

Declaration of Conformity

The following in-vitro diagnostic medical device (software as medical device)

Name of product: **Ikaros** Release Version: 6.3

fulfill the applicable requirements of the Directive 98/79 EC of 27 Oct. 1998, Annex III, "Other (General)".

This declaration covers all options and configurations of the above products delivered by MetaSystems

Manufacturer: **MetaSystems Hard & Software GmbH**

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The MetaSystems products are in-vitro diagnostic medical devices according to

GMDN CT1250 Analyser software IVDs

This declaration of conformity is issued under the responsibility of MetaSystems Hard & Software GmbH and valid until 26.05.2026.

Altlusheim, 2022-05-12



Dr. Andreas Pfuhl
(Managing Director)

CE DECLARATION