

# 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

#### Document Metadata

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1. Peter Guldentops (amlpo) on 2014-04-10 09:30 AM CET

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