MEDI PLAST

EC Declaration of Conformity

Manufacturer	**Mediplast AB**
Address	**Bronsåldersgatan 2** **213 76 Malmö** **Sweden**
Product group	**Vastrip**
Classification	**Class IIa**
Assessment route	**Annex II, MDD 93/42/EEC**
Notified body	**Intertek Semko AB identification No. 0413**
EC certificate	**41311518**

Mediplast AB hereby declares that the product group Vastrip, enclosing the below listed products, fulfills applicable requirements of the Swedish Medical Device Act SFS 1993:584 and Swedish Regulation LVFS 2003:11, enforcing the European Medical Device Directive 93/42/EEC.

Product name	***Product ID***
67321	Vastrip2+
67322	Vastrip Special

Johan Bongstorp, Managing Director
Malmö 2020-08-17