DECLARATION OF CONFORMITY

Product name / Trade name	REF IS-5100	BASIC UDI-DI: 5060169690738XP
IDS-iSYS 17-OH Progesterone	GMDN: 58346	EMDN: W0102050107
Intended Purpose	For *In Vitro* Diagnostic Use. For Laboratory Professional Use. The IDS-iSYS 17-OH Progesterone assay is an in vitro diagnostic device intended for the quantitative determination of 17-OH Progesterone in human serum on the IDS system. Results are to be used as an aid in the diagnosis and monitoring of various disorders of the adrenal glands in adult and paediatric population.	

	EC REP
Immunodiagnostic Systems Limited, 10 Didcot Way, Boldon Business Park, Boldon, Tyne and Wear, NE35 9PD, UK SINGLE REGISTRATION NUMBER: GB-MF-000015851	Immunodiagnostic Systems SA 101 Rue Ernest Solvay B-4000 Liege Belgium SINGLE REGISTRATION NUMBER: BE-AR-000015342

RISK CLASS: ☐ A ☒ B ☐ C ☐ D

CLASSIFICATION RULE (ANNEX VIII) : Rule 6

CONFORMITY ROUTE: ☒ ANNEX IX Full Quality System (Class B, C & D)

☐ ANNEX I & II+III *(non-sterile Class A)*

Name of the Notified Body / Identification: TÜV Rheinland LGA Products GmbH 0197

CE Marking Date: **23rd June 2026**

EU CERTIFICATE No.: **HX 2333083-1**

COMMON SPECIFICATIONS: Not Applicable

We, as the manufacturer of the device(s) take sole responsibility for and hereby declare that the above-mentioned product(s) meet(s) the provisions of the following Regulation(s)/Directives

- Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices

Date: 23rd June 2026

Place: UK

Signed on behalf of Immunodiagnostic Systems Limited

M Henderson
Group RA Manager & PRRC

Date: 23rd June 2026

Place: UK

Signed on behalf of Immunodiagnostic Systems Limited

Mohammed Adil
RA Manager (Specialist)

Ver.06