Norwegian
Medicines Agency

To Whom it May Concern

NO-23/27676

Date: 04.12.2023

Certificate of Free Sale

The Norwegian Medicines Agency as competent authority for medical devices hereby declares that the following manufacturer has its registered place of business in Norway:

For **Easy Roller AS**
SRN: NO-MF-000010301
Hagebyvegen 40
3734 Skien
Norway

We have been notified that the following medical devices bear the CE-mark in accordance with Regulation (EU) 2017/745/Regulation (EU) 2017/746:

Product Name	*Certificate number*	*Basic UDI-DI*
EasyRoller	**N/A**	**709006211EASYROLLER2R**
EasyRoller Pool	**N/A**	**709006211EASYROLLER2R**

The precondition for medical devices to be CE-marked in accordance with Regulation (EU) 2017/745 on medical devices and Regulation (EU) 2017/746 on in vitro diagnostic medical devices is to meet the essential requirements for safety. A CE-marked medical device may be marketed within the European Union/European Economic Area without any approval from the Norwegian Medicines Agency, and may be exported without any restrictions.

This certificate is valid until 04. December 2025

Raymond Ludvigsen
Senior Advisor

NORWEGIAN MEDICINES AGENCY

Jørgen Bryn
Consultant

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Please state our reference.
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