



Livinguard AG  
Bahnhofstrasse 12, 6300 Zug,  
Switzerland

21/09/2020

Dear business partners,

Based on requests to provide official documentation of the submissions made on 1<sup>st</sup> July 2020, Livinguard AG writes to inform you that the products listed below have been confirmed by Swissmedic as **Class I Medical Device** according to Art. 6 MedDO. We confirm that the products below can continue to be sold in Switzerland and the European Union as Class I Medical Device.

The notification number by Swissmedic is **CH-202007-0004**.

The CE Marking, the wording "Medical Device Class I" and "compliant with EN 14683:2019 Medical Face Mask Type I (or Type II)" must be displayed on the packaging of the products.

**Livinguard PRO MASK** (without valve) - SKU #FM-1-NV

**Livinguard ULTRA MASK** (without valve) - SKU #FM-2-NV

We declare the above-mentioned products fulfill the European Regulation MDR 2017/745 and EN 14683:2019 Medical Face Mask Type I (PRO MASK without valve) or Type II (ULTRA MASK without valve).

Sincerely,

5.1.2e

Livinguard AG

---

**Livinguard AG**  
Bahnhofstrasse 12, 6300 Zug, Switzerland