

DECLARATION OF CONFORMITY

We,

Name and address of manufacturer: *Penta Arzneimittel GmbH
Werksstrasse 3
92551 Stulln
Germany*

declare on our own responsibility that

the medical device

Name **EyeMedica® Gereizt + Gerötet
Augen-Tropfen**

Class **IIb**

Manufacturing Instructions **XVII/MP19 - OSD**

meets all applicable requirements of the Directive
93/42/EEC Annex II. – excluding Section 4.

Applied standards (optional): -

Other normative documents (optional): -

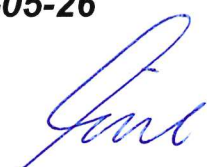
Name, address and identification
number of Notified body(ies): *mdc medical device certification
GmbH
Kriegerstrasse 6
70191 Stuttgart
Germany
(0483)*

Conformity assessment procedure: *93/42/EEC, annex II-excl. Sec.4*

Registration no.: **D1061000047**

Validity: **2024-05-26**

Place, date
Stulln, 20.05.2021



Legally binding signature

Name and function **C. Wilhelm
Head of Regulatory Affairs
Medical Devices** **Penta Arzneimittel GmbH
Werksstraße 3
92551 Stulln**