

European Union Medical Device Regulation (EU MDR)

The European Union Medical Device Regulation (EU MDR) establishes rules about the presence of certain substances in medical devices for the purpose of improving the quality, safety and reliability of medical devices.

This letter certifies that HandyTube's stainless steel products do not contain any of the substances regulated under the requirements of the EU Medical Device Regulation.

We thank you for your business.

Respectfully,



Rebecca Reimer
Environmental, Health and Safety Manager

Date: March 2, 2023