champion.

2019 December 10

EC DECLARATION OF CONFORMITY

Champion Manufacturing Inc.
2601 Industrial Parkway
Elkhart IN 46516 USA

Matt Weismiller, President/CEO
Rhonda Hurd, Director of Quality/Regulatory Affairs

Declares that the medical device(s) described hereafter

54 Series, 59 Series, Continuum, Passage – PF, 250 Series, 251 Series, 85 Series, 89 Series, Ascent, Ascent II, Concord, Passage – PS, 56 Series, 58 Series, 86 Series, Ascent XL, Ascent II XL, 54P Series, 59P Series, 56P Series,58P Series, 85P Series, 89P Series, Ascent, Ascent II, 86P Series, Ascent XL, Ascent II XL, Elevate

Having a classification of (1) using Annex IX rule (12), is in conformity with the essential requirements and provisions of Council directive 93/42/EEC, per Annex VII and is designed and manufactured under a quality management system that is self-certified.

I, the undersigned, hereby declare that the device specified above conforms to Directive 93/42/EEC.

Signature, Quality/Regulatory

Rhonda Hurd
Director of Quality and Regulatory Affairs
On behalf of: Champion Mfg. Inc.

Signature, President/CEO

Matt Weismiller
President/CEO
On behalf of Champion Mfg. Inc.