Declaration of Conformity

Manufacturer:	Becton Dickinson and Company 7 Loveton Circle Sparks, Maryland 21152, USA Tel: +1.410.316.4000 Fax: +1.410.316.4499
Authorized Representative:	Benex Limited Pottery Road, Dun Laoghaire Co. Dublin, Ireland Tel: +353.1.202.5222
Conformity Assessment Procedure:	Directive 98/79/EC of the European Parliament and of the Council, Annex III of Directive 98/79/EC
Product:	

REF	Product Name
222371	Difco Haemophilus Influenzae Antiserum Poly, Contains Types a, b, c, d, e, f. 1 mL
222501	Difco Haemophilus Influenzae Antiserum Type a, 1 mL
222361	Difco Haemophilus Influenzae Antiserum Type b, 1 mL
227891	Difco Haemophilus Influenzae Antiserum Type c, 1 mL
227901	Difco Haemophilus Influenzae Antiserum Type d, 1 mL
227911	Difco Haemophilus Influenzae Antiserum Type e, 1 mL
227921	Difco Haemophilus Influenzae Antiserum Type f, 1 mL

We hereby declare that the above mentioned products comply with the European In Vitro Diagnostic Medical Devices Directive and its relevant transposition into national laws of the member states into which we place the devices. This declaration is issued under the sole responsibility of Becton, Dickinson and Company.
Date: 07-Feb-2022
Name and Authority: Anne Zavertnik Vice President Regulatory Affairs, IDS
Signature: 

Technical File Number: BALTER222371

RECORD REVISION HISTORY TABLE	
Revision	**Description of Changes**
A	Initial Release