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

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Werksstraße 3
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