EC DECLARATION OF CONFORMITY

We, Ralon Medical Equipment Co., Ltd, Xilian Development Zone, Jinsha Danzao, Nanhai Foshan, Guangdong, China, hereby declare under our own responsibility that the following products:

Product Name: MOKA Rollator

Product Code: SA1303601

Meet the provisions of the Council Directive 93/42/EC and the essential requirements which apply to them.

The abovementioned devices have been classified as class I devices according to rule I

This declaration is supported by the Quality Management System certification:

No. Q60 711570 005, EN ISO 13485:2016 + AC:2016 Medical devices- Quality management systems issued by TÜV SÜD PRODUCT Service GmbH-Zertifizierstelle- Ridierstraße 65, 80339 München, Germany

This declaration is based on the conformity assessment of products to the requirements of Annex VII

This Declaration is valid for all products concerned bearing the CE mark and manufactured by the above entitled "Manufacturer"

Foshan China

Issue place and date (14/11/2019)

<Stamp>

European Authorized Representative:

«Manufacturer's Signature

<Name>

<Position>Pr

Registered Address:

Obelis s.a.

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Representative: Mr. Gideon ELKAYAM (CEO)

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