## UNITED STATES PATENT AND TRADEMARK OFFICE

BEFORE THE PATENT TRIAL AND APPEAL BOARD

MEDTRONIC, INC. AND MEDTRONIC VASCULAR, INC.

Petitioner,

v.

## TELEFLEX LIFE SCIENCES LIMITED,

Patent Owner.

Case IPR2020-01341 U.S. Patent No. 8,142,413

Case IPR2020-01342 U.S. Patent No. 8,142,413

Case IPR2020-01343 U.S. Patent No. RE 46,116

Case IPR2020-01344 U.S. Patent No. RE 46,116

SUPPLEMENTAL DECLARATION OF STEPHEN JON DAVID BRECKER, MD, FRCP, FESC, FACC



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		2.	After removing Kontos's proximal funnel, a POSITA would have been motivated to maximize the usable real estate, but even if no further modifications were made, Kontos would not have a "problem gap."			
		3.	After removing Kontos's proximal funnel, Kontos's support catheter 10 would remain coaxially aligned with the guide catheter			
	D.	expe diffe	OSITA would have been motivated with a reasonable ectation of success to achieve the no-more-than-one-french erential between inner diameters of the guide catheter and 12.			
	E.	My opinion remains unchanged: the claims that recite a two-inclined side opening are obvious				
		1. 2.	Kontos + Ressemann and Kontos + Ressemann + Takahashi55 Kontos + Ressemann + Takahashi + Kataishi			
V.		eflex's evidence regarding secondary considerations does not alter my nion that the challenged claims are obvious				



#### I. Overview

- 1. I have been retained by Robins Kaplan LLP on behalf of Medtronic, Inc. and Medtronic Vascular, Inc. ("Medtronic") as an independent expert to provide my opinions concerning U.S. Patent Nos. 8,142,413 and RE46,116 (the "Teleflex Patents") in IPR2020-01341, IPR2020-01342, IPR2020-01343, and IPR2020-01344.
- 2. I set forth the information below as a supplement to my original declarations, as Teleflex raised new issues in their Patent Owner Responses on which I had not previously been given an opportunity in these IPRs to offer testimony. Having considered Teleflex's arguments, and the testimony of Teleflex's declarants, my opinions on the invalidity of the Teleflex Patents remain the same.
- 3. I have reviewed additional material in conjunction with my analysis set forth herein. This includes the Final Written Decisions in IPRs on related Teleflex patents: IPR2020-00126, IPR2020-00127, IPR2020-00128, IPR2020-00129, IPR2020-00130, IPR2020-00132, IPR2020-00134, IPR2020-00135, IPR2020-00136, IPR2020-00137 and IPR2020-00138 ("Related IPRs"). A list of

<sup>&</sup>lt;sup>1</sup> Citations to exhibits refer to exhibits filed in IPR2020-01341, unless noted otherwise. I understand that most of Teleflex's and Medtronic's exhibits are numbered consistently across all four IPRs.



these materials includes everything cited in this declaration, and the materials disclosed in my original declarations.

## II. Claim Construction

- A. "interventional cardiology device"
- 4. The opinion in this section generally relates to at least IPR2020-01341 and IPR2020-01342.
- 5. I am aware that when the Board instituted the IPR2020-01341 petition it determined that the limitation "interventional cardiology device" refers to "at least two types of the devices selected from the group that includes, but is not limited to, guidewires, balloon catheters, stents, and stent catheters." IPR2020-001341, Paper 11 ("I.D.") at 16. This limitation appears in independent claim 1 of the '413 patent.
- 6. At institution, the Board additionally determined that the claims do not require that more than one of guidewires, stents, stent catheters, and balloon catheters be simultaneously insertable into and through the lumen of the claimed coaxial guide catheter. *Id.* Additionally, it determined that Medtronic demonstrated where Itou discloses every limitation of claim 1. *Id.* at 24.



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