

# EU Quality Management System Certificate

Regulation (EU) 2017/746, Annex IX Chapter I and III

## IVDR 741870 R000

**Manufacturer:** Canterbury Scientific Limited

**Address:**

71 Whiteleigh Avenue  
Addington  
Christchurch  
8011  
New Zealand

**Single Registration Number:** NZ-MF-000001778

**EU Authorised Representative:** Emergo Europe B.V.

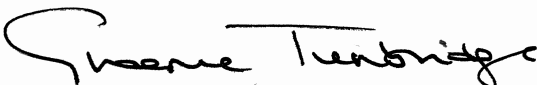
**Address:**

Westervoortsedijk 60  
6827 AT Arnhem  
The Netherlands

**Scope:** See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/746, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class D devices, and self-test, near-patient test and companion diagnostic devices an Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):



Graeme Tunbridge, Senior Vice President Medical Devices

First Issue Date: **2023-02-15**

Current Issue Date: **2023-02-15**

Starting Validity Date: **2023-02-15**

Expiry Date: **2028-02-14**

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### Class C devices

W0101 – Clinical Chemistry.  
IVP 3007 In vitro diagnostic devices which require knowledge regarding immunoassays.

### Intended purpose

Control materials with quantitative assigned values for glycated haemoglobin

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### Certificate History

*(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)*

Date	Reference Number	Action
Current	3346873	Issued



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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.

This certificate was issued electronically and is bound by the conditions of the contract.