CYTOTEST®

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MANUFACTURER'S EC DECLARATION OF CONFORMITY

CytoTest declares that the device meets the provisions of the Council Directive 98/79/EC for in vitro diagnostic medical devices.

Product Identification:

Generic Device Term	Commercial Name	Catalogue Reference Number
DNA in situ Hybridization Reagents	DNA Fish Probe Kit	CT-PACxxx
	Chromosome Counting Probes	CT-CCPxxx
	Locus Specific Probes	CT-LSPxxx
	Subtelomere Probes	CT-STPxxx

Device Classification:	All Other IVDs: General IVD; Self-Certified
Manufacturer's Name:	CytoTest, Inc
Business Address:	1395 Piccard Drive Suite 308 Rockville, MD 20850
European Authorized Representative:	Obelis S. A. Bd. General Wahis 53 B-1 030 Brussels Belgium
CE Mark First Applied:	2016
Route to Compliance:	Annex III, self-declared
Responsible Person:	Reinhard Ebner, President and CEO

Date: 12/13/2022

Reinhard Ebner
President and CEO

DELAWARE 2015
CYTOTEST INC.
File No. 5861157