Declaration of Conformity

CE

AGFA HEALTHCARE N.V.

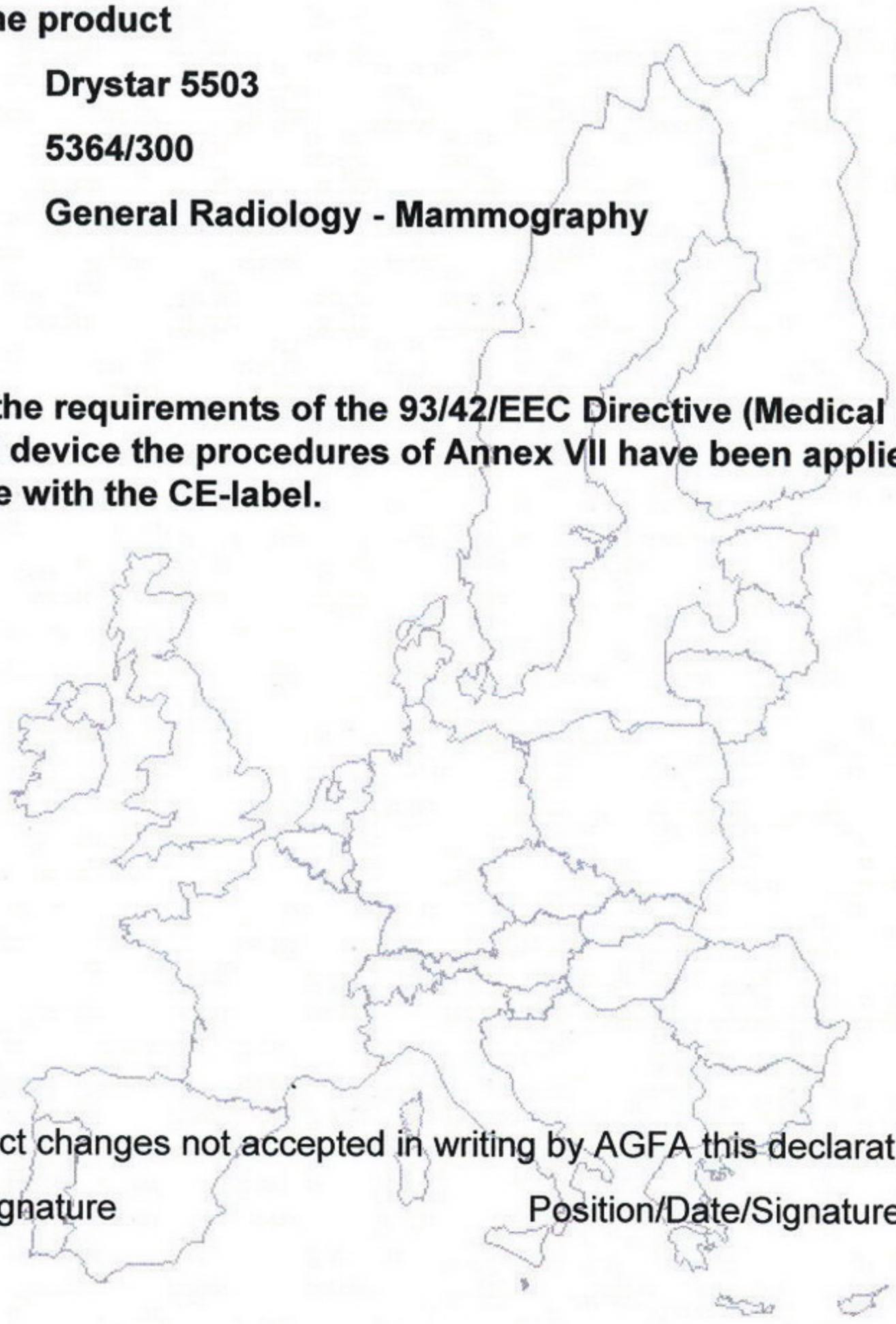
Septestraat 27, 2640 Mortsel, Belgium

declares that the product

Name: **Drystar 5503**

Type/model: **5364/300**

Application: **General Radiology - Mammography**



complies with the requirements of the 93/42/EEC Directive (Medical Device).
For this Class I device the procedures of Annex VII have been applied in order to mark the device with the CE-label.

In case of product changes not accepted in writing by AGFA this declaration will expire.

Position/Date/Signature

30 MAART 2007

Jan Langmans
Manager Global Regulatory Affairs Agfa HealthCare

Position/Date/Signature

30 MAART 2007

Stephen R. Abbott Ph. D.
Vice President, Global Quality Assurance & Regulatory Affairs Agfa HealthCare

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