

Declaration of Conformity



AGFA HEALTHCARE N.V.

Septestraat 27, 2640 Mortsel, Belgium

declares that the product

Category: X-ray film, medical, screen

Name/Type:	DRYSTAR DT 1 B	DRYSTAR TS 2 O
	DRYSTAR DT 1 C	DRYSTAR TS 2 C
	DRYSTAR DT 2 B	DRYSTAR TS 2 CF
	DRYSTAR DT 2 C	DRYSTAR DT 1.000 B
	DRYSTAR DT 5 B	DRYSTAR DT 1.000 C
	DRYSTAR DT 10 B	DRYSTAR DT 5.000 B
		DRYSTAR DT 5.000I B

Application: General Radiology

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.
This declaration is valid maximum for 5 years after the signature date.

Position, Signature & Date

02 APR. 2014

Paul Merckx
Director Agfa HealthCare Quality Assurance & Regulatory Affairs
Business Division Imaging



Details as of PDF Creation Date

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