

## EU DECLARATION OF CONFORMITY (DoC)

|   |  |
|---|--|
| <b>Manufacturer:</b>  | Becton, Dickinson and Company<br>BD Biosciences<br>155 North McCarthy Boulevard<br>Milpitas, California 95035 USA  |
| <b>Manufacturer SRN:</b>  | US-MF-000017797  |
| <b>Authorised Representative:</b>   | Becton Dickinson Ireland Ltd.<br>Donore Road, Drogheda<br>Co. Louth, A92 YW26 Ireland  |
| <b>Authorised Representative SRN:</b>   | IE-AR-000007610  |
| <b>Product:</b>   | <div>BD® CD14 (MΦP9) FITC <span>REF</span> 345784</div> <div>BD® CD14 (MΦP9) PE <span>REF</span> 345785</div> <div>BD® CD14 (MΦP9) PerCP <span>REF</span> 345786</div> <div>BD® CD14 (MΦP9) APC <span>REF</span> 345787</div> <div>BD® CD14 (MΦP9) APC-Cy7 <span>REF</span> 333951</div>   |
| <b>Basic UDI-DI:</b>  | 038290OHQPGOVBFN   |
| <b>Risk Class and Rule:</b>   | Class C, Annex VIII, Rule 3(g) & Rule 3(h)   |
| <b>Intended Purpose:</b>  | <p>CD14 (MΦP9) is intended for in vitro diagnostic use in the identification of cells expressing the CD14 antigen in peripheral blood, using a BD FACSLytic™ flow cytometer.</p> <p><u>Clinical Applications</u></p> <p>Expression of the CD14 antigen in the characterization of subjects having, or suspected of having, hematological neoplasia.</p> <p>CD14 (MΦP9) is a qualitative reagent intended for laboratory professional use only.</p> |
| <b>Notified Body:</b>   | BSI Group The Netherlands B.V.<br>Say Building, John M. Keynesplein 9,<br>1066 EP Amsterdam, Netherlands<br>Notified Body Number: 2797   |
| <p>We, as the manufacturer of the device(s) take sole responsibility for and hereby declare that the above-mentioned product(s) meet(s) the provisions of the following Directives/ Regulation(s):</p> <ul style="list-style-type: none"> <li>Regulation (EU) 2017/746 on In vitro Diagnostic Medical Devices.</li> </ul> |  |

## Conformity Assessment Route:

|  |   |
|--|---|
| <input type="checkbox"/> ANNEX IX Technical File Examination     | EC CERTIFICATE No.:<br>EC Certificate Expiration Date:                              |
| <input checked="" type="checkbox"/> ANNEX IX Full Quality System | EC CERTIFICATE No.: 728780 R000<br>EC Certificate Expiration Date: 2025-December-16 |
| <input type="checkbox"/> ANNEX X Type Examination                | EC CERTIFICATE No.:<br>EC Certificate Expiration Date:                              |
| <input type="checkbox"/> ANNEX XI Production Quality System      | EC CERTIFICATE No.:<br>EC Certificate Expiration Date:                              |
| <input type="checkbox"/> ANNEX I & II+III                        | N/A   |

## Common Specifications (CS):

Common Specifications have not been issued for this product.

| Authorised Signatory:    |  |
|--------------------------|--|
| <b>Name &amp; Title:</b> | Mirna Dipano, Vice President, Regulatory Affairs   |
| <b>On behalf of:</b>     | Becton, Dickinson and Company<br>BD Biosciences<br>155 North McCarthy Boulevard<br>Milpitas, California 95035 USA  |
| <b>Place of Issue:</b>   | Milpitas, California USA   |
| <b>Date of Issue:</b>    | Please refer to Signature section.   |
| <b>Signature:</b>        | <div> DocuSigned by:<br/> <br/>  Signer Name: Mirna Dipano<br/> Signing Reason: I approve this document<br/> Signing Time: 30-Jun-2023   8:46:45 AM PDT<br/> 6166B70768DA44C1A0CA7A9871E2F40A </div> |

## DECLARATION OF CONFORMITY Revision History:

| Version: | Detailed Change Description:   |
|----------|--|
| A        | Original release.  |
| B        | Update to Corporate Template CBI-058 FRM24 Revision 04. Updated Risk Class & Rule section by replacing “Class C, Annex VIII Rule 3” with “Class C, Annex VIII, Rule 3(g) & Rule 3(h)” to support global registration, and removed the Template Revision History section. Included the detailed address of the Notified Body. Updated the Date of Issue section in the Authorised Signatory table from date generated from DocuSign to “Please refer to Signature section.” |
| C        | Updated grammar and formatting. Updated the legal manufacturer address and PRRC.   |