

IN THE CIRCUIT COURT FOR THE
FOURTH JUDICIAL CIRCUIT AND
FOR NASSAU COUNTY, FLORIDA

ROBERT HOGAN,

Plaintiff,

vs.

No. 06-CA-44

BAPTIST MEDICAL CENTER -
NASSAU, INC., a Florida corporation,

Defendant and
Third-Party Plaintiff,

vs.

AMERICAN CANCER SOCIETY and
GRAY GABLE, NASSAU VILLAGE
VOLUNTEER FIRE DEPARTMENT, INC.,

Third-Party Defendants.
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VIDEOTAPED EVIDENTIARY DEPOSITION OF

ROBERT NORMAN HARDEN, M.D.

VOLUME II

September 12, 2007  
5:00 p.m.

446 East Ontario Street, Suite 1011  
Chicago, Illinois

Stacee L. Jackson, Notary Public and Certified Shorthand Reporter  
within and for the County of Will, State of Illinois

APPEARANCES:

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APPEARED ON BEHALF OF THE PLAINTIFF;  
COKER, SCHICKEL, SORENSON & DANIEL  
AARON SPRAGUE, ESQUIRE  
135 East Bay Street  
Jacksonville, Florida 32202  
(904) 356-6071  
.  
APPEARED ON BEHALF OF THE DEFENDANT AND  
THIRD-PARTY PLAINTIFF VIA SPEAKERPHONE;  
SAALFIELD, SHAD, JAY, LUCAS & STOKES, P.A.  
CLEMENTE J. INCLAN, ESQUIRE  
P.O. Box 41589  
Jacksonville, Florida 32202  
(904) 355-4401  
.  
APPEARED ON BEHALF OF THE THIRD-PARTY  
DEFENDANT AMERICAN CANCER SOCIETY VIA  
SPEAKERPHONE.  
FULMER, LEROY, ALBEE, BAWMANN & GLASS  
THOMAS TOLLEFSEN, ESQUIRE  
4720 Salisbury Road  
Jacksonville, Florida 32256  
(904) 562-1020

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APPEARANCES: (CONT.)

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APPEARED ON BEHALF OF THE THIRD-PARTY  
DEFENDANTS GRAY GABLE AND NASSAU VILLAGE  
VOLUNTEER FIRE DEPARTMENT, INC.  
QUINTAIROS, PRIETO, WOOD & BOYER, P.A.  
TERESA A. ARNOLD-SIMMONS, ESQUIRE  
One Independent Drive  
Suite 1650  
Jacksonville, Florida 32202  
(904) 354-5500

.  
ALSO PRESENT:

MR. MICHAEL DESCHAMP, THE VIDEOGRAPHER.  
. . . . .

1 Videotaped Evidentiary Deposition of

2 Robert Norman Harden, M.D.

3 Volume II

4 September 12, 2007

5 THE VIDEOGRAPHER: This is tape  
6 No. 1 in the video -- videotaped deposition  
7 of Norman Harden (sic), M.D., in the matter  
8 of Robert Hogan, plaintiff, versus Baptist  
9 Medical Center-Nassau, Incorporated, a  
10 Florida corporation, defendant and  
11 third-party plaintiff, versus American  
12 Cancer Society and Gray Gable, Nassau  
13 Village Volunteer Fire Department,  
14 Incorporated, third-party defendants, before  
15 -- before the matter of the court in the  
16 Circuit Court, Fourth Judicial Circuit in  
17 and for Nassau County, case No. 06 C A 44,  
18 Division A.

19 This deposition is being held at  
20 446 East Ontario Street, Chicago, Illinois,  
21 60611, on 9/12/07. Approximately the time  
22 is 17:19:09. My name is Michael Deschamp  
23 and I am your videographer. The court  
24 reporter is Stacey Jackson. Counsel,  
25 please introduce yourselves and

1                   affiliations, and the witness will be  
2                   sworn.

3                   MR. SPRAGUE: My name is Aaron  
4                   Sprague. I represent Robert Hogan, the  
5                   plaintiff.

6                   MS. ARNOLD-SIMMONS: My name is  
7                   Teresa Arnold-Simmons, and I represent Gray  
8                   Gable and Nassau Village Volunteer Fire  
9                   Department, Inc.

10                  MR. INCLAN: I'm Clemente Inclan  
11                  appearing on behalf of Baptist Hospital  
12                  Nassau.

13                  MR. TOLLEFSEN: I'm Thomas  
14                  Tollefsen appearing on behalf of third-party  
15                  defendant American Cancer Society.

16                  THE COURT REPORTER: Doctor,  
17                  would you please raise your right hand?

18                  (WHEREUPON, the witness was  
19                  duly sworn.)

20                  ROBERT NORMAN HARDEN, M.D.,  
21                  called as a witness by the Third-Party  
22                  Defendants Gray Gable and Nassau Village  
23                  Volunteer Fire Department, Inc., having been  
24                  first duly sworn, was examined and  
25                  testified as follows:

1 DIRECT EXAMINATION

2 BY-MS.ARNOLD-SIMMONS:

3 Q. Good afternoon, Dr. Harden. Would  
4 you please state your full name for the  
5 record, please?

6 A. Robert Norman Harden.

7 Q. And where are we currently  
8 today?

9 A. This is the Center for Pain  
10 Studies at the Rehabilitation Institute of  
11 Chicago.

12 Q. So we're in Chicago, Illinois?

13 A. Correct.

14 Q. Would you tell us briefly what  
15 your occupation is?

16 A. I'm a physician who works in the  
17 academic field. As such, I'm a clinician,  
18 researcher, administrator, manager, many  
19 hats.

20 Q. Could you take us through your  
21 educational background starting with your  
22 college degree?

23 A. Yes. I went to Emory and the  
24 University of Georgia for my undergraduate  
25 degree, got a degree in bachelor of

1 science. Then I began a PhD program in  
2 neurophysiology which was cut short because  
3 I was accepted to medical school; went to  
4 medical school, went to the Medical College  
5 of Georgia, did my internship and combined  
6 residency at the Medical University of  
7 South Carolina, did fellowship work back to  
8 Emory in the Pain Control Rehabilitation  
9 Institute of Georgia with a specialty in  
10 pain management. In the course of that  
11 fellowship I also spent time at Cornell and  
12 Sloan-Kettering University -- I'm sorry --  
13 Sloan-Kettering Cancer Center in New York.

14 Q. Let me --

15 A. And then.

16 Q. Let me stop you for a minute  
17 and ask you --

18 A. Yes.

19 Q. -- what year did you graduate  
20 from medical school?

21 A. 1984, wasn't it.

22 Q. Okay. And what did you do your  
23 residency in, what specialty?

24 A. It was actually a combined  
25 residency in internal medicine and

1                   neurology.

2                   Q.       And then you were telling us  
3                   about what happened when you were a fellow  
4                   in pain control in 1989?

5                   A.       Yes. I was -- did a fellowship  
6                   in pain control at Emory with rotations at  
7                   Cornell and Sloan-Kettering.

8                   Q.       Have you had any occasion to be  
9                   at the NIH for any reason?

10                  A.       Yes. I did observerships at the  
11                  NIH on three occasions.

12                  Q.       After medical school did you  
13                  begin your practice?

14                  A.       Well, no. Of course, after my  
15                  fellowship I began practice.

16                  Q.       Right, excuse me. Can you tell  
17                  us where you began your practice and what  
18                  type of practice?

19                  A.       Yes. I was on the staff of the  
20                  Medical University of South Carolina in the  
21                  departments of neurology and the department  
22                  of physical medicine rehabilitation. And I  
23                  was sort of the pain management guy at  
24                  Medical University of South Carolina. I  
25                  was the director of the pain control and



1 rehabilitation section, and I founded the  
2 headache program and a RSD program at  
3 Medical University of South Carolina.  
4 After working there for approximately four  
5 or five years, I was recruited to come  
6 here to Chicago and have worked ever since  
7 here as the director of the Center for  
8 Pain Studies. I was the founder of the  
9 Chronic Pain Care Center here and still  
10 serve as an attending physician in that --  
11 in that clinic as well.

12 Q. Currently what does your practice  
13 consist of from a clinical versus research  
14 versus teaching capacity?

15 A. As of the last three years I  
16 shifted my attention more to research. But  
17 I do still spend time in the clinics in  
18 that old clinic that I founded. And I see  
19 general chronic pain in that clinic but do  
20 have a tendency to see more complex  
21 regional pain syndrome than most doctors  
22 simply because that's an area of interest  
23 and specialization. But then I spend most  
24 of my time now as a researcher not only in  
25 the area of complex regional pain syndrome

1 but in other neuropathic conditions,  
2 spending a lot of time with post-amputation  
3 pain, spinal cord injury pain, post-stroke  
4 pain specifically.

5 Q. Prior to -- to going in or  
6 spending more time in research did you have  
7 a significant clinical practice?

8 A. Yes.

9 Q. How many years have you actually  
10 been in more of a clinical practice?

11 A. Approximately 18 years.

12 Q. What exactly is the Chronic Pain  
13 Care Center here in Chicago?

14 A. Well, the Chronic Pain Care  
15 Center is our clinical operation, and its  
16 sister organization, if you will, is the  
17 Center for Pain studies. So the one does  
18 the clinical work and the other does the  
19 research, but there's considerable overlap  
20 in terms of personnel, patients' emphasis,  
21 et cetera.

22 Q. Are either of these entities  
23 associated with the Rehabilitation Institute  
24 of Chicago?

25 A. They both are.

1           Q.       Okay. Tell us a little bit  
2           about the Rehabilitation Institute of  
3           Chicago, what type of organization that is,  
4           the facility.

5           A.       Well, it is a free-standing  
6           rehabilitation hospital. It is not for  
7           profit. It is affiliated with Northwestern  
8           University. And I am an associate  
9           professor at Northwestern University as all  
10          our doctors have an academic affiliation  
11          with Northwestern. We are fiscally  
12          different, separate from Northwestern but,  
13          of course, academically affiliated. The  
14          Rehab Institute of Chicago is the largest  
15          rehabilitation facility in the world. It's  
16          been ranked No. 1 in the nation for the  
17          last 15 years or so. I think we're  
18          actually up to 17 years now.

19                 So it's -- you know, it's a  
20          great place to work. I certainly enjoy  
21          the opportunity to try to serve  
22          rehabilitation patients. And complex  
23          regional pain syndrome remains an area of  
24          intense interest for us in terms of our  
25          research and clinical operation.

1 Q. Are you board certified, Dr.  
2 Harden?

3 A. Yes, I am.

4 Q. In what area?

5 A. Pain management.

6 Q. As part of your training being a  
7 pain management doctor I guess the focus is  
8 on pain?

9 A. Yes.

10 Q. Is there a component of your  
11 training that deals with psychiatric or  
12 psychological issues?

13 A. Yes.

14 Q. Can you explain to us what  
15 training you've had related to that?

16 A. Well, I have formal rotations  
17 that were required in my residency. I  
18 took a lot of extra training in psychiatry  
19 because I find it very interesting. And  
20 my forte has always been psychopharmacology,  
21 so a real interest in the drugs or the  
22 chemistry and how it worked. Since  
23 practicing, of course, as a pain management  
24 doctor it's necessary that a good pain  
25 management doctor understand and embrace

1 psychiatric diagnoses, learn how to make  
2 these diagnoses and, more importantly, learn  
3 how to participate and engage in  
4 psychotherapy.

5 We do some of the cognitive  
6 behavioral-based psychotherapy, but a lot of  
7 the psychotherapy we do is -- is -- are  
8 issues that will not be received well  
9 unless they come from the doctor. In  
10 other words, a lot of the things that I  
11 suggest or help patients to embrace or  
12 endorse -- they can only hear it from the  
13 doctor.

14 So we work hand in hand with  
15 psychologists and, of course, I take a  
16 great interest in who the psychiatrists in  
17 our centers are. As a matter of fact, I  
18 hired all of them that are working in our  
19 center now. I work very closely with them  
20 throughout. Obviously, since my role is  
21 primarily of psychopharmacology, of course  
22 that heavily overlaps with what they're  
23 interested in and what they're doing with  
24 their talk therapy. So that's my current  
25 role in the field of psychopharmacology, if

1                   you will, psychiatry and, of course,  
2                   psychology.

3                   Q.           Okay. Do you diagnose  
4                   psychiatric conditions such as depression?

5                   A.           Yes.

6                   Q.           Do you diagnose psychoschematic  
7                   pain disorders?

8                   A.           Yes.

9                   Q.           Okay. Do you prescribe  
10                  antidepressants?

11                  A.           Yes, I do.

12                  Q.           Do you have to be a psychiatrist  
13                  to treat psychiatric conditions?

14                  A.           No.

15                  Q.           How about to diagnose them?

16                  A.           No.

17                  Q.           Has your practice dealt with the  
18                  psychiatric issues of pain and how it  
19                  causes pain?

20                  A.           Yes.

21                  Q.           How do psychiatric issues explain  
22                  pain?

23                  A.           Explain pain?

24                  Q.           Uh-huh.

25                  A.           Well, I mean I guess the

1 question is if somebody has a primary  
2 psychiatric diagnosis, can that manifest as  
3 pain complaints? I'm sorry. I'm -- is  
4 that what you mean?

5 Q. Yes, sir.

6 A. Yes. As a matter of fact, we  
7 very frequently see that -- that people  
8 that -- really their primary diagnosis is  
9 depression or anxiety. They will manifest  
10 the subjective complaint of pain as a more  
11 socially acceptable complaint to get them  
12 in to see the doctor. So we embrace that,  
13 diagnose the depression or the anxiety, and  
14 treat it as such. And -- and in those  
15 cases, of course, the pain complaint  
16 usually diminishes or goes away.

17 Q. As a pain doctor, then, do you  
18 have to consider psychological or  
19 psychiatric issues in your treatment and  
20 diagnosis of patients with RSD, also known  
21 as CRPS?

22 A. Yes, ma'am, absolutely. It's 80  
23 plus percent of our patients with that --  
24 with that -- what the psychiatrist would  
25 call Axis 1 diagnosis of CRPS have an Axis

1           2 diagnosis, such as depression or anxiety.  
2           It is ubiquitous in the field and must be  
3           diagnosed and treated.

4           Q.        Have you, in fact, written  
5           articles related to -- dealing with pain  
6           and psychological status?

7           A.        Yes, ma'am.

8           Q.        And I'm going to embarrass you a  
9           little bit here, Doctor, and ask you to  
10          kind of toot your own horn about your  
11          awards. Have you received any awards  
12          related to your position as a physician?

13          A.        Yes, ma'am.

14          Q.        Okay. Could you tell us what  
15          those are?

16          A.        Well, there's a few. For  
17          instance, Castle Connolly does national  
18          doctors and local doctors. So I've been  
19          the top doc. I've been mentioned as top  
20          local doctor in the field of pain  
21          management. I've received teaching awards;  
22          for instance, the Golden Apple. I was  
23          considered to be teacher of the year at  
24          RIC one year. I've received awards from  
25          philanthropic organizations for my work in



1                   this area, particularly the Reflex  
2                   Sympathetic Dystrophy Association. There's  
3                   a few things there.

4                   Q.        Okay. Are you on any editorial  
5                   boards?

6                   A.        Yes.

7                   Q.        Which boards are you on?

8                   A.        I'm actually on about a dozen,  
9                   maybe -- I'm senior editor on the Journal  
10                  Pain Medicine. I'm a senior editor of the  
11                  Journal Pain Practice. I have -- I am  
12                  section editor of the American Pain Society  
13                  Bulletin. And then there's multiple  
14                  journals that I serve as -- on the  
15                  editorial board and certainly many more  
16                  that I'm a reviewer.

17                  Q.        Okay. Do you have any  
18                  professional -- are you a member of any  
19                  professional associations related to your  
20                  position as a physician, pain management  
21                  physician?

22                  A.        Yes, ma'am.

23                  Q.        What would those be? Just name  
24                  a couple of them for us.

25                  A.        Well, internationally I belong to

1 the International Association for the Study  
2 of Pain. I have sporadically been a  
3 member of the World Institute of Pain which  
4 is sort of the second big international  
5 organization. Nationally I'm a long-term  
6 member of the American Pain Society. I am  
7 also a member of the American Academy of  
8 Pain Medicine. There's several.  
9 Regionally I'm the -- actually have been  
10 the president of Midwestern Pain Society  
11 and still -- still remain a member of  
12 that.

13 I'm on the Headache. I'm in  
14 Headache Societies. I am in Physical  
15 Medicine and Rehabilitation Societies. I'm  
16 actually an honorary member of the American  
17 Academy of Physical Medicine and  
18 Rehabilitation which makes me the only --  
19 the only neurologist and the only doctor --  
20 nonphysiatrist doctor to actually hold that  
21 post. They've had three or four before,  
22 but I think both of those are deceased  
23 now. So I -- I may be the only living  
24 honorary doctor for the AAPMR. It was  
25 actually quite a honor and I -- I felt

1 touched that they felt -- felt they need  
2 to come out and grab me and pull me in to  
3 their club because normally I couldn't join  
4 that.

5 Q. Are you on any boards or  
6 committees related to reflex sympathetic  
7 dystrophy?

8 A. Yes, I am.

9 Q. Are you involved at all with the  
10 Reflex Sympathetic Dystrophy Syndrome  
11 Association of America?

12 A. Yes, I am.

13 Q. Can you tell us what your  
14 involvement is with that organization?

15 A. I'm the -- one of the chairman  
16 of their clinical advisory board with Dr.  
17 Rodge (phonetic).

18 Q. And what are your duties related  
19 to that?

20 A. I'm the scientific liaison. When  
21 they have questions or their members have  
22 questions about -- scientific questions,  
23 specifically diagnosis, drugs, treatments,  
24 what works, what doesn't work, what their  
25 -- what the status of the evidence about

1 particular treatments, for instance, that's  
2 my role.

3 Q. Have you written in your career  
4 as a physician articles related to complex  
5 regional pain syndrome also known as reflex  
6 sympathetic dystrophy?

7 A. Yes, ma'am.

8 Q. Have you written any books  
9 related to those symptoms?

10 A. I've edited books, yes, ma'am.

11 Q. Okay. And who were those books  
12 generally written for, what's the audience?

13 A. The audience is a specialty  
14 scientific audience in most cases. The two  
15 -- the two books I wrote with the  
16 International Association for the Study of  
17 Pain is -- was written for a very  
18 specialty pain management audience.  
19 However, some of the -- some of the  
20 ramifications of that which was the  
21 diagnosis of CRPS was meant for any doctor  
22 who would presume to make a diagnosis of  
23 CRPS, sort of guidelines that they could  
24 use to make that diagnosis properly.

25 Another book that I had edited

1 was about treatment guidelines for this.  
2 And, again, this was less for the specialty  
3 pain management doctors who are supposed to  
4 know about that. They're supposed to keep  
5 up with the literature but more for the  
6 other doctors, the primary care doctors and  
7 the secondary care doctors that would be  
8 called on to treat CRPS in the community.  
9 This is meant to be a practical pragmatic  
10 guide so that they could organize their  
11 thinking about how to properly treat the  
12 disease.

13 Q. And those are the treatment  
14 guidelines for complex regional pain  
15 syndrome?

16 A. That's correct.

17 Q. Do you make presentations on  
18 complex regional pain syndrome?

19 A. Yes, ma'am.

20 Q. Okay. Are you making any  
21 presentations the week of September 24th?

22 A. The week of September 24th, no,  
23 I'm not. I'll be in -- I'll be in  
24 Holland. I will be serving as a senior  
25 advisor to the Dutch who are trying to

1 organize their effort in terms of diagnosis  
2 and treatment of CRPS. I won't be making a  
3 formal presentation, but I'm sure I'll have  
4 a lot of comments about what they're up  
5 to.

6 Q. And is your trip to Holland is  
7 -- that is why we are taking your video  
8 today?

9 A. Yes.

10 Q. Because you will not be able to  
11 be with us in Florida on that day?

12 A. No, ma'am. Unfortunately, I'll  
13 be in Europe.

14 Q. So, I take it, based on  
15 everything you -- you've told us that the  
16 research, diagnosis, and treatment of RSD  
17 is a large part of your practice?

18 A. Yes, ma'am.

19 Q. I'm going to hand you a copy of  
20 what was provided to us as your CV. It  
21 looks like it's 35 pages long and ask you  
22 if that is a copy of your CV?

23 A. Yes, ma'am.

24 Q. And was I correct? It's 35  
25 pages long?

1 A. Let me see what that version is.

2 Q. Okay.

3 A. That's July 18th. There's a  
4 newer one that's 36 pages but, yes.

5 Q. The newer version, what  
6 additional things would have been added to  
7 that that's not on this version?

8 A. Well, really all the research  
9 that we've done or presented since  
10 obviously July, but mostly poster  
11 presentation and abstracts at scientific  
12 meetings that have happened since then. As  
13 a senior researcher, I get mentioned on an  
14 awful lot of research for the -- for my  
15 young colleagues out there that they're --  
16 that they're working on different projects  
17 and things. I coordinate and direct their  
18 efforts, but they're very energetic and  
19 there's a lot.

20 Q. Dr. Harden, could you explain to  
21 the jury generally exactly what is reflex  
22 sympathetic dystrophy which we also use  
23 another term for that, I think, complex  
24 regional pain syndrome?

25 A. Well, this is basically a

1 disease or a syndrome that has been around  
2 since the dawn of time. We certainly see  
3 reference to this in some of the very,  
4 very old Samarian tablets, for instance.  
5 So as far as we know it's been around  
6 since people have been injured. It is a  
7 syndrome that is characterized in general  
8 by pain and a disturbance of the autonomic  
9 nervous system. You need to have both, and  
10 that is the -- sort of pivot of this  
11 disease and what it is.

12 It was actually well  
13 characterized by a doctor in the 18 --  
14 late 1800's with civil war veterans or, as  
15 we say in the south, war between the  
16 states. So Silas Weir Mitchell gave really  
17 good diagnostic information back in the  
18 1800's, but it seems like there hadn't been  
19 an awful lot of movement in terms of  
20 making that diagnosis properly until  
21 recently. But basically it's a disease  
22 that is characterized by pain, disturbance  
23 of the autonomic nervous system which is  
24 the automatic nervous system that controls  
25 the tone of blood vessels. It controls



1           sweating. It is associated with central  
2           features that manage pain, and it's  
3           remotely related to other changes you can  
4           see in the peripheral like changes in skin  
5           and hair. So all of these features as a  
6           collection are what we use to make the  
7           diagnosis now.

8           Q.       What happens to the autonomic  
9           nervous system when someone is injured?  
10          Give us a little science lesson, I guess,  
11          on that if you would.

12          A.       Well, in an acute situation if  
13          somebody is injured, their sympathetic  
14          nervous system immediately goes into play  
15          and clamps down the blood vessels so  
16          somebody doesn't, for instance, bleed to  
17          death. As time passes normally the  
18          sympathetic nervous system resets. The  
19          pain begins to subside as healing occurs.  
20          In CRPS something goes wrong, and the  
21          sympathetic nervous system goes haywire.  
22          It starts to sort of act on its own  
23          without any -- any useful further input  
24          from the pain system.

25                    The pain begins to increase we

1           feel due to a relationship between the  
2           damage and the periphery and the changes in  
3           the central nervous system which feed back  
4           to the periphery. And it forms these very  
5           negative feedback loops between the central  
6           nervous system and the periphery unit, and  
7           it becomes self-maintaining. Now -- in  
8           excess of what you would expect with normal  
9           healing and normal recovery.

10           Q.       Is the diagnosis of CRPS  
11           controversial?

12           A.       That's an interesting question. I  
13           don't think it's controversial. I mean I  
14           think that for years there was sort of a  
15           chaotic idiosyncratic approach to the  
16           diagnosis meaning that, you know, whatever  
17           doctor sat down in front of a typewriter  
18           in those days could type out a criterion,  
19           and people would begin to use that. But  
20           lately we have begin to -- begun to apply  
21           the principals of the validation of a  
22           diagnostic criteria to this disease.

23                       Interestingly, this is one of  
24           the first pain diseases that that's ever  
25           been done. It was actually done well for

1 psychiatric disease for years and then done  
2 pretty well for headache, and now finally  
3 we're trying to get it into other -- other  
4 areas of chronic pain. Complex regional  
5 pain syndrome represents the very first one  
6 that a statistical derivation was done of a  
7 diagnostic criteria, and we have that  
8 extent now.

9 Q. Is there disagreement in the  
10 medical community as to, then, how to make  
11 the diagnosis? Was that part of the  
12 problem?

13 A. There's not disagreement. There  
14 may be ignorance. There may be an  
15 unwillingness to embrace the criteria that's  
16 accepted in the field but, you know, pretty  
17 much doctors in the field -- pain  
18 management doctors are in good and general  
19 agreement about how to make the diagnosis  
20 and what criteria should be used.

21 Q. Have you studied the criteria  
22 and help establish criteria to make the  
23 diagnosis of CRPS with more specificity?

24 A. Yes.

25 Q. And can you explain to the jury

1           what we mean by specificity as it relates  
2           to the diagnostic criteria?

3           A.       Well, there's really two features  
4           of a good criteria. The one is  
5           sensitivity meaning that it doesn't leave  
6           anybody out. If somebody has it, they're  
7           going to get caught by this net, if you  
8           will, of the criteria. Specificity refers  
9           to the ability of that criteria to exclude  
10          other disease; in other words, somebody  
11          comes in with a collection of signs and  
12          symptoms that is CRPS. You want a  
13          criteria that is sensitive enough to  
14          capture those with a high frequency, but  
15          you also want a criteria that is specific  
16          enough to not include things like other  
17          neuropathic disease or headache or  
18          postherpetic neuralgia or diabetic  
19          neuropathy.

20                 You want to make sure that your  
21          criteria is specific enough to rule those  
22          out. Those are other diagnoses. They  
23          need to -- doctors need to make the other  
24          diagnoses and treat them as such whereas  
25          you really want to identify properly the

1 people with CRPS with sensitivity and  
2 specificity so you're treating the right  
3 thing. You're not over treating or you're  
4 not under treating or you're not using  
5 treatments that are unsafe or, you know,  
6 irrelevant to the prognosis of the patient.

7 Q. And is one of the reasons the  
8 diagnostic criterion are necessary for CRPS  
9 because there's not one test that can  
10 diagnosis the condition like an X-ray or  
11 something?

12 MR. SPRAGUE: Objection, leading.

13 THE WITNESS: Okay. You're  
14 correct. There is no gold standard test.  
15 It would be nice if we had, you know, a  
16 simple objective test that we could give  
17 somebody, you know, a blood test that we'd  
18 get a number and say you got it. It's  
19 not that kind of a syndrome. It is a  
20 collection of signs which are to things  
21 that a doctor observes and symptoms which  
22 are complaints that the patient would have;  
23 things like pain and swelling, et cetera.

24 BY MS. ARNOLD-SIMMONS:

25 Q. Have you published any articles

1 or materials related to the diagnostic  
2 criterion and their specificity?

3 A. Yes.

4 Q. Are you familiar with diagnostic  
5 criteria as propounded by the IASP?

6 A. Yes, ma'am.

7 Q. Can you explain to us what the  
8 IASP is?

9 A. The International Association for  
10 the Study of Pain; that's the big  
11 international organization that dedicates  
12 itself to the study of -- and the  
13 promulgation of research in the area of  
14 pain management. So the IASP is huge in  
15 my field. They're the big guys. So they  
16 -- they have meetings. They have a lot of  
17 sociologic efforts in regards to educating  
18 patients, educating doctors, but they also  
19 involve themselves in, of course, publishing  
20 a journal. And it is considered the best  
21 journal in pain management. It's the  
22 Journal of Pain. And they publish books  
23 -- actually a series of books. I happen  
24 to have one of them here about the  
25 diagnosis and the treatment of pain and

1 research and pain. So that's -- that's  
2 who the IASP are, the International  
3 Association for the Study of Pain.

4 Q. And have you been involved with  
5 the ISA -- IASP? Excuse me.

6 A. Yes, I have.

7 Q. Now, can you explain to us the  
8 IASP criteria? And if you need to, there  
9 is a dry erase board behind you that we  
10 could move so you can write those down.

11 A. Okay. Since this is important,  
12 I like to make sure I get this correct.  
13 And if you don't mind, I'll just carry it  
14 with me. But I'll be happy to --

15 MS. ARNOLD-SIMMONS: If you'll  
16 give the videographer one second to adjust  
17 so we can see the dry erase board.

18 THE VIDEOGRAPHER: Going off the  
19 record, the time is 17:46:42. We are off  
20 the record.

21 (WHEREUPON, a discussion was had  
22 off record.)

23 THE VIDEOGRAPHER: Back on the  
24 record in the deposition of Norman Harden  
25 (sic). The time is 17:48:14. Counsel,

1                   you may proceed.

2                   BY MS. ARNOLD-SIMMONS:

3                   Q.        Dr. Harden, I think I asked you  
4                   to list for us the diagnostic criterias  
5                   propounded by the IASP.

6                   A.        Yes.  There's actually four  
7                   criterion or four steps that are used to  
8                   make this diagnosis.  So the first  
9                   criterion says the presence of an  
10                  initiating noxious event or cause of  
11                  immobilization.  So this is -- noxious  
12                  event, something bad happened, and  
13                  immobilization.  The second criteria says  
14                  continuing pain, allodynia, or hyperalgesia  
15                  with which the pain is disproportionate to  
16                  any inciting event.  So it's pain.

17                  Allodynia means something that is  
18                  normally an innocuous stimulus such as a  
19                  light touch is now very painful up to and  
20                  including things like even the presence of  
21                  clothing or even an air conditioner coming  
22                  on and blowing across the affected part in  
23                  extreme cases.  Hyperalgesia simply means  
24                  sort of the same thing where there's a  
25                  sensory phenomenon that is now very, very



1 sensitive. So hyperalgesia which is  
2 disproportionate to any inciting event  
3 meaning that if somebody gets a needle  
4 stick and just to get a shot -- of course  
5 that hurts when it happens. But if the  
6 pain is intense and the pain is long  
7 lasting we would say it is  
8 disproportionate.

9 Q. Okay.

10 A. And then it says the third  
11 criterion is evidence at some time of edema  
12 or swelling, changes in skin, blood flow,  
13 or abnormal sudomotor activity in the  
14 region of pain. Now sudomotor is, again,  
15 back to that sympathetic nervous system.  
16 This is how the nervous system controls  
17 sweating. So when it says sudomotor that  
18 equals sweat. Now remember I said that  
19 the sympathetic nervous system also controls  
20 blood flow to a part. So this, again, is  
21 the sympathetic nervous system. And edema,  
22 swelling may or may not be related to the  
23 sympathetic nervous system.

24 Q. What is the fourth criteria?

25 A. It says this diagnosis is

1 excluded by the existence of conditions  
2 that would otherwise account for the degree  
3 of pain and dysfunction. In other words,  
4 you have to make sure that there's no  
5 other better explanation; in other words,  
6 not another diagnosis that explains the  
7 signs which is what the physician observes  
8 or the symptoms which is what the patient  
9 portrays. So no better diagnosis or  
10 explanation for the sign and symptoms.

11 Q. Okay. When were the IASP  
12 criteria adopted?

13 A. They were adopted by a consensus  
14 group that met in Orlando Florida in 1994.

15 Q. And have there been any problems  
16 with the IASP criteria?

17 A. Yes. There have been big  
18 problems. Now this group when they got  
19 together in '94, their charge was to -- to  
20 try to clean this up, clean up some of the  
21 chaos and confusion that had surrounded  
22 this diagnosis prior to. And one of the  
23 ways that they did that was to say we're  
24 going to -- we're going to stop using the  
25 term reflex sympathetic dystrophy. The

1 point being there may or may not be  
2 reflexes involved. We don't know that.  
3 The sympathetic nervous system must be  
4 involved, but it may be an innocent  
5 bystander.

6 So RSD implied that the  
7 sympathetic nervous system caused the  
8 problem. It may not be the cause. And  
9 probably, most importantly, dystrophy only  
10 occurs in about 10 percent of cases. So  
11 the old name RSD or reflex sympathetic  
12 dystrophy was considered misleading. So  
13 the first thing that this consensus group  
14 did -- and I might add it took them a  
15 long time to do this. They had to meet  
16 two times before Orlando to agree on a  
17 name. But they agreed on the term complex  
18 regional pain syndrome as a general name,  
19 sort of a general overarching theme that  
20 would be very sensitive, that would bring  
21 everybody under the umbrella, if you would.  
22 Nobody would be left out in the cold or in  
23 the rain.

24 So their charge was to create a  
25 very broad, very general name and a very

1 broad and very general diagnostic criteria  
2 that would make sure that nobody was left  
3 out. And they did that by creating a list  
4 here that was very, very open, very  
5 general, sort of loose. They did that  
6 with the full understanding that they would  
7 create a very sensitive diagnostic criteria  
8 with the understanding that the research  
9 would be done to validate this empirically  
10 so that we would improve the specificity as  
11 time passed.

12 Q. Okay.

13 A. So what we have with the IASP  
14 criteria is a very broad, very general  
15 diagnosis that really a lot of people can  
16 get under this umbrella. The specificity  
17 was supposed to come with research over  
18 time, empirical validation research.

19 Q. Through the use of this broad  
20 criteria have there been experienced  
21 problems by pain management?

22 A. Yes, a lot of over diagnosis;  
23 many people would come in with something  
24 else. Somebody would sort of superficially  
25 go through this, not thoughtfully go

1 through this and they would conclude that  
2 they had RSD or CRPS -- to use the proper  
3 term -- and then they would charge off  
4 with very expensive treatments, very  
5 inappropriate treatments, and in some cases  
6 very dangerous treatments.

7 Certainly there's a lot of the  
8 more experimental treatments that are quite  
9 dangerous. They have death and morbidity  
10 associated with those. So, you know, it's  
11 a real problem if you use a criteria like  
12 this and you bring too many people into  
13 the fold and begin treating all of those  
14 people as if they truly had the disease;  
15 so a lot of over diagnosis. There has  
16 also been abuses in -- obviously in this  
17 field of people that because this is so  
18 loose, they can easily make this diagnosis.  
19 And they can actually make it on the basis  
20 of a subjective report of the patient. So  
21 that a patient that comes in and says, you  
22 know, I had a noxious event; I have this  
23 pain and allodynia which is subjective --  
24 that's my report. And I had edema and  
25 blood flow changes, swelling, and, you

1 know, it was -- the limb got cold one  
2 time, and it turned blue. And it was  
3 sweating. They can say that that happened  
4 weeks ago or months ago. So to use this  
5 criteria you would have to say, well, maybe  
6 that's CRPS. And then it's the doctor's  
7 absolute responsibility to look at this and  
8 say is this what it is or are there other  
9 possibilities that could explain this?

10 But there have been a lot of  
11 abuses. Either doctors that, you know,  
12 take this as gospel in the sense that they  
13 feel like this is a real good tight  
14 specific criteria, which it is not, or  
15 patients that -- for instance, the  
16 psychiatric patient that can't come in and  
17 say I'm depressed, I'm crying all the time.  
18 But they can come in to a doctor. They  
19 feel comfortable saying I have pain. It's  
20 horrible. My hand turns blue. And the  
21 doctor looks at it and says, well, your  
22 hand is not blue now. They say, well, but  
23 on the basis of this we can accept  
24 historical symptoms.

25 So there were a lot of problems

1           that came in directly due to this point  
2           that -- I know it's very technical but  
3           this point that, you know, it's very  
4           sensitive, embraces many people that have  
5           pain and anything going wrong with them but  
6           not specific. See, the problem is that we  
7           didn't -- for a long time we didn't really  
8           get back and improve on the specificity of  
9           this so we can now say this is really what  
10          you have. You don't have something else.  
11          It's just sort of sliding in under the  
12          umbrella to use the analogy I keep using.

13           Q.       So the I -- can using the IASP  
14           criteria -- you can make a diagnosis  
15           through a patient report alone without a  
16           physician seeing signs or symptoms?

17           A.       That's correct.

18           Q.       Have other criteria --

19           A.       Well, I want to add to that  
20           too.

21           Q.       Sure.

22           A.       That, of course, ignores Criteria  
23           4 because it's still the absolute  
24           responsibility of the physician to do a  
25           good history to do a good physical

1 examination. So they're going to have data  
2 that will allow them to answer Criterion  
3 No. 4 which is -- you know, there's other  
4 possibilities. And really good practice  
5 requires that they list the other  
6 possibilities as well. To not go into a  
7 situation and say, well, somebody said this  
8 was CRPS, so I guess that's what this is  
9 and to focus on that as the only  
10 possibility.

11 Q. Okay.

12 A. Okay.

13 Q. Since 1994 have other criteria  
14 been developed to address these issues?

15 A. Yes.

16 Q. Can you tell us what those  
17 criteria are?

18 A. Did you want me to sit down  
19 or...

20 Q. Well, I'm going to ask you to  
21 write those events on the board, if you  
22 can tell us what they are.

23 A. Well, the -- remember the charge  
24 was that these guys were going to create  
25 this broad loose umbrella diagnostic



1 criteria because they trusted that research  
2 would march ahead and that we would do  
3 what's called empirical validation of these  
4 criteria, meaning we would go to the  
5 patients. We would run the statistics and  
6 we would improve on that specificity -- I'm  
7 sorry -- the, the -- yeah, the specificity  
8 feature. So we actually did that. We --  
9 we empirically validated these criteria.

10 Q. When you say we, were you  
11 involved in that?

12 A. Yes. I was.

13 Q. I'm sorry. I didn't mean to  
14 interrupt you.

15 A. We being a team of  
16 biostatistician and clinicians and actually  
17 I like to include my patients in this. I  
18 feel like our patients helped us a lot to  
19 try to understand what was going on. But  
20 we collected all the signs and all the  
21 symptoms that had ever been mentioned in  
22 any literature about this in a very  
23 methodical way. And then we went back and  
24 said what signs and what symptoms are  
25 specific to this diagnosis and what signs

1           and what symptoms don't help us to rule  
2           out other disease; in other words, to  
3           develop a statistical process where we  
4           could now start to talk about improving the  
5           specificity.

6                     In fact, the specificity for  
7           making the diagnosis -- in other words, the  
8           ability to rule out other disease is less  
9           than a coin flip using this criteria. In  
10          other words, it's less than a doctor just  
11          saying, well, do they have it or not and  
12          flipping a coin. It's 40-percent specific.  
13          It's essentially 100-percent sensitive but  
14          it's only 40-percent specific. It's less  
15          good than a coin flip. So using the  
16          empirical validation, internal validation,  
17          external validation, factor analysis,  
18          cluster analysis, all these biostatistical  
19          technologies that are -- that are well  
20          proven. They were proven in psychiatry and  
21          proven in headache and now applied to  
22          improving criteria for all pain -- should  
23          be applied for all pain for all diagnoses  
24          in medicine for that matter.

25                     But we applied this to this, and

1 we did create a criteria which was  
2 considerably more specific, and basically  
3 twice as specific. So now we were  
4 achieving specificity -- can achieve  
5 specificity of 80 percent or better which  
6 is great for any type of diagnostic test;  
7 for X-rays or MRI's or whatever. A  
8 specificity of 80 percent is good. A  
9 specificity of 40 percent is terrible. So  
10 somewhere in there is where you want to  
11 be. And we retained good sensitivity I  
12 might add on the basis of this empirical  
13 derivation of a new criteria.

14 Q. And what year was the new  
15 criteria developed?

16 A. Well, it was over time. It  
17 obviously takes a lot of time and a lot of  
18 work to do this. But we had been doing  
19 this work in the late 90's and published  
20 the two similar articles about this in  
21 1999, the internal and external validation.  
22 Then we took that statistical derivation of  
23 the research criteria and clinical criteria  
24 and then we subjected that to consensus  
25 meaning we went back to the research

1 community and the IASP sponsored a -- a --  
2 what's called a closed workshop, basically  
3 a think tank of all the international  
4 experts we could get ahold of; researchers,  
5 clinicians, interventionalists, PhD's,  
6 everybody that was involved either in  
7 trying to understand how the disease worked  
8 or how to treat the disease or how to  
9 diagnosis the disease and put these people  
10 in a room actually in the city of Budapest  
11 which is why it's called the Budapest  
12 criteria.

13 And they sat down and very  
14 critically assessed what the statistics were  
15 telling us and said does this make sense  
16 in terms of reality? Does this make sense  
17 in terms of really diagnosing these people  
18 that we see in our clinics all the time?  
19 And to make a long story short after four  
20 days of hard work and a lot of jumping up  
21 and down and yelling and screaming, if you  
22 will, but -- but -- but, finally, at the  
23 end of the day we were in agreement and we  
24 said yes.

25 Q. And what you --

1           A.       These statistically derived  
2           criteria are good. They do apply and we  
3           should embrace these. We should adopt  
4           these as the criteria for making the  
5           diagnosis.

6           Q.       And what year was that?

7           A.       That would have occurred around  
8           about 2001 I believe.

9           Q.       And so the more -- and so I  
10          understand, is the more specific diagnostic  
11          criteria --

12          A.       Yes.

13          Q.       -- called the Budapest criteria?

14          A.       Correct.

15          Q.       And what's written on the board  
16          right now is not the Budapest criteria?

17                   (Indicating.)

18          A.       No. This is the IASP. This is  
19          the loose, very sensitive, not very  
20          specific criteria with the less than coin  
21          flip specificity in terms of ruling out  
22          other disease when somebody walks into your  
23          office.

24          Q.       Can you -- I don't know if  
25          you --

1           A.       This is the one that we were  
2           meant to improve on.  And that's -- that's  
3           what the charge with the crew that put  
4           this together.  They said, look, we don't  
5           have to worry about this right now.  We  
6           want to get something going so that  
7           research can kick in and improve on this.  
8           They had no intention on this going forward  
9           as a long-term solution to the real problem  
10          of diagnosing this.

11          Q.       Can you set out for us on the  
12          board then what are the Budapest diagnostic  
13          criteria?  And we're talking about clinical  
14          criteria.  Is that right?

15          A.       Well, there's two.  Let me  
16          explain.  In research you need to have the  
17          maximum degree of specificity, meaning that  
18          you don't want anybody in your research  
19          paradigms that don't have CRPS because  
20          you're doing such thing as trying to  
21          determine if drug A works or therapy B  
22          works.  So you can't have people that  
23          don't have the disease in there or your  
24          results are obviously completely  
25          compromised.  So in research we have to

1           have a maximum degree of specificity.

2                       However, because as you move up  
3           in terms of higher and higher specificity  
4           -- and we -- the Budapest research criteria  
5           goes up above 90 percent specificity which  
6           is extraordinary in medicine. Because the  
7           sensitivity started to fall off what  
8           happened was a lot of people that had  
9           previously had a diagnosis of CRPS would  
10          now go undiagnosed if we use that. So  
11          some of the clinicians in the group,  
12          including myself, said, well, maybe this is  
13          too hard. This is too hard a cut. Let's  
14          drop back and create a corollary criteria  
15          that's very similar. There's actually only  
16          one difference, as I'll go into.

17                      Let's create a corollary criteria  
18          that's more for clinical use people who  
19          have the disease will still retain good  
20          specificity. We'll still retain much  
21          better sensitivity than we would get with a  
22          research criteria. And let's do that and  
23          create a clinical criteria that is now  
24          statistically derived and consensus  
25          validated by the IASP group here and

1 include that as well. So, in other words,  
2 clinically as is relevant in this case we  
3 want to use the clinical -- the Budapest  
4 clinical diagnostic scheme, okay.

5 (Indicating.)

6 Q. Okay.

7 A. Are we clear?

8 Q. We are. Could you set out then  
9 for us what the Budapest clinical  
10 diagnostic criteria are?

11 A. Okay. Now -- okay, this is  
12 what the statistics tell us. We looked at  
13 the patients and they now -- we say these  
14 signs and these symptoms. So it's going  
15 to be similar to IASP, but there will be  
16 some differences.

17 Q. Okay.

18 A. So basically the signs and  
19 symptoms that we saw in our patients fell  
20 into four categories. One of the  
21 categories was sensory. And that's pain.  
22 And that is this thing, allodynia, I talked  
23 about. Innocuous stimuli is now painful or  
24 painful stimuli is now very, very painful.  
25 It is vasomotor. And that's the



1           sympathetic nervous system acting on the  
2           vessels. In other words, if you have the  
3           sympathetic nervous system active, it causes  
4           vasoconstriction which when you look at a  
5           limb, it's going to cause that limb to be  
6           blue colored. It's going to cause that  
7           limb to be cooler relative to a part that  
8           doesn't have the sympathetic activity and  
9           vasoconstriction. So vasomotor just simply  
10          means the size and the diameter of the  
11          blood vessels which controls the heat and  
12          the color of the limb -- of the skin.

13           Q.        Okay.

14           A.        Then there was sudomotor. Now  
15          everybody knows that's sweat. But also was  
16          included in that was edema.

17           Q.        By edema what do we mean?

18           A.        Swelling. And then there's  
19          motor. Now, this is something that's new  
20          because we talked about these symptoms  
21          before in terms of the IASP criteria and  
22          just basically defining what the disease  
23          is. But the new one -- this is what the  
24          patients told us and the statistics require  
25          that we include in the new criteria was

1 motor changes. And this means power,  
2 strength. It also means things that are  
3 wrong with the motor system such as tremor.  
4 People you can actually see with this  
5 disease have these strange tremor, and  
6 sometimes they have contractions of the  
7 muscles or contractions sometimes long term  
8 of the tendons. They have these -- these  
9 -- these very awkward postures that you  
10 see.

11 So these all fall under this  
12 category of motor. Now -- so the  
13 statistics say, yes, pain is clearly in  
14 there. So Criterion 1 of the new criteria  
15 -- the Budapest criteria requires that you  
16 have pain and, of course, pain that is  
17 disproportionate with what you would expect.  
18 If somebody has a fracture, certainly it's  
19 going to hurt. But as the healing occurs  
20 and the fracture resolved the pain goes  
21 away. So if somebody has pain six months  
22 or a year later after the fracture is  
23 resolved, that is disproportionate.

24 So we still had the  
25 disproportionate, disproportionate pain.

1 But the thing we lose on this was the  
2 immobilization. Everybody knows that  
3 immobilization is terrible with CRPS and  
4 the treatment guidelines at least -- so we  
5 took that out. It didn't -- it was always  
6 a nonmandatory criteria anyhow. So that  
7 was gone -- statistically gone and gone by  
8 consensus by the Budapest group.

9 (Indicating.)

10 Q. Okay.

11 A. Criterion No. 2 requires that we  
12 have -- I'm sorry. At this point it  
13 doesn't -- it doesn't fall under the 1, 2,  
14 3, 4 kind of thing. But basically under  
15 No. 2 we required, for a clinical diagnosis  
16 of CRPS by the Budapest criteria, that we  
17 had at least one -- now, this is -- this  
18 is -- remember I said that there's signs  
19 and symptoms.

20 Q. Right.

21 A. Patients come in and tell us  
22 symptoms. So the Budapest clinical  
23 requires that there is at least a symptom  
24 in three of these four categories.

25 Q. And so we understand, what is a

1 symptom?

2 A. It's pain or it's I can't move  
3 my arm or it's my hand turned blue.

4 Q. Are symptoms something that  
5 patients report?

6 A. Correct.

7 Q. Okay. How is that different  
8 from a sign?

9 A. A sign is something that a  
10 physician observes and documents.  
11 Obviously, the physician can observe  
12 symptoms -- observe signs but not document  
13 them, but then they're useless in terms of  
14 moving forward with diagnostic criteria or  
15 certainly court where you need it to be  
16 documented in the record. But -- so they  
17 have to have symptoms of what the patient  
18 tells us is going on with them, and they  
19 have to have three of four; in other  
20 words, at least one -- one symptom in all  
21 -- in three of these four categories, okay.

22 Q. Okay.

23 A. Three requires that there be two  
24 signs. This means it's something that the  
25 physician observes; the color changes, the

1 temperature changes, the allodynia, motor  
2 changes like the dystonia in two, so signs  
3 in two of four categories. So you have to  
4 have Criterion 2 now which is three or  
5 four symptoms. You have to have Criterion  
6 3 which is two or four signs. And then  
7 four we still have no matter explanation,  
8 no other diagnosis -- no other diagnosis,  
9 okay. And all of these, of course, are  
10 required now to make the clinical diagnosis  
11 under the Budapest criteria. Now, if you  
12 do that -- you'll notice that there's not  
13 a huge change from what we were doing with  
14 the IASP except we're now requiring the  
15 doctor see at least two of these  
16 quasi-objective signs in the lab or in the  
17 clinic and record those as well as three  
18 of four symptoms.

19 So they require that they have  
20 to have that. To achieve that specificity  
21 they have to have that to retain that  
22 sensitivity and, of course, it -- it  
23 improves the diagnostic acumen, if you  
24 will, by 100 percent because we go from a  
25 40-percent -- I'm sorry -- .4 specificity

1 to .8 specificity, 100-percent improvement.

2 Q. Doctor, I think you might retake  
3 your seat, and I think we may need to  
4 change out the tape.

5 A. Okay.

6 THE VIDEOGRAPHER: This is the  
7 end of tape No. 1 in the deposition of  
8 Norman Harden (sic). The time is 18:14:16.  
9 We are off the record.

10 (WHEREUPON, a discussion was had  
11 off record.)

12 THE VIDEOGRAPHER: Back on the  
13 record. This is the beginning of tape No.  
14 2 in the deposition of Norman Harden (sic).  
15 The time is 18:19:47. Counsel, you may  
16 proceed.

17 MS. ARNOLD-SIMMONS: Thank you.

18 BY MS. ARNOLD-SIMMONS:

19 Q. Dr. Harden, you had just set out  
20 for us the Budapest criteria, and I want  
21 to make sure I -- I understand -- or the  
22 jury understands. Can you tell us what  
23 you mean by sensory component of a patient  
24 report?

25 A. This is subjective presentation

1 by the patient; when it's symptoms it mean  
2 things like I have pain, it's burning, it's  
3 stinging, it's electrical. If it is signs,  
4 it is more in the hands of the  
5 investigator such as they will -- they will  
6 use a light touch or a brush or a light  
7 touch of their hand to see if that causes  
8 pain now which is allodynia. They will  
9 use a pin and actually do light pin  
10 pricks, and the patient will say, you know,  
11 that hurts, of course, with a pinprick.  
12 But if it hurts intensely and it hurts for  
13 a long way, that's hyperpathia. So those  
14 are the components, both signs and  
15 symptoms, that we would be looking for in  
16 the diagnosis of CRPS.

17 Q. What about for the vasomotor  
18 area? What are the symptoms that a  
19 physician would look for to comply with  
20 that part of the criteria?

21 A. Well, the -- the symptoms the  
22 patient will portray was -- either  
23 temperature changes in the limb or color  
24 changes in the limb; both of which reflect  
25 the amount of blood flow to the -- the

1           subcutaneous tissues. So symptoms; the  
2           patient would say it turns blue or it  
3           turns red or it's cooler or it's hotter  
4           than the unaffected side. Of course, the  
5           signs that the doctor would be looking for  
6           would be to observe, obviously, for color  
7           and to lay hands on the patient and  
8           actually feel for temperature differences.  
9           This is an area where there are some tests  
10          that -- although they're not required for  
11          the diagnostic criteria, they do corroborate  
12          and they do quantitate the result.

13           Q.       How about for the area of the  
14          criteria called the sudomotor? How -- what  
15          would you as a physician look for -- for  
16          the patient report as to the sudomotor  
17          being a component?

18           A.       The patient would say, for  
19          instance, the affected part or the affected  
20          region is drenched in sweat. You know, I  
21          can't keep it dry. I change my socks five  
22          times a day or it's dripping sweat when  
23          I'm trying to grab something and it's  
24          slippery. The signs, obviously, again would  
25          be a physician observing these sweat



1 changes. And then there are ways to  
2 quantitate that again. There is a very,  
3 very research-based test called the  
4 Quantitative Sudomotor Axon testing which we  
5 don't need to go into because it's strictly  
6 research.

7 Q. How about the area of motor as  
8 part of the criteria? What would a  
9 physician look for to see a patient report?

10 A. In terms of the symptoms the  
11 patient would say I'm weak. I can't move  
12 as fluidly as I did. Or in the case of  
13 dystonia they'll say it locks into this  
14 uncomfortable odd posture, and I can't get  
15 it out of that. Or they'll say it jerks  
16 around on its own and, of course, the  
17 symptoms would be formal strength testing  
18 by the doctor. The range of motion can be  
19 measured by the physician in terms of  
20 normal range or comparing it to the  
21 opposite side. Dystonia and the tremors  
22 can be documented recorded, you know. Of  
23 course, in our labs we actually film that  
24 or take a picture of that.

25 The test that can corroborate

1           that impression, of course, are -- there's  
2           a lot of formal testing of motor such as  
3           EMG or electromyography. There's strength  
4           testing with devices to measure the  
5           strength of the patient, the sustained  
6           strength so...

7           Q.       And you may have already told us  
8           and maybe I just forgot, but what is the  
9           fourth criteria under the Budapest criteria?

10          A.       The fourth criteria is that  
11          there's no better explanation for what you  
12          see, meaning signs or symptoms. This comes  
13          into your office, and you don't have a  
14          better way to explain that. There's no  
15          diagnosis that could equally well explain  
16          this or better explain it than CRPS.

17          Q.       Now, the things you have listed  
18          for the Budapest criteria, are those -- do  
19          the criteria require that those be seen all  
20          during one visit?

21          A.       The symptoms have to be  
22          portrayed and documented, and the signs  
23          have to be seen and documented at the  
24          visit.

25          Q.       Okay. What if a another

1 physician documents that they've seen  
2 something. What type of finding is that?

3 A. That doesn't have any place in  
4 the criteria. It's -- you know, it is the  
5 responsibility of a doctor that presumes to  
6 make a diagnosis and treat this disease  
7 that they assume responsibility for  
8 documenting this and recording this, however  
9 they prefer to do so. But it's clearly  
10 their responsibility. And it's no good  
11 that, you know, it was observed, you know,  
12 months ago by doctor X because obviously  
13 that can't be corroborated. That can't be  
14 -- you know, that can't be attested to by  
15 the physician -- who's now -- it's that  
16 doctor's responsibility to make the  
17 diagnosis. It's that doctor's  
18 responsibility to elicit the signs and  
19 symptoms necessary to meet the criteria --  
20 the established criteria for the diagnosis.

21 Q. As part of the Budapest criteria  
22 are there objective findings that are  
23 required?

24 A. That's really an interesting  
25 question, but objective, to me, as a

1 scientist means that it's -- it's something  
2 that's measured and reported by a machine.  
3 It's not subject to bias, meaning it's not  
4 the patient's subjective opinion or their  
5 portrayal of something. It's not even the  
6 doctor who in their opinion now -- if they  
7 see a blue extremity -- and it always  
8 should be relative to the other side or  
9 the unaffected side which, of course, it --  
10 it -- although it's a trained medical  
11 professional, and they should be able to  
12 observe and record accurately, they're, of  
13 course, biased at this point by what the  
14 patient has told them because we get the  
15 history first. So they're already biased  
16 before they even go in to do the exam.  
17 But they do measure and record hopefully  
18 reliably, hopefully truthfully into the  
19 record, and that's -- that's what's  
20 required for the criteria. There are no  
21 fully objective tests for CRPS. If we had  
22 that, we would have the -- you know, we  
23 would the gold standard, and we wouldn't  
24 need to do a syndromic diagnosis anymore.  
25 We would have it.

1           Q.       Are there any objective tests to  
2           measure temperature change in the  
3           extremities?

4           A.       Yes.

5           Q.       What is that?

6           A.       The best test for measuring  
7           temperature in the extremity in this  
8           context -- because you can't do points.  
9           You can't just say, well, it was hot here  
10          or cold here because we know that  
11          temperature across the surface of an  
12          extremity is -- is -- is very changeable.  
13          It goes from hot to cold in a patchy  
14          distribution and then there's gradations  
15          where, for instance, your hands are cooler  
16          than your shoulder.

17                    So it's no good to measure a  
18                    spot. You have to measure the whole limb  
19                    at once. And there's really only one  
20                    technology that can do that. And it is  
21                    quantitative infrared telethermography.  
22                    It's basically a camera that measures the  
23                    -- the infrared radiation or emitted heat  
24                    off of a surface, and then the computer  
25                    now will assess the average temperature of

1 one side versus the other side. And then  
2 you can say this -- you know, this side --  
3 the affected region is hotter or the  
4 affected region is colder. You can say --  
5 you know, of course now because it's fully  
6 objective and coming off a machine off a  
7 computer, you can say it's 1.23 degrees  
8 centigrate higher or lower on the affected  
9 part versus the unaffected part. But  
10 that's really the only technology that  
11 allows us to fully objectively measure  
12 temperature.

13 Q. How about edema? Is there a  
14 way to objectively measure edema in a  
15 person?

16 A. Yes. It's called volume immetry  
17 and basically it's a big bucket full of  
18 water. And the patient puts the unaffected  
19 part into that water and see how much  
20 water they displace. You can measure that  
21 in cubit centimeters. Then they take the  
22 affected part and put it in, and you can  
23 see, if it's swollen, it's going to  
24 displace more water. And then you can say  
25 uneffected versus affected there was a

1 displacement of 42 centimeters square more  
2 water by the affected part; the conclusion  
3 being it's bigger.

4 Now, to get back and say that  
5 that's actually edema or swelling is a  
6 trick because I mean it could be that the  
7 unaffected side is smaller or atrophy. So  
8 this has to be in the context of a  
9 clinical examination where you can say  
10 that, you know, the left side is the  
11 unaffected side and it is normal in  
12 strength, power, range, all that. And then  
13 you go to the affected side and say it  
14 looks to me to be swollen to have edema.  
15 You know, I will push on it and it's kind  
16 of boggy. There seems to be fluids there.  
17 And by volume immetry, this technology I'm  
18 talking about, there's a 40-centimeters  
19 cubed more displacement. That is the most  
20 objective technology that can be used to  
21 measure this. So this is -- this is a  
22 bedside test. However, you know, it's not  
23 -- it's not required for the criteria. It  
24 is the best way of doing this.

25 Q. Can you also measure edema using

1 a tape measure?

2 A. Yes. Now -- again, you know,  
3 that has to be very carefully documented;  
4 what -- you know, where you put the tape  
5 measure on. Obviously you have to again  
6 compare the unaffected to the affected  
7 side. And, you know, you have to say it's  
8 37 centimeters down from the -- the elbow  
9 crease on the unaffected side, the same  
10 place on the -- and then you can compare  
11 side to side. Of course, that only gets  
12 align as opposed to a whole affected part.  
13 So it's not nearly as good as the volume  
14 immetry. And this is something that -- it  
15 has to be one or the other really that is  
16 required to make the bedside diagnosis of  
17 CRPS.

18 Q. Let me ask you a question. As  
19 a physician how important is documentation  
20 to you?

21 A. It's critical, absolutely  
22 critical, not only in terms of being able  
23 to convince my colleagues that I did it  
24 right and that this is what I saw but to  
25 communicate to myself because I'll come in



1 to see the patient the next time -- I mean  
2 I see hundreds of patients. I won't  
3 remember exactly all the features. So I  
4 can look for change, either change bad or  
5 change good; in other words, they're  
6 getting better, they're getting worse. I  
7 need to change my approach. I need to  
8 change my therapy. Without documentation  
9 all is lost unless you have a picture  
10 perfect memory which I do not which most  
11 doctors do not. That would be a rare  
12 commodity.

13 Q. Doctor, have you reviewed certain  
14 materials in the case we're here today  
15 about?

16 A. Yes, I have.

17 Q. Have you reviewed the bylaws of  
18 Baptist Medical Center?

19 A. Yes, I did, a piece of those  
20 bylaws, the portion about credentially.

21 Q. Have you reviewed a copy of the  
22 complaint in this matter?

23 A. Yes, I have.

24 Q. Have you reviewed any deposition  
25 transcripts in this case?

1 A. Yes, I have.

2 Q. Have you reviewed the deposition  
3 transcripts of Dr. Robert Hogan?

4 A. Yes, ma'am.

5 Q. Have you reviewed the deposition  
6 transcripts of Dr. Dennis George?

7 A. Yes, ma'am.

8 Q. Have you reviewed the deposition  
9 transcript of Dr. George Mada (phonetic)?

10 A. Yes, ma'am.

11 Q. Have you reviewed the deposition  
12 transcript of Dr. Anthony Kirkpatrick?

13 A. Yes, ma'am.

14 Q. Have you also reviewed medical  
15 records?

16 A. Yes, ma'am.

17 Q. Have you reviewed medical records  
18 from Dr. George Mada?

19 A. Yes, ma'am.

20 Q. From Dr. Ullah?

21 A. Yes, ma'am.

22 Q. Dr. Dana Twiggs?

23 A. Yes.

24 Q. Dr. Bruce Steinberg?

25 A. Yes, I have.

1 Q. Dr. Anthony Kirkpatrick?

2 A. Yes, ma'am.

3 Q. Dr. Gregory Hartly?

4 A. I'm sorry. Who?

5 Q. Gregory Hartly. I think he  
6 might be a dentist actually.

7 A. Oh, yes, yes, those huge dental  
8 records. Yes, I did review those.

9 Q. Have you reviewed records from  
10 the Institute of Pain Management and First  
11 Coast Pain Management which would be Dr.  
12 George?

13 A. Yes, ma'am.

14 Q. Have you reviewed records from  
15 Health South?

16 A. Yes, ma'am.

17 Q. Have you reviewed records from  
18 Dr. Drunani (phonetic)?

19 A. Junani (phonetic).

20 Q. Drunani.

21 A. Oh, yes, I have with a D. Thank  
22 you.

23 Q. Sorry. Have you reviewed  
24 medical records from Baptist Medical  
25 Center-Nassau?

1           A.       Yes, ma'am.

2           Q.       Have you reviewed medical records  
3 -- excuse me. Have you reviewed a report  
4 from Dr. Halprin related to a medical  
5 examination done on Dr. Hogan?

6           A.       Yes, ma'am.

7           Q.       Okay. Having reviewed those  
8 documents, do you have an opinion as it  
9 relates to this case?

10          A.       Yes, ma'am.

11          Q.       Okay. And can you tell us what  
12 your opinion is to a reasonable degree of  
13 medical certainty as it relates to this  
14 case?

15          A.       Specifically in regards to the  
16 diagnosis of complex regional pain syndrome,  
17 I cannot state to a reasonable degree of  
18 medical certainty that on the basis of this  
19 record I would make that diagnosis.

20          Q.       Okay. And why can you -- why  
21 is there insufficient information in the  
22 medical records and documentation you  
23 reviewed to make that diagnosis?

24          A.       At no place in this record in  
25 no single occasion by no doctor is there

1 sufficient signs and symptoms to make this  
2 -- the more specific Budapest criteria in  
3 this case.

4 Q. And which criteria did you use  
5 for your opinion?

6 A. Well, I always use the more  
7 specific criteria which is the Budapest  
8 clinical criteria which is obviously the  
9 relevant criteria in this case.

10 Q. And is that the criteria you  
11 normally use in your practice?

12 A. Yes. That's the criteria I  
13 always use in my practice.

14 Q. You have of course reviewed the  
15 medical records of Dr. Anthony Kirkpatrick,  
16 I believe?

17 A. Yes, ma'am.

18 Q. Could you bring those out for  
19 us? And what I would like you to do is  
20 explain to us why his records are  
21 insufficient for a diagnosis of RSD.

22 A. Okay. Well, in general I can  
23 say I went through all his records such as  
24 they are. And in no case did I find, as  
25 I say, three of the four symptoms and two

1 of the four signs in the same record at  
2 the same time. In fact, in aggregate if I  
3 put all his records together, I still don't  
4 have enough signs and symptoms to meet the  
5 Budapest clinical criteria, okay.

6 Q. Okay.

7 A. So do you want me to elaborate  
8 on that?

9 Q. Yes, if you would.

10 A. Okay. So, for instance...

11 Q. There you go.

12 A. Thank you. I have a -- I have  
13 a note here dated 1/18/2006. And Criterion  
14 1 requires pain. And he does report pain.  
15 Criterion 2 requires -- remember symptoms.  
16 In this -- in this note I have -- he  
17 mentions dysesthesia but says that he does  
18 not have allodynia.

19 Q. Why is that important?

20 A. Well, it's critical. Remember  
21 that allodynia is one of those signs and  
22 symptoms that the statistics led us to as  
23 being necessary for the criteria. So  
24 dysesthesia is not mentioned as a criterion  
25 for CRPS. Allodynia is. And he actually

1           says in this that he does not have  
2           allodynia. So obviously that's -- that  
3           doesn't count.

4                     He mentions modeling to bluish  
5           discoloration as a symptom. Modeling is --  
6           is not real clear. That means sort of a  
7           -- a particulated pattern, sort of a  
8           geographic, you know, in some places blue,  
9           some places white. So that's sometimes --  
10          certainly saying it's CRPS, but it's not  
11          part of the criteria but more -- more to  
12          the point more bluish. So he does record  
13          that symptom. So that's one of three that  
14          we need. And then he goes on to say that  
15          there was some swelling. So that was two  
16          of four where we need three of four. So  
17          it doesn't meet the symptoms in this  
18          letter.

19                    Q.       Did he have any recording in his  
20          documentation of meeting any signs that he  
21          personally saw?

22                    A.       Yes. And he calls this in this  
23          note objective, but that means physical  
24          examination. And he says that it's  
25          swollen. So it has edema. He says that it

1 has a bluish discoloration. And that's it.  
2 And then he goes on to talk about skin  
3 appears to be bluish. He says that again.  
4 And then he talks about the patient's gait  
5 which is irrelevant to his upper  
6 extremities. So in this case he -- let me  
7 see. We've got two; edema, blue. He's  
8 got -- he's got two signs that he  
9 documents.

10 Q. Okay.

11 A. Two of four so he meets the  
12 signs but not the symptoms on this note.

13 Q. Is there -- as a clinician who  
14 diagnoses RSD is there anything you look at  
15 related to asymmetry as it relates to this  
16 condition?

17 A. This is a critical issue and a  
18 critical lack, I think, in terms of either  
19 Dr. Kirkpatrick's exam or his documentation  
20 is he doesn't say. I mean he says it's  
21 -- the skin appears to be bluish. What  
22 skin? I mean is he talking about the  
23 affected part? Is he talking about the  
24 affected part relative to the unaffected  
25 part so is it in general? I mean he



1           didn't even really mention that that is  
2           specifically in regard to -- let's see.  
3           Yes, he does mention upper extremity and  
4           lower extremity, but he doesn't -- he  
5           doesn't bother to tell us that this --  
6           that the affected part is blue relative to  
7           the unaffected side. And that's critical  
8           because all of these tests require that it  
9           be a distinct difference from the  
10          unaffected side.

11                        So to not mention that this is  
12          asymmetrical and this is different from  
13          side to side is pretty much a critical  
14          lack. In terms of a hard cut in terms of  
15          looking for signs and symptoms you would  
16          have to essentially discard everything that  
17          he says because he does not mention the  
18          symmetry or the asymmetry specifically in  
19          regards to the criteria.

20                        Q.        In your review of these records  
21          was there any physician that actually  
22          looked at all of the diagnostic criteria  
23          under Budapest and looked at the  
24          symmetrical considerations that you just  
25          told us about?

1           A.       Some -- some doctors do refer to  
2           the symmetry or asymmetry but, again -- as  
3           I say, I went through these records in  
4           great detail and no doctor gives enough  
5           signs or enough symptoms to make the  
6           Budapest clinical criteria. Also, several  
7           doctors don't mention what body part  
8           they're referring to when they say it's  
9           blue or cold or swollen which again -- you  
10          know, in terms of at least being a  
11          research scientist I would have to say, you  
12          know, I don't know what they're talking  
13          about. But that throws it pretty much out  
14          anyhow. So, A, they don't have signs and  
15          symptoms. B, they don't mention that  
16          critical piece which is symmetry or  
17          asymmetry in regards to affected part  
18          versus unaffected part.

19          Q.       Does the fact that a physician  
20          may have some of the signs or symptoms in  
21          their notes -- is that sufficient for the  
22          diagnosis of CRPS or RSD?

23          A.       No. Well that's why there is a  
24          criteria, and the answer is of course no.  
25          We wouldn't have bothered to go through all

1           that trouble if it was okay for people to  
2           say, well, you know, I think this is it.  
3           And I've got a couple things here. It has  
4           to be more rigorous with that or we go  
5           back to the chaos of the 60's and 70's in  
6           terms of making this diagnosis.

7           Q.       And can you make the diagnosis  
8           just by looking at all the records together  
9           over a continuum of time using the Budapest  
10          criteria?

11          A.       But that's not acceptable in  
12          terms of Budapest criteria. It has to be  
13          all done at the time of the exam and  
14          documented at the time of the exam.  
15          However, I can tell you that an aggregate  
16          of all the records would not allow me to  
17          make on the basis of any doctor that saw  
18          him, a diagnosis -- a clinical diagnosis of  
19          CRPS by the Budapest clinical criteria.

20          Q.       Have you reviewed the medical  
21          examination as done by Dr. Halprin?

22          A.       Yes, I have.

23          Q.       Did Dr. Halprin -- tell me your  
24          opinion as to -- as to your review of Dr.  
25          Halprin's report as to his documentation of

1                   criterias or list of CRPS?

2                   MR. SPRAGUE: I'm going to be  
3                   -- I'm going to object as being not the  
4                   proper subject of expert testimony.

5                   THE WITNESS: I'm sorry. I  
6                   didn't hear.

7                   MR. SPRAGUE: I just made an  
8                   objection for the record.

9                   BY MS. ARNOLD-SIMMONS:

10                  Q.       Here. Let me rephrase the  
11                  question. Is there any physician whose  
12                  report that you reviewed who followed the  
13                  Budapest criteria and listed them all in  
14                  his report?

15                  A.       Yes. Remember I said that  
16                  there's nobody that gave sufficient signs  
17                  and symptoms to make the diagnosis of CRPS.  
18                  The only doctor that actually reviewed  
19                  sufficient data to say he asked and  
20                  answered the question of signs and asked  
21                  and answered the question of symptoms  
22                  thoroughly was Dr. Halprin. But Dr.  
23                  Halprin, if you look at what he did, he  
24                  has -- he answers no to enough questions.  
25                  He answers not based on criteria to enough

1 things that he concludes, and I conclude  
2 that he does not support a diagnosis of  
3 CRPS. In other words, he does sufficient  
4 signs and he does sufficient symptoms for  
5 me to say Dr. Hogan does not have CRPS.  
6 But he's the only doctor that's bothered to  
7 record and document all of those signs and  
8 symptoms.

9 MR. SPRAGUE: I'm going to  
10 object again as not the proper basis of  
11 expert testimony and move to strike that  
12 entire last answer.

13 BY MS. ARNOLD-SIMMONS:

14 Q. Okay. Dr. Harden, in reviewing  
15 -- or in your knowledge and expertise as a  
16 pain management physician would using the  
17 IASP criteria be appropriate for a  
18 diagnosis of CRPS?

19 A. Well, we should use -- use the  
20 newer better criteria. We should use the  
21 consensus base statistically derived  
22 criteria, but clearly some doctors go back  
23 to the older looser criteria and use that.  
24 I don't use it but certainly people --  
25 usually people outside of the pain field

1 would use the IASP criteria.

2 Q. Okay. Have you reviewed the  
3 medical records of Dr. Steinberg in this  
4 case?

5 A. Yes, I have.

6 Q. Did -- in reviewing those  
7 medical records did you see whether the  
8 documentation was sufficient for diagnosis  
9 of RSD/CRPS?

10 A. No. He did not meet the  
11 criterion for signs or the criterion for  
12 symptoms. Actually, he didn't even come  
13 close.

14 Q. Did anyone in your review of  
15 these materials work Dr. Hogan up for a  
16 neuropathy?

17 A. Some people sort of did a  
18 glancing blow of that possibility, but  
19 nobody actually got down and said, oh,  
20 well, I think this may be an unusual  
21 neuropathy, not CRPS. We don't have support  
22 for that. But it may be an unusual  
23 neuropathy. Certainly we can explain the  
24 signs and symptoms such as they were such  
25 as were reported on the basis of

1 neuropathy. Therefore, it's very logical  
2 that we have to entertain that possibility.  
3 We have to rule that out. Because it's a  
4 different diagnosis. It's a different  
5 disease and it's treated differently, et  
6 cetera.

7 Q. In people who actually have RSD  
8 or CRPS are those people counseled at all  
9 as to the use of the affected part?

10 A. Yes. The treatment guidelines  
11 are very clear about that. We want them  
12 to use the affected part as much as  
13 possible. In fact, so much so that we  
14 counsel our patients if you cannot get to  
15 essentially normal use of the part, we  
16 cannot help you. You will never feel  
17 better. So it's a critical issue that  
18 they use the affected part as much as they  
19 can; either passive range of motion or  
20 lifting or carrying or actually getting  
21 back to, you know, complete vocational use  
22 of the affected part. It's critical.

23 Q. Would you encourage a patient  
24 diagnosed with RSD or CRPS of the upper  
25 extremity to use a sling on that extremity?

1           A.       Absolutely not. That would be  
2           dead wrong in terms of -- of their  
3           prognosis. You discourage the use of slings  
4           and immobilizing, casts or splints. So  
5           that's -- that's -- that's -- remember all  
6           the way back to the old IASP criteria.  
7           They said immobilization was a cause of  
8           CRPS. Thus, immobilization is a bad thing.  
9           You know, clearly you want to keep them  
10          moving and active with the affected part.

11          Q.       We've talked about the criteria  
12          as it relates to signs and symptoms but  
13          the fourth criteria for the Budapest  
14          criteria is what again?

15          A.       No better explanation, no other  
16          diagnosis that explains the signs, symptoms  
17          as you do see.

18          Q.       And when you're looking at  
19          documentation, what does that criteria  
20          comport with?

21          A.       Well, it needs -- basically it  
22          means that even if you do have signs and  
23          symptoms to make a Budapest research  
24          criteria, the tightest cut, if you conclude  
25          that those signs or symptoms are created on



1 the basis of a different diagnosis, it's  
2 your responsibility to -- to entertain  
3 that, to prove that, and then to -- to  
4 make that diagnosis and treat it.

5 So, you know, even -- even if  
6 you do have all these Criterion 1, 2, 3  
7 but then fail 4 because you have a better  
8 explanation -- you know, something like a  
9 diabetic neuropathy as an example of what  
10 we use as another disease to test the  
11 specificity of the diagnostic criteria. So  
12 if you -- if you make Criteria 1, 2, 3  
13 but fail criteria 4 you don't have it. So  
14 it is necessary. Again, it's the  
15 responsibility of any doctor that holds  
16 himself out as a specialist or a pain  
17 management doctor who's going to diagnosis  
18 and treat CRPS to at least consider other  
19 possibilities.

20 They can't conclude that because,  
21 you know, the doctor before me said it was  
22 RSD, therefore, I'm just going to look for  
23 signs and symptoms and go along. It's  
24 their responsibility to create a list of  
25 other possibilities. We call that a

1 differential diagnosis.

2 Q. Okay.

3 A. That's a very serious error in  
4 medical systems to not do your own history  
5 and physical; in other words, to just  
6 simply go by what other people have said.  
7 And it's also a very serious error to not  
8 keep an open mind which is what happens if  
9 you just, you know, say, well, you know,  
10 the last three doctors have said X.  
11 Therefore, I conclude it's X without  
12 entertaining the possibility of Y and Z.

13 So you have to generate a  
14 differential diagnosis. You have to test  
15 the differential diagnosis, meaning, you  
16 know, if you think it may be CRPS, you got  
17 to entertain the possibility that it could  
18 be an unusual neuropathy or rheumatologic  
19 disease or psychiatric disease or  
20 sociologic disease, if you will; things  
21 like, you know, secondary gain and -- and  
22 -- and begin to at least go through some  
23 sort of process to rule those out and rule  
24 CRPS in. And that -- that should actually  
25 go on at every single visit but over time

1 -- I mean the first visit you should have  
2 an extensive differential diagnosis. This  
3 is just good practice. Everybody is taught  
4 this in medical school.

5 Q. And so we understand, what  
6 exactly do you mean because some of us  
7 might not -- by differential diagnosis what  
8 is that?

9 A. Just means all the possible  
10 explanations of what you see; it's your  
11 responsibility to, as a doctor -- to say,  
12 well, it probably is this but I can't  
13 absolutely say it's not this, this, or  
14 this. And so you list them all. And you  
15 say, you know, No. 1 is what I really  
16 think it is. So I'm going to run some  
17 tests to try to corroborate that diagnosis  
18 but, No, 2 that's serious. I need to find  
19 out if that's the case because that's  
20 totally different. It needs to be treated  
21 differently.

22 And No. 3 and 4, you know,  
23 well, you know, that's gee whiz. But, you  
24 know, they have some features of that.  
25 And there's a test that's going to tell me

1           for sure whether they have that or not.  
2           So at the first visit you really do have  
3           to have a differential. That's good  
4           medical practice. And you need to begin the  
5           process over time of saying, well, I've  
6           ruled out No. 4. I've ruled out No. 3.  
7           I can't rule out No. 2. So I'm going to  
8           have to do another test, and I still hold  
9           No. 1 as the best explanation. So over  
10          time you finally get to the point where  
11          you can now in good faith and in good  
12          practice say, yeah, this is what the  
13          diagnosis is. And this is what we're  
14          going to spend our money and spend our  
15          time and our effort to treat.

16           Q.       And was diagnose -- diagnostic  
17          criteria No. 4, the Budapest criteria, was  
18          it addressed in any of the treating  
19          physicians of Dr. Hogan?

20           A.       Some of the doctors mentioned  
21          other diagnoses as a possibility, but  
22          they're very few. And actually the only  
23          other diagnosis that I recall from my  
24          review is -- I think Dr. George mentioned  
25          depression and anxiety as another diagnosis.

1 I don't know whether he was mentioning that  
2 as differential point, meaning this is an  
3 explanation for all the signs and symptoms  
4 I see, not CRPS, or whether he was saying  
5 it was comorbidity of the CRPS. He's not  
6 clear about that. He just throws it out  
7 and gives the diagnostic code for that  
8 so...

9 Q. Based on your review of the  
10 treating physicians' records in this case  
11 do you have an opinion on a differential  
12 diagnoses that would apply in this case?

13 A. Yes.

14 Q. Okay. And what is your opinion?

15 A. The opinion from the record --  
16 of course, I've never seen the gentleman.  
17 But if I were the doctor and I were  
18 getting the history and I were getting the  
19 tests and I were getting the physical exam  
20 that I saw in this gentleman's case, I  
21 would have to immediately conclude  
22 neuropathy.

23 Q. And why is that?

24 A. Because a lot of the signs and  
25 symptoms that he does show -- since, of

1 course, I've concluded that it's  
2 inconsistent with a diagnosis of CRPS --  
3 would be consistent with neuropathy. A  
4 specific type of neuropathy or a general  
5 type of neuropathy is unclear. But then  
6 you have to do the test and you have to  
7 entertain that and go forward through the  
8 process of ruling in, ruling out which in  
9 the case of most neuropathies requires that  
10 you do EMG testing or nerve conduction  
11 testing, you know, electrodiagnostic test  
12 that's pretty quick and pretty easy to rule  
13 in and rule out that possibility.

14 Q. What else would be on your  
15 differential diagnosis?

16 A. Well, throughout the record  
17 there's a lot of talk about his mood and  
18 certainly doctors and doctors specifically  
19 that should make that diagnosis have  
20 diagnosed him as being depressed. There's  
21 also diagnoses made of anxiety. I would  
22 base on the history that I read and the  
23 symptoms that I read make a diagnosis of  
24 certainly anxiety, depression probably. But  
25 that's another criteria. See, that's the

1 diagnostic and statistic manual revision 4  
2 of the psychiatrist. They have a very  
3 formal criteria for making these diagnoses.

4 And based on their criteria I  
5 see signs and symptoms throughout the  
6 record that you could make a diagnosis of  
7 depression and anxiety. And at least two  
8 of the doctors in this record also  
9 corroborate that and make a diagnosis of  
10 depression. And then Dr. George also makes  
11 a diagnosis of anxiety. So that's a  
12 differential possibility. Do patients with  
13 depression and anxiety have, for instance,  
14 changes in temperature? Well, anxiety  
15 absolutely cause vasoconstriction especially  
16 in the context of panic attack. And it  
17 causes intense vasoconstriction and bluish  
18 discoloration, cool extremities, modeling of  
19 the skin, all of these things we're seeing.

20 One of the principal symptoms  
21 mentioned by depressed patients -- and this  
22 is 60 plus percent of depressed patients  
23 will say that they have pain of some sort.  
24 So pain itself is something that is very  
25 frequently endorsed by depressed patients

1 and anxious patients as well. So although  
2 it seems odd, can you have physical  
3 manifestations of depression and the answer  
4 is yes. Can you have physical  
5 manifestations of anxiety? And the answer  
6 is absolutely yes especially in terms of  
7 some of the things that you see because  
8 anxiety activates the sympathetic nervous  
9 system. So you can see how that would be  
10 an overlap and must be entertained as a  
11 differential diagnostic possibility.

12 Q. Let me --

13 A. Especially -- I'm sorry.  
14 Especially in a case where a gentleman is  
15 portraying historically symptoms and signs  
16 of anxiety and is being treated now for  
17 depression and anxiety.

18 Q. Can life stressers be a cause of  
19 depression and anxiety?

20 A. Yes.

21 Q. In somebody like Dr. Hogan would  
22 failing your surgical board --

23 MR. SPRAGUE: Objection --

24 BY MS. ARNOLD-SIMMONS:

25 Q. -- be --



1 MR. SPRAGUE: -- more  
2 prejudicial than probative.

3 MS. ARNOLD-SIMMONS: Can I  
4 finish the question first, please?

5 MR. SPRAGUE: Okay. Go ahead.

6 BY MS. ARNOLD-SIMMONS:

7 Q. I think you had told me that  
8 life stressers can be a cause of depression  
9 and anxiety?

10 A. Yes, ma'am.

11 Q. Okay. Can a surgeon failing  
12 their boards on four different occasions be  
13 a stresser that could be a cause of  
14 depression and/or anxiety?

15 MR. SPRAGUE: Objection, more  
16 prejudicial than probative.

17 MS. ARNOLD-SIMMONS: You can  
18 answer.

19 THE WITNESS: Yes, ma'am.

20 BY MS. ARNOLD-SIMMONS:

21 Q. Okay. Can being substantially  
22 in debt either through defaulted student  
23 loans or tax liens be the type of stresser  
24 that can cause depression or anxiety?

25 MR. SPRAGUE: Objection, more

1 prejudicial than probative.

2 THE WITNESS: Yes, ma'am.

3 BY MS. ARNOLD-SIMMONS:

4 Q. How about being involved in  
5 litigation such as there's a judgment  
6 against someone -- is that type of stresser  
7 that can cause depression and anxiety?

8 MR. SPRAGUE: Objection, more  
9 prejudicial than probative.

10 THE WITNESS: Yes, ma'am.

11 MR. INCLAN: Excuse me. This is  
12 Clemente. I cannot hear Aaron's objection.  
13 I just need to know what they are for the  
14 record, please.

15 MS. ARNOLD-SIMMONS: The  
16 objections are more prejudicial than  
17 probative.

18 MR. INCLAN: Okay. Thank you.

19 BY MS. ARNOLD-SIMMONS:

20 Q. Okay. Can the possibility of  
21 losing one's job and one's privileges to  
22 practice be the type of stresser that can  
23 cause depression and anxiety?

24 MR. SPRAGUE: Objection, more  
25 prejudicial than probative.

1 THE WITNESS: Yes, ma'am.

2 BY MS. ARNOLD-SIMMONS:

3 Q. Those things I just listed for  
4 you; failing the boards, the tax liens,  
5 lawsuits, the possibility of losing medical  
6 privileges, and the defaults on student  
7 loans, have you -- do you have any  
8 knowledge of that as it relates to this  
9 case?

10 MR. SPRAGUE: Objection, more  
11 prejudicial than probative.

12 THE WITNESS: I have -- I do  
13 have some information provided in the  
14 record about certain of those things, yes,  
15 ma'am.

16 BY MS. ARNOLD-SIMMONS:

17 Q. I think I interrupted you to ask  
18 you more questions about depression as a  
19 cause of pain -- as part of the  
20 differential diagnosis. Was there anything  
21 else you needed to tell us about depression  
22 and anxiety as part of the differential  
23 diagnosis?

24 A. No. Well, except for the fact  
25 that -- that -- we -- we -- anxiety and

1 depression are actually seen in CRPS. But  
2 the anxiety thing I think I underlined  
3 this, but it's critical that you entertain  
4 that as a possibility because it -- it  
5 plugs right into the same type of neuroaxis  
6 that is involved in CRPS. So, you know,  
7 it really would be inappropriate, bad  
8 medicine, I guess, to not -- to not  
9 entertain that as a differential possibility  
10 at least early on and then rule it in,  
11 rule it out, call it a comorbidity, what  
12 you need to do, but to treat it.  
13 Obviously, that's what we're all about here  
14 is to try to determine treatment targets.

15 So if his disease, if his  
16 problem, if his principle diagnosis is  
17 anxiety, that would be a huge error to not  
18 get that and to go off and do things and  
19 -- and treatments appropriate to CRPS  
20 because those would be dangerous and  
21 inappropriate in somebody who has anxiety  
22 disorder.

23 Q. Can someone be depressed and not  
24 be getting treatment for it?

25 A. Yes, ma'am. Yes, ma'am, of

1 course.

2 Q. All right. You told us so far  
3 that on your differential diagnosis in this  
4 case would be neuropathy and psychological  
5 causes such as depression and anxiety?

6 A. Well, that's not all the  
7 psychological causes but those are the two  
8 preeminent. I mean those are -- those are  
9 what the -- the psychiatrists call Axis 2.

10 Q. Okay. What other things would  
11 go on your differential diagnosis as it  
12 relates to Dr. Hogan?

13 A. Well, a big one in this field  
14 you need to entertain the possibility that  
15 there is secondary gain.

16 Q. Why is that a big one in this  
17 field? Explain that to us.

18 A. Because the -- as you say  
19 stresses and strain and expectation --  
20 basically we are animals like the pigeons  
21 in the skitter box. And if you give the  
22 pigeons or us positive reenforcement, it  
23 tends to propetuate behaviors. If you give  
24 us negative reenforcement, it extinguishes  
25 the behavior. You know, if you give the

1 pigeons food pellets and they keep doing  
2 that. You give them electrical shock and  
3 they stop doing those behaviors.

4 But, you know, I don't want to  
5 equate us to pigeons but we're not much  
6 different because things that reenforce  
7 behaviors like love, affection, money, the  
8 hope of money, getting out of work, getting  
9 out of military service; these sort of  
10 things can be positively reenforcing. And  
11 then, of course, negative reenforcing --  
12 you have to go back to work. You have to  
13 work in spite of some discomfort. You  
14 know, you still have to pass the test or  
15 go forward with, you know, life and pay  
16 taxes. You know, all of those things,  
17 they're very negative.

18 You know, if one were to say,  
19 you know -- you know, I started to have  
20 this pain that something's going wrong and  
21 I don't -- but they get a lot of positive  
22 reenforcement to continue that behavior and  
23 negative reenforcement to get better, it  
24 tends to -- to perpetuate those behaviors.  
25 So secondary gain is one sociologic piece

1           that can perpetuate illness, behavior -- is  
2           very serious in this business because  
3           that's actually a treatment target. We  
4           have to break all that up. So secondary  
5           gain or, as the psychiatrists say in the  
6           DSM-IV, malingering, are important  
7           considerations. And they -- again, they  
8           have a criteria. They have a diagnostic  
9           approach that can be methodically used to  
10          rule in, rule out these possibilities.

11          Q.       As a pain management physician  
12          in the diagnosis and treatment of CRPS, do  
13          you have to assess whether a patient may  
14          have issues related to secondary gain in  
15          your diagnosis and treatment?

16          A.       Yes. It's absolutely essential  
17          that we diagnosis people on biologic  
18          grounds but also psychologic grounds and  
19          sociologic grounds. We need to understand  
20          if there's lawsuits, if there's  
21          compensation, if there's disability, also if  
22          there's spouses that are saying, you know,  
23          honey, stay in bed. Don't go to work.  
24          I'll take out the garbage. You're sick.  
25          And that, as you can see, is positive

1           reenforcement for illness behavior. You  
2           know, you have to tell spouses don't do  
3           that just like you have to say settle the  
4           lawsuit. You know, let's get on with  
5           life.

6           Q.        What did you see in this case  
7           that would cause you to put secondary gain  
8           under your differential diagnosis?

9           MR. SPRAGUE:  Objection, lack of  
10          predicate, lack of qualification; more  
11          prejudicial than probative.

12          THE WITNESS:  Well, it's the  
13          context of our interaction here. This is  
14          a lawsuit. So in context this is all  
15          about compensation and money. As it comes  
16          out there's a -- there's a lot of negative  
17          reenforcers for getting well and going back  
18          to work in Dr. Hogan's case. There are  
19          positive reenforcers for staying sick; not  
20          just money but, unfortunately, in this case  
21          there's -- you know, there's opioid  
22          medications. There's drugs that -- that  
23          come with staying sick in this regard that  
24          some people find to be very -- very  
25          positive, very reinforcing, euphoric.



1 Euphorigenicity of the opioids of course is  
2 well known as you can see from street  
3 value and diversion and all of that.

4 BY MS. ARNOLD-SIMMONS:

5 Q. Even if you were not acting as  
6 an expert witness in this case but if you  
7 were treating Dr. Hogan, would you be  
8 forced to evaluate those secondary gain  
9 issues in your treatment and diagnosis?

10 A. Absolutely. I mean we would  
11 have -- we would take a very critical view  
12 of these and not -- not to be mean but  
13 simply to say, you know, if we want to  
14 help this guy get on with his life, we're  
15 going to have to clean up some of the  
16 sociology. We're going to have to make it  
17 seem to be positive to get well and to go  
18 back to work and to get on with life as  
19 opposed to staying sick and, you know,  
20 taking more and more drugs and stronger and  
21 stronger drugs, et cetera.

22 So you have to you have to  
23 diffuse the sociologic reenforcers, meaning  
24 you have a talk with the support group and  
25 say, you know, encourage him to get out

1                   and use the limb. Don't encourage him to  
2                   stay in bed and not -- not work. You  
3                   know, the secondary gain money, you don't  
4                   encourage people to settle quickly or  
5                   inappropriately or without advice of counsel  
6                   but, you know, clearly let's do this.  
7                   Let's get this done. Let's settle this  
8                   case. Let's get that out of your hair so  
9                   that you can get on with your life.

10                                 And, you know, psychologically  
11                   you treat the depression. You treat the  
12                   anxiety. You try to create an operant  
13                   paradigm so it is the best thing to do to  
14                   get better and get on with life. That's --  
15                   that's a critical part of any pain  
16                   management doctor's repertoire. It has to  
17                   be part or they're just not doing the job.  
18                   For them to treat only biomedical issues  
19                   they're doomed to failure anyhow. If they  
20                   don't treat the psychology, they're doomed  
21                   to failure. If they don't treat the  
22                   sociology, they're not only doomed to  
23                   failure, they're doomed to tremendous  
24                   frustration and -- and, you know, a sense  
25                   of just not moving forward here.

1           Q.       Okay. You had listed on your  
2 differential so far neuropathy,  
3 psychological causes, specifically depression  
4 and anxiety, secondary gain. Is there  
5 anything else you would include on your  
6 differential diagnosis in this particular  
7 case?

8           A.       Well, yeah. Unfortunately, Dr.  
9 Hogan has gotten on some very high doses  
10 of opioid medication, and the opioids --  
11 it's very paradoxical. Opioids, of course,  
12 are good pain-fighting drugs acutely. But  
13 as -- as you take these medicines since  
14 over time, they actually cause a resetting  
15 of the nervous system. So now people  
16 become very sensitized to their environment;  
17 in other words, things that were not  
18 painful now become painful or allodynia.

19                   They have pains -- things that  
20 are normally painful like, you know,  
21 basically sitting too long and getting  
22 ischemia of your behind now become  
23 intensely painful to a patient because they  
24 lose that natural ability to modulate pain  
25 or to screen out pain because the opioids

1 -- the drugs have now come in and the --  
2 and the natural -- the dogenous opioid  
3 system has gone to sleep. So now anything  
4 that hurts is going to hurt ten times  
5 worse. And actually documented in the  
6 literature -- not only in rats, cats, and  
7 dogs but now in people that opioid  
8 medication cause allodynia, caused  
9 hyperpathia.

10 Now, opioid medications also  
11 cause depressed mood. Opioid medications  
12 also cause clumsiness and, you know,  
13 inability to do fine motor tasks. So --  
14 So, you know, in this case I see that --  
15 I see him coming in on high dose opioids.  
16 I would have to suggest that maybe the  
17 drugs, the opioids, are at least part of  
18 the problem, if not the problem in this  
19 case now. So that's -- that's a big issue  
20 and it has to be. Again, any good pain  
21 management doctor has to entertain that  
22 possibility, has to mention it as a -- as  
23 at least a possibility that must be  
24 considered and ruled in, ruled out, if that  
25 can be accomplished.

1 Q. Have you reviewed the medications  
2 which Dr. Hogan is on as a pain management  
3 specialist and as a specialist with an  
4 expertise in pharmacotherapy?

5 A. Yes, ma'am.

6 Q. Okay. In reviewing his  
7 medications is his medication regimen  
8 appropriate in this particular case?

9 A. Well, I don't think so. I mean  
10 I think he's on too much opioids obviously  
11 for the reasons we just talked about. In  
12 our clinics we use opioids very sparingly  
13 in these cases and use them short term.  
14 We like to use the opioids so we can get  
15 moving, but once moving and feeling better  
16 we try to withdraw those drugs and get rid  
17 of those instead of, unfortunately, in this  
18 case where it's more and more and more and  
19 stronger and stronger and stronger seemingly  
20 without end. So he's on too much opioid.

21 Q. Do you have an opinion to a  
22 reasonable degree of medical certainty  
23 whether the amount of opioids he is on  
24 could be causing his hyperalgesia and pain?

25 A. I have -- that -- you phrase

1           that interestingly. I am not sure but I  
2           do believe it could be a cause. It should  
3           be included in the differential and it must  
4           be addressed. I am sure of that. I am  
5           100 percent sure of that. I'm not sure  
6           that that's what's causing his signs and  
7           symptoms, and the only way to know that is  
8           as his treating doctor -- stop the opioids  
9           and see what happens.

10          Q.       Let me ask you, have you also  
11          written papers related to the effects of  
12          nicotine and caffeine on CRPS/RSD?

13          A.       I have written one paper, but I  
14          have done research in that area, yes,  
15          ma'am.

16          Q.       And can you tell us is caffeine  
17          a factor -- caffeine use a factor in  
18          someone with CRPS and RSD?

19          A.       Caffeine is a secondary factor.  
20          We did show a trend with caffeine to  
21          worsen CRPS because it works on the  
22          sympathetic axis and causes  
23          vasoconstriction.

24          Q.       How about smoking?

25          A.       Now, that was strongly positive

1 as a negative factor in CRPS. Nicotine is  
2 a powerful vasoconstrictor. It also causes  
3 alteration of pain modulatory systems and  
4 rewards systems. So it -- it's really a  
5 bad idea in CRPS clearly in itself. It  
6 can mimic some of the signs and symptoms  
7 that we require for the criteria.

8 Specifically smoking can cause  
9 vasoconstriction so much so that in -- in  
10 extreme cases -- what's called Berger's  
11 disease -- smoking can cause  
12 vasoconstriction on to auto amputation.  
13 People who are addicted continue to smoke,  
14 and they get this clamping down -- this  
15 vasoconstriction in the residual limbs so  
16 that they actually start to lose fingers,  
17 hands, arms, if they can't quit smoking.  
18 So it clearly is a very negative habit to  
19 continue, if you have a diagnosis of CRPS.

20 Q. I think you already told us that  
21 the criteria that's no longer used is the  
22 IASP criteria. Is that correct?

23 A. Not by pain specialists at  
24 least.

25 Q. In reviewing this chart was

1           there any physician that made the diagnosis  
2           that you could make the diagnosis using  
3           this less specific and outdated criteria?

4           A.       Yes.  It's my recollection that  
5           Dr. George in one note on one occasion  
6           actually would have recorded enough signs  
7           and enough symptoms in the format of the  
8           old IASP criteria to make that diagnosis.

9           Q.       You, of course, treat patients  
10          with RSD/CRPS on a regular basis?

11          A.       Yes, ma'am.

12          Q.       Do you have RSD patients who  
13          work?

14          A.       Yes, ma'am.

15          Q.       Does the diagnosis of RSD  
16          necessarily mean someone cannot work?

17          A.       No, not at all.  In fact,  
18          remember when I said we encourage them to  
19          keep active, keep using the affect part.  
20          Therefore, we actually encourage work as  
21          part of the therapy in effect.

22          Q.       Does the diagnosis of RSD mean  
23          that someone is going to be on narcotics  
24          the rest of their life?

25          A.       No, ma'am.



1           Q.       Being an expert in the areas of  
2 pain management and RSD, can you tell us  
3 what are the usual treatment options when  
4 somebody is diagnosed with RSD/CRPS?

5           A.       Because it's a complicated  
6 disease that has biopsychosocial aspects it  
7 requires a team, what we call an  
8 interdisciplinary team, meaning a team  
9 working together. They work towards the  
10 same goal. They talk. They discuss the  
11 case. They discuss progress and problems  
12 that the people are having and the plan  
13 for the future. And they adapt and  
14 modify. They're very flexible in their  
15 appropriate.

16                   But it should include physicians  
17 for diagnosis critically, monitoring  
18 medications. Oftentimes, the physicians are  
19 the leaders of these teams but not  
20 necessarily. I worked on a team where a  
21 psychologist was on the leader and I was  
22 the medical director. It requires  
23 psychologists absolutely. You can't proceed  
24 without psychology in terms of evaluation  
25 for the psychosocial part but treatment --

1 cognitive behavioral treatments, biofeedback,  
2 et cetera.

3 It is essential that you have  
4 physical therapy and occupational therapy,  
5 again, to keep them moving to pursue the  
6 goal of functional restoration. It can  
7 involve groups like recreational therapy.  
8 Certainly skilled nursing educators are  
9 critically important on teams in my  
10 opinion. And then I think in a lot of  
11 cases vocational rehabilitation plays a big  
12 part and then other team players. I mean  
13 you can get into chaplaincy and social  
14 workers, et cetera, if that seems to be  
15 appropriate.

16 Q. As a physician who treats  
17 patients with RSD/CRPS and a physician who  
18 does pain management do you make  
19 recommendations whether patients should go  
20 back to work or not?

21 A. Yes, I do.

22 Q. Okay. Do you have an opinion  
23 in this case to a reasonable degree of  
24 medical certainty as to whether Dr. Hogan  
25 can return to work?

1           A.       Yes, I do.

2           Q.       And what is your opinion?

3           A.       I am sure to a reasonable degree  
4 of medical certainty and far beyond that 50  
5 percent require -- 51-percent requirement  
6 that he can work. You know, as a  
7 rehabilitationist I have amputees that have  
8 no arm. So worse case scenario as if he  
9 were an amputee and had no arm, he can  
10 work. I mean there's tens of thousands of  
11 jobs that can be done by an amputee, much  
12 less somebody with some pain and hopefully  
13 the encouragement to use the limb such as  
14 would be in this case.

15          Q.       Do you have an opinion whether  
16 he can work in the medical field?

17          A.       Again, I'm certain he could.  
18 Again, worse case scenario -- let's say  
19 that he could never use the arm again. He  
20 can still function as a -- for instance, a  
21 radiologist. Radiologists sit in a quiet  
22 room and observe the films as they come  
23 across and -- and then make a report. So  
24 he wouldn't have to have his arm to do  
25 that. And actually there's many, many

1 different fields of medicine that he  
2 doesn't need to have both arms.

3 Now, of course, if we could get  
4 him moving and using actively the limb  
5 again and could show dexterity and show  
6 strength, then there's no real reason he  
7 couldn't work as a surgeon again. But  
8 that's more a choice in regards to, you  
9 know, how much the pain impacts his ability  
10 to do fine manipulation, et cetera. So,  
11 you know, yes, absolutely, he can work in  
12 the medical field.

13 Your next question is going to  
14 be how about as a surgeon and I would --  
15 I don't know to a reasonable degree of  
16 medical certainty that he can do that. I  
17 really don't have enough information. See,  
18 if I'm asked to make that judgment, you  
19 know, the first thing I do is a functional  
20 capacity evaluation, a complete occupational  
21 therapy evaluation, a complete vocational  
22 rehabilitation evaluation. Then I deliver  
23 the treatments that are appropriate to  
24 optimize his ability to return to the  
25 original job which is, of course, always

1 the target. And only then using all that  
2 data and all that information would I  
3 assess whether or not this job was  
4 realistic. So, unfortunately, that's never  
5 been done here.

6 Q. As a pain management specialist  
7 are you familiar with a treatment option  
8 called sympathectomy?

9 A. Yes, ma'am.

10 Q. Can you tell me why you're  
11 familiar with it?

12 A. Well, it used to be done. It  
13 used to be considered a very -- this is  
14 considered a very, you know, third-tier  
15 kind of intervention. I guess to use the  
16 lay terminology it would be kind of Hail  
17 Mary, when all else fails kind of stuff.  
18 And people did it for a long time  
19 remembering that, you know, it looks like  
20 the sympathetic nervous system is  
21 dysfunctional. So one way to look at it  
22 is, well, you know, if it doesn't work,  
23 well, let's destroy it. To me it's like  
24 if it doesn't work well, how much sense  
25 does it make to cut it out, you know.

1                   So basically my opinion that it  
2                   doesn't make any sense to completely  
3                   destroy a part of the nervous system that's  
4                   not functioning well has been corroborated  
5                   by the literature and, in fact, it's not  
6                   considered standard of care and not  
7                   appropriate to do sympathectomy now in CRPS  
8                   patients. There's still doctors that do  
9                   it. There's still doctors that make a good  
10                  living doing it, and there's still  
11                  countries that it may be a part of their  
12                  algorithm. But clearly in the United  
13                  States amongst pain management doctors  
14                  nobody is really doing sympathectomies any  
15                  more.

16                 Q.           Based on your review of these  
17                 records would a sympathectomy be appropriate  
18                 in this case to a reasonable degree of  
19                 medical certainty?

20                 A.           No. And I say that because  
21                 sympathetic blocks are no longer effective  
22                 in his case, and if the sympathetic blocks  
23                 don't work, it clearly is not indicated to  
24                 do a surgical manipulation.

25                 Q.           Are you familiar with a

1 treatment modality for CRPS involving spinal  
2 cord stimulators?

3 A. Yes.

4 Q. And can you tell me what your  
5 experience is with the use of spinal cord  
6 stimulators?

7 A. Well, it's -- you know, this is  
8 -- this is even further down the line than  
9 the surgical sympathectomy in some context.  
10 I guess it's more main stream now. But it  
11 is experimental. And I say that because  
12 all know it's commonly done, there has  
13 never been the appropriate scientific  
14 research to prove that it works. This is  
15 very unfortunately where you got, you know,  
16 doctors that are putting in these devices  
17 without the support of research to say that  
18 it's effective.

19 I do a lot of pharmaceutical  
20 research, drugs. And to convince the FDA  
21 that something works, I have to have three  
22 huge randomized control trials. In terms  
23 of spinal cord stimulators there's only  
24 been one small. And everybody in the  
25 field agrees, including the doctors that

1           conducted this, that it's a highly flawed  
2           trial. So one small flawed trial that  
3           early on suggested that spinal cord  
4           stimulators work. But when they analyzed  
5           it six months and a year later, it didn't  
6           really look like it helped that much.

7                         That's essentially the sum and  
8           total of the high level science that  
9           supports this. Now, of course, there's a  
10          lot of doctors that are saying, gee whiz,  
11          I put this in Mr. Smith and he felt  
12          better. That's called an anecdote, you  
13          know, and that doesn't compel anybody in --  
14          in -- in medicine or in my field that  
15          something works. There's open label trials  
16          where they just put in the spinal cord  
17          stimulator in a whole bunch of people and  
18          ask them do you feel better and they say  
19          yeah.

20                        The reason that's flawed is  
21          because we know there's a huge placebo  
22          response. 50 percent of people you start  
23          monkeying around with their back and doing  
24          surgery, even if you do saline injections  
25          or do a sham operation are going to say I



1           feel better; in other words, a placebo  
2           response. That's why we have to do  
3           randomized control trials to prove that  
4           something works. This has not been done  
5           for spinal cord stimulators yet. There's  
6           no excuse for it, but it has not been  
7           done.

8                         So the point is would I  
9           recommend this for in any case or in this  
10          case specific, and I would have to say,  
11          no, I would not. I don't believe in the  
12          things. I really think that when the  
13          science is finally done, it's going to show  
14          they don't work. And it's a waste of time  
15          and money. But it's still in there as an  
16          experimental approach. In other words,  
17          after everything else has been tried and  
18          done and exhausted, you can find a doctor,  
19          you know, on any street corner almost  
20          that's willing to do this thing because  
21          it's very, very highly compensated. I mean  
22          it's upwards of \$100,000 to do this and it  
23          takes 15 minutes to put it in. So I mean  
24          there's certainly people that are willing  
25          to do this, not me, but certainly there's

1 doctors that will do this whether it's  
2 proven or not. And it is not proven. It  
3 is experimental. So I'm sorry. That was  
4 a long answer.

5 Q. That's okay. I think we  
6 probably need to take a break to change  
7 out the tape again.

8 A. Okay.

9 THE VIDEOGRAPHER: This is the  
10 end of tape No. 2 in the deposition of  
11 Norman Harden (sic). The time is 19:22:56  
12 and we are off the record.

13 (WHEREUPON, a discussion was had  
14 off record.)

15 THE VIDEOGRAPHER: Standby. Back  
16 on the record; the beginning of tape No. 3  
17 in the deposition of Norman Harden (sic).  
18 The time is 19:29:35. Counsel, you may  
19 proceed.

20 BY MS. ARNOLD-SIMMONS:

21 Q. Dr. Harden, let me clear up a  
22 couple things I forgot to do earlier. Are  
23 you being paid for your time and review in  
24 acting as an expert witness in this case?

25 A. Yes, ma'am.

1 Q. And what is your rate for  
2 review?

3 A. It's \$450 for record review.  
4 It's \$650 for deposition or court time.

5 Q. And have you generated a bill as  
6 it relates to this case?

7 A. Yes, I have.

8 Q. And do you have an estimate  
9 about what the total is going to be for  
10 your fees related to this case?

11 A. An estimate for the total?

12 Q. Yes, sir.

13 A. I think we computed today it  
14 would be about \$7,000.

15 Q. Have you ever testified in court  
16 as an expert witness prior to your  
17 appearance here today?

18 A. No, I have not.

19 Q. Have you reviewed cases in the  
20 past as a potential expert witness?

21 A. Yes, I have.

22 Q. Have you ever given a deposition  
23 as an expert witness prior to today?

24 A. No, I have not.

25 Q. Earlier we had -- I was asking

1           you about your differential diagnosis in  
2           this case based on everything you had  
3           reviewed. Did we go through everything on  
4           your differential diagnosis?

5           A.       Well, those to me are the  
6           prominent things on the differential.  
7           These are things that, if I were to see  
8           this case, I would really focus on. There  
9           is a half dozen other possibilities. I  
10          mean there's rheumatologic disease.  
11          There's, you know, things like carpal  
12          tunnel that should be entertained, but to  
13          me they're less likely in regards to  
14          explaining signs, symptoms, and what's going  
15          on here.

16          Q.       And, Dr. Harden, do you have an  
17          opinion to a reasonable degree of medical  
18          certainty whether Dr. Hogan's medical  
19          records support a diagnosis of RSD using  
20          the Budapest criterion in this case?

21          A.       Yes, I do.

22          Q.       Okay. And what is that opinion?

23          A.       I don't think that the records I  
24          see support that diagnosis to a reasonable  
25          degree of medical certainty.

1 MS. ARNOLD-SIMMONS: Dr. Harden,  
2 I believe that's all the questions I have  
3 at this time but other counsel may have  
4 questions for you, please.

5 THE WITNESS: Okay.

6 CROSS EXAMINATION

7 BY-MR.SPRAGUE:

8 Q. Dr. Harden, now that you're only  
9 seeing patients about 10 percent of the  
10 time -- or 10 percent of your professional  
11 time. Is that correct?

12 A. Yes, sir.

13 Q. In fact, you only see patients  
14 one day a week. Is that right?

15 A. Yes, sir.

16 Q. Now, you mentioned or -- when a  
17 patient comes in to see you with chronic  
18 pain and RSD is suspected, you would want  
19 to take a history from that person,  
20 correct?

21 A. Yes, sir.

22 Q. And you would want to do that  
23 even if you had other medical records to  
24 review, correct?

25 A. Yes, sir.

1 Q. In fact, that's critical to you  
2 in making your diagnosis. Isn't that  
3 correct?

4 A. Yes, sir.

5 Q. But you didn't do that in this  
6 case?

7 A. No, sir. I'm not a treating  
8 physician. I'm asked to review the records  
9 as an expert.

10 Q. So that's a no?

11 A. Yes, sir. I'm sorry, you're  
12 correct. No, I did not do that in this  
13 case.

14 Q. Okay. You would also want to  
15 do a physical examination on that -- on  
16 that patient, wouldn't you, Doctor?

17 A. Yes, sir. If I were to treat  
18 -- if I were to be responsible, that would  
19 be important.

20 Q. Okay. And doing a physical  
21 examination is also crucial to you in  
22 arriving at your diagnosis when you're  
23 seeing a patient. Is that correct?

24 A. Yes, sir.

25 Q. But you didn't do that in this

1 case either?

2 A. No, sir.

3 Q. Okay. And there's also certain  
4 tests that you might like to do when  
5 you're entertaining a diagnosis of RSD or  
6 trying to rule out RSD --

7 A. Yes, sir.

8 Q. -- correct?

9 And you didn't do those either?

10 A. No, sir. That's not my role  
11 here today.

12 Q. Yeah. You are -- in fact, your  
13 role is as a hired expert?

14 A. Yes, sir.

15 Q. Okay. And you were hired by  
16 the counsel for Gray Gables, one of the  
17 third-party defendants in this case,  
18 correct?

19 A. I prefer the term "independent  
20 expert." You know, certainly I -- I like  
21 to maintain my integrity and answer  
22 truthfully. So I'm not hired in the sense  
23 I'm going to say what they want me to say.  
24 I will say the truth and I will say it  
25 clearly and loudly in a case like this.

1 MR. SPRAGUE: Okay. I'm going  
2 to move that as violative of the motions  
3 in limine and nonresponsive. And I'm going  
4 to ask my question again.

5 THE WITNESS: Yes, sir.

6 BY MR. SPRAGUE:

7 Q. You're actually a hired expert.  
8 Isn't that correct?

9 MS. ARNOLD-SIMMONS: Objection,  
10 asked and answered.

11 BY MR. SPRAGUE:

12 Q. My question is --

13 A. You know, I'm really not  
14 familiar with this term. I've never been  
15 called or asked to be a hired expert. I  
16 am an expert witness. I am an expert  
17 reviewer. Yes, I am those things.

18 Q. Yes. And you were hired by one  
19 of the defendants?

20 A. Correct.

21 Q. Okay. You've never seen Dr.  
22 Hogan?

23 A. No, sir.

24 Q. Never examined him?

25 A. No, sir.



1 Q. Never spoken to him?

2 A. No, sir.

3 Q. And yet you're still comfortable  
4 making the opinions that you've rendered in  
5 this case?

6 A. Absolutely. I mean I'm asked to  
7 review these records and then form opinions  
8 about these records. That's appropriate.

9 Q. And how much is the defendant  
10 paying you?

11 A. \$450 an hour to review the  
12 records and \$650 an hour in this context.

13 Q. And that's \$7,000?

14 A. Approximately 7,000 all said and  
15 done.

16 Q. And you mentioned that you've  
17 acted as an expert in other cases. It's  
18 true, Doctor, that five out of six times  
19 you've consulted on behalf of defendants?

20 A. That's correct.

21 Q. I want to ask you some questions  
22 about complex regional pain syndrome in  
23 general. You would agree, Doctor, that  
24 complex regional pain syndrome, otherwise  
25 known as RSD, is -- or can be

1 characterized by severe and relentless pain?

2 A. The criteria says that it's pain  
3 -- excuse me -- pain that is  
4 disproportionate to what you would expect  
5 on the basis of a lesion.

6 Q. Okay. My question was, RSD can  
7 be characterized by severe and relentless  
8 pain. Is that correct, Doctor?

9 A. Certainly in some cases I've  
10 heard that, yes.

11 Q. Okay. You also would agree with  
12 me, Doctor, that minor injuries can cause  
13 complex regional pain syndrome?

14 A. Yes, sir.

15 Q. Now, as far as treatment for RSD  
16 you would agree, Doctor, that opioids may  
17 be effective as far as maintenance of pain  
18 relief in patients with RSD?

19 A. Actually, I believe I opined  
20 that we use those drugs acutely, but we  
21 rarely would entertain using those drugs as  
22 maintenance. Specifically in your question  
23 you asked me about maintenance.

24 Q. Doctor, Doctor, you edited a  
25 manual called Complex Regional Pain Syndrome

1 Treatment Guidelines?

2 A. I did.

3 Q. And there is a section in there  
4 about pharmacotherapy?

5 A. Yes.

6 Q. In that section about  
7 pharmacotherapy was written by you?

8 A. Yes, it was.

9 Q. And you, in fact, wrote the use  
10 of opioids for general chronic pain  
11 management is still subject to some  
12 controversy, but they may have value as  
13 both a rescue and maintenance treatment for  
14 CRPS?

15 A. Correct. And the operative  
16 word, the important word is "may." There  
17 is no definition that these should be used  
18 or have to be used. It's just an  
19 occasional use. It's a possibility.

20 Q. Okay. But they --

21 A. Not in my clinics, not in my  
22 clinics, but I have to entertain the  
23 possibility that other doctors may choose  
24 to use these drugs.

25 Q. But they may have value,

1 correct?

2 A. Correct.

3 Q. Okay. And also Methadone may  
4 have specific value in the treatment of  
5 CRPS. Is that correct?

6 A. Yes, sir. That's possible.

7 Q. And in -- you would agree with  
8 me that opioids may be an appropriate  
9 treatment for intractable pain due to CRPS.  
10 Is that correct?

11 A. Possible, yes.

12 Q. The subject of a spinal cord  
13 stimulator -- again, in this manual you  
14 edited a spinal cord stimulator is included  
15 in the evasive treatment algorithms that's  
16 set forth in this manual. Isn't that  
17 correct?

18 A. By Dr. Burton, yes. I did not  
19 write that section.

20 Q. Yeah. But you edited the  
21 manual?

22 A. That's correct.

23 Q. Okay. Also you're the clinical  
24 affairs director of the RSD Association of  
25 America?

1 A. That's correct.

2 Q. And dorsal column stimulators,  
3 otherwise known as spinal cord stimulators,  
4 are listed as a possible treatment for RSD  
5 on the RSD Association of America website.  
6 Isn't that correct?

7 A. That's correct.

8 Q. All right. I want to talk to  
9 you about the -- hold on. You would agree  
10 with me, Doctor, that RSD can render  
11 someone unable to work. Is that correct?

12 A. That's possible, yes.

13 Q. It also can render them unable  
14 to take care of the activities of daily  
15 living, correct?

16 A. That's a possibility, yes. I've  
17 certainly seen that.

18 Q. And the pain and other  
19 psychological factors that are associated  
20 with RSD can lead to losses in the areas  
21 of families, social acquaintances, and jobs?

22 A. Yes, sir, that can happen.

23 Q. The criteria that were set for  
24 the diagnosis of RSD in 1994 by the  
25 International Association for the Study of

1 Pain, which we've already discussed, those  
2 criteria were codified by one of committees  
3 of the IASP. Is that correct?

4 A. Yes, sir, a committee of two.

5 Q. The Budapest criteria have not  
6 been codified by the IASP. Is that  
7 correct?

8 A. That's not correct. The IASP  
9 actually published the book that mentions  
10 these criteria. The IASP put together the  
11 consensus group in Budapest and, in other  
12 words, they sponsored that work and  
13 published that work. So, you know, I  
14 would say that certainly the IASP is behind  
15 this effort and supporting this effort.

16 Q. Okay. Doctor, listen to me  
17 closely.

18 A. Okay.

19 Q. The question I asked is, have  
20 the Budapest criteria been codified by the  
21 Taxonomy Committee of the IASP?

22 MS. ARNOLD-SIMMONS: Objection to  
23 the form. That's not the question.

24 THE WITNESS: That's not what  
25 you asked me. You asked me if the IASP

1 has codified and I would say, yes,  
2 absolutely. They sponsored this work. They  
3 published this work. The Taxonomy  
4 Committee which has not met since the  
5 Budapest consensus group certainly has not  
6 commented on that one way or the other.  
7 In fact, that group such as it is has  
8 asked us to do this consensus work. That  
9 is specifically why we had the consensus  
10 group. That is specifically why we are  
11 revalidating the criteria at this moment.  
12 So the IASP Taxonomy Committee is very  
13 interested in this, and they're asking us  
14 to do a couple more steps before they do,  
15 as you say, codify.

16 BY MR. SPRAGUE:

17 Q. So the answer to my question is  
18 no?

19 A. The Taxonomy Committee has not  
20 -- has not mentioned -- I'm sorry. They  
21 have not published a new diagnostic  
22 criteria book since 1994. So, of course,  
23 they haven't -- they don't have a new  
24 criteria.

25 Q. So the answer to my question is

1 no?

2 MS. ARNOLD-SIMMONS: Objection to  
3 the form, asked and answered.

4 THE WITNESS: They have not  
5 printed another diagnostic criteria since  
6 1994. That's the answer to the question.  
7 You know, to say that they codified or  
8 approved or adopt is essentially irrelevant.  
9 They just simply have not published another  
10 criteria.

11 BY MR. SPRAGUE:

12 Q. Okay. Do you remember when your  
13 deposition was taken earlier today, Doctor?

14 A. Yes.

15 Q. And you were sworn to tell the  
16 truth?

17 A. Yes.

18 Q. And I asked that exact same  
19 question?

20 A. And you asked the exact same  
21 question two different ways at that time as  
22 well.

23 Q. And you answered no?

24 A. I answered that the IASP has  
25 sponsored and published the work of the



1 Budapest criteria and that the Taxonomy  
2 Committee has not -- you know, they haven't  
3 written another criteria. They haven't had  
4 an opportunity to codify or not codify  
5 since 1994. So we live with the 1994 IASP  
6 criteria. That doesn't mean that they  
7 support or do not support. In fact, the  
8 head of the IASP Taxonomy Committee  
9 specifically has told us if we accomplish  
10 this goal, which we did, and another goal,  
11 which we're doing, that he will publish  
12 this new criteria in the next book that  
13 comes out.

14 Q. Okay. Doctor, again, you're the  
15 clinical affairs director for the RSDAA?

16 A. Yes, sir, Reflex Sympathetic  
17 Dystrophy Association of America, yes.

18 Q. On that organization's website  
19 under the heading of what is RSD the 1994  
20 criteria are set forth. Is that correct?

21 A. That's correct.

22 Q. The Budapest criteria are not  
23 set forth, correct?

24 A. I'm sorry, the website?

25 Q. Yeah.

1           A.       You're correct.  However, there's  
2           a publication of the RSDAA that does  
3           mention the Budapest criteria.  It's in  
4           their book about the treatment guidelines.

5           Q.       But under the heading what is  
6           RSD the Budapest criteria are not to be  
7           found there?

8           A.       Correct, on the website under  
9           that heading, you're correct.

10                   (Indicating.)

11           Q.       Now, as to your differential  
12           diagnosis the list that you gave;  
13           neuropathy, depression, anxiety, secondary  
14           gain, opioids, you're not in a position as  
15           you sit here today to say within a  
16           reasonable degree of medical probability  
17           that any of these are in fact the cause of  
18           Dr. Hogan's problems?

19           A.       That's correct.

20                   MR. INCLAN:  Objection.

21                   THE COURT REPORTER:  Clemente?

22                   MR. INCLAN:  Yes.

23                   BY MR. SPRAGUE:

24           Q.       Okay.  Also, as to Dr. -- the  
25           treatment for Dr. Hogan you are not in a

1 position as you sit here today to set  
2 forth a treatment plan for Dr. Hogan within  
3 a reasonable degree of medical probability.  
4 Is that correct?

5 A. Certainly on the basis of not  
6 having, I think, a proper diagnosis it  
7 would be -- it would be improper for me to  
8 even discuss treatment. You need to  
9 diagnosis before you treat. He hasn't been  
10 diagnosed in my opinion.

11 Q. And you haven't seen him?

12 A. I have not seen him.

13 Q. So you don't have any opinions  
14 as to what the proper treatments for this  
15 man is?

16 MR. INCLAN: Objection.

17 THE WITNESS: Well, no, that's  
18 not correct. I mean there's features. If  
19 I can accept the opinions and the signs  
20 and symptoms that are mentioned in the  
21 records, certainly I would say that he  
22 needs aggressive treatment for depression.  
23 I can say that unequivocally unless my  
24 colleagues are absolutely misportraying his  
25 situation. So I can say that, although I

1           have not seen him, you're correct. I've  
2           not seen him. I have not diagnosed him  
3           myself. But if my colleagues are  
4           presenting the signs and symptoms properly,  
5           I would say that, you know, there's a very  
6           serious depression problem that needs  
7           aggressive treatment. So I would say  
8           absolutely if these records are correct to  
9           a reasonable degree of medical certainty,  
10          somebody needs to treat this man's  
11          depression aggressively.

12           BY MR. SPRAGUE:

13           Q.       Now, then -- you did a records  
14           review in this case?

15           A.       Yes, sir.

16           Q.       Okay. Now, as to the Budapest  
17           criteria which you wrote on the board  
18           didn't you leave out something, Doctor?

19           A.       What are you referring to? I  
20           didn't write everything. I was trying to  
21           make it fit on the board.

22           Q.       I think you left out trophic  
23           changes?

24           A.       Correct. That's -- again, that  
25           comes under -- that fell under the -- let

1 me make sure I get this right. It's  
2 important so I get it right. The trophic  
3 changes are under the motor. Motor and  
4 trophic are clustered together  
5 statistically.

6 (Indicating.)

7 Q. And you didn't discuss what  
8 trophic changes are?

9 A. That's correct. You're right.

10 Q. Okay. What are trophic changes?

11 A. They are changes in the  
12 integument, meaning changes in skin, nails,  
13 hair. Also not mentioned is a trophic  
14 change -- are changes in bony integrity  
15 osteopenia as it's called.

16 Q. So changes to the appearance of  
17 the skin would qualify as a trophic change?

18 A. Yes, sir.

19 Q. And that would include if the  
20 skin appeared to transition as being shiny,  
21 correct?

22 A. Shiny is considered to be an  
23 early trophic change. It's not a hard  
24 trophic change such as scaliness, horniness,  
25 thickness. But certainly shiny skin, if it

1 is different from the unaffected side, is  
2 relevant.

3 Q. Okay. Now, Dr. Twigg observed  
4 hyperesthesia to the left third finger.  
5 That would be a sign of sensory -- a sign  
6 of complex regional pain syndrome in the  
7 sensory category, would it not, Doctor?

8 A. I'm sorry. I don't -- repeat  
9 the question because I can't find the  
10 record right this second.

11 MS. ARNOLD-SIMMONS: Here it is,  
12 Doctor.

13 THE WITNESS: Oh, I'm sorry.

14 MS. ARNOLD-SIMMONS: Here you  
15 go.

16 (Indicating.)

17 BY MR. SPRAGUE:

18 Q. Dr. Twigg observed  
19 hyperesthesia --

20 A. Yes.

21 Q. -- in the left third finger?

22 A. Yes.

23 Q. That would be a sign of RSD  
24 under the sensory category, correct, Doctor?

25 A. That is a sign, yes.

1 Q. Dr. Hogan complained to Dr.  
2 Twigg of left hand weakness. That would  
3 be a symptom under the motor category,  
4 would it not, Doctor?

5 A. That's correct.

6 Q. Dr. Twigg also observed that  
7 left handgrip strength was decreased. That  
8 would be a sign of RSD under the motor  
9 category, would it not, Doctor?

10 A. This actually could be a symptom  
11 and a sign because the patient -- it  
12 requires patient participation, patient  
13 compliance with what the doctor is asking  
14 the patient to do. So it is in the hands  
15 -- if you'll pardon the pun -- of the  
16 patient as to what -- you know, whether or  
17 not they actually try as hard as they can.

18 Q. Okay. Sign or symptom it's a  
19 -- it's consistent with RSD, correct?

20 A. Correct.

21 Q. She also observed skin color as  
22 shiny in the appearance of the left hand.  
23 That would be a sign of RSD under the  
24 trophic category, would it not, Doctor?

25 A. Yes.

1 Q. Dr. Steinberg observed that the  
2 left hand was cold. That would be a sign  
3 of RSD under the vasomotor category of the  
4 criteria, correct, Doctor?

5 A. Yes.

6 Q. Okay. Moving to Dr. George, Dr.  
7 George observed discoloration in the left  
8 hand. That would be a sign of RSD in the  
9 vasomotor category, correct?

10 A. Yes, sir.

11 Q. He also observed that it was  
12 noticed -- the left hand was noticeably  
13 cooler. That's also a sign of RSD under  
14 the vasomotor category?

15 A. Yeah. That's not a separate --  
16 I mean that's not a separate category.  
17 That's still one single category.

18 Q. It's one category but it's  
19 another sign in that category?

20 A. That's correct.

21 Q. He also observed that the left  
22 upper extremity was pale. That's also a  
23 sign in the -- of RSD in the vasomotor  
24 category?

25 A. Same category, yes, yet another



1                   embellishment to that category.

2                   Q.       He also observed -- made a  
3                   finding of hyperalgesia and allodynia.  
4                   That's a sign of RSD in the sensory  
5                   category, correct?

6                   A.       Correct.

7                   Q.       He also has found weakness in  
8                   the upper -- the left upper extremity or  
9                   that grip strength was reduced. And,  
10                  again, as we discussed, this would be a  
11                  sign or symptom of RSD in the motor  
12                  category, correct?

13                  A.       Correct.

14                  MR. INCLAN: Mr. Sprague, can  
15                  you tell us what date you're referring to?

16                  MR. SPRAGUE: I'm skipping  
17                  around.

18                  MR. INCLAN: Okay.

19                  BY MR. SPRAGUE:

20                  Q.       Dr. Ullah, he observed left  
21                  upper extremity edema, correct?

22                  A.       Yes.

23                  Q.       And that is a sign of RSD,  
24                  correct?

25                  A.       That's correct.

1 Q. He also observed decreased range  
2 of motion as a sign of RSD in the motor  
3 category, correct?

4 MR. INCLAN: Objection.

5 BY MR. SPRAGUE:

6 Q. Correct?

7 MS. ARNOLD-SIMMONS: Objection.

8 THE WITNESS: If it's  
9 asymmetric, yes.

10 MR. SPRAGUE: Okay.

11 THE COURT REPORTER: If it's...

12 THE WITNESS: Asymmetric meaning  
13 side-to-side differences.

14 (Indicating.)

15 BY MR. SPRAGUE:

16 Q. Dr. Ullah in that same note,  
17 Doctor, also observed that the right upper  
18 extremity was perfectly normal, did he not?

19 MR. INCLAN: Objection.

20 THE WITNESS: Correct.

21 BY MR. SPRAGUE:

22 Q. So this is asymmetrical, correct?

23 A. Correct. I decided to start  
24 saying that because many of the doctors  
25 that you mentioned before did not bother to

1 discuss symmetry, and this doctor actually  
2 did.

3 Q. Okay. So -- and he also  
4 discussed purple discoloration of the left  
5 upper extremity, correct?

6 A. That's correct.

7 Q. And that is a sign of RSD,  
8 correct?

9 A. That's correct.

10 Q. Okay. Now, we have had some  
11 discussion about Dr. Kirkpatrick. I want  
12 you to assume that Dr. Kirkpatrick's whose  
13 deposition has been taken in this case and  
14 he's testified that his notes regarding his  
15 observations of the left upper extremity  
16 are all in comparison to the right upper  
17 extremity and should be taken to indicate  
18 asymmetry. Can you make that assumption  
19 with me, Doctor?

20 A. I will make that assumption.  
21 It's certainly not in his records so.

22 Q. But, again, assuming that he's  
23 testified that way in his deposition, he  
24 has observed swelling in the left upper  
25 extremity. That would be a sign of RSD,

1 correct?

2 A. That's correct.

3 Q. Decreased strength in the left  
4 upper extremity, that would also be a sign  
5 of RSD, correct?

6 MR. INCLAN: Objection.

7 MS. ARNOLD-SIMMONS: Objection.

8 BY MR. SPRAGUE:

9 Q. Correct?

10 A. I'm sorry. What was the  
11 question again? You distracted me.

12 (Indicating.)

13 Q. He also -- he also observed  
14 decreased strength in the left upper  
15 extremity, and that would also be a sign  
16 of RSD, correct?

17 A. If we assume that he is speaking  
18 in reference one side to the other, yes,  
19 you're correct.

20 Q. And, again, we're making that  
21 assumption based on his deposition.

22 A. Yes.

23 Q. He testified of -- or to  
24 swelling of the left upper extremity. That  
25 would be a sign of RSD, correct?

1 A. That is correct.

2 Q. And also bluish discoloration;  
3 that would also be a sign of RSD, correct?

4 A. Now, it's important to note, of  
5 course, this is across many notes, many  
6 context, but you are correct. That is a  
7 sign.

8 Q. Okay. So we have Dr. George,  
9 Dr. Twigg, Dr. Ullah, Dr. Steinberg, and  
10 Dr. Kirkpatrick who all have observed signs  
11 of RSD in this man, correct?

12 A. Yes.

13 Q. And you noted that Dr. George  
14 has in your opinion observed enough to meet  
15 the criteria of the 1994 IASP criteria,  
16 correct?

17 A. That's correct.

18 Q. And that is --

19 A. On one occasion.

20 Q. And that is the -- that is the  
21 criteria that is currently codified by the  
22 Taxonomy Committee of the IASP, correct?

23 A. Yes, the Taxonomy Committee.

24 MR. SPRAGUE: I don't have any  
25 other questions.

1 (Indicating.)

2 MS. ARNOLD-SIMMONS: Mr. Inclan?

3 MR. INCLAN: Yes.

4 EXAMINATION

5 BY-MR. INCLAN:

6 Q. Hello, Doctor. This is Clemente  
7 Inclan on behalf of Baptist Hospital. I  
8 just want to clarify that as counsel for  
9 the plaintiff was just asking you questions  
10 regarding different entries in the medical  
11 records for various physicians, was he  
12 reviewing with you notes from one  
13 particular entry or was he taking you  
14 through an entire medical record for each  
15 individual physician?

16 A. It is -- it's clear that --  
17 that he was aggregating signs and symptoms  
18 across multiple notes for multiple doctors.

19 Q. What does aggregating mean for  
20 somebody who doesn't understand what that  
21 means, like myself?

22 A. Well, for instance, for Dr.  
23 Kirkpatrick he had gone through all of his  
24 notes and selected the signs across all of  
25 those notes, different notes, different

1 days.

2 Q. Okay. Regarding counsel for the  
3 plaintiff's questioning regarding Dr. -- Dr.  
4 George, did he ask you about signs and  
5 symptoms that cover more than one note?

6 A. I believe so, yes.

7 Q. Same question with regards to  
8 Dr. Kirkpatrick's entries?

9 A. Yes, sir.

10 Q. Okay. Doctor, let me ask you  
11 to assume that Dr. Hogan, the plaintiff in  
12 this case, at or about the time that --  
13 well, at or about the time of November  
14 where this dunking booth incident occurred  
15 -- let me ask you to assume that at that  
16 time he had a tax lien with the IRS of  
17 \$150,000. Let me also ask you to assume,  
18 Doctor, that he had a judgment entered  
19 against him for a previous job that he had  
20 a Lake City Medical Center whereby the  
21 judgment against him was for over \$90,000.

22 Let me also ask you to assume,  
23 Doctor, that at the same time Dr. Hogan  
24 was behind on his student loans. Let me  
25 ask you to assume also that he was behind

1 in paying the rent for his office at  
2 Baptist-Nassau. Let me ask you to assume  
3 also that Dr. Hogan at the time was aware  
4 of the fact that his ability to maintain  
5 his staff privileges at Baptist-Nassau were  
6 dependent upon his ability to become board  
7 certified as a general surgeon and that in  
8 fact he had taken and not passed the board  
9 exam on four previous occasions.

10 Let me ask you to assume also,  
11 Doctor, that all of these stressers were on  
12 Dr. Hogan at or about the time of this  
13 dunking booth. Can you say more likely  
14 than not whether or not those stressers can  
15 affect an individual, such as Dr. Hogan, to  
16 become depressed based on your training and  
17 experience?

18 MR. SPRAGUE: Okay. Objection,  
19 more prejudicial than probative, lack of  
20 predicate, and beyond the scope of this  
21 expert's qualifications.

22 THE WITNESS: Yes. Absolutely,  
23 the sociologic and psychologic stressers  
24 clearly could precipitate depression and  
25 anxiety in terms of the literature and in



1 terms of common medical knowledge and  
2 psychiatric knowledge.

3 MR. INCLAN: Okay. Thank you,  
4 Doctor.

5 MS. ARNOLD-SIMMONS: Is that  
6 all, Mr. Inclan?

7 MR. INCLAN: (No response.)

8 MR. TOLLEFSEN: Yeah, Doctor,  
9 this is Tom Tollefsen. I think Mr. Inclan  
10 answered -- everything I'd have for you, so  
11 I don't think I have any questions for  
12 you.

13 EXAMINATION

14 BY-MS.ARNOLD-SIMMONS:

15 Q. Dr. Harden, I just have a couple  
16 of follow-ups for you. The medical records  
17 that you were asked about by attorney  
18 Sprague; Twiggs, George, Ullah, any of  
19 those use the Budapest criteria which is  
20 what is necessary to make a diagnosis of  
21 RSD?

22 A. No.

23 MR. SPRAGUE: Objection.

24 BY MS. ARNOLD-SIMMONS:

25 Q. Looking at Dr. Ullah's record,

1 do you have that in front of you?

2 A. Yes, I do.

3 Q. If not -- I can -- if you would  
4 look at the last page, does he assess  
5 lower extremities in the last page of his  
6 report?

7 A. Yes, he does.

8 Q. And does he find any problem  
9 with the lower extremities?

10 MR. SPRAGUE: Objection, beyond  
11 the scope of cross.

12 THE WITNESS: No. The quote is  
13 full range of motion of all joints, no  
14 active swelling, no edema, good pedal  
15 pulses, no varicosity of leg veins in terms  
16 of the lower extremities, both sides.

17 BY MS. ARNOLD-SIMMONS:

18 Q. Now, I would ask you to look at  
19 Dr. Twiggs' records that you were asked  
20 about earlier. That's in your hand right  
21 there.

22 A. In my hand?

23 Q. (Indicating.)

24 A. Okay. Thank you.

25 Q. Looking at her note, does she

1 make the comparison from left to right as  
2 required under the Budapest criteria or as  
3 required for diagnosis?

4 A. In one case she -- she does  
5 talk about the left hand slight hypothenar  
6 wasting, hyperesthesia of the distal left  
7 third finger which would have to  
8 theoretically be compared to the right  
9 hand. But she doesn't say that. She  
10 doesn't specifically compare or mention  
11 asymmetry, but she does at least mention  
12 left. And then she goes on to talk about  
13 left hand; so only in the case of motor  
14 abnormalities. Again, motor abnormalities  
15 are contingent on patient's effort and  
16 patient's compliance with the direction of  
17 the doctor.

18 MS. ARNOLD-SIMMONS: Okay. Thank  
19 you. That's all.

20 MR. INCLAN: Aaron, are you  
21 going to ask more?

22 MR. SPRAGUE: Hold on.

23 MR. INCLAN: While Aaron is  
24 looking, Teresa, can you look at your  
25 BlackBerry?

1 MS. ARNOLD-SIMMONS: Yeah.

2 MR. SPRAGUE: Yes, I do have a  
3 question.

4 EXAMINATION

5 BY-MR.SPRAGUE:

6 Q. Dr. Harden, were you given the  
7 report of Dr. Dansinger (phonetic) in this  
8 case?

9 A. Dansinger?

10 Q. Dr. Dansinger.

11 A. I don't believe I have that.  
12 There was a D -- a doctor whose name began  
13 with D, but I think it was D-w, something,  
14 Drewin (phonetic), no, not Dansinger.

15 Q. So you're unaware of any of his  
16 opinions as they relate to the cause of  
17 Dr. Hogan's psychological problems, if any?

18 A. I don't have those records. No,  
19 sir.

20 MR. SPRAGUE: Okay. No further  
21 questions.

22 (Indicating.)

23 EXAMINATION

24 BY-MS.ARNOLD-SIMMONS:

25 Q. Doctor, one further question. In

1           your experience as a pain management  
2           specialist treating patients with RSD have  
3           you had any experience with patients who  
4           were factitious or manufacturing RSD?

5           A.       Yes.

6           Q.       Okay. And can you tell me how  
7           that can happen?

8           A.       I had one specific case that  
9           comes to mind where a lady was putting a  
10          tourniquet on her extremity prior to coming  
11          into the clinic. Putting a tourniquet on  
12          for approximately 30 minutes will cause  
13          swelling, discoloration, pain, and, of  
14          course, mimicks some of the signs and, of  
15          course -- well, specifically the signs of  
16          CRPS. We actually caught her doing that,  
17          of course, and that was the cause of  
18          unraveling a sociologic issue now with her.  
19          But it's actually fairly easy to create or  
20          feign the -- some of the symptoms of --  
21          and some of the signs of CRPS.

22                   MS. ARNOLD-SIMMONS: Okay. Thank  
23                   you. That's all I have.

24                   MR. SPRAGUE: Nothing.

25                   MS. ARNOLD-SIMMONS: I think

1                   that's it.

2                   THE VIDEOGRAPHER: This is the  
3                   end of tape No. 3 which concludes the  
4                   deposition of Norman Harden. The time is  
5                   20:07:20 and we are off the record.

6                   (WHEREUPON, a discussion was  
7                   had off record.)

8                   THE COURT REPORTER: Everybody's  
9                   order is the same?

10                  MS. ARNOLD-SIMMONS: Yes.

11                  MR. SPRAGUE: Yes.

12                  MR. INCLAN: Yes.

13                  MR. TOLLEFSEN: Yes.

14                  MR. SPRAGUE: And I would like  
15                  to order that video if I could get that by  
16                  Friday.

17                  NO EXHIBITS MARKED

18                  NO EXHIBITS ATTACHED

19                  .  
20                  .  
21                  .  
22                  .  
23                  .  
24                  .  
25                  .

CERTIFICATE OF REPORTER

1  
2 .  
3 I, STACEE L. JACKSON, Certified  
4 Shorthand Reporter for the State of  
5 Illinois, do hereby certify that the  
6 foregoing was reported by stenographic and  
7 mechanical means, which matter was held on  
8 the date, and at the time and place set  
9 out on the title page hereof and that the  
10 foregoing constitutes a true and accurate  
11 transcript of same.

12 I further certify that I am not  
13 related to any of the parties, not am I an  
14 employee of or related to any of the  
15 attorneys representing the parties, and I  
16 have no financial interest in the outcome  
17 of this matter.

18 I have hereunder subscribed my  
19 hand on the 14th day of September, 2007.

20 .  
21 \_\_\_\_\_  
22 STACEE L. JACKSON, CSR  
23 .  
24 .