CONFIDENTIAL
SOLELY FOR REGULATORY PURPOSES

Date: 08 February 2019

TO WHOM IT MAY CONCERN

Manufacturer's declaration

We Contipro a.s. as a manufacturer of the product Hyiodine® hereby declare that the product is classified as a class III Medical Device in the Czech Republic. Therefore neither manufacturing licence, GMP certificate nor other separate certification is required for its manufacturing and placing of the product into the market.

Yours faithfully

Pavel Kušnierik
Regulatory affairs specialist
Contipro a.s.

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