



KONICA MINOLTA



# EU DECLARATION OF CONFORMITY

**Manufacturer**

Name KONICA MINOLTA, INC.  
Address 1 Sakura-machi, Hino-shi, Tokyo, 191-8511, Japan  
Single Registration Number JP-MF-000008214

**TỔNG GIÁM ĐỐC**  
*Trần Thị Mỹ Duyên*

**declares, sole responsibility, that the following product**

Generic Device Group: Laser Imager  
Type: LASER IMAGER  
Model (Product Name): DRYPRO MODEL 873  
Basic UDI-DI: 45601419200002589  
Intended Purpose: A device is intended for use in the acquisition and process of radiographic images of human anatomy. It is intended to replace radiographic film/screen system in general-purpose diagnostic procedures.  
Classification: Class I, Rule 1, according to Annex VIII of REGULATION (EU) 2017/745  
Serial Number: from 0993-61235 to 0993-99999 (A2GP)  
Including Printlink5-IN

**referred to in this declaration conforms with the following EU law(s):**  
REGULATION (EU) 2017/745 and Directive 2011/65/EU

**and conforms with the following standard(s):**  
EN ISO 13485:2016, EN ISO 14971:2012, EN 1041:2008+A1:2013,  
prEN ISO 15223-1:2020, EN 60601-1:2006+A1:2013, EN 60601-1-2:2015,  
EN 60601-1-6:2010, EN 60825-1:1994+A1:2002+A2:2001, EN 62366:2008,  
EN 62304:2006 for REGULATION (EU) 2017/745,  
EN IEC 63000:2018 for Directive 2011/65/EU

**and that this declaration is valid upon approval for release of each product.**

**EU Representative**

Name Konica Minolta Business Solutions Europe GmbH  
Address Capellalaan 65, 2132 JL, Hoofddorp, The Netherlands  
Single Registration Number NL-AR-000002026

**Signed for and on behalf of manufacturer:**

Tokyo Japan, 2021-08-24  
(Place and date of issue)  
HAJIME NOZAWA  
General Manager,  
Quality Assurance Operations  
Healthcare Business Unit  
Healthcare Business Headquarters  
(Name, function)

(Signature of equivalent authorized by the manufacturer)