



EU Quality Management System Certificate

Regulation (EU) 2017/746, Annex IX Chapter I and III

IVDR 744391 R000

Manufacturer: The Binding Site Group Ltd

Address:

8 Calthorpe Road Edgbaston Birmingham B15 1QT United Kingdom

Single Registration Number: UK-MF-000036707

EU Authorised Representative: The Binding Site Ireland Limited

Address:

First floor, 43-49 Sir John Rogerson's Quay Dublin 2 Ireland

Scope: See attached Device Schedule

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/746, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class D devices, and self-test, near-patient test and companion diagnostic devices an Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

First Issue Date: 2023-05-04 Starting Validity Date: 2024-11-04

Current Issue Date: **2024-11-04** Expiry Date: **2028-05-03**

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.

This certificate was issued electronically and is bound by the conditions of the contract.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80 Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK. A Member of the BSI Group of Companies.





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Device Schedule: Class D, C and B devices

Class C devices	Intended purpose
W0102 - Immunochemistry (Immunology)	Immunochemistry (Immunology) devices for in vitro
IVP 3007 - In vitro diagnostic devices which require knowledge regarding	detection of proteins as an aid in diagnosis and
immunoassays	monitoring of monoclonal gammopathies,
	inflammatory conditions, kidney disease and
	haematological malignancies.
Class B devices	Intended purpose
IVR 0608 – Devices intended to be used for screening, determination or	Immunochemistry (Immunology) devices intended
monitoring of physiological markers.	to be used for screening, determination or
	monitoring of physiological markers
IVR 0506 – Other devices intended to be used to determine markers of	Immunochemistry (Immunology) devices intended
infections / immune status	to be used to determine markers of infections /
	immune status

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Certificate History

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate. Verification@bsigroup.com)

Date	Reference Number	Action
2023-05-04	3387963	Issued
Current	30299791	Supplemented – addition of device category Class B IVR 0506 Supplemented – addition of SRN

First Issue Date: **2023-05-04**

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