

## EC Declaration of Conformity



We hereby declare that the product described below conforms to all applicable requirements of Council Directive 98/79/EC for in vitro diagnostic medical devices.

**Legal Manufacturer:** Siemens Healthcare Diagnostics Inc.  
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Tarrytown, NY, 10591-5097, USA

**Place of Manufacture:** ThermoFisher Scientific  
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Middletown, VA, 22645-0307, USA

**EC Authorized Representative:** Siemens Healthcare Diagnostics Ltd.  
Sir William Siemens Square  
Frimley, Camberley, GU16 8QD, UK

**Product Name:** ADVIA 120/2120/2120i CBC TIMEPAC

**Catalogue Number (REF):** 09826813

**Siemens Material Number (SMN):** 10312269

**Legacy Product Code:** T01-3620-52

**Classification:** General IVD

**Conformity Assessment Route:** ANNEX III

**Document Control Number:** DoC\_ADVIA 120/2120/2120i CBC TIMEPAC

**Version:** 1.0

*This declaration of conformity is issued under the sole responsibility of Siemens Healthcare Diagnostics Inc.  
This declaration supersedes any declaration issued previously for the same product.*

**Signature:**

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**Matthew Gee**  
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2015-11-12

**Date**  
[YYYY-MM-DD]