

25 October 2017

Blount Swain
Ability Dynamics
1236 W. Southern,
Suite 101
Tempe, AZ 85282
USA

Dear Blount:

I am writing to inform you that today, we have notified the Dutch Competent Authority*.

With this notification, Ability Dynamics has met the requirements of Article 14 of the Medical Devices Directive, 93/42/EEC for the following devices:

- **RUSH Chopart Plate**

As of today and without any further notice from the respective Competent Authorities, Ability Dynamics can consider the respective devices as officially notified.

If you have any questions, please do not hesitate to contact me.

Yours sincerely,



Rene van de Zande
Director
Emergo Europe

**MDD notification with the Dutch Healthcare Inspectorate grants you access to 28 EU countries and 4 others:*



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28 EU: Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, and United Kingdom.

Additional: Switzerland and the EEA – European Economic Area (Iceland, Liechtenstein, Norway)