CE Conformity Declaration

In accordance with the annex n° VII of the application decree n° 45 dated February 24th 1997 of the Italian Government (annex VII of the European Directive 93/42/EEC) the undersigned Mr. Sergio Battara (Quality Assurance Manager) of EUROSIREL S.p.A., legal seat at Settimo Milanese,MI - ITALY, Via Volta 23 ensures and declares that the following Medical Device:

- Plasters 6cm x 2cm (material: woven non woven brown colour with pre-cut system, polybag 350 pcs, without covering paper)
- Plasters 6cm x 1,7cm (material: foil different colours with pictures , with pre-cut system, polybag 350 pcs, without covering paper)

Manufaktured for: MEDIPOS P&P, s.r.o., 675 52 Lipník č.p.44 , rezidence adress : Vinohradská 156/2222, 130 00 Praha 3, IČO: 469 94 742 DIČ: CZ46994742

belong to Class I and fulfil the applicable rules of the decree.
The above mentioned medical device is not sterile.
I undertake to fulfil the general requirements of the annex I.
On Your demand EUROSIREL S.p.A. has manufactured in its facility of Settimo Milanese, via Volta, 23, Milano - ITALY.
All the technical files in accordance with annex VII are at Your disposal and at disposal of the competent authorities.

Settimo Milanese, 05/10/23

Sergio Battara
Quality Assurance Manager