Declaration of Conformity

Agfa NV

Septestraat 27, 2640 Mortsel, Belgium

declares that the product

Name/Type:	**DRYSTAR DT 1 B**	**DRYSTAR DT 1 C**
	DRYSTAR DT 2 B	**DRYSTAR DT 2 C**
	DRYSTAR DT 5 B	**DRYSTAR DT 5 C**
	DRYSTAR DT 10 B	**DRYSTAR DT 10 C**
	DRYSTAR DT 1.000 B	**DRYSTAR DT 1.000 C**
	DRYSTAR DT 5.000 B	**DRYSTAR DT 5.000 C**
	DRYSTAR DT 5.000I B	**DRYSTAR DT 5.000I C**
	DRYSTAR DT 2 B XL	**DRYSTAR DT 2 C XL**
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

14 -02- 2019

Paul Merckx
Head of Quality Assurance & Regulatory Affairs
Agfa NV

AGFA