

IN THE COURT OF COMMON PLEASCUYAHOGA COUNTY, OHIO

STANLEY T. MYSLIWIEC,
et al.,

Plaintiffs,

- vs -

JUDGE E. GALLAGHER
CASE NO. 339022

STANLEY R. GAHRING, M.D.,
et al.,

Defendants.

- - - -

1 Deposition of NATHAN LEVITAN, M.D., taken as
1 if upon cross-examination before Kenneth F.
1. Barberic, a Registered Professional Reporter and
1: Notary Public within and for the State of Ohio,
14 at the offices of Reminger & Reminger, Seventh
15 Floor, 113 St. Clair Building, Cleveland, Ohio, a
16 8:10 a.m., on Wednesday, March 3, 1999, pursuant
17 to notice and/or stipulations of counsel, on
18 behalf of the Plaintiffs in this cause.

- - - -

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1 APPEARANCES:

2 Harlan M. Gordon, Esq.

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6 On behalf of the Plaintiffs;

7 John R. Scott, Esq.

8 Reminger & Reminger

9 113 St. Clair Building

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(215) 687-1311,

10 On behalf of the Defendants

11 Dr. Gahring, Luebbers &

12 West Side Pathology.

MR. GORDON: Ken, will you mark
that?

(Thereupon, Plaintiff's Exhibit
Levitan-1, Notice of Deposition, was mark'd for
purposes of identification.)

NATHAN LEVITAN, M.D., of lawful age,
called by the Plaintiffs for the purpose of
cross-examination, as provided by the Rules of
Civil Procedure, being by me first duly sworn, a;
hereinafter certified, deposed and said as
follows:

CROSS-EXAMINATION OF NATHAN LEVITAN, M.D.

BY MR. GORDON:

Q. Good morning again. My name is Harley Gordon.
Seated next to me is my partner, Morey Heller.
We represent the plaintiffs in the case and this
morning because you have been identified as an
expert in this case. I'll be asking you
questions primarily in the area of your opinions
and the bases of your opinions and then also
other areas that might develop.

During the course of my questioning,

Dr. Levitan, please make sure you understand the question before you answer it. Do you understand that, sir?

4 A. I do.

5

6 the question, the question is not clear to you,
please don't answer and tell me to rephrase it.
7 Do you understand that?

A. Yes.

1 Q. If you want to take a break for whatever reason,
1 please tell me and we'll stop these proceedings.
1 Okay?

1 A. Yes.

14 Q. First of all, could you give us your full name
15 and home address?

16 A. Nathan Levitan. 2980 East Belvoir Oval, Shaker
15 Heights.

18 A. And what is your professional address?
19 University Hospitals of Cleveland, 11100 Euclid
20 Avenue, Cleveland, 44106.

21 Q. Mr. Scott was kind enough to forward to us a copy
22 of your curriculum vitae. Let me hand it to
23 you. Is that curriculum vitae which I have
24 handed to you, there is a mark at the bottom
25 February 18th, 1999, is that your current

1 curriculum vitae?

A. I actually have had some additional publications since this version

4 Q. Okay. Other than the additional publications, is
5 there any other additions to this curriculum vitae which would bring it current and up to date?

A. Based on a very brief and cursory review of this, I don't believe so.

11 Q. Could you take more than a brief and cursory
12 review then?

12 A. Since the completion of this version, as I
13 mentioned, there are some additional
14 publications, I have a somewhat different title^e
15 at the cancer center and I have one additional
16 degree.

17 Q All right. Let's go over each of them then.
7 What is your additional title at the cancer
18 center?

19 I'm currently the medical director of clinical
20 cancer programs for the cancer center and
21 University Hospitals Health System, which is to
22 say that I'm in charge of overseeing cancer
23 patient care.

24 When you talk about the cancer center, are you
25

1
2 talking by about the Ireland Cancer Center?
3 Correct.

4 A.

5 Q. When did you assume that position?

6 A. Two years ago.

7 Q. Okay. You also indicated that you have received
8 an additional degree?

9 A. I earned an MBA as of May of 1996 at the
10 Weatherhead School.

11 Q. Okay. And then the last item is additional
12 publications. And how many would those be?

13 A. I can't recall exactly, but perhaps three or four
14 additional publications.

15 Q. Do any of those additional publications relate in
16 any way to the issues in this case?

17 A. They do not relate to penile carcinoma.

18 Q. Okay. What generally do they relate to?

19 A. As I recall, I wrote a chapter on rehabilitation
20 from cancer in a textbook, I have written an
21 article on cancer and deep vein thrombosis and
22 pulmonary embolism, and I have written a couple
23 of articles on lung cancer, something on
24 esophageal cancer.

25 Q. Okay. Could you please tell me what publication
you authored a chapter for?

A. There's a textbook on rehabilitation medicine and

I was asked to write the chapter pertaining to cancer and rehabilitation.

Q. And could you tell us the author in the title of that publication?

A. I just received a copy of the textbook last week. I believe that the editor is -- you know, I can't remember. I can provide that information, for you very easily following my return to my office this morning.

Q. Okay. Offhand, would you have a more current curriculum vitae than the one we have here now?

A. I don't have it with me, but I would be pleased to provide it to you.

A. Okay.

MR. SCOTT: Off the record.

- - - -

(Thereupon, a discussion was had off the record.)

- - - -

Q. Dr. Levitan, while we're talking about publications, are there any publications identified in your curriculum vitae and those that you have supplemented to your curriculum vitae this morning that relate to any of the issues in this case?

A.

Q.

A.

or discuss specifically carcinoma of the penis.

Q. Okay. Do any of the articles or publications that you authored relate to dysplasia, carcinoma in situ, invasive cancer, generally speaking?

A. I can't specifically recall. That is not the focus of any of those articles.

Q. Okay. Where were you born?

A. Near Boston.

Q. Are you presently licensed to practice medicine in the State of Ohio?

A. Yes.

Q. Also I note that you have a license in Massachusetts?

A. Yes.

Q. Has your license to practice medicine ever been revoked or suspended for whatever reason?

A. No. And I should clarify that I have since

allowed my license in Massachusetts to lapse.
have not renewed it. ^I
a

Q. So as we sit here today the only license you have
to practice medicine is in the State of Ohio?

A. Correct.

6 Q. Now, you are board certified in hematology and
7 oncology?

8 A. Correct.

9 Q. And that is one board certification, is that
correct?

A. Two.

I. Two separate?

I A. Yes.

1 Which one, is there any order in which you take
1 those?

1 It makes no difference which, in which order
1 those exams are taken.

1 Okay. How did you take the exams, in what
1 order?

20 A. I frankly would have to look at my curricu um
21 vitae. It's been a long time.

22 Q. Okay.

23 A. My curriculum vitae indicates that I was boarded
24 in internal medici in 1983 in medical oncol
25 in 1985 and in h atology in 1986

1 Q. Okay. Did you pass your boards on, for internal
2 medicine on the first attempt?

3 A.

4 Q. Did you pass your boards for medical oncology on
the first attempt?

A. Yes, sir.

Q. Did you pass your boards in hematology on the
first attempt?

A. Yes.

1 2. And referenced in your curriculum vitae is that
1 you are board eligible for blood banking. What
1 does that mean?

1. 1. During the course of my postgraduate training I
14 had enough exposure to blood blanking, which
15 overlaps considerably with hematology, such that
1 I could have taken those boards if I had wanted
1 to.

1 Q. At the present time what is your practice in
1 terms of specialty?

2 4. I practice general medical oncology currently,

2 Q. And how long have you been doing that?

2: 1. I completed my training, as my curriculum vitae
2: indicates, in 1986 and I have been practicing
24 general medical oncology since that time.

25 1. Do you have any subspecialty or special interest

in the area of medical oncology?

A. At the Ireland Cancer Center I'm the director of the thoracic malignancy research program.

Although in terms of patient care I see all types of patients.

Q. With cancer?

A. Yes.

Q. Okay. And solely with cancer?

A. Are you asking whether I see patients with any hemologic disorders?

Q. Let's take that out. The patients that you do see from -- well, I should phrase it this way, do you do general internal medicine?

1. There is a lot of general internal medicine in cancer care.

Associated with the cancer treatment, is that it's So the internal medicine that I practice is, occurs because cancer patients develop general medical problems.

Q. Okay. So, in other words, you don't do general internal medicine in and of itself?

A. Patients do not come to me for general medical care without either a diagnosis of cancer or a suspicion of cancer or a hematologic disorder.

Q. And has your practice been primarily in the

1 1990's at the University Hospitals of Cleveland?
2 A. I moved from Boston to the University Hospitals
3 of Cleveland in 1991.

4 Q. Okay. Are you employed by the hospital? Or let
5 me put it this way, what business relationship,
6 since you have an MBA, do you have with the
7 Ireland Cancer Center?

8 A. I am employed full time at the Ireland Cancer
Center. The actual structure of my compensation
1 is more complicated because I'm employed by a
1 private practice based at the hospital. I would
1 be pleased to go into that in more detail if
1 you're interested.

14 Q. What's the name of the private practice group?

15 A. University Physicians, Incorporated.

16 Q. Okay. And you also -- strike that.

17 Do you also teach at the medical school?

18 A. Yes.

19 a. Now, handing you what has been marked Exhibit 1,
20 this is a Notice of Deposition that we forwarded
21 to Mr. Scott regarding certain information which
22 we would like relative to your involvement in
23 this case.

24 Have you seen this Notice of Deposition
25 before?

1 A. I don't believe so.

2 Q. Could you turn to the second page, please?

3 A. (Thereupon, the witness so complied.)

4 Q. We asked in Request Number 1 for your up to date
5 curriculum vitae and you indicated you'll forward^d
6 that to us?

7 A. Yes, sir.

8 Q. Number 2 of the Notice of Deposition requests
9 your complete file, including your personal
10 notes. Did you bring your complete file here
11 today?

A. I have it here.

13 Q. Which consists of three pieces of paper?

14 A. Correct. Plus the records, which I didn't bring¹⁹
15 with me.

16 MR. GORDON: Okay. Why don't we
17 mark these:
- - - -

18
19 (Thereupon, Plaintiff's Exhibit
20 Levitan-2, Dr. Levitan's report, was mark'd for
21 purposes of identification.)
- - - -

22
23 (Thereupon, Plaintiff's Exhibit
24 Levitan-3, Dr. Levitan's notes, were mark'd for
25 purposes of identification.)

Q. Exhibit 2, am I correct, is a copy of your report^{rt} consisting of two pages?

A. Correct.

Q. Dated December 14th, 1998, is that correct?

A. Correct.

Q. The report has some writing here, it says depo
of?

A. I recently received a copy of the deposition of
Mr. Mysliwiec. I added that because my report
summarizes those records that I have reviewed andnd
at that time I had not yet received that
deposition for review.

Q. Okay. And when did you receive the deposition of
Mr. Mysliwiec?

A. Yesterday.

Q. All right. Have you reviewed or read any other
depositions in this case?

A. Everything that I've read for this case is
indicated here.

Q. So you've read the treatment records and what did
that consist of?

A. That consisted of all the records that were sent

1 to me by Mr. Scott and I did not break those
2 down. Perhaps, Mr. Scott, you could help me with
3 a breakdown of those.

4 MR. SCOTT: We gave, if this is all
5 right with you, all the medical records, all
the records of Dr. Kasick and Gahring and
all the pathology reports, all of the
records of Dr. Spirnak, records of all
treaters up through the present.

1 Q. Does that sound correct to you?

1 A. Yes.

1 Q. Okay. Did you in any way summarize those record
1 or take any notes relative to your review of
14 those records?

1 A. I have, my style is to prepare notes as you see
1 here and I have simply flagged important pages
1 such as pathology reports in the record.

1 Q. Okay. Is there any reason why you didn't bring
1 those records today?

2 A. I wasn't aware that, that that was requested.

2 MR. GORDON: We're just going to
2 make a request to have an opportunity to
23 look at those records.

24 MR. SCOTT: That's okay, Harley.
25 I'll be happy to do that. I frankly did not

1 tell the doctor to bring anything.

Q. Okay. And then Exhibit 3 is what?

4 A. Is notes that I've made to summarize important
4 dates that may be raised for discussion.

Q. And when did you prepare Exhibit 3?

6 A. I don't recall the exact date. But this was
E prepared following my review of the records
E several months ago.

10 Q. And would you have reviewed those records before
10 you authored your report of December 14th, 1998?

11 A. Yes, sir.

12 2. And these notes, Exhibit 3, would have been
1 prepared then before your report on December
1 14th, 1998?

1 A. Correct.

1 2. So in terms of this case, so I can understand
1 specifically what you looked at, we know about
1 the treatment records. Then in terms of
1 depositions, you've only read Mr. Mysliwiec's
2c deposition?

21 A. Correct.

22 And did you ask to read his deposition?

23 I simply read those records that were sent to me
24 by Mr. Scott.

25 Q. Okay. Do you have any understanding why

3 Q. So as we, at the present time you have not read

5 Dr. Luebbers, is that correct?

6

7

8

9 A. Correct.

10 Q. In addition, you have not read the depositions of
11 any of the expert witnesses in this case?

13 reviewed are listed in the letter in front of
14 you.

15 Q. Okay. You looked at the reports of Dr. Nuovo and
16 Dr. Lee?

17 A. As I indicated, all the records that I reviewed
18 are in the letter in front of you.

19 Q. Do you know Dr. Lee?

20 A. I don't believe so

21 Q. Okay. Give me one moment. Let me read your
22 notes.

23 In your notes you state, I think you also
24 indicated this in your report, that Dr. Kasick
25 stated in his deposition he wanted to refer him

to Dr. Gahring or another urologist. Where did you get that information if you didn't read Dr. Gahring's deposition?

A. Well, I must have. I stand corrected. That must have been included in those notes. As I told you, I read these records several months ago. I did not re-read the records prior to today. I rely on my report to prepare for a deposition.

Q. In looking at these notes in Exhibit 3, the treatment records that you reviewed, was that through April of '97?

4. Yes.

2. Have you seen any additional records relating to Mr. Mysliwiec since April of 1997?

1. Not that I recall.

3. Now, have you conducted any medical research in regard to this case?

No.

Q. Do you intend to use any demonstrative exhibits during your trial testimony?

21 A. I do not.

Q. Have you discussed this case with any individual other than Mr. Scott and/or any attorney in the Reminger & Reminger law firm?

A. No.

1 Q. Have you -- strike that.

2 Do you know Dr. Luebbers?

3 A. No.

4 Q. Do you know Dr. Gahring?

5 A. No.

6 Q. Do you know Dr. Kasick?

7 A. No.

8 Q. Do you know Dr. Spirnak?

9 A. No.

10 Q. Do you practice pathology?

11 A. No.

12 Q. Do you practice urology?

13 A. My practice is limited to medical oncology.

14 Though as a medical oncologist I care for
15 patients with urologic malignancies though I'm
16 not a urologist, I'm not a surgeon, I don't do
17 the type of procedures that urologists do.

18 Q.

19 other than as it relates to cancer?

20 A.

21 only a urologist would be qualified to carry
22 out.

23 Q. Okay. The patients that you do see that have
24 urological problems are those that relate
25 specifically to cancer?

1 A. Correct

2 Q. In other words, you see those patients that
3 already have cancer associated with the
urological tract, is that correct?

A. That either have cancer or suspected cancer.

Q. Okay. Do you treat patients with lesions of the
penis that have not been diagnosed as cancer or
pre-cancer?

4. I have seen such patients, though my care for any
10 such patient would be in close association with
11 other physicians such as urologists and
dermatologists.

1 Q Okay. So the patients that you do see who would
1 have skin lesions on the penis would be based
1 upon a referral from a urologist or
1 dermatologist?

1 A The referral could come from anyone. Though all
1 patients that are referred to me either have an
1 established diagnosis of cancer or a suspicion of
2 cancer.

2 Q. Okay. Just to clarify one point, the patients
22 that you do see with skin lesions, let's say of
23 the penis, are referred to you?

24 A. Every patient who comes to me is either referred
25 by another physician or is self referred.

Q. Do you practice dermatology?

A. My answer to that question would be analogous to the previous line of questioning.

Q. No. Do you practice surgery?

A. Ditto. My answer would be analogous to the previous line of questioning.

Q. Do you practice surgery?

A. No. I work closely with surgeons.

Q. Do you know what a Mohs' procedure is?

10 A. I do.

1 Q. What is a Mohs' procedure?

1 A. A Mohs' procedure is a technique practiced
1 generally by dermatologists in which a thin layer
1 of skin is removed in an effort to eliminate a
1 malignancy and there is an interim analysis
1 during the course of the procedure to determine
1 if additional tissue needs to be removed.

18 2. And to your knowledge has the Mohs' procedure
19 been utilized in the treatment of penile cancer?

20 A. I do not know the answer to that.

21 1. Do you know whether any topical chemotherapy
22 cream is associated or used with penile cancer?

23 A. I've never been involved in the treatment of
24 penile cancer with that modality.

25 Q. Do you know whether laser treatment is used in

1 conjunction with penile cancer or a pre-cancerous
2 condition?

3 MR SCOTT: Oops That's two
4 questions I mean to clarify I'm sorry
5 It is not really two questions, but I want
6 to make sure that the doctor, I want to make
7 sure the doctor heard You were talking
8 about cancer initially.

9 MR. GORDON: I'll break it down
10 to you.

11 MR SCOTT: Okay

12 Q Do You know whether a Mohs' procedure is used in
13 conjunction with the diagnosis of dysplasia or
14 carcinoma in situ relative to a penile lesion?
15 A That is outside my area of expertise I do not
16 know the answer to that

17 Q Similarly, do you know whether topical
18 chemotherapy treatment is utilized in terms of
19 dysplasia or carcinoma in situ identified in a
20 penile lesion?

21 A I have never used such a treatment in following
22 patients with penile carcinoma

23 Q Do you know whether that's used in the treatment
24 of such a condition?

25 A In my experience I'm unaware of it

1 Q. By reading the literature are you aware that this³
type of treatment is used?

MR. SCOTT: You're assuming it is?

MR. GORDON: Yeah.

4 A. I have not been involved in such treatments and I
6 do not know.

7 Q Okay. And do you know whether laser therapy is
8 used in a situation where you have dysplasia or
9 carcinoma in situ identified in a penile lesion?

1 A. It is my understanding that surgeons are using
1 laser excision in most areas of the body. The
1 experience that I've had with penile carcinoma
1 has involved more traditional surgical
1 techniques.

1 Q. And those are what?

1 A. Excisional biopsy and other conventional surgical
1' procedures to treat cancer.

1t 2. When you say other conventional surgical
1 treatments, what are those as it relates to the
2 treatment of penile cancer?

2: A. Surgical procedures that involve surgical
2: instruments that are conventional, that are
2: scalpels and knives and so forth.

24 Q. The long and short of it, to remove the cancer
25 and the surrounding tissue?

A. Correct.

Q. And the excisional biopsy is done by a urologist dermatologist and/or surgeon?

A. Correct.

Q. And the surgical procedures are done by a urologist and/or surgeon?

A. Correct.

Q. Do you know if dermatologists do surgical procedures relative to penile cancer?

1 A. They'll do superficial procedures, but generally
1 not complex procedures.

1 Q. Do you receive or read publications directed to
1: urologists?

14 A. I do not.

15 Q. DO you read -- strike that.

16 Do you receive any publications directed to
17 dermatologists?

18 A. I read general medical oncology publications
19 exclusively and internal medicine publications.

20 Q. What textbooks do you from time to time look at
21 in the area of oncology?

22 A. I use a whole variety of textbooks.

23 Q. Okay.

24 A. Including DeVita, including Haskell, I frequently
25 use computer literature searches, Harrison's

1 Textbook of Internal Medicine and others, the
2 authors of which I can't immediately recall.

3 Q. Is DeVita's publication considered a reliable
4 authority?

5 A. One of many.

6 Q. Is Haskell's publication considered a reliable
7 authority?

8 A. One of many.

9 Q. And is Harrison's publication considered a
10 reliable authority?

11 A. One of many. There is no single supreme
12 authority among those textbooks.

13 Q. What has your experience been in terms of penile
14 cancer in terms of how many cases you see a
15 year?

16 A. Penile cancer is a rare disorder. No medical
17 oncologist can specialize in this area. In my
18 career I have seen perhaps a dozen cases.

19 Q. And your career would be approximately how many
20 years?

21 A. Just over twelve years.

22 Q. Did you see any penile cancer cases during your
23 internship, residency and/or fellowship?

24 A. I'm sure I did.

25 Q. Okay. So that would increase the amount?

1 A. Correct.

2 Q. When is the last time you were involved in the
3 treatment of a patient who had penile cancer?

4 A. December.

5 Q. And what role do you play in terms of the
6 treatment of a patient with penile cancer?

7 A. My involvement is to assist with decisions
8 regarding the need for chemotherapy, the need for
9 radiation therapy and prognostic assessment.

10 Q. Prognostic assessment means what?

11 A. Helping to determine the prognosis of the patient
12 with such a cancer.

13 Q. That means what the future would foretell for the
14 patient?

15 A. And accordingly what treatments would be needed.

16 Q. Prognostic assessment would be assessing the
17 potential life expectancy of a patient, is that
18 correct?

19 A. Correct.

20 Q. Do you also do staging of penile cancer?

21 A. Staging is a very simple process that is
22 determined by data provided by the surgeon and
23 pathologist.

24 Q. Are you saying then that you rely upon this
25 staging of the surgeon and/or pathologist?

1 A. I'm familiar with the staging categories of
2 penile carcinoma and the data used to determine
3 the stage are provided by the surgeon and the
4 pathologist.

5 Q. Okay. Do you independently stage penile cancer?

6 A. Whenever I see a patient I review the available
7 data and determine the stage and if for any
8 reason I disagree with the staging assigned by
9 other physicians I would so indicate.

10 Q. Have you done any studies in the area of the
11 treatment and prognosis of patients who have
12 penile cancer?

13 MR. SCOTT: Any studies, you say?

14 MR. GORDON: Yes.

15 MR. SCOTT: I'm sorry. I wasn't
16 listening. I apologize.

17 A. All the studies that I have done have been
18 published. And we've discussed the relevance of
19 my publications to penile carcinoma.

20 Q. So have you done any studies in that area?

21 A. I believe that I have answered your question.

22 Q. I don't think you have. I mean yes or no, have
23 you done any studies in the area of penile
24 cancer?

25 A. I haven't done any studies that specifically

1 focused on penile carcinoma.

2 Q. Okay. I notice you are looking at your watch.
3 What time do you need to leave?

4 A. I need to leave here by ten o'clock.

5 MR. SCOTT: What time is it? Almost
6 9:00?

7 MR. GORDON: Yeah.

8 - - - -

9 (Thereupon, a discussion was had off
10 the record.)

11 - - - -

12 Q. Have you consulted with the Reminger & Reminger
13 firm before?

14 A. I have done expert witness work for them before,
15 yes.

16 Q. Have you done expert consultations with Mr. Scott
17 before?

18 A. Yes.

19 Q. On how many occasions?

20 A. I don't recall the number.

21 Q. Okay.

22 A. A small number.

23 Q. Have you ever acted as a consultant for a
24 patient?

25 A. Yes.

1 Q. On how many occasions?

2 A. Several.

3 Q. In the greater Cleveland area?

4 A. Some in the greater Cleveland area and I believe
5 one in New England.

6 Q. All right. And in terms of the cases that you
7 have participated in, either by review or giving
8 a report or deposition, going to trial, what
9 would the percentage be of plaintiff versus
10 defendant, patient versus medical care provider?

11 A. There is a fair representation of both. There
12 may be a slight majority pertaining to defense of
13 the physicians involved, though there is
14 significant representation on both sides.

15 Q. Could you give me the percentage?

16 A. I can't.

17 Q. How many cases do you review a year?

18 A. I would guess perhaps six to eight cases a year.

19 Q. And how much do you charge an hour for your
20 review?

21 A. 250.

22 Q. And what do you charge for a deposition?

23 A. It's the same.

24 MR. SCOTT: \$7,000.

25 Q. And trial?

1 A The same

2 Q Have you ever testified in a case involving
3 penile cancer, carcinoma in situ and or carcinoma of
4 a penile lesion?

5 A I have never testified regarding carcinoma of the
6 penis. I cannot recall whether issues of
7 penile cancer and so forth in other cancers have come
8 up. I believe that probably have.

9 Q I was. I overheard one of these. What role do you
10 play in the treatment of a patient who is
11 diagnosed with penile cancer of a penile lesion?

12 A As a medical oncologist I'm often involved in
13 coordinating the care of cancer patients who are
14 seeing multiple specialists. A patient with
15 penile cancer would not specifically require services
16 such as chemotherapy that are particular to a
17 medical oncologist. But my involvement would be
18 as a coordinator of care.

19 Q Which means what as a practical matter?

20 A Which means following such patients periodically
21 and reviewing reports of other physicians to make
22 sure that the care provided is appropriate.

23 Q Okay. What role do you play in a patient who is
24 diagnosed with carcinoma in situ of the penis?

25 A The same. A medical oncologist, as an internist,

1 plays an important role to make sure there is
 2 appropriate follow-up and that the appropriate
 3 specialists are involved in that patient's care.

4 Q For what purpose?

5 A As I said, it's very important that the patient
 6 has coordinated medical care and the medical
 7 oncologist generally often plays the role of
 8 coordinating such care.

9 Q In your experience, have you seen patients who
 10 have had dysplasia or moderate dysplasia which
 11 resulted in a laser procedure?

12 MR. SCOTT: I'm sorry. Just at that
 13 stage or after it developed into cancer?

14 MR. GORDON: No. Just at that
 15 stage.

16 A As I believe I've said, in the dozen plus cases
 17 that I have seen involved in the surgical
 18 procedures have not involved laser treatment
 19 Q Okay that would apply both to dysplasia a,
 20 moderate dysplasia and/or carcinoma in situ?

21 A. Correct.

22 Q. Then when you have invasive cancer your role
 23 would be what of the penis?

24 A Precisely the same, determining what type of
 25 intervention or procedure is multi-specialists.

1 following that patient over time and coordinating
2 care

3 Q Okay In terms of the decision whether or not a
4 Mohs' procedure or use a topical chemotherapy
5 treatment for dysplasia or carcinoma in situ.
6 That decision would be made by the urologist or
7 the dermatologist?

8 A. Correct.

9 Q. Not you?

10 A. Correct.

11 Q. And similarly the decision whether to use a laser
12 procedure or surgical intervention of, of a
13 penile lesion with dysplasia or carcinoma in
14 situ, that would be the decision of the urologist
15 and/or dermatologist or a consulting surgeon?

16 MR SCOTT: If those modalities are
17 used at that stage?

18 MR. GORDON: Right.

19 A Correct.

20 Q Now, with respect to your report, Exhibit B,
21 have -- let me phrase

22 Am I correct that all your opinions are
23 contained in Exhibit 2?

24 A That's a very broad question

25 Q And it's purposely that way. Because at this

1 stage of these proceedings I'm entitled to find
2 out your opinions and the bases for your
3 opinions. What's what I'm asking, are all your
4 opinions then contained in Exhibit 2?

5 A. I can't at the moment recall any major points
6 that I have to make that are not included in that
7 document

8 Q. Okay. Then do you have an opinion as to what the
9 accepted standard of care was for Dr. Luebbers in
10 reading the pathology slides?

11 A. Could you restate your question?

12 Q. Okay. Do you have an opinion, to a reasonable
13 degree of medical probability, as to the standard
14 of care required of Dr. Luebbers in reading the
15 pathology slides in February of '93?

16 A. I'm not sure how to answer that question.

17 Q. And why is that?

18 A. When one asks a question about standard of care
19 one could list a hundred principles pertaining to
20 pathologic practice. It would be more helpful to
21 me if you could ask a specific question regarding
22 her practice and I could answer as to whether or
23 not that is within the accepted standard of care
24 in my opinion

25 Q. And how do you feel you are qualified to answer what

1 the accepted standard of care of a pathologist is
2 when though you don't practice pathology?

3 4 I could only comment within my expertise as a
4 medical oncologist

5 MR SCOTT: Harlow, if it helps in
6 any way, I am not going to be asking him any
7 questions about any pathology, that is what
8 features of a pathology slide might be
9 consistent with what conditions

10 Q From your understanding, is it the accepted
11 standard of care for a pathologist to accurately
12 interpret tissue on a slide?

13 MR. SCOTT: I object to that. I
14 don't know what you mean be accurately
15 A. I would concur.

16 MR SCOTT: And that's a legal
17 question and it's not appropriately a
18 question me, appropriate legal standard is
19 what is reasonable.

20 Q. Can you answer the question?

21 4 It is my understanding that a pathologist is
22 trained in the interpretation of tissue specimens
23 and it is his or her obligation to apply that
24 expertise in pathologic evaluation.

25 Q And going beyond that, wouldn't you agree that

1 the accepted standard of care of a pathologist is
2 to appropriately read the tissue that's presented
3 for interpretation?

4 MR SCOTT: I object to that because
5 that's not a proper question as to what is
6 appropriate. You may think appropriate --

7 Q Well, let me put it this way --

8 MR SCOTT: There needs to be a
9 definition of appropriate

10 Q Let me put it this way, assuming hypothetically
11 that tissue presented to a pathologist contains
12 tissue and cells that are either moderate
13 dysplasia or carcinoma in situ. Do you agree that
14 accepted standard of care would be for a
15 pathologist to identify those conditions?

16 MR SCOTT: Objection.

17 A Recognizing that pathologic interpretations are
18 not black and white and that there is often
19 considerable room for interpretation among
20 different pathologists, I would expect a well
21 trained pathologist to render an opinion that is
22 within the range of interpretation that is
23 determined to be correct in accordance with a
24 community standard.

25 Q In this case we have the Plaintiff's expert.

Dr. McCart~~X~~, reading the slides of ebruary of 1993 and he identified morphologic dyslasia. Were you aware of that?

A Yes

Q You were also aware that in July of 1996 the tissue of the February of '93 biopsy was re-read by two pathologists at Metro and that identified morphologic dysplasia, is that correct?

A I have seen those reports

Q Okay In addition, were you aware that during the deposition of Dr. Blath he indicated that he had two pathologists review the slides and that identified morphologic dysplasia?

A I don't specifically recall that detail, but I believe you.

Q. Okay. We'll take that out.

Did you also, were you also aware that Dr. Barnard Ackerman, a dermatopathologist, reviewed the slides and indicated that signs of superficial squamous cell carcinoma were present?

A I was not aware of that

Q All right Let me mark this.

- - -

(Thereupon, Plaintiff's Exhibit

Lewitan-4, r. Ackerman's report, was marked for

1 purposes of identification.)

2

3 MR. GORDON: Back on the record.

4 Q. You have now read Dr. Ackerman's report, Exhibit
5 4, is that correct?

6 A. Yes.

7 Q. And the first time you have seen Dr. Ackerman's
8 report is today?

A. Correct.

1 Q. You were never provided then Dr. Ackerman's
1: report or his opinions before today?

12 A. Not as far as I can recall.

Q. Do you know Dr. Ackerman?

A. No.

1 Q. Now, then to go back with what we were
1 discussing, Dr. Ackerman in his view of the
1 slides finds superficial squamous cell carcinoma,
1 is that correct?

1 A. His report so indicates.

2 Q. Okay. So we have Dr. McCarty identifying
2 moderate dysplasia, two pathologist at Metro
2: identifying moderate dysplasia and we have
23 Dr. Ackerman identifying superficial squamous
24 cell carcinoma, is that correct?

25 A. As you demonstrated.

1 Q Day And then we have four pathologists reading
2 these slides to contain moderate dysplasia or
3 superficial squamous cell carcinoma, is that
4 correct?

5 A As you've demonstrated

6 in comparison to Dr. Lubbers' interpretation of
7 basically representative atypia?

8 MR SCOTT: Now that's an improper
9 question there is not only Dr. Lubbers,
10 there's Dr. Conover, there's Dr. Al-Kaisi,
11 there's Dr. Nuovo, all of whom are
12 pathologists and all of whom agree that
13 there is only atypia

14 MR GORDON: Who is Dr. Conover?

15 MR SCOTT: He is another
16 pathologist who read this pathology all
17 those say atypia So you don't have quite
18 the balance that you've indicated. But now,
19 Harlow, let's march on. We have 50 minutes
20 and this is not a subject that he's going to
21 be talking on.

22 Q So in terms of whether Dr. Lubbers deviated from
23 the accepted standard of care in the
24 interpretation of these slides, you don't have
25 any opinion?

1 AS a medical oncologist I certainly cannot
2 interpret pathology slides

3 Q Okay Then do you have an opinion, to a
4 reasonable degree of medical certainty and/or
5 probability, what was present in the penis of
6 Mr. Mysliwiec in February of 1993, was it
7 dysplasia, carcinoma in situ or a benign
8 condition?

9 I'm not qualified to make such a determination.

10 Q And in rendering your opinions does it matter
11 what was present in terms of whether it was
12 moderate dysplasia, carcinoma in situ or a benign
13 condition at that time?

14 A It would be more helpful to me if you could ask
15 me a specific question and I will tell you
16 whether that distinction is important

17 Q And why do you say that?

18 A When you ask me a question like in rendering my
19 opinions, that's extremely vague But I would be
20 happy to answer a specific question and indicate
21 whether that distinction is important

22 Q Okay. Assuming hypothetically that -- let me
23 back track.

24 What do you understand superficial squamous
25 cell carcinoma to mean?

1 A Again, it's a very vague question

2 Q When you read Dr. Ackerman's report to indicate
3 superficial squamous cell carcinoma, what did
4 that mean to you?

5 MR. SCOTT: Well, he says carcinoma in
6 situ? Signs of superficial squamous cell

7 A. When a pathologist determines that frank
8 carcinoma is present, as a medical oncologist I
9 would indicate the importance of making every
10 effort to remove that carcinoma

11 Q. Why?

12 A. The primary treatment for penile carcinoma, as
13 with most cancers, is to perform complete
14 surgical removal.

15 Q. Now, based upon what Dr. Ackerman says, signs of
16 superficial squamous cell carcinoma, would that
17 mean to you frank carcinoma?

18 MR. SCOTT: We, we don't know what it
19 means, Harlow, and if you want to give him
20 that hypothetical, if in the event there was
21 frank carcinoma is that invasive cancer,
22 that's what you are saying?

23 MR. GORDON: No, I'm not saying
24 that I'm just being very limited

25 MR. SCOTT: This doctor can't really

interpret what another doctor is saying by that definition as seems to me

A Perhaps I can answer your question this way. If I have a patient with a penile lesion and frank carcinoma identified therein I would recommend surgical excision of the lesion

Q. All right. Then I'm saying in terms of what you have just said, when there is a reference or signs of superficial squamous cell carcinoma, would you then on the basis of that pathological diagnosis recommend excision?

MR SCOTT: Well, we don't know what he means by that we don't know what Dr Ackerman means by that is it invasive cancer, metastatic, is it simply sarcoma in situ, is it dysplasia? You can't take that person's definition and then ask the doctor how to apply it.

Q. Can you answer my question?

A Again, best say I can answer your question is if a patient under my care has a lesion on the penis which is determined to be frank carcinoma, that excision is the appropriate treatment is in fact it hasn't already been carried out

Q All right Assume hypothetically that there is

present carcinoma in situ of the penile lesion,
 what would your recommendation be in terms of
 treatment?

MR. SCOTT: The doctor as answered
 that question.

A. Again the same answer as I provided a minute ago,
 surgical excision is the appropriate treatment if
 it has not already been carried out.

Q Okay. Now, if you have moderate dysplasia
 identified in a penile lesion, what is your
 recommended course of treatment?

A If it is possible to resect the area that
 involved dysplasia that is moderate. If it is
 a small area then either excision or very
 close follow-up would be indicated depending upon
 the details of the lesion.

Q. So with respect to a pathologic diagnosis of a
 lesion of the penis which has moderate dysplasia,
 the preferable recommendation is to have
 excision, is that correct?

A It is feasible to perform an excision and it
 isn't excessively disfiguring, excision is
 preferable. But there are many cases because of
 the nature of the organ where very close
 follow-up is also appropriate.

1 Q Why is excision preferable?

2 A If there's --

3 Q Under those circumstances?

4 A As with any part of the body, if there is a
5 localized area of dysplasia that can be easily
6 removed that is the preferred course. But,
7 again, if it is an extensive area, as is often
8 the case, close follow-up is needed and excision
9 may not be feasible,.

10 Q. Okay. Excision is preferable because you can
11 remove the moderate dysplasia and that reduces
12 the risk of developing invasive cancer, is that
13 correct?

14 A. Correct

15 Q. Okay.

16 A. If it is present in a very localized area.

17 Q. Okay. In this opinion, in this case do you have
18 an opinion as to whether there is any moderate
19 dysplasia of a localized area which would require
20 an excision as the course of treatment?

21 MR. SCOTT: In this case?

22 MR. GORDON: Yes.

23 A. As I understand it, the area that was excised in
24 February of 1993 was interpreted by some
25 pathologists as showing dysplasia, by other

1 pathologist as not out in pit er case excision
 2 was performed So regardless of whether or not
 3 dysplasia was present, the appropriate surgical
 4 procedure was completed

5 Q And are you saying by that excision all the
 6 dysplastic cells were removed at that time, to a
 7 reasonable degree of medical certainty?

8 A Well, I can't answer that question because
 9 there's not a uniform agreement that dysplasia
 10 was present at all.

11 Q Assuming moderate dysplasia was present, were all
 12 the cells involving moderate dysplasia removed?

13 A I'm unable to answer that question.

14 Q Do you know whether the, do you know what I mean
 15 when I say margins in terms of a pathological
 16 specimen?

17 I do.

18 Q Do you know whether, in terms of the February of
 19 '98 specimen whether margins were identified?

20 A I am not aware of pathologist reports which
 21 specifically indicate a positive margin following
 22 review of this lesion.

23 Q Okay And if you don't have the report
 24 involving -- let me rephrase the question.

25 If you don't have margins identified you

1 can't determine whether the dysplastic cells or
2 the cancerous cells are beyond the area of
3 margin, is that correct?

4 MR. SCOTT: Objection.

5 A. Would you rephrase your question?

6 Q. If you don't have the margins identified one
7 cannot then determine whether the dysplastic or
8 cancerous cells are still present beyond the
9 margin, is that correct?

10 A. Well, it's very important to differentiate
11 between frank cancer and dysplasia. When frank
12 carcinoma is present obtaining negative margins
13 is of greater importance because, as I've said,
14 dysplastic lesions particularly in an area such
15 as the penis will often always be closely
16 followed.

17 MR. GORDON: Could you repeat the
18 answer, Ken?

19 - - - -

20 (Thereupon, the requested portion of
21 the record was read by the Notary.)

22 - - - -

23 A. I believe I said will often be closely followed.

24 Q. Is carcinoma in situ of the penis considered
25 frank carcinoma?

1 A. Carcinoma in situ refers to carcinoma that has
2 not invaded through a basement membrane. But it
3 is a lesion that should be surgically excised.

4 Q. All right. Then are you saying that carcinoma in
5 situ is frank carcinoma?

6 A. Carcinoma in situ is a category of carcinoma.

7 Q. All right. But --

8 A. I'm not sure what frank means.

9 Q. That's what you told me, what you mentioned in
10 terms of frank carcinoma. That's the terminology
11 that I'm using based on what you have said. So I
12 just want to find out whether carcinoma in situ
13 is considered by you to be frank carcinoma?

14 A. Carcinoma in situ is a category of carcinoma and
15 for that reason whenever possible should be
16 treated with complete surgical excision.

17 Q. All right. But going back, you mentioned the
18 terminology frank carcinoma in your deposition
19 testimony. I just want to clarify. Do you
20 consider carcinoma in situ as frank carcinoma?

21 A. Perhaps you could re-read the sentence where I
22 used that term and then I can clarify that for
23 you.

24 Q. I don't know. You used the terminology frank
25 carcinoma. I didn't use the terminology.

1 A Well, again, if you would re-read the sentence
2 where I used that term I will clarify it for you.
3 We'll have to do that then at another time

4 Then is moderate dysplasia considered
5 carcinoma?

6 A No.

7 Q All right. Now, if you have moderate dysplasia
8 the decision whether to do excision or close
9 follow-up, is that up to the urologist or
10 urologist following the patient?

11 A I would say this, the decision concerning the
12 feasibility of resecting an area of dysplasia
13 depends upon the details of the anatomy and
14 generally the urologist or urologist would
15 make what determination

16 Q Okay where do you -- strike that
17 What's your basis that if you do have
18 dysplasia it is preferable to excise the lesion
19 and/or have close follow-up?

20 MR SCOTT: Are you talking about a
21 slight, mild, I mean mild, moderate, severe
22 cancer in situ? What kind of dysplasia are
23 you talking about?

24 Q You mentioned that if you have moderate dysplasia
25 the preferable approach is excision or you can

1 follow-up Where did you get that, or
 2 what is the basis of your opinion that that's
 3 that is the treatment for that type of condition?

4 A There's a basic principle in medical oncology
 5 that various parts of the body can, as a result
 6 of exposure to some carcinogens that you may or
 7 may not be able to identify. Some susceptible
 8 to transformation into carcinomas very often.
 9 very often large areas become dysplastic As an
 10 example, large masses of the mucous membrane,
 11 the mouth, the throat, esophagus, the lung, the
 12 esophix and other areas can become dysplastic
 13 Which means that there's a risk that carcinoma
 14 can develop very often it's simply impossible
 15 to surgically remove large areas of dysplasia and
 16 in such cases, which are extremely common, close
 17 follow-up is the form rather than excision.

18 Q Okay And I'm asking you what do you base that
 19 course of treatment on? Is that identified in
 20 any textbook, any literature?

21 A It's a very individualized decision.

22 Q What is based upon what the treating physician
 23 feels is appropriate?

24 A. Based upon the details of the pathology, the
 25 anatomy and the opinion of the surgeon involved

1 Q But is there any literature, publications that
 2 indicate if you do have moderate dysplasia, this
 3 is the course of treatment if you have moderate
 4 dysplasia?

5 A No It has to be individualized
 6 or similarly if you have carcinoma in situ this
 7 is the course of treatment for carcinoma in situ
 8 in a lesion of the penis, is there any literature
 9 that identifies that course of treatment?

10 4 As I believe I've explained a few minutes ago, it
 11 is generally accepted in medical oncology that a
 12 carcinoma in situ should be removed surgically
 13 whenever possible That requirement is of
 14 greater importance than an area of dysplasia
 15 which can be followed

16 Q Okay When plaintiff's urology expert,
 17 Dr. Blath, has given his opinion that the
 18 standard of care for the treatment of dysplasia
 19 of a penile lesion includes excision, laser
 20 treatment, or treatment or topical anti-cancer
 21 cream. Do you disagree with that?

22 MR SCOTT: Or monitoring the record
 23 reflect.

24 4 Am I responding to your comments and Mr Scott's
 25 comments?

Q. Just my comment.

A. As I've said, an effort should be made to remove the area of dysplasia if that is feasible.

Otherwise, close follow-up is often undertaken.

Q. Well, the question is more specific. Do you disagree with Dr. Blath's opinion that the standard of care for treating moderate dysplasia in a penile lesion is excision, laser treatments, Mohs' procedure or use of topical anti-cancer cream?

1 A. Am I to understand his comments don't leave room
1 where close follow-up is necessary? And is he
1 suggesting that close follow-up in certain cases
1 is contrary to the standard of care?

1 Q. No, he's not saying that. He's saying that the
1 initial treatment is excision, laser treatments,
1 Mohs' procedure or topical anti-cancer cream. Do
1 you disagree with that?

15 4. I agree with that with the addition that in many
20 cases such treatment is not possible and close
21 follow-up is also within the standard of care.

2% Q. Okay. The standard of care of a urologist and/o
23 dermatologist, is that what you are saying?

24 A. Yes.

25 Q. Okay. And what does that close follow-up

entail?

A. This is highly variable depending upon the details of the situation. But the urologist and/or dermatologist should see that patient at a reasonably frequent interval to make sure that there is not evidence of progression from one type of lesion to carcinoma.

Q. And what is the frequency?

A. Again, that's highly variable depending upon the situation. Every few months.

Q. In your practice then do you, if there is moderate dysplasia or carcinoma in situ you direct the urologist or dermatologist to excise the lesion?

A. I believe that I've answered that question extremely comprehensively regarding the role.

MR. SCOTT: And repeatedly.

Q. But do you direct, I'm changing the question, you are presented with -- strike that.

If you had been presented with -- strike that. I have to backtrack.

Have you seen cases in which there was moderate dysplasia of a penile lesion?

A. Yes.

Q. And how many have you seen?

1 A. I can't recall.

2 Q. And what was the treatment in those cases?

3 A. The treatment is individualized based on whether
4 excision is feasible or whether follow-up is
5 necessary in place of complete excision.

6 Q. Do you know what indeed was done in those cases?

7 A. I can't recall the specifics of each case.

8 Q. And how many cases of moderate dysplasia have you
9 seen?

10 A. Again, I believe you just asked me that
11 question. I refer you to the record. Perhaps
12 you can read my response.

13 Q. Can you be so kind, to move this thing along
14 could you tell me your response?

15 A. Well, I believe that I've answered that question
16 and I would ask you to re-read my response.

17 MR. SCOTT: He has answered that.

18 He believes he has seen moderate dysplasia
19 in the twelve cases or whatever.

20 Q. Have you seen carcinoma in situ in a penile
21 lesion?

22 A. Yes.

23 Q. How many?

24 A. I can't recall the specific number.

25 Q. And what was the course of treatment?

1 A The course of treatment is consistent with that
2 which I have already explained to you

3 Q And have you seen invasive cancer in a penile
4 lesion?

5 A. I have.

6 Q. And how many cases?

7 A. Again, I can't recall the specific number.

8 Q. And what was the course of treatment?

9 A. The course of treatment was consistent with that
10 which I have already described to you

11 Q Excision?

12 A Correct

13 Q Okay. Now, dysplasia is a pre-cancerous
14 condition is that correct?

15 MR SCOTT: Objection

16 A Dysplasia can but now does not necessarily
17 progress to carcinoma

18 Q. So are you saying that dysplasia is not a
19 pre-cancerous condition?

20 MR SCOTT: Objection Harley!

21 Q. Yes or no, that's all.

22 MR SCOTT: No, it doesn't take a
23 Yes or no answer Those conditions that
24 progress to cancer obviously are
25 pre-cancerous Those that don't ever and

1 regress obviously are not. I mean compare on.
2 let's go on.

3 MR. GORDON: No. It's very
4 important

5 Q Are you saying that dysplasia is not considered a
6 pre-cancerous condition?

7 A As I explained, dysplasia can but not does not
8 always progress to carcinoma

9 Q But that still doesn't answer my question.

10 MR. SCOTT: Yes it does

11 Q Is dysplasia considered a pre-cancerous
12 condition, yes or no?

13 MR. SCOTT: Harlan has answered
14 that question. He's not required to give
15 you yes and no. And he's answered fully.
16 Some dysplasias do, some don't

17 MR. GORDON: That's a different
18 issue.

19 MR. SCOTT: It's not when those
20 don't then they can't be pre-cancerous

21 Q In the medical community is dysplasia considered
22 a pre-cancerous condition?

23 A. The best way that I can answer your question, the
24 clearest and most thoughtful way I can answer
25 your question is to say dysplastic lesions can

1 but don't necessarily progress to carcinoma.

2 Q. You still haven't answered my question.

3 MR. SCOTT: Yes, he has.

4 Q. In the medical community is dysplasia considered
5 a pre-cancerous condition?

6 A. Sir, I believe that to the best of my ability
7 I've answered your question

8 Q. Dysplasia is an abnormal condition, is that
9 correct?

10 A. What do you mean by abnormal?

11 Q. That means different from normal. Do all people
12 have dysplasia? That's what I'm saying. Is
13 dysplasia a normal or abnormal condition?

14 A. Dysplasia is a condition that can progress to
15 carcinoma and therefore needs to be either
16 excised or closely followed.

17 Q. I know. You mentioned that before. I just want
18 to know is dysplasia a normal or abnormal
19 condition?

20 A. I don't use terms like normal and abnormal
21 because I think they're extremely vague.

22 Q. Is dysplasia considered a benign condition?

23 A. Dysplasia is a change that does not represent
malignancy. If we are classifying all lesions as
25 benign versus malignant then I guess you could

call dysplasia benign. But I actually think in medical terminology that would be misleading. Dysplasia is a term describing changes in the tissue which could but do not always lead to carcinoma.

Q. Because there is a potential that dysplasia can develop into cancer would you agree with the diagnosis of dysplasia of a, of tissue of the penis that a treating physician has to be more vigilant than if it was considered a benign tissue?

1: MR. SCOTT: Objection.

1: A. I believe that I've repeatedly on multiple occasions in the past hour and a half stated that dysplastic lesions in any part of the body need to be either excised or carefully followed because of the potential to progress to carcinoma.

19 Q. So, again, with the diagnosis of dysplasia, moderate dysplasia, does the vigilance of index of suspicion of a treating physician increase in any way?

23 MR. SCOTT: Objection. You are not giving enough facts. But go ahead, doctor, answer if you can.

1 A. I would, I would give the same answer that I've
2 given to you a dozen times before this morning.
3 I don't understand how I have failed to answer
4 that question for you.

5 Q. Okay. Your opinion is that the biopsy of
6 February of '93 did not show any evidence of
7 invasive cancer?

8 A. I believe I said to you that as a medical
oncologist I'm not able to interpret pathology
1 slides.

1 Q. But in your report you said this biopsy,
1 referring to the February of '93 biopsy, showed
1 no evidence of invasive cancer. Do you still
1 stand by that?

1 A. I believe that in saying that it showed no
1 evidence of invasive cancer I was basing that
1 determination on the pathology report rendered at
1 that time. I was not indicating that I had
1 personally reviewed those slides and made a
2 judgment as to how they should be interpreted.

21 Q. All right. But, in any event, you still stand by
2.2 your statement this biopsy of February of '93
23 showed no evidence of invasive cancer?

24 MR. SCOTT: Based on what the report
25 says?

MR. GORDON: Right.

Q Is that correct?

A I'm only making reference to a pathology report. I am not indicating my opinion as to the pathologic diagnosis because as a medical oncologist I can't review pathology slides. Q. But you are assuming for purposes of your opinions that there was no evidence of invasive cancer?

MR. SCOTT: For purposes of what opinion?

MR. GORDON: Of his opinions in this case.

MR. SCOTT: What opinions specifically?

MR. GORDON: Any of these opinions.

MR. SCOTT: Well, all right.

A I'm only rendering the report of the pathologist.

Q What I'm trying to get at, is there any information that you have gleaned to indicate that there was invasive cancer present in February of '93?

A Upon my review of the medical records pertaining to the care of Mr. Melnick there is no

indication from the single pathology report dated February, 1993 that there was invasive cancer present.

Q. Now, then assuming hypothetically indeed that either moderate dysplasia was present or carcinoma in situ was present in February of 1993, okay?

A. Okay.

Q. And Dr. Luebbers reported out, among other things, reparative atypia but did not report out the moderate dysplasia or the carcinoma in situ in her report, okay?

A. Okay.

Q. Would you agree that the physicians that were treating Mr. Mysliwiec and who were aware of the report were misled by the information in her

MR. SCOTT: Objection. What you say

-- first of all, what you are saying is not even answerable. I mean, if in fact in your hypothetical the true diagnosis was dysplasia and the doctors were not told of dysplasia, then ergo the doctors did not know of dysplasia, so that's a tautology. What's the point of this question?

MR. GORDON: By asking the question and if I repeat the question through Ken you can see the point of the question.

MR. SCOTT: No, It is just, if they did not know there was dysplasia then, and it was there then of course the pathologist did not tell the clinicians that it was there and the clinicians did not become aware. That's obvious. So I mean why are we even --

MR. GORDON: There is a further point if you are listening.

MR. SCOTT: All right. I will try to listen again.

A. Why don't you repeat the question for me.

Q. It is too complicated now, but I'll repeat it.

A. Okay.

Q. Assuming hypothetically indeed that the interpretation of the slides should have been moderate dysplasia or carcinoma in situ and Dr. Luebbbers did not report that in her report, the physicians relying upon her report were misled in terms of the diagnosis of the penile tissue, is that correct?

MR. SCOTT: Now I'm going to object

because you know from the record that both of those doctor have said that when if the interpretation was dysplasia it would make no difference. So it's unfair to kind of ambush this witness in that manner. You're talking about the facts of this case, are you not?

Q Can you answer the questions?

A I can answer the question explaining the principles of management of such lesions which are, as I've stated, if there is carcinoma present every effort should be made to ensure that complete excision has been accomplished. If there is dysplasia present and the area of dysplasia has been removed, then follow-up is appropriate.

If there is dysplasia present and it is not feasible to remove all of that area then follow-up is appropriate.

Q. Would you agree that treating physicians rely upon the interpretations of the pathologist in terms of making treatment decisions?

A. Yes.

Q. Okay And you do that?

A. Yes.

Q. Now, in Paragraph 5 you say 'In my opinion, regardless of whether or not dysplasia was represented in this specimen, the appropriate course of action was to follow the patient closely for evidence of new or non-healing lesions suggestive for malignancy.' Is that your statement?

A. You've read that correctly.

Q. Okay. Now, you just indicated, though, that if there was dysplasia present the course of treatment would have been excision and then follow-up, is that correct?

MR. SCOTT: No, that's not.

A. I don't believe that's what I said. I believe I said, again --

Q. You said if there's moderate dysplasia or carcinoma in situ the preferable approach is excision, is that right?

MR. SCOTT: No. You are misstating his testimony. He said repeatedly that if it was cancer in situ he would have excised it if possible, with moderate dysplasia excision and follow-up, following up is always an option.

A. That's correct.

Q. But you didn't identify the fact that one of the

1 modes of treatment is excision in your report, is
2 that correct?

3 A. As I recall in writing this report I was making
4 reference to the fact that an excisional biopsy
5 had been performed, i.e., the area in question
6 had been surgically removed. For this reason
7 whether or not dysplasia was present in this area
8 it had already been removed and subsequent
follow-up was appropriate.

10 And then you state further, "The presence or
absence of dysplasia on the original biopsy
specimen from February of '93 would not have
altered the nature of the patient's subsequent
close follow-up." Would you explain that for
me?

16 MR. SCOTT: I thought we just did.

17 A. As I have said, following the removal of a lesion
from the penis or any other part of the body
follow-up of that area is necessary because other
lesions can certainly develop adjacent to it.
Hence, after this excisional biopsy was completed
follow-up was the appropriate course and it
appears was carried out.
Are you testifying that it doesn't matter to a
clinician whether the pathologist reports out

1 reparable atypia or moderate dysplasia as it
2 relates to treatment decisions by the clinician?

3 MR. SCOTT: You mean in this case
4 where there has been an excisional biopsy
5 done?

7 A. Well, in this case if you look at the records you
8 see that there were areas of inflammation in
9 multiple locations on the penis and inflammatory
10 changes of the penis can be precursors of
11 carcinomas and may also actually be pre-cancerous
12 changes and for that reason such a patient
requires periodic follow-up by a dermatologist
and/or a urologist.

15 So this patient had physical signs that were
16 indications of a risk for penile carcinoma
17 regardless of whether dysplasia was or wasn't
18 present in that specimen and those physical
19 findings merited periodic follow-up as I believe
20 he received.

21 Q. Are you saying then, to a reasonable degree of
22 medical -- let me withdraw that.

23 Is it your opinion then that Dr. Gahring
24 complied with the accepted standard of care, to a
25 reasonable degree of medical probability?

MR. SCOTT: Objection.

A To the extent that I as a medical oncologist can comment on urologic standards, it appears that this patient had reasonable follow-up visits. I am not able to assess his specific actions in evaluating subsequent lesions because that's out of my area of expertise.

Q. Okay. Similarly, do you have an opinion, to a reasonable degree of medical certainty and/or probability, whether Dr. Kasick complied with the accepted standard of care?

A As a medical oncologist it appears that this patient had periodic follow-up for occurrence of penile lesions following the initial excision and as a medical oncologist that appears to be the appropriate course of care.

Q Would you defer to a board certified urologist in determining what the accepted standard of care was for Dr. Gahring?

A I would defer in evaluating a specific lesion.

Q Such as here?

A If you were to show me a picture of a specific lesion on a penis and ask me do you think that merits topical antibiotics or surgical excision, I would defer to a urologist or dermatologist.

Q Would you prefer to a doctor certification
dermatologist as to the accepted standard of care
that Dr. Kasick should have complied with?

4 Well the determination of standard of care, and
I would ask for Mr. Scott's legal help because I
think it is somewhat of a legal issue, as a
medical oncologist I can say that close follow-up
is necessary and I can delineate the principles
of care for such a patient. What I can't do as a
medical oncologist is render an opinion about a
specific lesion and whether it should be handled
surgically.

Q Do you have an opinion, to a reasonable degree of
medical certainty, as to whether according to the
accepted standard of care re-biopsy should have
been done sometime after February of 1993?

4 Again, that would be a question of judgment based
on the appearance and suspiciousness of a
particular lesion and that would be a decision to
be made by the dermatologist and/or urologist
Q That would be outside of your expertise?

A Yes.

Q Okay Now, in terms of your opinions, in your
report you do not address or give any opinion
regarding timing or progression of this lesion.

is that correct?

I don't believe --

Let me ask you the question.

Do you have an opinion as to what stage the cancer was in in 1993, 1994, 1995, to a reasonable degree of medical certainty and of probability?

MR SCOTT: Well, I'm going to

object because your question assumes that the patient had cancer in 1993. It is not in the record about that and in 1994 and when parts of 1995

Q. All right. Let me go to that then.

Do you have an opinion, to a reasonable degree of medical certainty and/or probability, whether the patient had cancer in 1993?

A. I think there's no way to make a determination of that.

Q. And why is that?

A. Well, I know that this cancer in December of 1996 was a T1N2 Stage 3. I know that cancers of this type develop over a long period of time. Over a period of a number of years it's not possible to determine with greater certainty exactly how many years that is. But I know that

1 cancers such as this are odd at that time of
2 diagnosis.

3 Q And would that be what time frame, years or
4 months?

5 A. Measured in years.

6 I should, I should clarify that by saying
7 that just because a cancer is old doesn't mean
8 that it was clinically detectable at an earlier
9 point in time but rather that pathologically
10 speaking cancer cells were present for a long
11 period of time. Pathologically speaking cancer
12 cells were present for a long period of time
13. Do you have an opinion, to a reasonable degree of
14 medical certainty and/or probability, whether
15 moderate dysplasia or cancer in situ was present
16 in 1993?

17 MR. SCOTT: We have already gone
18 through that. The doctor said he doesn't
19 know any more than what's contained in the
20 reports.

21 I have no information in that regard.

22 Q Do you have an opinion, to a reasonable degree of
23 medical certainty and/or probability, as to
24 whether dysplasia and/or carcinoma in situ was
25 present in 1994?

1 A. In 1994?

2 Q. Yes.

3 MR. SCOTT: I thought we have gone
4 through that as well.

5 A. I have no information to make a determination
6 either way.

7 Q. Do you have an opinion, to a reasonable degree of
8 medical certainty, as to whether in 1994 --

9 MR. SCOTT: 5 you mean. I
10 thought you were going to the next year. Go
11 on.

12 MR. GORDON: There is a logic to
13 what I am doing.

14 MR. SCOTT: Go ahead. Let me hear
15 your logic.

16 Q. Do you have an opinion, to a reasonable degree of
17 medical certainty and/or probability, as to
18 whether cancer was present in 1994? And I'm
19 referring to obviously in his penis.

20 A. As I've explained, the only reference point that
21 we can really use is December of 1996 and as I've
22 told you we can go back in time and say that the
23 age of such cancers is generally measured in
24 years rather than months, but I'm not able to
25 tell you that the first neoplastic cell occurred

1 in 1995 or 1994 or 1993, et cetera.

2 Q. Then in 1996 we have metastatic penile cancer, is
3 that correct?

4 A. In 1996 we have involvement of inguinal nodes.
5 The term metastatic is used in many different
6 ways, but to state it more correctly we know that
7 we have N2 disease in 1996.

8 Q. Okay. So you wouldn't use the term metastasize?

9 MR. SCOTT: I think what the
10 doctor is saying is that people use it
11 differently, but in this instance it would
12 obviously --

13 A. The term often is used to refer to tissue spread
14 and therefore I use the term very carefully so as
15 not to be misleading.

16 Q. So in September of 1996 there was spread of the
17 cancer to the nodes?

18 A. Well, precisely we know that in December,

19

20

21 December.

22 Q.

23

24, growing for some period of time?

25 A. Correct.

1 Q Okay And to reach that condition in December of
2 1996 the cancer would have been present for at
3 least a year, is that correct?

4 A Correct.

5 Q And to reach the condition in December of 1996
6 the cancer would have been present, would you
7 agree, at least two years?

8 MR SCOTT: Objection. He's

9 answering that question.

10 A I believe that I've explained it, that you can
11 try to press for a determination on my part as to
12 exactly how many years old that cancer is and I
13 believe I've explained it at except for saying
14 that it is years old based on what we know of the
15 natural history of such a cancer. I would be
16 unable to determine with greater accuracy exactly
17 what year the cancer began.

18 Q Okay In your report you do not address the
19 issue in terms of survivability of Mr. Myalistic.
20 is that correct?

21 A. Correct.

22 Q. And I assume therefore you don't have any
23 opinions in that regard since you didn't include
24 it in your report, is that correct?

25 A Well, there are data regarding predictive care

1 rates for this cancer on a stage by stage basis.

2 Q. Okay. But in this case you didn't discuss in
3 your report his survivability, is that correct?

4 A. As I recall, I have not included any specific
5 statistics regarding his predictive five year
6 survival in my report.

7 Q. Okay. And therefore can I assume you have no
8 opinion that you intend to offer in this case in
9 that regard?

10 MR. SCOTT: Objection.

11 Q. Because you didn't include it in your report?

12 A. When you ask me that open-ended question the
13 reason that I requested clarification is because
14 one could ask me 5,000 questions about this case
15 to which I would hopefully provide an answer. I
16 certainly haven't included 5,000 paragraphs in my
17 report. So to ask if all of my opinions are
18 included is an extremely confusing question. I
19 would be happy to answer any additional questions
20 that you have for me regarding this case.

21 Q. I just asked you, I assume you do not intend to
22 offer an opinion in terms of survivability
23 because you have not rendered, not identified
24 that in your report?

25 MR. SCOTT: Harlan, I'm going to ask

1 him at the trial of this case about his
2 opinions about survivability

3 MR. GORDON: Although it is not in
4 his report? I'm surprised.

5 MR. SCOTT: Don't be and go ahead
6 and finish your deposition.

7 Q. Is there any reason why you did not include your
8 opinion as to survivability in your report?

9 A. Again, allow me to clarify that I believe it
10 would be unreasonable to ask me a question to
11 which I have a response, a question of your
12 determination, and then ask me why it wasn't
13 included in my report. I have no way of reading
14 your mind to know exactly which questions you
15 would like to ask me and therefore I couldn't
16 have preempted this session by including all the
17 answers to your questions in my report

18 Q. Well, was that on a different process, though
19 The reports, according to our rules that we, that
20 we follow, indicate that the opinions of the
21 expert should be in his report. That's why. So
22 therefore let me ask you the question, were you
23 asked to give your opinion before December 14th,
24 1998 regarding this patient's survivability?

25 A I can't recall.

1 Q. And if you were asked would you have included
2 that in your report?

3 A. I don't believe that I was given specific
4 instructions as to which points should be
5 addressed in my report and, and also in the
6 malpractice cases that I have been involved in I
7 don't believe I've ever been told that any answer
8 to any question that I may be asked is expected
9 to be included in my report.

10 Q. Well, be that as it may, were you asked to --
11 strike that.

12 If you had been asked to give an opinion
13 regarding survivability before December 14th you
14 would have included that in your report?

15 MR. SCOTT: Objection.

16 A. I don't recall whether I was asked that so I
17 can't really answer your question.

18 Q. Okay. Well, do you have an opinion, to a
19 reasonable degree of medical certainty and/or

21 as of today?

22 A. I can tell you that we know as of December of
23 1996 that he had a T1N2 Stage 3 cancer. We
24 know that the predicted cure rate for someone
25 with a cancer in Stage 3 is somewhere in the

1 range of 40 to 50 percent We know that he had a
2 T1N2, which is on the earlier end of the spectrum
3 of Stage 3 disease So I would predict that as
4 of December of 1993 he had approximately a 50
5 percent likelihood of cure from his penile
6 carcinoma

7 Your question was where do we stand today
8 This is now more than two years later. All that
9 I can tell you based on the data with which I'm
10 familiar concerning penile carcinoma is that the
11 survival curve plateaus at about five years.

12 meaning that if someone is diagnosed from at five
13 years they're probably cured and also that the
14 survival curve drops off fairly rapidly over the
15 first two years, which is to say that a great
16 many of the occurrences happen within the first
17 two years So the best I could do in terms of
18 predicting this outcome right now is to say that
19 he has a better than 50-50 chance of being cured
20 of his cancer.

21 MR SCOTT: We're going to pass to
22 run.

23 MR GORDON: If he has to leave he
24 has to leave.

25 MR. SCOTT: Go ahead.

1 A. I can take five more minutes.

2 MR. GORDON: It's going to take
3 longer.

4 Q. You would agree that the earlier you diagnose
5 penile cancer the better the prognosis, is that
6 correct?

7 A. Well, I would agree that there is a definite
8 correlation between cancer stage and prognosis
9 and if a patient is diagnosed with a Stage 1
10 cancer he has a higher likelihood of cure at five
11 years than a Stage 2 and so forth, Stage 3 and
12 Stage 4.

13 Q. So assume hypothetically Mr. Mysliwiec's cancer
14 was in a Stage 1 condition, let's say two years
15 before 1996 and now it's in a Stage 3, you would
16 agree that based upon the delay of the diagnosis
17 there is a reduction in the survivability, is
18 that correct?

19 A. Well, you're making a lot of assumptions about
20 this specific case which I think are, are
21 questionable. But I can tell you that in general
22 if a penile cancer or a lung cancer or a breast
23 cancer can be diagnosed at an earlier point in
24 time and at an earlier stage the predicted five
25 year survival which correlates with stage would