EU DECLARATION OF CONFORMITY

MANUFACTURER:	DIESSE DIAGNOSTICA SENESE SPA STRADA DEI LAGHI 39 53035 MONTERIGGIONI (SI), ITALY
SINGLE REGISTRATION NUMBER	IT-MF-000013311
AUTHORIZED REPRESENTATIVE:	NOT APPLICABLE
PRODUCT: CODE:	**CHORUS HERPES SIMPLEX 1 IgG RECOMBINANT** **81023**
INTENDED PURPOSE:	CHORUS HERPES SIMPLEX 1 IgG RECOMBINANT (REF 81023) is an immunoassay kit for automated qualitative detection of IgG class antibodies against Herpes simplex virus (Type 1). The test is performed in human serum, using a disposable device applied on the Chorus and Chorus TRIO instruments. The kit is intended to detect the exposure to Herpes simplex virus (Type 1) infection as an aid to the relative diagnosis. It must be used by professional laboratory users only.
BASIC UDI-DI	803389132CHORUSHSV00BE
UDI-DI	08033891328689
RISK CLASS:	CLASS C
CLASSIFICATION RULE:	RULE 3a
CONFORMITY ASSESSMENT ROUTE:	ANNEX IX (CHAPTER I AND III, SECTION 4.4 TO 4.8)

WE HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCT IS IN CONFORMITY WITH **REGULATION (EU) 2017/746** OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL ON IN VITRO DIAGNOSTIC MEDICAL DEVICE. ALL SUPPORTING DOCUMENTATION IS RETAINED UNDER THE PREMISES OF THE MANUFACTURER.

REFERENCE TO ANY CS APPLIED:	NOT APPLICABLE
NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH ZERTIFIZIERSTELLE RIDLERST. 65 – 80339 MÜNCHEN GERMANY No. 0123
(EU) CERTIFICATE:	V12 056726 0004 Rev.00
REVISION:	0
PLACE, DATE OF ISSUE:	MONTERIGGIONI, 28 JUNE 2022

EXPIRY DATE:	2027-06-26

THE PRESENT EU DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE AND EXCLUSIVE RESPONSIBILITY OF THE MANUFACTURER.

SIGNATURE:

CHIARA MUZZI
PERSON RESPONSIBLE FOR THE REGULATORY COMPLIANCE

THIS IS THE CERTIFIED COPY OF THE ORIGINAL DOCUMENT STORED IN ARCHIVE OF DIESSE DIAGNOSTICA SENESE SPA

ISSUED: MONTERIGGIONI, 2022-06-28

MAGDALENA STOCZKO
REGULATORY SUPERVISOR