EXHIBIT 5

TO THE DECLARATION OF BRIAN J. NISBET IN SUPPORT OF DEFENDANTS' MOTION FOR SUMMARY JUDGMENT



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9	UNITED STAT	ΓES DISTRICT COURT
11	SOUTHERN DISTRICT OF (CALIFORNIA – SAN DIEGO DIVISION
12		
13	NUVASIVE, INC., a Delaware corporation	Case No. 3:18-CV-00347-CAB-MDD
14	Plaintiff,	[Assigned to Courtroom 4C – Honorable Cathy Ann Bencivengo]
15	V.	[Magistrate: Hon. Mitchell D. Dembin]
16 17 18	ALPHATEC HOLDINGS, INC., a Delaware corporation and ALPHATEC SPINE, INC., a California corporation, Defendants.	HIGHLY CONFIDENTIAL – ATTORNEYS' EYES ONLY SUBJECT TO PROTECTIVE ORDER
19		REBUTTAL REPORT OF BARTON L.
20		SACHS, M.D., M.B.A., F.A.C.P.E.,
21		F.A.C.H.E.
2223		Complaint Filed: February 13, 2018
23 24		Jury Trial Demanded
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27 28		EXHIBIT Ex 002 Barton Sachs 01/11/2021
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doctrine of equivalents analysis. Dr. Youssef's conclusory analysis simply erases meaningful structural limitations from the claims. I understand that in any case, that is not a proper doctrine of equivalents analysis and Dr. Youssef has failed to meaningfully set forth any such analysis here.

255. For the reasons discussed above, the Battalion™ Lateral Spacer does not infringe claim 1 of the '334 patent.

(a) Dependent Claims 16 and 18

256. Claims 16 and 18 depend from independent claim 1. Alphatec's BattalionTM Lateral Spacer does not infringe these claims at least for the same reasons as described above with respect to independent claim 1.

(b) Dependent Claim 18

- 257. Claim 18 additionally requires that the "maximum lateral width of said implant is approximately 18 mm." It is my opinion that certain configurations of the BattalionTM Lateral Spacers do not meet this limitation, literally or under the doctrine of equivalents.
- 258. The following configurations of the Alphatec's Battalion™ Lateral Spacers do not have a "maximum lateral width ... [of] ... approximately 18 mm."

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16 mm width, 30 mm length, 6 mm height (parallel and lordotic)
22 mm width, 45 mm length, 8 mm height (parallel and lordotic)
22 mm width, 45 mm length, 10 mm height (parallel and lordotic)
22 mm width, 45 mm length, 12 mm height (parallel and lordotic)
22 mm width, 50 mm length, 8 mm height (parallel and lordotic)
22 mm width, 50 mm length, 10 mm height (parallel and lordotic)
22 mm width, 50 mm length, 12 mm height (parallel and lordotic)
22 mm width, 50 mm length, 14 mm height (parallel and lordotic)
22 mm width, 55 mm length, 8 mm height (parallel and lordotic)
22 mm width, 55 mm length, 10 mm height (parallel and lordotic)

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22 mm width, 55 mm length, 12 mm height (parallel and lordotic)
22 mm width, 60 mm length, 8 mm height (parallel and lordotic)
22 mm width, 60 mm length, 10 mm height (parallel and lordotic)
22 mm width, 60 mm length, 12 mm height (parallel and lordotic)

259. Dr. Youssef states—without any explanation—that "Alphatec offers a version of the BattalionTM Lateral Spacer that is 22 mm wide, which is approximately 18 mm." (Youssef Rpt. at Ex. D at 69 (emphasis added).) The 22 mm wide BattalionTM Lateral Implant, which is 22% wider than the 18 mm Battalion™ Lateral Spacer, does not have a maximum lateral width of "approximately 18 mm." In fact, Dr. Youssef's opinion that the 22 mm wide BattalionTM Lateral Spacer falls within the scope of this claim only confirms that this claim lacks reasonable certainty and renders the claim indefinite.

260. In fact, Dr. Youssef offers no written description or intrinsic evidence to support that the claims cover the above-listed configurations of the Alphatec's BattalionTM Lateral Spacers. Specifically, the '334 patent only discloses implants "having a width ranging between 9 and 18 mm, a height ranging between 8 and 16 mm, and a length ranging between 25 and 45 mm." ('334 patent at 2:17–21.) Per this disclosure, the above-listed configurations of the BattalionTM Lateral Spacers having a 22 mm width, 6 mm height, 50 mm length, 55 mm length, and 60 mm length are categorically excluded from the claim scope. Dr. Youssef's opinion that these implants are covered by the claimed dimensions appears to be based on a subjective assessment and there is no objective criteria by which a POSA would conclude that these implants would fall within the scope of this claim.

Further, Dr. Youssef states in a conclusory fashion and without the required analysis or explanation that this limitation is also infringed under the doctrine of equivalents. (Youssef Rpt. at Ex. D at 74.) Dr. Youssef's generalized statement alleging infringement under the doctrine of equivalents is unsupported, and lacks

particularized testimony and argument regarding the insubstantiality of any differences between the claimed invention and the accused implants that I understand is required in a proper doctrine of equivalents analysis. Dr. Youssef's conclusory analysis simply erases meaningful structural limitations from the claims. I understand that in any case, that is not a proper doctrine of equivalents analysis and Dr. Youssef has failed to meaningfully set forth any such analysis here.

262. Accordingly, these configurations of the BattalionTM Lateral Spacer do not infringe claim 18 of the '334 patent.

2. TranscendTM LIF PEEK Spacer

263. In my opinion, Alphatec's TranscendTM LIF PEEK Spacer does not infringe claims 16 and 18 of the '334 patent for the reasons set forth below.

(a) Independent Claim 1

- 264. Below I have reproduced independent claim 1 of the '334 patent. I understand that NuVasive is not asserting this claim, but asserted claims 16 and 18 depend from independent claim 1.
 - 1. A spinal fusion implant of non-bone construction positionable within an interbody space between a first vertebra and a second vertebra, said implant comprising:

an upper surface including anti-migration elements to contact said first vertebra when said implant is positioned within the interbody space, a lower surface including anti-migration elements to contact said second vertebra when said implant is positioned within the interbody space, a distal wall, a proximal wall, a first sidewall and a second sidewall, said distal wall, proximal wall, first sidewall, and second sidewall comprising a radiolucent material;

wherein said implant has a longitudinal length greater than 40 mm extending from a proximal end of said proximal wall to a distal end of said distal wall;

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