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This is to certify that

Certificate: 181030177P
Date of release: 30-10-2018

Willem van de Biggelaar

Has participated in the extensive EU MDR (2017/745) training:

- Module 1: Introduction, timelines, scoping, definitions, Eudamed, implementing/delegated acts
- Module 2: Classification, Conformity Assessment routes, QMS
- Module 3: GSPR's, Labeling, UDI
- Module 4: Clinical evaluation, PMCF, PMS, Vigilance / FSCA,
- Module 5: Technical documentation, Notified Body assessment, Health Institutes

Each Module consisted of 8 hours plenary training.

By Pieter de Vries
Principal Consultant Medical Device Project B.V.