MINISTRY OF HEALTH, LABOUR AND WELFARE
GOVERNMENT OF JAPAN
2-2, KASUMIGASEKI 1-CHOME, CHIYODA-KU, TOKYO 100-8916

CERTIFICATE

It is hereby certified that the following medical devices marketed by KONICA MINOLTA, INC., 2-7-2 Marunouchi, Chiyoda-ku, Tokyo, 100-7015, Japan are manufactured under our supervision as stipulated in the Pharmaceuticals, Medical devices and Other Therapeutic Products Act of Japan and are authorized to be marketed in Japan.

Medical devices:

1. LASER IMAGER DRYPRO MODEL 832
2. LASER IMAGER DRYPRO MODEL 873

Manufacturing Site and Address:

KONICA MINOLTA TECHNOPRODUCTS CO., LTD.

2-2-1 Hirosedai, Sayama-shi, Saitama, 350-1328, Japan

For legalization by the foreign consul in Japan,
this is to certify that the Seal affixed to this document
is genuine.
Tokyo, DEC. 20 2018 T. TANAKA
Official
Ministry of Foreign Affairs
(Consular Service Division)

No. 4065

Tokyo, date NOV. 30. 2018

Kiyohito Nakai
Director, Medical Device Evaluation Division
Pharmaceutical Safety and Environmental Health Bureau
Ministry of Health, Labour and Welfare