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Oncosia Scientific

CE Declaration of Conformity

according to the Regulation (EU) 2017/745 on Medical Devices (MDR), Annex IV

We, the manufacturer

Oncosia Scientific GmbH
Dorfstr. 19
91602 Duerrwangen
Germany
Basic UDI: PP11873ONCOPREVIA31 / IFA

declare herewith under our sole responsibility that the following medical device

OncoPrevia™

in the category

of risk class **I (one)** (according to rule 1 and 13 set out in annex VIII of Regulation (EU) 2017/745)

CE-marked with

CE

to which this declaration relates, meets all the provisions of Regulation (EU) 2017/745 of the European Parliament and of the Council on Medical Devices which apply to it through conformity assessment procedure annex II and III of the Regulation (EU) 2017/745.

Place, Date, Signature of CEO	Dürrwangen, 12.12.2024	Nico Schürrle
Place, Date, Signature of PRRC	Dürrwangen, 12.12.2024	Nico Schürrle