

EU DECLARATION OF CONFIRMITY

Manufacturer: Huka B.V.

Address: Münsterstraat 13, 7575 ED Oldenzaal, The Netherlands

SRN: Single Registration Number NL-MF-000003494

Product type: Tricycle

Device name: City L

Basic UDI: 8720589794030

Risk class: Class 1

Conformity assessment route: Huka uses the following procedures for the CE marking of

their products according to Regulation MDR 2017/745:

Class 1: according to (EU) 2017/745 Annex VIII

This declaration of conformity is issued under the sole responsibility of Huka. We hereby declare that the medical device (s) specified above comply with the provisions of Regulation (EU) MDR 2017/745 for medical devices.

This declaration is supported by the certified quality system according to ISO 13485: 2016 issued by "het Keurmerkinstituut" and the relevant sections of the following harmonized standards: EN-ISO 14971: 2012 and EN 12182: 2012.

All supporting documentation is kept on the manufacturer's premises.

Signed on behalf of Huka: Place and date:

Oldenzaal, 21 May 2021

Rob Lotgerink Managing Director

Spenia !



Form Name: MDR CE Declaration of Conformity ENG v20210521