


EU Declaration of Conformity

(N° dc90137ben)

WE THE MANUFACTURER

| | |
|--|---|
| Name | HORIBA ABX SAS |
|  Address | Parc Euromédecine Rue du Caducée BP 7290 34184 Montpellier Cedex 4 FRANCE |
| Single Registration Number | FR-MF-000000320 |

TAKE SOLE RESPONSIBILITY FOR AND HEREBY DECLARE THAT THE PRODUCT(S)

| | |
|-------------------|------------------------------|
| Device category | Hemostasis Reagent |
| Product name | Yumizen G CLEANER |
| Models | 1300036420 |
| Basic UDI-DI | 361023ymz_g_cleanerP2 |
| Country of origin | HUNGARY |

Intended Use

Yumizen G CLEANER is a washing solution used for daily maintenance and cleaning of Yumizen G Line fully automated coagulation analyzers during routine use.
For *in vitro* diagnostic use only.

MEET(S) THE PROVISIONS OF THE FOLLOWING DIRECTIVES, REGULATIONS, STANDARDS AND COMMON SPECIFICATIONS

| | |
|----------------------------------|--|
| Regulations | Regulation (EU) 2017/746 on <i>in vitro</i> diagnostic medical devices Risk Class: A <input checked="" type="checkbox"/> B <input type="checkbox"/> C <input type="checkbox"/> D <input type="checkbox"/> |
| IVDR conformity assessment route | <input checked="" type="checkbox"/> ANNEX II + ANNEX III + ANNEX IV (<i>Class A devices excluding sterile devices</i>) |
| Common Specifications | Not applicable |

Montpellier, France
2022/09/06

Claire MALLIÉ
Quality & Regulatory Affairs Junior
Director / PRRC

