# 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:

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| 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:

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

Authorised representative:

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| Name: | SYSMEX EUROPE GMBH |
| Address: | Bornbarch 1, 22848 Norderstedt |
| Country: | Germany |
| Authorised officer: | Kohei Sumitani |
| Position: | Managing Director |
| Date: | September 13, 2013 |
| Place: | Germany |

This certificate was issued under sole responsibility of:

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| Authorised officer: | Keiji Fujimoto |
| Position: | Executive Officer |
| Date: | September 2nd, 2013 |
| Place: | Japan |

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