

Budapest, 17th February 2023.

## **DECLARATION OF CONFORMITY 07-2022**

We undersigned **MY MED KFT**, with head office addressed in Budapest, Vendel utca 11, declare under its own responsibility that the medical device "My Filler" (Type: Soft, Medium, Strong, Man Max, Glips, and HA Refill), Lot Number O003, N082, N083, N041, N082 and N039, risk class III, GMDN code 59131, according to rule 8 to the Directive 93/42/EEC (enforced in Italy by Legislative Decree 46/97, and further amendments), as amended by Directive 2007/47 / EC (enforced in Italy with Legislative Decree 37/10),

- comply with essential requirements and dispositions of the Directive 93/42/EEC and further amendments, as the Technical File no. FT 002 MM retained by the Company;
- is manufactured according to the Quality System which satisfies the requirements of Annex II of above mentioned Decree, as per Certificates n. EPG-0218-19 and QCT-0118-19 dated 14/05/2019 issued by Notify Body Istituto Superiore di Sanità, Notify Body 0373, viale Regina Elena 229 - 00161 - Roma (RM), Italy.

**My Med Kft**  
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