

# Declaration of Conformity

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**MANUFACTURER**

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Hygiene360 AG  
Büelstrasse 17  
8330 Pfäffikon ZH  
Switzerland

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**AUTHORIZED REPRESENTATIVE**

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Hygiene360 s.r.o.  
Vinohradská 2828/151  
Žižkov  
130 00 Prague 3  
Czech Republic

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**IDENTIFICATION AND DESCRIPTION OF THE DEVICE**

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PROSEPT® Spray (Ready-to-use solution for the quick and comprehensive disinfection of non-invasive medical devices):

Item Code	Trade Name	Description of Delivery Form
OD-041005	PROSEPT® Spray	250 ml bottle
OD-041013	PROSEPT® Spray	1 litre bottle with flip top cap
OD-041025	PROSEPT® Spray	5 litre canister

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**RISK CLASS OF THE DEVICE**

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Class IIa (according to the classification rules in Annex IX of the Council Directive 93/42/EEC concerning medical devices)

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**CONFORMITY ASSESSMENT PROCEDURE**

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Annex II (excluding Section 4 of the Council Directive 93/42/EEC concerning medical devices)

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**STANDARDS APPLIED**

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EN ISO 13485:2016, EN ISO 14971:2019, ISO 10993-1:2018, EN 62366-1:2015, EN 14885:2018, EN ISO 21530:2004, EN 1041:2008+A1:2013, EN ISO 15223-1:2016

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**NOTIFIED BODY**

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DNV Product Assurance AS  
Veritasveien 3  
1363 Høvik  
Norway

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**CE MARK AFFIXED**

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**CE**  
2460

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**AUTHORIZED SIGNATORY**

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This Declaration of Conformity is issued under the sole responsibility of Hygiene360 AG. We hereby declare that the above-mentioned device(s) meet the provisions of the Council Directive 93/42/EEC concerning medical devices. This declaration is supported by the Quality System approval to EN ISO 13485, issued by DNV Product Assurance AS Notified Body Number 2460. All supporting documentation is retained at the premises of the manufacturer.

Name: Juerg Suter  
Designation: Chief Operating Officer  
Place of Issue: Pfäffikon ZH  
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