MEDI PLAST

# EC Declaration of Conformity

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| ***Manufacturer*** | **Mediplast AB** |
| ***Address*** | **Bronsåldersgatan 2**<br>**213 76 Malmö**<br>**Sweden** |
| ***Product group*** | **IV-fixation** |
| ***Classification*** | **Class 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 IV-fixation, 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*** |
|---|---|
| Transparent IV fixation, 5 x 5,7 cm | 60702101 |
| Transparent IV fixation, 7 x 8,5 cm | 60702102 |
| Transparent IV fixation, 10 x 12 cm | 60702103 |
| Transparent IV fixation, 8,5 x 11,5 cm | 60702104 |
| Transparent IV fixation, 10 x 15,5 cm | 60702105 |
| Transparent IV fixation, 6 x 7 cm | 60702106 |
| Transparent IV fixation, 8,5 x 10,5 cm | 60702107 |
| Transparent IV fixation, 7 x 9 cm | 60702108 |

*Johan Bongstorp, Managing Director*
*Malmö 2020-07-29*