KONICA MINOLTA

EU DECLARATION OF CONFORMITY

Manufacturer

Name	KONICA MINOLTA, INC.
Address	1 Sakura-machi, Hino-shi, Tokyo, 191-8511, Japan

declares, sole responsibility, that the following product

GMDN term:	Diagnostic-imaging multiformat camera
GMDN code:	35580
Generic Device Group:	Laser Imagers
Type:	LASER IMAGER
Model:	DRYPRO MODEL 832
Classification:	Class I, Rule 12, 93/42/EEC
Serial Number:	From 0922-52376 to 0922-99999
Including:	Printlink5-IN, Lispl-832

referred to in this declaration conforms with the following EU law(s) :

COUNCIL DIRECTIVE 93/42/EEC, confirmed by the procedure of its Annex VII, and Directive 2011/65/EU

and conforms with the following standard(s):

EN ISO 13485:2012+AC:2012, EN ISO 14971:2012, EN 1041:2008, EN 980:2008, EN 60601-1:2006, EN 60601-1-2:2007, EN 60601-1-6:2010, EN 60825-1:1994+A1:2002+A2:2001, EN 62366:2008, EN 62304:2006 for DIRECTIVE 93/42/EEC,
EN 50581:2012 for Directive 2011/65/EU

and that this declaration is valid upon approval for release of each product.
The manufacturer will keep on file for review the technical documentation.

EU Representative

Name	Konica Minolta Business Solutions Europe GmbH
Address	Hoogoorddreef 9, 1101 BA Amsterdam, The Netherlands

Signed for and on behalf of manufacturer:

Tokyo Japan, 2017-10-01
(Place and date of issue)
YASUSHI YAMANAKA
General Manager,
Quality Assurance Operations
Healthcare Business Unit
Healthcare Business Headquarters
(Name, function)

(Signature of equivalent authorized by the manufacturer)

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