

EC Declaration of Conformity

Application of Council Directive:

- 98/79/EC of 27 October 1998 on In Vitro Diagnostic Medical Devices
- 1999/5/EC of 9 March 1999 on radio equipment and telecommunications terminal equipment and the mutual recognition of their conformity

Means of conformity:

The following product is in conformity with;

- Directive **98/79/EC** based on the test results using harmonised standards in accordance with Article 5 of the Directive
- Directive **1999/5/EC** based on the test results using harmonised standards in accordance with Article 5 of the Directive

Product identification:

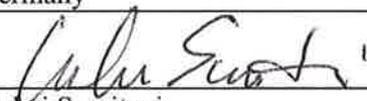
Product name: Automated Hematology Analyzer XN series
System name: XN-1000, XN-2000, XN-3000, XN-9000
Module name: XN-10, XN-20, XN-11, XN-21
Accessories: SA-10, SA-20, BT-40, CV-50, CV-60, CV-70, ST-40, ST-41, ST-42, TU-40, RR-10, SA-30, WG-40, WG-50, WG-60, SA-01

Manufacturer:

Name: SYSMEX CORPORATION
Address: 1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073
Country: Japan

Authorised representative:

Name: SYSMEX EUROPE GMBH
Address: Bornbarch 1, 22848 Norderstedt
Country: Germany

Authorised officer: 
Kohéi Sumitani

Position: Managing Director
Date: September 13, 2013
Place: Germany

This certificate was issued under sole responsibility of:

Authorised officer: 
Keiji Fujimoto

Position: Executive Officer
Date: September 2nd, 2013
Place: Japan