MEDI PLAST

EC Declaration of Conformity

Manufacturer	**Mediplast AB**
Address	**Bronsåldersgatan 2** **213 76 Malmö** **Sweden**
Product group	**Sterile bandages for fixation and support**
Classification	**Is**
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 Adhesive Non-woven Dressings, 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.

Ref. No	Type/Model	Size/Length
607027S	Primalast sterile	4 cm x 5 m
607028S	Primalast sterile	6 cm x 4 m
607029S	Primalast sterile	8 cm x 4 m
607030S	Primalast sterile	10 cm x 4 m
607032S	Primalast sterile	15 cm x 4 m
607034S	Primalast sterile	12 cm x 4 m
6050401S	DanaUniversal sterile	4 cm x 5 m
6050402S	DanaUniversal sterile	6 cm x 5 m
6050403S	DanaUniversal sterile	8 cm x 5 m
6050404S	DanaUniversal sterile	10 cm x 5 m
6050406S	DanaUniversal sterile	15 cm x 5 m
6050301S	Danalastic sterile	6 cm x 5 m
6050303S	Danalastic sterile	10 cm x 5 m

Ref. No	Type/Model	Size/Length
6050304S	Danalastic sterile	12 cm x 5 m

PGL 127(4)

Johan Bongstorp, Managing Director
Malmö 2021-05-28