

EU Quality Management System Certificate

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

IVDR 747622 R000

Manufacturer: Cytocell Limited

Address:

Oxford Gene Technology
418 Cambridge Science Park
Milton Road
Cambridge
CB4 0PZ
United Kingdom

Single Registration Number: GB-MF-000016893

EU Authorised Representative: Sysmex Europe SE

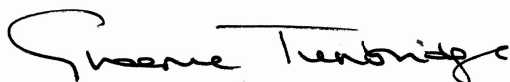
Address:

Bornbarch 1
22848 Norderstedt
Germany

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 Medical Devices

First Issue Date: **2023-03-09**

Current Issue Date: **2023-03-09**

Starting Validity Date: **2023-03-09**

Expiry Date: **2028-03-08**

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Device Schedule: Class C devices

| Class C devices | Intended purpose |
|--|---|
| W0106 – Genetic testing | Chromosomal analysis device for the detection of prenatal trisomy 13 & 21. |
| IVP3004 – In vitro diagnostic Devices which require knowledge of chromosomal analysis | |
| W0106 – Genetic testing | Chromosomal analysis devices for the detection of acquired cancer-related chromosome alterations. |
| IVP3004 – In vitro diagnostic Devices which require knowledge of chromosomal analysis | |

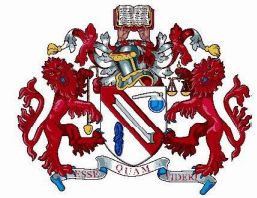
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Certificate History

(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 |
|---------|------------------|--------|
| Current | 3413836 | Issued |



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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.