UNITED STATES PATENT AND TRADEMARK OFFICE

BEFORE THE PATENT TRIAL AND APPEAL BOARD

NUVASIVE, INC. Petitioner

v.

WARSAW ORTHOPEDIC, INC. Patent Owner

Patent Number: 8,444,696 Issue Date: May 21, 2013 ANATOMIC SPINAL IMPLANT HAVING ANATOMIC BEARING SURFACES

Case IPR2013-00395

WARSAW'S MOTION FOR OBSERVATION REGARDING CROSS-EXAMINATION OF DR. BRANTIGAN

EXHIBITS

WARSAW2001	Affidavit of Mr. Luke Dauchot.
WARSAW2002	Affidavit of Mrs. Nimalka Wickramasekera.
WARSAW2003	U.S. Patent No. 4,834,757 to Brantigan
WARSAW2004	U.S. Patent No. 5,425,772 to Brantigan
WARSAW2005	Declaration of Dr. Charles L. Branch, Jr., M.D.
WARSAW2006	Curriculum vitae of Dr. Charles L. Branch, Jr., M.D.
WARSAW2007 '430 patent.	Comparison of claim 1 of the '696 patent and claim 1 of the
WARSAW2008 '430 patent.	Comparison of claim 4 of the '696 patent and claim 4 of the
WARSAW2009 April 7, 2014.	Deposition transcript of Dr. John W. Brantigan, M.D. taken
WARSAW2010	Declaration of Lori Ferrell, CPA, CGMA.
WARSAW2011	CLYDESDALE® Spinal System Product Information.

DOCKET A L A R M Find authenticated court documents without watermarks at <u>docketalarm.com</u>. Warsaw Orthopedic, Inc. ("Patent Owner") submits this motion for observation regarding cross-examination of Dr. Brantigan, the reply declarant of Petitioner NuVasive, pursuant to the Board authorization provided via email communication dated June 18, 2014. In lieu of taking an additional deposition of Dr. Brantigan, the parties have agreed to Patent Owner's use of the prior deposition testimony (taken April 7, 2014) of Dr. Brantigan. Warsaw submits the following observations based on Dr. Brantigan's prior deposition testimony.

Observation 1

In Exhibit 2009 (at 81:2 to 82:20; and 86:15 to 87:7), Dr. Brantigan concedes that adding ratchetings such as the nubs 122 of Brantigan '035 to the intermediate ridges 68, 68a of the spinal disk implant 50 of Senter would cause resistance to insertion thereof between the vertebrae 22a and 22b. This testimony is relevant to the reply declaration of Dr. Brantigan at paragraphs 8 and 9 because Dr. Brantigan now ignores the downside (specifically identified in Senter) of adding ratchetings such as the nubs 122 to the intermediate ridges 68, 68a. Such resistance to insertion is undesirable – Senter indicates that the ridges 68, 68a are "preferably smooth, without serrations, to permit it to be surgically implanted." (Patent Owner's Response at 36:1-5; and Senter (Ex. 1007) at 11:30-31.) Furthermore, Dr. Brantigan in paragraph 8 of the reply declaration considers adding ratchetings such as nubs 122 to the intermediate ridges 68, 68a would be a

"belt-and-suspenders" approach, but such an approach is contrary to the express teachings of Senter.

Observation 2

In Exhibit 2009 (at 92:22-24), Dr. Brantigan indicates that, when placing bone in an opening of the modified spinal disk implant 50 of Senter, "you would put as much bone in there as you could," and "[t]hat bone would be tending to bulge out beyond the confines of the implant." This testimony is relevant to the reply declaration of Dr. Brantigan at paragraph 11 because Dr. Brantigan now asserts that "[o]ne of skill in the art would also have known before June 1995, as is known today, that spinal fusion implants do not need to be loaded to the very absolute top of the fusion aperture, in order to facilitate bone growth through the implant from one adjacent vertebra to the other." During insertion between the vertebrae 22a and 22b, the bone filling the opening of the modified spinal disk implant 50 of Senter would be subject to dislodgement. (Patent Owner's Response at 44:12-15.) Patent Owner submits that Dr. Brantigan's changing testimony is relevant to his credibility and to whether one of ordinary skill would modify the spinal disk implant 50 of Senter as suggested by Petitioner NuVasive.

Observation 3

In Exhibit 2009 (at 94:1-22), Dr. Brantigan indicates that the portion of the intermediate ridges 68, 68a of the spinal disk implant 50 of Senter removed to

provide the opening therethrough would be filled with bone-growth promoting material, and that a portion of the bone-growth promoting material that now occupies the cut-away of the intermediate ridge 68 would extend above the posterior ledge 60. Regarding the bone-growth promoting material that extends above that posterior ledge 60, Dr. Brantigan in Exhibit 2009 (at 96:7-8) indicates that "I believe that some of the bone would be scraped off." This testimony is relevant to the reply declaration of Dr. Brantigan at paragraphs 13 and 14 because Dr. Brantigan now disparages the dislodgement problem identified by Patent Owner. The bone-growth promoting material filling the cut-away of intermediate ridge 68 would not be protected during insertion of the modified spinal disk implant 50 between the vertebrae 22a and 22b. (Patent Owner's Response at 44:12-15.) As seen in Petitioner NuVasive's modified Fig. 3 of Senter (Petitioner's Corr. Petition '395 at page 14), the abrupt transition of the intermediate ridge 68 with the posterior ledge 60 does not provide such protection. Furthermore, Dr. Brantigan (reply declaration at paragraph 13) now asserts that, since Medtronic, Inc. (Patent Owner's parent company) does not identify dislodgement problems associated with the Clydesdale implant, the dislodgement problems identified by Patent Owner regarding the modified spinal disk implant 50 of Senter are not actually a problem. In making these assertions, Dr. Brantigan does not acknowledge his previous indication ("I believe that some of the bone

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