

Annex to declaration of accreditation (scope of accreditation)  
Normative document: EN ISO/IEC 17021-1:2015  
Registration number: **C 637**

of **Kiwa Dare B.V.**

This annex is valid from: **23-06-2022** to **01-07-2026**

Replaces annex dated: **23-02-2022**

### Location(s) where activities are performed under accreditation

#### Head Office

Vijzelmolenlaan 7  
3447 GX  
Woerden  
The Netherlands

Standard / Normative document	Certification scheme <sup>1</sup>
ISO 13485	Medical Devices - Quality Management Systems - Requirements for regulatory purposes for the scopes: <ul style="list-style-type: none"><li>- Active (non-implantable) medical devices<ul style="list-style-type: none"><li>- General active medical devices</li><li>- Devices for imaging</li><li>- Monitoring devices</li><li>- Devices for radiation therapy and thermotherapy</li><li>- Active (non-implantable) medical devices other than specified above</li></ul></li></ul>

This annex has been approved by the Board of the  
Dutch Accreditation Council, on its behalf,

J.A.W.M. de Haas

<sup>1</sup> If no date or version number is mentioned for a normative document, the accreditation concerns the most current version of the document or scheme.

<sup>1</sup> If there is a reference to a code starting with NAW, NAP, EA or IAF, this concerns a scheme mentioned on the [RvA-BR010-lijst](#).