Declaration of Conformity

Manufacturer:
Leggero, LLC
2802 S. Water Street
Burnet, Tx USA 78611 Ph: +1 (512) 756-4700

EC Authorized Representative:
MDSS GmbH
Schiffgraben 41
30175 Hannover, Germany
Tel: +49 511 6262 8630+

We, Leggero, LLC, declare that the Dyno is in conformity with the applicable provisions of 2017/745 European Union (EU) Medical device Regulation of the European Parliament and, if applicable, with any other relevant Union legislation that provides for the issuing of an EU declaration of conformity. The Dyno has been tested and been found in full compliance with EN 12183:2009 and EN 12182:1999 by Berlin Cert as of May 18, 2011.

Dyno Part Numbers: 3001-1001 and 3001-1002

Class of Device: Class 1 medical device. Rule 1.

SRN: DE-AR-000005430

Basic UDI-DI #: ++B5143001VN

Description of Device: Rehab pushchairs and pediatric positioning systems are appropriate for children who are limited to a seated position and require mobility and positioning assistance. Individuals who need such assistance can have a wide variety of diagnosis limiting their ability to achieve mobility without such a system.

"We explicitly designate MDSS GmbH, Schiffgraben 41, 30175 Hannover, Germany to act as our sole Authorized Representative in the European Union for the above indicated products."

Place of Issue:
Leggero, LLC
2802 S. Water Street
Burnet, Tx USA 78611

Attestation made by the following Leggero LLC Personnel:

Gabriela Romero Cedillo	Gabriela R. Cedillo	7/5/2021
(Name) Authorized Representative	General Manager, Leggero LLC	Date of Issue