patient has a suspicious-looking
9 mammogram, they're seen by everybody.

10 Q Okay. Is it a breast center of some
11 type?

12 A It is

13 Q Okay. Can you, for my education, just
14 go through the materials you have reviewed in
15 this case?

16 A Certainly.

17 Q You have -- let me interrupt for a
18 second. Do you have a stack in front of you?

19 A I do.

20 Q All right. If you could just go
21 through them and list for me what you have, I'd
22 appreciate it.

23 A Certainly. Give me one moment. I
24 reviewed the office records of Dr. Roberta Brown,
25 the office records of Dr. John Dorsky, the Lake

'1 Hospital System records, the office records of
2 Dr. Green, the deposition of Dr. Brown, the --
3 I'm not sure what you call it, but the report
4 from Dr. Levitan, the brief report from the other
5 physician.

6 Q Dr. Resnick?

7 A Yes, Dr. Resnick. And I believe that's
8 everything.

9 Q Did you read Dr. Green's deposition?

10 A I did not see Dr. Green's deposition.
11 Hang on one -- yes, I'm sorry, I did see Dr.
12 Green's deposition.

13 Q Do you know Armin Green at all?

14 A I know the name. That's it.

15 Q Okay. Did you read Mary Ann Feathers's
16 deposition?

17 A I did not see Mary Ann Feathers's
18 deposition, no.

19 Q Okay. Do you have any notes, either
20 handwritten or typewritten, within your file?

21 A I do not.

22 Q Okay. How do you keep track of the
23 information as you're reviewing it?

24 A I -- when I give my first report, I
25 review it and immediately write a report, and

1 then I've spent the last two evenings
2 re-reviewing it for this deposition.

3 Q All right. But you don't take any
4 notes during that process?

5 A I do not. I did bring a copy of the
6 AJCC's handbook for cancer staging to address the
7 issue of survival statistics.

8 Q You knew I was going to ask you about
9 that, huh?

10 A I did.

11 Q Okay. We'll get to that in a minute.
12 Did you bring any other literature with you?

13 A No, just that.

14 Q Okay. Did you do any other type of
15 literature search --

16 A No,

17 Q -- prior to today's deposition?

18 A No, I did not.

19 Q Okay. Do you have any letters or
20 correspondence from Donna Kolis in your file?

21 A I do.

22 Q Okay. Can you just tell me what the
23 dates of those are?

24 A Sure. In no particular order, I have
25 one from February 15th, 2001. That is the

1 transmittal of the first set of materials; one
2 from August 26 of 2001; one from January 24th of
3 2002; one from -- actually I take it -- I have
4 two from January 24th of 2002.

5 Q Donna was working hard that day.

6 A She was. One from March 20th of 2002,
7 and one from April 22nd of 2002. And that, I
8 believe, is it.

9 a Okay. If you could do me a favor
10 before you leave today, just have the court
11 reporter make a copy of those --

12 A Will do.

13 Q -- for me. Thanks.

14 Have we gone through all the materials
15 you have reviewed in this case and that are
16 contained within your file?

17 A Yes, we have.

18 Q The report I have that was furnished to
19 me by Mrs. Kolis is dated April 13th, 2001. Do
20 you have any additional reports other than that
21 one?

22 A No. That's it.

23 Q Okay. Was there a draft sent to Mrs.
24 Kolis before this copy was produced?

25 A No. This was it.

'1 Q All right. This is the first, the only
2 report?

3 A Yes, it is.

4 Q Did you say yes? I'm sorry.

5 A Yes, I did.

6 Q Dr. Ozer, I'd like to go to your report
7 and talk about your opinions in this case. First
8 of all, let's talk about your opinions related to
9 the standard of care.

10 It's my understanding that, in your
11 opinion, Dr. Brown should have sent Mrs. Feathers
12 for surgical evaluation in February of 1999.

13 A That's correct. I think the
14 appropriate response for a dominant mass is one
15 of two things: either an immediate referral to a
16 surgeon or very close follow-up and repalpation
17 of that mass and remammography.

18 Q What do you mean by a dominant mass?

19 A A dominant mass, in women who have
20 fibrocystic changes of the breast, they may feel
21 lumpy both to the patient and to the examining
22 physician. But in this case, the patient reports
23 a new, quote, dominant mass, unquote; in other
24 words, a mass that she has identified is
25 different and distinct, and those warrant close

'1 follow-up.

2 Q And your opinion about the necessity of
3 close follow-up is irregardless of the fact that
4 Dr. Brown could not palpate that mass?

5 A That's correct. Particularly given its
6 size, it's at the lower limit of what has been
7 documented in the literature for physicians to be
8 able to detect, and therefore, it is quite
9 understandable that a physician might miss it.

10 At that point, it warrants referral of
11 the patient to a breast surgeon who is more
12 experienced in detecting such masses or,
13 alternatively, a very careful follow-up to detect
14 whether it's growing or not.

15 Q What's your understanding of what the
16 size of the mass was in February of 1999?

17 A Well, the patient describes it as pea
18 sized.

19 Q Okay. What's the size of a pea?

20 A That would roughly be half a
21 centimeter.

22 Q And the literature does document that
23 masses of that size can be difficult for
24 physicians to detect?

25 A That's correct. Suzanne Fletcher, who

1 was at UNC when I was there, did a rather elegant
2 study in which they made breast models and put
3 different size masses in those breast models and
4 were able to demonstrate that physicians, in some
5 cases, could not detect half centimeter to one
6 centimeter lesions, but half a centimeter was the
7 point at which they began to become detectable.

8 Q Assuming the accuracy of that study and
9 that opinion, is it possible that, even if Mrs.
10 Feathers had been referred to a surgeon, the
11 surgeon would not have been able to palpate the
12 mass?

13 A That is a possibility.

14 Q How can we know one way or the other
15 whether or not the surgeon would have been able
16 to palpate the mass?

17 A Well, we can't know one way or the
18 other.

19 Q And if a surgeon could not palpate the
20 mass, he or she couldn't take any further steps;
21 i.e., a needle biopsy or anything along those
22 lines?

23 A Well, the appropriate response is that,
24 number one, you're correct, a needle biopsy
25 cannot be done unless the mass can actually be

1 palpated; but number two, if the patient is
2 complaining about it, you sit with the patient --
3 and this can either be the surgeon or the general
4 practitioner -- and say we can't find it but
5 you're feeling something, we recommend self-exam
6 and we recommend you come back in two months, a
7 different time of your period, and we will, if
8 necessary -- and we will re-examine you and we
9 may repeat a mammography in three months.

10 Q Does the standard of care under these
11 circumstances require a follow-up visit in two
12 months?

13 A Well, I hesitate to say specifically
14 two months, but certainly the standard of care
15 does require that degree of close follow-up.

16 Q Okay. Well, what's the outline of when
17 the standard of care would require Mrs. -- or
18 Dr. Brown to -- let me start all over again. I'm
19 getting jumbled here.

20 What does the standard of care require
21 as relates to follow-up for Mrs. Feathers?

22 A Two to three months would be
23 appropriate and within --

24 Q Three months for a follow-up visit?

25 A Correct.

1 Q How about a mammogram?

2 A Same thing.

3 Q As we sit here today, can you say to a
4 reasonable degree of medical probability that had
5 Mrs. Feathers returned in two to three months for
6 a follow-up, which would bring us to April or May
7 of 1999, that the mass would have been palpable
8 or would have been evidenced on mammogram?

9 A Well, the mass clearly grew in the 16
10 months, and the tumor, itself, was 1.7
11 sonometers, and the mass was significantly larger
12 than that, which is fairly typical for the growth
13 of breast cancers.

14 So, yes, I think that within a
15 reasonable degree of medical probability, it
16 would have been palpable very shortly after
17 February of '99.

18 Q And detectable on mammogram?

19 A It was not detectable on mammogram
20 until rather late. I believe she had a mammogram
21 in October of '99 that was read as normal. And
22 again, there's a 15 percent false negative rate,
23 so that's not -- not surprising. But the
24 mammogram, nonetheless, should have been repeated
25 at that second follow-up visit.

1 Q Is there ever a situation where a
2 patient detects a mass, the physician does not,
3 and referral to a surgeon is not necessary?

4 A Well, you're implying that the patient
5 is -- does not have a cancer in that situation.
6 And the problem is that we don't know that the
7 patient doesn't have a cancer.

8 If the physician cannot palpate the
9 mass, it is not, as I said, imperative that the
10 patient immediately be referred to a surgeon, but
11 it is imperative, if that physician chooses not
12 to refer, that the physician then assumes the
13 responsibility of that very close follow-up.

14 Q Okay. So while a surgical consult may
15 not be necessary, under these circumstances, you
16 must always have close follow-up every two to
17 three months?

18 A Correct.

19 Q Do you have any other opinions
20 regarding the standard of care in this case?

21 A No, I do not.

22 Q Okay. So just so I can summarize, make
23 sure I'm clear, it's your opinion that either
24 Dr. Brown either should have sent Mrs. Feathers
25 for surgical evaluation or should have arranged

1 for closer follow-up every two to three months
2 with a follow-up visit and mammogram?

3 A Correct.

4 Q Okay. Now, in your report, you state
5 that it's your belief that the tumor identified
6 in June of 2000 is the same mass that was
7 reported by Mrs. Feathers in February of 1999.

8 A That's correct.

9 Q What's your basis for that opinion?

10 A It's in the same location, the patient
11 feels it's the same mass throughout, and it is
12 very close to an area of DCIS when it's
13 ultimately diagnosed. So clearly that area of
14 the breast is involved in the development of this
15 tumor.

16 Q All right. I must be -- I'm confused
17 about this. Are you -- are you familiar with
18 Dr. Brown's note where she describes the mass as
19 existing at 11:00?

20 A Correct.

21 Q And on a mammogram in June of 2000,
22 it's reported the -- the mass is reported at
23 2:00?

24 A Correct. Now, there I would differ
25 with Dr. Levitan, although he says it's in a

1 different quadrant, which is technically correct
2 because of the way that we normally draw a
3 compass, it's actually within the same quadrant.
4 11:00 and 2:00 are only three hours apart.

5 Q There's four hours in each quadrant?
6 Is that what you're saying?

7 A Correct.

8 Q But regardless of the quadrant
9 analysis, don't those describe a mass in two
10 separate areas on the breast?

11 A Not necessarily. We have the same
12 problem with specific localization that we do
13 with the ability to palpate a mass. What feels
14 different to the patient -- what the patient
15 perceives may not be what the physician
16 perceives, and it may not be what is true in
17 reality.

18 The tumor may be starting at 2:00 but
19 growing medially and, therefore, actually be a
20 little bit to the -- to the right of where it's
21 initially detected.

22 Finally, the mass, itself, may -- may
23 actually be spreading over a larger area than
24 what the pathology is showing.

25 What I mean by that is that the way --

'1 the way you're detecting the mass is to rub the
2 breast back and forth. And so -- I've not
3 examined the patient, I don't know whether she
4 has large breasts or small breasts, but you can
5 appreciate that breasts are mobile and the tumor,
6 itself, is mobile. So as you move your -- your
7 fingers across the breast, you may feel it in a
8 different area than it actually is located.

9 In any event, I think the likelihood,
10 if you use the term more likely than not, more
11 likely than not, what the patient felt, continued
12 to think she felt, and what was ultimately
13 diagnosed were one in the same lesion.

14 Q And your basis for that is that it's
15 within the different quadrant and there's medical
16 or anatomic explanations as to why you could have
17 two different descriptions?

18 A That's correct.

19 Q You mentioned DCIS in your basis for
20 your opinion that it's the same tumor. I'm not
21 sure I understand how that fits in.

22 A Well, DCIS is a precursor, and more of
23 this tumor is composed of DCIS than actual
24 infiltrating carcinoma. That tends to be a
25 relatively non-lumpy lesion, although it can

1 appear as a lump. It can be more difficult to
2 detect. And she probably had more of an element
3 of DCIS in February of '99 than she had in June
4 and July of 2000.

5 Q Is it possible that the tumor
6 identified in June of 2000 is a different mass
7 than that was reported in February of 1999?

8 A Anything's possible, but I think it's
9 less likely than not.

10 Q Let's talk about mammograms for a
11 minute. I think you stated earlier that there's
12 a 15 percent false negative rate in mammograms?

13 A Correct.

14 Q Is there a particular type of mass
15 that's less detectable with mammograms?

16 A Lobular carcinoma and DCIS may be a
17 little more difficult to detect.

18 Q Have you seen the mammograms in this
19 case?

20 A I have not seen the mammograms, only
21 the written reports.

22 a I take it you don't have any criticisms
23 of the mammography techs or the radiologist who
24 interpreted the mammograms?

25 A No, I do not.

1 Q Let's turn to the survival rates that
2 you speak about in your report, if you would,
3 Dr. Ozer.

4 A Sure.

5 Q Now, you brought with you what
6 publication today?

7 A This is the -- it's a pocket version,
8 but has the same information, the AJCC Staging
9 Manual for all tumors. And I can get the court
10 reporter to make a copy, but breast cancer begins
11 on page 161.

12 Q Okay. What edition is that?

13 A This is the undefined edition. Hang on
14 a moment.

15 Q Now, how does that -- the -- the AJCC
16 is the staging manual?

17 A Correct.

18 Q Does the NCCN just relate to treatment
19 guidelines?

20 A That's right. NCCN publishes treatment
21 guidelines. The AJCC staging system is the one
22 -- it's the American Joint Committee on Cancer,
23 and it's the one that we all adhere to.

24 I, for example, chair the cancer
25 committee of the university, and we have to meet

1 their staging guidelines and use their staging
2 system.

3 And the only identification here is it
4 says copyright 1993. It doesn't have a date
5 other than that or a volume other than that.
6 Q I'd appreciate it, as you offered
7 earlier, if you could have the court reporter
8 just make a copy of the applicable pages for me.

9 A I'll do that.

10 Q Now, you've read Dr. Levitan's report
11 in this case?

12 A Yes, I have.

13 Q Does he follow the AJCC staging manual?

14 A Yes, he does, although I would dispute
15 his survival statistics by a hair. But
16 otherwise, he does.

17 Q All right. Well, let's talk about the
18 survival statistics. And I assume you're taking
19 what you've listed in your report straight out of
20 the staging manual?

21 A I am.

22 Q All right. And it's your opinion that
23 when she was diagnosed in June of 2000, her
24 cancer was Stage IIA; correct?

25 A That's correct.

1 Q And then that stage, based upon the
2 type of tumor it was, gives her a greater than 80
3 percent five-year survival rate?

4 A It gives -- yes, it gives her a 80 --
5 at five years, it's 85 percent according to this
6 document.

7 Q Okay. Because your report -- and I'm
8 not trying to dance on a pin head here, but your
9 report says 80 percent five-year survival.

10 A Correct.

11 Q But 85 percent would be more accurate?

12 A 85 percent would be the more precise
13 number, that's correct.

14 Q That was a better way to put it. Now,
15 Dr. Levitan uses 88 percent. You don't agree
16 with that?

17 A I won't argue over 1 or 2 percent. The
18 only point I'll make is that for years four, five
19 and six from diagnosis, the differential between
20 Stage 1 and Stage IIA is 10 percent. And the
21 numbers provided for five years are 95 percent
22 for Stage I and 85 percent for Stage II. So he's
23 a tad optimistic or pessimistic, depending on
24 your point of view, when he says 3 to 5 percent.
25 I'm saying it's closer to 10 percent.

1 Q All right. Now, for a Stage I cancer,
2 you've stated that the survival rate is greater
3 than 95 percent for five years.

4 A Correct.

5 Q Is that the most precise number?

6 A Yes, that is more precise.

7 Q Okay. And you're taking issue somewhat
8 with Dr. Levitan's differential between the
9 survival rates for those two stages of cancer?

10 A Yes. I mean, he says 3 to 5, I'm
11 saying 10. That wouldn't matter much in lots of
12 measurements, but if it's the amount of time you
13 spend at home with your wife, it would be a big
14 difference.

15 Q Okay. Now, Dr. Levitan figures,
16 though, that he lists -- for the survival rates
17 for a Stage I cancer, he lists 91 percent at five
18 years, 85 percent at 10 years. Do those not come
19 from the staging manual?

20 A I couldn't tell you where they come
21 from.

22 Q Okay. And that's why you take issue
23 with them?

24 A Correct.

25 Q All right. Do you know Dr. Levitan at

1 all?

2 A I know of him, yes.

3 Q Have you and he been on opposite sides
4 of the fence before in a medical malpractice
5 expert situation?

6 A That, I really couldn't tell you.

7 Q Okay. Just curious. Do you agree with
8 Dr. Levitan's statement that the prognosis for a
9 patient with a single microscopically involved
10 lymph node approximates that of a patient with a
11 negative node?

12 A I would agree with that statement, with
13 the caveat that every time you add a lymph node,
14 your prognosis is impacted just slightly. But I
15 think that's what he means when he says
16 approximates.

17 Q Okay. If Mrs. Feathers had been
18 diagnosed in 1999, it's your opinion that she
19 would have had a Stage I cancer; right?

20 A Correct.

21 Q All right. Assuming that to be true,
22 diagnosing -- why I can't get that -- February of
23 1999 -- I can't get that date in my head.
24 Assuming a diagnosis in February of 1999 Stage I,
25 would she still have needed a lumpectomy?

1 A Yes.

2 Q Chemotherapy?

3 A I believe she would of needed
4 chemotherapy. There are some who might not
5 recommend that; however, the fact that she's a
6 rather -- rather low on her ER/PR expression, and
7 more importantly the fact that she's HER-2
8 positive, to me is sufficient to warrant
9 chemotherapy. So, yes, I would have recommended
10 chemotherapy for her as a Stage I.

11 Q Okay. Radiation as well?

12 A Yes.

13 a Tamoxifen therapy?

14 A Yes.

15 Q So to kind of speak in lawyer terms
16 here, this delay -- I'll have to say alleged
17 delay in diagnosis resulted simply in a 10
18 percent differential in her survival rate?

19 A With the -- if you took out the simply,
20 I would agree with that statement.

21 Q Yeah. That's not -- let me -- let me
22 put it in -- take out that word simply.

23 So looking at this case on a whole, the
24 alleged delay in diagnosis of the breast cancer
25 for approximately a year or 14 months led solely

1 to a decrease -- a 10 percent decrease in the
2 survival rate?

3 A That's correct.

4 Q Do you follow the NCCN guidelines when
5 it comes to treatment regimens for patients?

6 A We certainly review them and use them
7 as a helpful guide. We don't blindly follow
8 them, no.

9 Q So they don't necessarily provide a
10 standard of care?

11 A They provide a framework but not a
12 standard of care.

13 Q Okay. Can we agree that the properties
14 of Mrs. Feathers's cancer did not change between
15 1999 and the year 2000?

16 A I would agree, again, in the more
17 likely than not context that that's true.

18 Q Okay. Do you have an opinion on how
19 long it took Mrs. Tumor -- Mrs. Tumor. It's been
20 a long day, Doctor.

21 Do you have an opinion on how long it
22 took Mrs. Feathers's tumor to grow to 1.7
23 centimeters?

24 A I think the natural history of breast
25 cancer can be anywhere from one to five years.

In this particular case, it's not a overly aggressive or rapidly growing tumor, but I think its origin almost certainly predates '99. I'm not certain whether it predates '98. But I think a ballpark would be an origin at some point in early '98, winter of '98.

7 ' If a physician will have difficulty
8 palpating a mass smaller than a half a
9 centimeter, would you expect a patient to be able
10 to palpate that size of a mass?

11 : Well, everyone varies quite a bit. I
12 think patients are more attuned to breast
13 self-examination than are physicians. And I
14 would estimate that the majority of masses that
15 we detect, "we" being -- that we see in the
16 clinic, rather, are brought to our attention by
17 patients rather than physicians being able to
18 pick up a mass when they do a physical exam.

19 So it wouldn't surprise me that a
20 patient would be able to detect a pea-sized mass
21 for the first time.

22 Q Just because they have a better sense
23 of their own body and the changes?

24 A Exactly.

25 Q Okay. Do you agree with the statement

1 that the highest risk of recurrence of breast
2 cancer is in the first two years?

3 A Actually, I -- I can give you the
4 numbers for that, and the answer to that is no.
5 It might be for Stage IV disease but it's not for
6 Stage 0, I or IIA if we look at the statistics.

7 For example, the majority of relapses
8 have occurred in year five rather than in year
9 one or two for those early stages.

10 Q What are you looking at? Are you
11 looking at the journal?

12 A I'm looking at the -- at page 165. And
13 these are the curves. And just to give you an
14 example of how a rapidly recurrent tumor would
15 be, if you look at Stage IV, which is metastatic,
16 these are survival, which is an equivalent to
17 relapse, there is a 60 percent roughly -- 62
18 percent survival after one year with metastatic
19 disease.

20 So that means that 38 percent have
21 relapsed and progressed and the patients have
22 died. The majority are dead after two years with
23 metastatic disease.

24 If you look now at Stage IIA, which is
25 what she's ultimately diagnosed with, her curve

1 demonstrates that after two years, the decrement
2 from 100 percent is, oh, 10 percent, maybe. So
3 10 percent of relapsed have progressed and died
4 in the first two years, and then the -- at five
5 years -- remember it's now down to 85 percent.

6 Q Right.

7 A So 50 percent of the total 15 percent
8 deaths is 7 percent, and the 7 percent has
9 occurred by about three years. So you have to go
10 to four years before the majority of patients are
11 relapsing and dying.

12 Q That curve you're looking at, does that
13 only consider patients who have a recurrence and
14 die within five years?

15 A Actually it considers -- it's an intent
16 -- what we call an intent to treat analysis which
17 is deaths from all causes which is what we use
18 for all our statistics. So it includes patients
19 that die of heart attacks and other things, all
20 thought to be related.

21 Q Does it include a patient that died of
22 a heart attack but didn't have a recurrence of
23 cancer?

24 A Yes, it does. That's how we do all our
25 statistics.

1 Q Okay. Because that statement I gave
2 you, that the highest risk of recurrence is
3 within the first two years, came from Dr. Green
4 during his deposition as Mrs. Feathers's treating
5 physician. I'm not quite sure where he gets that
6 statistic as well.

7 A I'm not sure, either. And I would
8 dispute that for this kind of a small, slowly
9 growing tumor.

10 What I tell my patients is that -- the
11 analogy I give them is that it's -- if you take
12 the gondola to the top of the mountain and a
13 group of bad skiers gets off, some of them are
14 going to fall early and some of them will make it
15 most of the way down, but there will be people
16 falling off all the way or falling all the way.

17 Q Dr. Ozer, I think I'm just about
18 through. I want to take a minute to talk to you
19 a little bit about your experience as an expert
20 in medical malpractice litigation.

21 A Certainly.

22 Q You served as an expert in the past
23 before?

24 A Yes, I have.

25 Q When did you first start doing this

1 type of work?

2 A The very first case I did was in 1984.

3 Q Do you know about how many cases you
4 review a year?

5 A I've now reviewed about 55, 58 cases,
6 and it's probably one every six weeks, I would
7 guess.

8 Q So you've done a total of 55 to 58
9 since 1984?

10 A Correct,

11 Q Any estimate on the breakdown of
12 plaintiff versus defense?

13 A About 80 percent have been for the
14 plaintiff and 20 percent for the defense.

15 Q What do you charge to review records?

16 A I charge an initial retainer fee of
17 \$1,500 and then \$350 an hour thereafter.

18 Q How about for deposition? I probably
19 should have asked this sooner. What's the charge
20 for deposition?

21 A I can give you any number I want now.

22 Q Oh, no. I'll pull an old transcript.

23 A It's \$1,500 as a flat fee for a
24 deposition because I have to block out the
25 afternoon.

'1 Q Okay. How about trial testimony?

2 A For trial testimony, I charge \$3,000 a
3 day, plus expenses.

4 Q Do you have plans to come -- actually
5 it's to Lake County, in Mentor, Ohio, in
6 September for the trial of this case?

7 A If I'm asked, I will.

8 Q Okay. I'm just going to take a minute
9 to take a look at my notes. I think I'm just
10 about through.

11 What's the risk of breast cancer in the
12 overall population?

13 A One in eight.

14 Q One in eight?

15 A Among women, it's one in eight.

16 Q Does that change depending on the
17 patient's age?

18 A It does. The lifetime risk is one in
19 eight. The majority of breast cancer occurs in
20 women over the age of 55 or 60.

21 Q What's the percent of one in eight?
22 Never mind.

23 Do you have an opinion whether or not
24 Mary Ann Feathers requires breast reconstruction?

25 A I saw a discussion in there regarding

1 that. The majority of women who have a
2 lumpectomy will -- let me take that back. It
3 varies a lot with the individual. It also varies
4 with the size of the woman's breasts.

5 Obviously if you have larger breasts
6 and you have a lumpectomy, the surgeon is able to
7 do a little bit of reconstruction at the time of
8 the lumpectomy without much of a difference.

9 If your breasts are very small, a
10 lumpectomy has a greater impact. And it depends
11 a lot on whether the woman doesn't like having
12 asymmetry. I've not seen her, I've not examined
13 her, I've not spoken to her. It would be her
14 call as to whether she needed reconstruction.

15 Q Is there a time frame you typically
16 wait before you do a reconstruction?

17 A It can be done immediately. However,
18 in someone who is getting chemotherapy and
19 radiation therapy, it is far preferable to wait
20 at least until all of that is complete and then
21 give them another four to six months before you
22 do reconstruction.

23 MS. REID: Dr. Ozer, I don't think I
24 have any more questions. I appreciate your time.

25 THE WITNESS: Certainly.

1 MS. REID: If you would just have the
2 court reporter make a copy of those letters and
3 the pages of the manual --

4 THE WITNESS: Will do.

5 MS. REID: -- I would appreciate it.

6 THE WITNESS: Okay.

7 MS. REID: Do you want to talk to Donna
8 for a minute?

9 MS. KOLIS: Doctor, are you going to
10 read the deposition or would you like to waive
11 the reading?

12 THE WITNESS: I'm happy with it if
13 you're okay with it.

14 MS. KOLIS: I'm fine with it. Just
15 indicate for the court reporter that I would like
16 a copy of the transcript.

17 THE WITNESS: Okay.

18 (Deposition concluded at 2:48 p.m.)
19
20
21
22
23
24
25

C E R T I F I C A T E

STATE OF OKLAHOMA)
) S S :
COUNTY OF OKLAHOMA)

I, ELIZABETH CATJDILL, CSR in and for
the State of Oklahoma, certify that HOWARD OZER,
M.D. was by me sworn to testify the truth; that
the above and foregoing deposition was taken by
me in stenotype and thereafter transcribed and is
a true and correct transcript of the testimony of
the witness; that the deposition was taken on
JULY 24, 2002 at 2:03 p.m. in Oklahoma City,
Oklahoma; that I am not an attorney for or a
relative of either party, or otherwise interested
in this action.

Witness my hand and seal of office on
this 31st day of July, 2002.

ELIZABETH CAUDILL
Oklahoma Certified Shorthand Reporter
Certificate No. 00161
-----Exp. Date: December 31, 2002-----

ELIZABETH CAUDILL, CSR, RMR, CRR
CSR No. 161

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DONNA TAYLOR-KOLIS
MICHAEL J. SKINDELL

February 15, 2001



Via Federal Express Standard Overnight

Howard Ozer, M.D., Ph.D.
1516 Camden Way
Oklahoma City, OK 73116

RE: Mary Ann Feathers

Dear Dr. Ozer:

Thank you for agreeing to review the matter of ~~Mary~~ Ann Feathers regarding a potential medical negligence claim. Enclosed for your review are the following records:

1. Family Care Associates (Drs. Hackett & Brown)
2. Armin Green, M.D. (Ireland Cancer Center)
3. John Dorsky, M.D.
4. Lake Hospital System

Also enclosed is a check in the amount of \$1,500.00 representing a retainer for your services rendered in this matter.

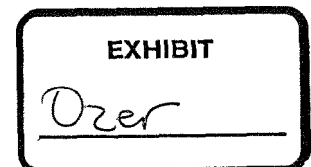
I would like to request that you call me upon completion of your review to discuss your opinions in this matter. If at all possible, I would like to know your opinions within 4 weeks of your receipt of this material.

Again, thank you for evaluating this matter, and I look forward to hearing from you.

Sincerely yours,

Donna Taylor-Kolis

DTK:vjw
Enclosures



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DONNA TAYLOR-KOLIS

August 26, 2001

Howard Ozer, M.D., Ph.D.
1516 Camden Way
Oklahoma City, OK 73116

RE: MaryAnn .Feathers

Dear Dr. Ozer:

In connection with the above-referenced matter, please call my office at your *earliest* convenience. The deposition of Dr. Brown will be going forward on September 11, 2001 and I would like to speak with you before that date regarding some issues to be addressed at the deposition.

Sincerely yours,



Donna Taylor-Kolis

DTK: sla

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January 24, 2002

Writer's Direct Dial
(216) 621-0070

Howard Ozer, M.D., Ph.D.
1516 Camden Way
Oklahoma City, OK 73116

RE: Mary Ann Feathers

Dear Dr. Ozer:

In connection with the above-captioned client, please be advised that this matter has been set for trial on **April 29, 2002 @ 8:30 am**. I need to make arrangements to go over your trial testimony with you should this case go forward. Upon receipt of this correspondence, please call me at the direct dial number listed above.

Sincerely yours,



Donna Taylor-Kolis

DTK:sla

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March 20, 2002

Howard Ozer, M.D., Ph.D.
1516 Camden Way
Oklahoma City, OK 73116

RE: MaryAnn Feathers

Dear Dr. Ozer:

The above-captioned lawsuit has now been re-set for **September 26, 2002**. Please have a staff member call my office to confirm that you would be available to testify on September 27th or the morning of September 30th.

Additionally, defense counsel would like to take your deposition in June of 2002 and I would ask that you or a staff member contact our office to advise us of your dates of availability.

Sincerely yours,



Donna Taylor-Kolis

DTK:sla
Enclosure

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DONNA TAYLOR-KOLIS

January 24, 2002

Howard Ozer, MD., Ph.D.
1516 Camden Way
Oklahoma City, OK 73116

RE: Mary Ann Feathers

Dear Dr. Ozer:

Enclosed please find the deposition testimony of Mary Ann Feathers' subsequent treating physician, Armin Green, M.D. I would like for you to read it to determine whether or not you are in agreement with the opinions that he has in this matter. when you concluded reading it, please give me a call so we can discuss the same.

Sincerely yours,


Donna Taylor-Kolis

DTK:jme
Enclosure

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April 22, 2002

Howard Ozer, M.D., Ph.D.
1516 Camden Way
Oklahoma City, OK 73116

RE: May Ann Feathers

Dear Dr. Ozer:

This case has yet again been reset for trial. The new trial date is September 16, 2002. At this point, the Defendant has not made an attempt to settle the claim and we do not know whether or not they will.

Enclosed please find the expert reports for Dr. Brown.

Counsel for Dr. Brown would like to take your discovery deposition sometime in late June, or early July. Please call my office with available dates.

Sincerely yours,


Donn Taylor-Kolis
a Taylor-Kolis

DTK/jme

HANDBOOK FOR **staging** OF Cancer

From the
MANUAL FOR STAGING OF CANCER,
Fourth Edition

American Joint Committee
on Cancer

J. B. Lippincott Company

BREAST

25

Breast

le bus
ssified

: desmoplastic variant also exists.
lanomas are identified according to site (e.g., mucosal, ocular, vaginal,
urethral). The staging classification described in this chapter applies
to those arising in the skin.

IOGRAPHY

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Clark WH Jr: The histogenesis and biological behavior of primary malignant melanoma of the skin. *Cancer Res* 29:705-717, 1969
Fitzpatrick AW, Rodriguez-Sains RS, Rigel DS, et al: "Small" melanomas: Evaluation of prognostic variables to diameter of superficial spreading melanomas. *J Dermatol Surg Oncol* 8:765-770, 1982

- C50.0 Nipple
- C50.1 Central portion
- C50.2 Upper-inner quadrant
- C50.3 Lower-inner quadrant
- C50.4 Upper-outer quadrant
- C50.5 Lower-outer quadrant
- C50.6 Axillary tail
- C50.8 Overlapping lesion
- C50.9 Breast, NOS

The following TNM definitions and stage groupings for carcinoma of the breast are the same for the AJCC and the UICC/TNM projects. This staging system for carcinoma of the breast applies to infiltrating and *in situ* carcinomas. Microscopic confirmation of the diagnosis is mandatory and the histologic type and grade of carcinoma should be recorded.

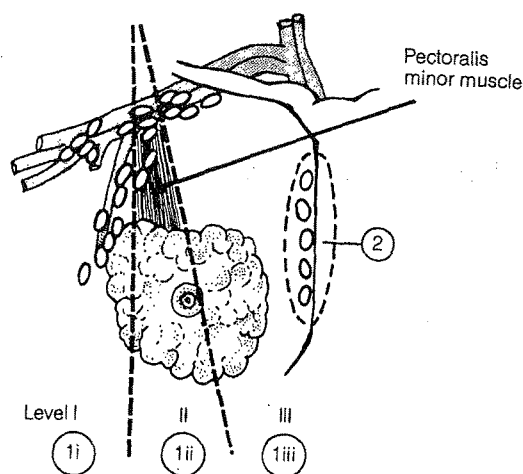
ANATOMY

Primary Site. Situated on the anterior chest wall, the mammary gland is composed of glandular tissue within a dense fibroareolar stroma. The glandular tissue consists of approximately 20 lobes, each of which terminates in a separate excretory duct in the nipple.

Regional Lymph Nodes. The breast lymphatics drain by way of three major routes: axillary, transpectoral, and internal mammary. Intramammary lymph nodes are considered with the axillary lymph nodes for staging purposes. Metastases to any other lymph nodes—including supraclavicular, cervical, and contralateral internal mammary nodes—are considered distant (M1). (Please refer to diagram.) The regional lymph nodes are:

- (1) Axillary (ipsilateral): interpectoral (Rotter's) nodes and lymph nodes along the axillary vein and its tributaries, which may be divided into the following levels:
 - (i) Level I (low-axilla): lymph nodes lateral to the lateral border of the pectoralis minor muscle
 - (ii) Level II (mid-axilla): lymph nodes between the medial and lateral borders of the pectoralis minor muscle and the interpectoral (Rotter's) lymph nodes
 - (iii) Level III (apical axilla): lymph nodes medial to the medial margin of the pectoralis minor muscle, including those designated as subclavicular, infraclavicular, or apical.

REGIONAL LYMPH NODES



Note: Intramammary lymph nodes are coded as axillary lymph nodes.

- (2) Internal mammary (ipsilateral): lymph nodes in the intercostal spaces along the edge of the sternum in the endothoracic fascia. Any other lymph node metastasis is coded as a distant metastasis (M1), including supraclavicular, cervical, or contralateral internal mammary lymph nodes.

Metastatic Sites. All distant visceral sites are potential sites of metastases. The four major sites of involvement are bone, lung, brain, and liver, but this widely metastasizing disease has been found in almost every remote site.

RULES FOR CLASSIFICATION

Clinical Staging. Clinical staging includes physical examination, with careful inspection and palpation of the skin, mammary gland, and lymph nodes (axillary, supraclavicular, and cervical), pathologic examination of the breast or other tissues, and imaging to establish the diagnosis of breast carcinoma. The extent of tissues examined pathologically for clinical staging is less than that required for pathologic staging (see Pathologic Staging). Appropriate operative findings are elements of clinical staging, including the size of the primary tumor and chest wall invasion and the presence or absence of regional or distant metastasis.

Pathologic Staging. Pathologic staging includes all data used for clinical staging and surgical resection as well as pathologic examination of the primary carcinoma, including not less than excision of the primary carcinoma with no tumor in any margin of resection by gross pathologic examination.

Breast

A case can be included in the pathologic stage if there is only microscopic, but not gross, involvement at the margin. If there is tumor in the margin of resection by gross examination, it is coded as TX, because the extent of primary tumor cannot be assessed. Resection of at least the low axillary lymph nodes (Level I)—that is, those lymph nodes located lateral to the lateral border of the pectoralis minor muscle—should be carried out. Such a resection ordinarily will include six or more lymph nodes. Metastatic nodules in the fat adjacent to the mammary carcinoma, without evidence of residual lymph node tissue, are considered regional lymph node metastases.

CLASSIFICATION

primary Tumor

The clinical measurement used for classifying the primary tumor (T) should be the one judged most accurate (e.g., physical examination or mammogram). Pathologically, the tumor size for classification (T) is a measurement of the invasive component. For example, if there is a large in situ component (4 cm) and a small invasive component (0.5 cm), the tumor is classified as T1a. The size of the primary tumor should be measured before any tissue is removed for special studies, such as for estrogen receptors.

Multiple Simultaneous Ipsilateral Primary Cancers

The following guidelines should be used when classifying multiple simultaneous ipsilateral primary (infiltrating, grossly measurable) carcinomas. These criteria do not apply to one grossly detected tumor associated with multiple separate microscopic foci.

1. Use the largest primary carcinoma to classify T.
2. Enter into the record that this is a case of multiple simultaneous ipsilateral primary carcinomas. Such cases should be analyzed separately.

Simultaneous Bilateral Breast Carcinomas

Each carcinoma is staged separately.

Inflammatory Carcinoma

Inflammatory carcinoma is a clinicopathologic entity characterized by diffuse brawny induration of the skin of the breast with an erysipeloid edge, usually without an underlying palpable mass. Radiologically, there may be a detectable mass and characteristic thickening of the skin over the breast. This clinical presentation is due to tumor embolization of dermal lymphatics. The tumor of inflammatory carcinoma is classified as T4d.

Paget's Disease of the Nipple

Paget's disease of the nipple without an associated tumor mass (clinical) or invasive carcinoma (pathologic) is classified as Tis. Paget's disease with a

nonstrable mass (clinical) or an invasive component (pathologic) is classified according to the size of the tumor mass or invasive component.

Skin of the Breast

Dimpling of the skin, nipple retraction, or any other skin change except those described under T4b and T4d may occur in T1, T2, or T3 without changing the classification.

Chest Wall

The chest wall includes the ribs, intercostal muscles, and serratus anterior muscle but not the pectoral muscle.

DEFINITION OF TNM

Primary Tumor (T)

Definitions for classifying the primary tumor (T) are the same for clinical and for pathologic classification. The telescoping method of classification can be applied. If the measurement is made by physical examination, the examiner will use the major headings (T1, T2, or T3). If other measurements, such as mammographic or pathologic, are used, the examiner can use the telescoped subsets of T1.

TX Primary tumor cannot be assessed

T0 No evidence of primary tumor

Tis Carcinoma *in situ*: intraductal carcinoma, lobular carcinoma in situ, or Paget's disease of the nipple with no tumor

T1 Tumor 2 cm or less in greatest dimension

T1a 0.5 cm or less in greatest dimension

T1b More than 0.5 cm but not more than 1 cm in greatest dimension

T1c More than 1 cm but not more than 2 cm in greatest dimension

T2 Tumor more than 2 cm but not more than 5 cm in greatest dimension

Tumor more than 5 cm in greatest dimension

Tumor of any size with direct extension to chest wall or skin

T4a Extension to chest wall

T4b Edema (including peau d'orange) or ulceration of the skin of the breast or satellite skin nodules confined to the same breast

T4c Both (T4a and T4b)

T4d Inflammatory carcinoma (See the definition of inflammatory carcinoma in the introduction.)

Note: Paget's disease associated with a tumor is classified according to the size of the tumor.

Regional Lymph Nodes (N)

NX Regional lymph nodes cannot be assessed (e.g., previously removed)

N0 No regional lymph node metastasis

BREAST CANCER

SURVIVAL ACCORDING TO AJCC STAGE

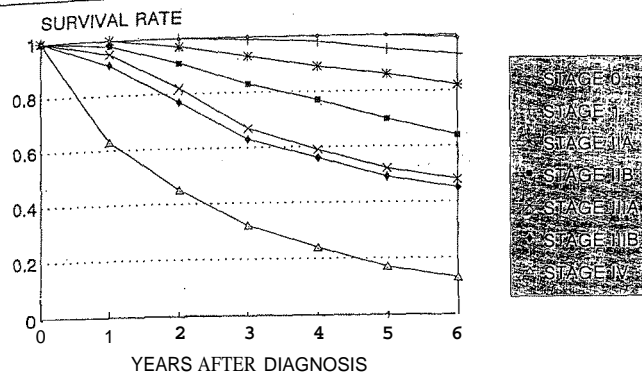


Fig. 25-1. Relative survival rates according to stage of disease. Data taken from 50,834 patients listed in the Surveillance, Epidemiology, and End Results Program of the National Cancer Institute. Patients were diagnosed between 1983 and 1987. Stage 0 represents 4,601 patients; Stage I, 16,519; Stage IIA, 14,692; Stage IIB, 8,283; Stage IIIA, 1,656; Stage IIIB, 1,389; and Stage IV, 3,694.

N1 Metastasis to movable ipsilateral axillary lymph node(s)

N2 Metastasis to ipsilateral axillary lymph node(s) fixed to one another or to other structures

N3 Metastasis to ipsilateral internal mammary lymph node(s)

Pathologic Classification (pN)

pNX Regional lymph nodes cannot be assessed (e.g., previously removed, or not removed for pathologic study)

pN0 No regional lymph node metastasis

pN1 Metastasis to movable ipsilateral axillary lymph node(s)

pN1a Only micrometastasis (none larger than 0.2 cm)

pN1b Metastasis to lymph node(s), any larger than 0.2 cm

pN1bi Metastasis in one to three lymph nodes, any more than 0.2 cm and all less than 2 cm in greatest dimension

pN1bii Metastasis to four or more lymph nodes, any more than 0.2 cm and all less than 2 cm in greatest dimension

pN1biii Extension of tumor beyond the capsule of a lymph node metastasis less than 2 cm in greatest dimension

pN1biv Metastasis to a lymph node 2 cm or more in greatest dimension

- pN2 Metastasis to ipsilateral axillary lymph node(s) that are fixed to one another or to other structures
- pN3 Metastasis to ipsilateral internal mammary lymph node(s)

Distant Metastasis (M)

- MX Presence of distant metastasis cannot be assessed
- M0 No distant metastasis
- M1 Distant metastasis (includes metastasis to ipsilateral supraclavicular lymph node(s))

STAGE GROUPING			
Stage 0	Tis	N0	M0
Stage I	T1	N0	M0
Stage IIA	T2	N0	M0
Stage IIB	T2	N1	M0
Stage IIC	T2	N2	M0
Stage IIIA	T3	N0	M0
Stage IIIB	T3	N1	M0
Stage IIIC	T3	N2	M0
Stage IVA	T4	N0	M0
Stage IVB	T4	N1	M0
Stage IVC	T4	N2	M0
Stage V	Tis	N0	M1
Stage VI	T1	N0	M1
Stage VII	T2	N0	M1
Stage VIII	T2	N1	M1
Stage IX	T2	N2	M1
Stage X	T3	N0	M1
Stage XI	T3	N1	M1
Stage XII	T3	N2	M1
Stage XIII	T4	N0	M1
Stage XIV	T4	N1	M1
Stage XV	T4	N2	M1
Stage XVI	T5	N0	M1
Stage XVII	T5	N1	M1
Stage XVIII	T5	N2	M1
Stage XIX	T6	N0	M1
Stage XX	T6	N1	M1
Stage XXI	T6	N2	M1
Stage XXII	T7	N0	M1
Stage XXIII	T7	N1	M1
Stage XXIV	T7	N2	M1
Stage XXV	T8	N0	M1
Stage XXVI	T8	N1	M1
Stage XXVII	T8	N2	M1
Stage XXVIII	T9	N0	M1
Stage XXIX	T9	N1	M1
Stage XXX	T9	N2	M1
Stage XXXI	T10	N0	M1
Stage XXXII	T10	N1	M1
Stage XXXIII	T10	N2	M1
Stage XXXIV	T11	N0	M1
Stage XXXV	T11	N1	M1
Stage XXXVI	T11	N2	M1
Stage XXXVII	T12	N0	M1
Stage XXXVIII	T12	N1	M1
Stage XXXIX	T12	N2	M1
Stage XL	T13	N0	M1
Stage XLI	T13	N1	M1
Stage XLII	T13	N2	M1
Stage XLIII	T14	N0	M1
Stage XLIV	T14	N1	M1
Stage XLV	T14	N2	M1
Stage XLVI	T15	N0	M1
Stage XLVII	T15	N1	M1
Stage XLVIII	T15	N2	M1
Stage XLIX	T16	N0	M1
Stage L	T16	N1	M1
Stage LI	T16	N2	M1
Stage LII	T17	N0	M1
Stage LIII	T17	N1	M1
Stage LIV	T17	N2	M1
Stage LV	T18	N0	M1
Stage LVI	T18	N1	M1
Stage LVII	T18	N2	M1
Stage LVIII	T19	N0	M1
Stage LVIX	T19	N1	M1
Stage LX	T19	N2	M1
Stage LXI	T20	N0	M1
Stage LXII	T20	N1	M1
Stage LXIII	T20	N2	M1
Stage LXIV	T21	N0	M1
Stage LXV	T21	N1	M1
Stage LXVI	T21	N2	M1
Stage LXVII	T22	N0	M1
Stage LXVIII	T22	N1	M1
Stage LXIX	T22	N2	M1
Stage LXX	T23	N0	M1
Stage LXXI	T23	N1	M1
Stage LXXII	T23	N2	M1
Stage LXXIII	T24	N0	M1
Stage LXXIV	T24	N1	M1
Stage LXXV	T24	N2	M1
Stage LXXVI	T25	N0	M1
Stage LXXVII	T25	N1	M1
Stage LXXVIII	T25	N2	M1
Stage LXXIX	T26	N0	M1
Stage LXXX	T26	N1	M1
Stage LXXXI	T26	N2	M1
Stage LXXXII	T27	N0	M1
Stage LXXXIII	T27	N1	M1
Stage LXXXIV	T27	N2	M1
Stage LXXXV	T28	N0	M1
Stage LXXXVI	T28	N1	M1
Stage LXXXVII	T28	N2	M1
Stage LXXXVIII	T29	N0	M1
Stage LXXXIX	T29	N1	M1
Stage LXXXX	T29	N2	M1
Stage LXXXXI	T30	N0	M1
Stage LXXXXII	T30	N1	M1
Stage LXXXXIII	T30	N2	M1
Stage LXXXXIV	T31	N0	M1
Stage LXXXXV	T31	N1	M1
Stage LXXXXVI	T31	N2	M1
Stage LXXXXVII	T32	N0	M1
Stage LXXXXVIII	T32	N1	M1
Stage LXXXXIX	T32	N2	M1
Stage LXXXXX	T33	N0	M1
Stage LXXXXXI	T33	N1	M1
Stage LXXXXXII	T33	N2	M1
Stage LXXXXXIII	T34	N0	M1
Stage LXXXXXIV	T34	N1	M1
Stage LXXXXXV	T34	N2	M1
Stage LXXXXXVI	T35	N0	M1
Stage LXXXXXVII	T35	N1	M1
Stage LXXXXXVIII	T35	N2	M1
Stage LXXXXXIX	T36	N0	M1
Stage LXXXXXX	T36	N1	M1
Stage LXXXXXXI	T36	N2	M1
Stage LXXXXXXII	T37	N0	M1
Stage LXXXXXXIII	T37	N1	M1
Stage LXXXXXXIV	T37	N2	M1
Stage LXXXXXXV	T38	N0	M1
Stage LXXXXXXVI	T38	N1	M1
Stage LXXXXXXVII	T38	N2	M1
Stage LXXXXXXVIII	T39	N0	M1
Stage LXXXXXXIX	T39	N1	M1
Stage LXXXXXXX	T39	N2	M1
Stage LXXXXXXXI	T40	N0	M1
Stage LXXXXXXXII	T40	N1	M1
Stage LXXXXXXXIII	T40	N2	M1
Stage LXXXXXXXIV	T41	N0	M1
Stage LXXXXXXXV	T41	N1	M1
Stage LXXXXXXXVI	T41	N2	M1
Stage LXXXXXXXVII	T42	N0	M1
Stage LXXXXXXXVIII	T42	N1	M1
Stage LXXXXXXXIX	T42	N2	M1
Stage LXXXXXXXx	T43	N0	M1
Stage LXXXXXXXxi	T43	N1	M1
Stage LXXXXXXXxii	T43	N2	M1
Stage LXXXXXXXxiii	T44	N0	M1
Stage LXXXXXXXxiv	T44	N1	M1
Stage LXXXXXXXxv	T44	N2	M1
Stage LXXXXXXXxvi	T45	N0	M1
Stage LXXXXXXXxvii	T45	N1	M1
Stage LXXXXXXXxviii	T45	N2	M1
Stage LXXXXXXXxix	T46	N0	M1
Stage LXXXXXXXxx	T46	N1	M1
Stage LXXXXXXXxxi	T46	N2	M1
Stage LXXXXXXXxxii	T47	N0	M1
Stage LXXXXXXXxxiii	T47	N1	M1
Stage LXXXXXXXxxiv	T47	N2	M1
Stage LXXXXXXXxxv	T48	N0	M1
Stage LXXXXXXXxxvi	T48	N1	M1
Stage LXXXXXXXxxvii	T48	N2	M1
Stage LXXXXXXXxxviii	T49	N0	M1
Stage LXXXXXXXxxix	T49	N1	M1
Stage LXXXXXXXxxx	T49	N2	M1
Stage LXXXXXXXxxxi	T50	N0	M1
Stage LXXXXXXXxxxii	T50	N1	M1
Stage LXXXXXXXxxxiii	T50	N2	M1
Stage LXXXXXXXxxxiv	T51	N0	M1
Stage LXXXXXXXxxxv	T51	N1	M1
Stage LXXXXXXXxxxvi	T51	N2	M1
Stage LXXXXXXXxxxvii	T52	N0	M1
Stage LXXXXXXXxxxviii	T52	N1	M1
Stage LXXXXXXXxxxix	T52	N2	M1
Stage LXXXXXXXxxxx	T53	N0	M1
Stage LXXXXXXXxxxxi	T53	N1	M1
Stage LXXXXXXXxxxxii	T53	N2	M1
Stage LXXXXXXXxxxxiii	T54	N0	M1
Stage LXXXXXXXxxxxiv	T54	N1	M1
Stage LXXXXXXXxxxxv	T54	N2	M1
Stage LXXXXXXXxxxxvi	T55	N0	M1
Stage LXXXXXXXxxxxvii	T55	N1	M1
Stage LXXXXXXXxxxxviii	T55	N2	M1
Stage LXXXXXXXxxxxix	T56	N0	M1
Stage LXXXXXXXxxxix	T56	N1	M1
Stage LXXXXXXXxxxix	T56	N2	M1
Stage LXXXXXXXxxxxxi	T57	N0	M1
Stage LXXXXXXXxxxix	T57	N1	M1
Stage LXXXXXXXxxxix	T57	N2	M1
Stage LXXXXXXXxxxix	T58	N0	M1
Stage LXXXXXXXxxxix	T58	N1	M1
Stage LXXXXXXXxxxix	T58	N2	M1
Stage LXXXXXXXxxxix	T59	N0	M1
Stage LXXXXXXXxxxix	T59	N1	M1
Stage LXXXXXXXxxxix	T59	N2	M1
Stage LXXXXXXXxxxix	T60	N0	M1
Stage LXXXXXXXxxxix	T60	N1	M1
Stage LXXXXXXXxxxix	T60	N2	M1
Stage LXXXXXXXxxxix	T61	N0	M1
Stage LXXXXXXXxxxix	T61	N1	M1
Stage LXXXXXXXxxxix	T61	N2	M1
Stage LXXXXXXXxxxix	T62	N0	M1
Stage LXXXXXXXxxxix	T62	N1	M1
Stage LXXXXXXXxxxix	T62	N2	M1
Stage LXXXXXXXxxxix	T63	N0	M1
Stage LXXXXXXXxxxix	T63	N1	M1
Stage LXXXXXXXxxxix	T63	N2	M1
Stage LXXXXXXXxxxix	T64	N0	M1
Stage LXXXXXXXxxxix	T64	N1	M1
Stage LXXXXXXXxxxix	T64	N2	M1
Stage LXXXXXXXxxxix	T65	N0	M1
Stage LXXXXXXXxxxix	T65	N1	M1
Stage LXXXXXXXxxxix	T65	N2	M1
Stage LXXXXXXXxxxix	T66	N0	M1
Stage LXXXXXXXxxxix	T66	N1	M1
Stage LXXXXXXXxxxix	T66	N2	M1
Stage LXXXXXXXxxxix	T67	N0	M1
Stage LXXXXXXXxxxix	T67	N1	M1
Stage LXXXXXXXxxxix	T67	N2	M1
Stage LXXXXXXXxxxix	T68	N0	M1
Stage LXXXXXXXxxxix	T68	N1	M1
Stage LXXXXXXXxxxix	T68	N2	M1
Stage LXXXXXXXxxxix	T69	N0	M1
Stage LXXXXXXXxxxix	T69	N1	M1
Stage LXXXXXXXxxxix	T69	N2	M1
Stage LXXXXXXXxxxix	T70	N0	M1
Stage LXXXXXXXxxxix	T70	N1	M1
Stage LXXXXXXXxxxix	T70	N2	M1
Stage LXXXXXXXxxxix	T71	N0	M1
Stage LXXXXXXXxxxix	T71	N1	M1
Stage LXXXXXXXxxxix	T71	N2	M1
Stage LXXXXXXXxxxix	T72	N0	M1
Stage LXXXXXXXxxxix	T72	N1	M1
Stage LXXXXXXXxxxix	T72	N2	M1
Stage LXXXXXXXxxxix	T73	N0	M1
Stage LXXXXXXXxxxix	T73	N1	M1
Stage LXXXXXXXxxxix	T73	N2	M1
Stage LXXXXXXXxxxix	T74	N0	M1
Stage LXXXXXXXxxxix	T74	N1	M1
Stage LXXXXXXXxxxix	T74	N2	M1
Stage LXXXXXXXxxxix	T75	N0	M1
Stage LXXXXXXXxxxix	T75	N1	M1
Stage LXXXXXXXxxxix	T75	N2	M1
Stage LXXXXXXXxxxix	T76	N0	M1
Stage LXXXXXXXxxxix	T76	N1	M1
Stage LXXXXXXXxxxix	T76	N2	M1
Stage LXXXXXXXxxxix	T77	N0	M1
Stage LXXXXXXXxxxix	T77	N1	M1
Stage LXXXXXXXxxxix	T77	N2	M1
Stage LXXXXXXXxxxix	T78	N0	M1
Stage LXXXXXXXxxxix	T78	N1	M1
Stage LXXXXXXXxxxix	T78	N2	M1
Stage LXXXXXXXxxxix	T79	N0	M1
Stage LXXXXXXXxxxix	T79	N1	M1
Stage LXXXXXXXxxxix	T79	N2	M1
Stage LXXXXXXXxxxix	T80	N0	M1
Stage LXXXXXXXxxxix	T80	N1	M1
Stage LXXXXXXXxxxix	T80	N2	M1
Stage LXXXXXXXxxxix	T81	N0	M1
Stage LXXXXXXXxxxix	T81	N1	M1
Stage LXXXXXXXxxxix	T81	N2	M1
Stage LXXXXXXXxxxix	T82	N0	M1
Stage LXXXXXXXxxxix	T82	N1	M1
Stage LXXXXXXXxxxix	T82	N2	M1
Stage LXXXXXXXxxxix	T83	N0	M1
Stage LXXXXXXXxxxix	T83	N1	M1
Stage LXXXXXXXxxxix	T83	N2	M1
Stage LXXXXXXXxxxix	T84	N0	M1
Stage LXXXXXXXxxxix	T84	N1	M1
Stage LXXXXXXXxxxix	T84	N2	M1
Stage LXXXXXXXxxxix	T85	N0	M1
Stage LXXXXXXXxxxix	T85	N1	M1
Stage LXXXXXXXxxxix	T85	N2	M1
Stage LXXXXXXXxxxix	T86	N0	M1
Stage LXXXXXXXxxxix	T86	N1	M1
Stage LXXXXXXXxxxix	T86	N2	M1
Stage LXXXXXXXxxxix	T87	N0	M1
Stage LXXXXXXXxxxix	T87	N1	M1
Stage LXXXXXXXxxxix	T87	N2	M1
Stage LXXXXXXXxxxix	T88	N0	M1
Stage LXXXXXXXxxxix	T88	N1	M1
Stage LXXXXXXXxxxix	T88	N2	M1
Stage LXXXXXXXxxxix	T89	N0	M1
Stage LXXXXXXXxxxix	T89	N1	M1
Stage LXXXXXXXxxxix	T89	N2	M1
Stage LXXXXXXXxxxix	T90	N0	M1
Stage LXXXXXXXxxxix	T90	N1	M1
Stage LXXXXXXXxxxix	T90	N2	M1
Stage LXXXXXXXxxxix	T91	N0	M1
Stage LXXXXXXXxxxix	T91	N1	M1
Stage LXXXXXXXxxxix	T91	N2	M1
Stage LXXXXXXXxxxix	T92	N0	M1
Stage LXXXXXXXxxxix	T92	N1	M1
Stage LXXXXXXXxxxix	T92	N2	M1
Stage LXXXXXXXxxxix	T93	N0	M1
Stage LXXXXXXXxxxix	T93	N1	M1
Stage LXXXXXXXxxxix	T93	N2	M1
Stage LXXXXXXXxxxix	T94	N0	M1
Stage LXXXXXXXxxxix	T94	N1	M1
Stage LXXXXXXXxxxix	T94	N2	M1
Stage LXXXXXXXxxxix	T95	N0	M1
Stage LXXXXXXXxxxix	T95	N1	M1
Stage LXXXXXXXxxxix	T95	N2	M1
Stage LXXXXXXXxxxix	T96	N0	M1
Stage LXXXXXXXxxxix	T96	N1	M1
Stage LXXXXXXXxxxix	T96	N2	M1
Stage LXXXXXXXxxxix	T97	N0	M1
Stage LXXXXXXXxxxix	T97	N1	M1
Stage LXXXXXXXxxxix	T97	N2	M1
Stage LXXXXXXXxxxix	T98	N0	M1
Stage LXXXXXXXxxxix	T98	N1	M1
Stage LXXXXXXXxxxix	T98	N2	M1
Stage LXXXXXXXxxxix	T99	N0	M1
Stage LXXXXXXXxxxix	T99	N1	M1
Stage LXXXXXXXxxxix	T99	N2	M1
Stage LXXXXXXXxxxix	T100	N0	M1
Stage LXXXXXXXxxxix	T100	N1	M1
Stage LXXXXXXXxxxix	T100	N2	M1

Note: The prognosis of patients with N1 is similar to that of patients with pN0.

HISTOPATHOLOGIC TYPE

Histologic types are as follows:

- Carcinoma, NOS (not otherwise specified)
- Intraductal (*in situ*)
- Invasive with predominant intraductal component
- Invasive, NOS
- Comedo
- Inflammatory
- Medullary with lymphocytic infiltrate
- Mucinous (colloid)
- Papillary
- Scirrhous
- Tubular
- Other

Breast

Lobular

- In situ
- Invasive with predominant *in situ* component
- Invasive

Nipple

- Paget's disease, NOS
- Paget's disease with intraductal carcinoma
- Paget's disease with invasive ductal carcinoma
- Other
- Undifferentiated carcinoma

HISTOPATHOLOGIC GRADE (G)

- GX Grade cannot be assessed
- G1 Well differentiated
- G2 Moderately differentiated
- G3 Poorly differentiated
- G4 Undifferentiated