

# MANUFACTURER'S DECLARATION OF CONFORMITY

## AUSTRALIAN THERAPEUTIC GOODS (MEDICAL DEVICES) REGULATIONS 2002

### DECLARATION OF CONFORMITY PROCEDURES

This is a declaration of conformity made under clause 6.6 of Schedule 3 to the Therapeutic Goods (Medical Devices) Regulations 2002.

**Manufacturer's Name:** ANSELL HEALTHCARE EUROPE N.V.

**Business Address:** Bld Internationelelaan 55,  
B-1070 Brussels,  
Belgium

**Medical device(s):** Micro-Touch® Blue Nitrile

**Classification:** Class I

**GMDN Code and Term:** 56286 - Nitrile examination/treatment glove, non-powdered, non-sterile

**Scope of Application:** All examination gloves from this manufacturer

Each kind of medical device to which the technical documentation applies complies with the applicable provisions of the essential principles and the classification rules before being supplied.

**Standards Applied:** EN 455 Parts 1, 2, 3, & 4 and ISO 11193-1

**Authorized Signatory:**

**Ansell Healthcare Europe NV**  
Riverside Business Park - Block J  
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