Certificate of Registration of Quality Management System to I.S. EN ISO 13485:2016

The National Standards Authority of Ireland certifies that:

CytoTest Inc.
9430 Key West Avenue
Suite 210
Rockville, MD 20850
USA

has been assessed and deemed to comply with the requirements of the above standard in respect of the scope of operations given below:

The design, manufacture, and distribution of DNA fluorescence in situ hybridization (FISH) probes for the diagnosis of chromosomal abnormalities.

Approved by:
Geraldine Larkin
Chief Executive Officer

Approved by:
Caroline Dore Geraghty
Director of Medical Devices
/ Head of Notified Body

Registration Number: MD19.4925
Certification Granted: March 31, 2015
Effective Date: February 13, 2019
Expiry Date: March 30, 2021

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