Remanufactured Electrosurgical Devices

Reducing the environmental impact of surgery at CUH and CUMH



HSE South West theatres at CUH and CUMH are sending some of their single use surgical devices for remanufacturing after use.

Electrosurgical devices are remanufactured instead of being disposed as healthcare risk waste after a single use.

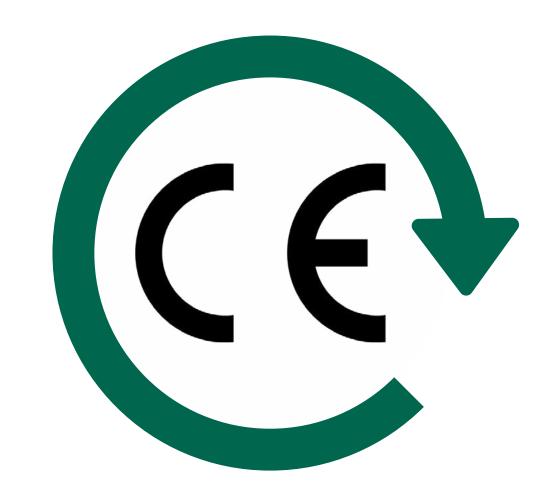




Remanufactured surgical devices are CE marked, denoting the same standard of functionality as new devices.

CUH and CUMH have been collecting single use electrosurgical devices for recycling for several years and some other public hospitals are also now setting up collections.

CUMH is working towards a purchase contract for remanufactured devices.





What is surgical device remanufacturing?

Remanufacturing of devices from original equipment manufacturers (OEMs) includes inspection; indelible marking; disinfection; cleaning; checks on functionality and cleanliness; packaging; and sterilisation. 100% function testing ensures the safety of the surgical device.

The CE Mark signifies that the manufacturer asserts that the electrosurgical device complies with all applicable EU health, safety, and environmental requirements in accordance with the EU Medical Devices Regulation 2017/745.

The remanufacturing process with a CE Mark ensures electrosurgical devices are fit for use once again with full functionality.







What are the benefits of recycling OEM devices & using remanufactured devices?

FINANCIAL BENEFITS

€1,200 per tonne reduced healthcare risk waste treatment costs



€3 - €8 procurement credit per recycled electrosurgical device, which can be used to purchase remanufactured devices

40-50% cost saving when purchasing a remanufactured device



ENVIRONMENTAL BENEFITS

33-50% lower carbon emissions with a remanufactured device compared to an OEM device



29% savings in resource use, mainly in plastic production and processing, and in packaging



Reduced carbon emissions from procurement of healthcare products and services





Is remanufacturing of devices permitted in Ireland?

Yes.

The EU Medical Devices Regulation 2017/745, enacted in 2021, permits the remanufacturing of 'single-use' devices where the reprocessor takes on full responsibility and liability as a new manufacturer i.e. CE marking.



Results to date



Over 600 surgical devices have been sent from the two hospitals for remanufacturing as of May 2025.



€2,800 procurement credithas been made to the
hospitals as payment
for devices sent for
remanufacturing.



A reduction in healthcare risk waste costs, with 130 kg of such waste being avoided.

Results to Date: HSE South West theatres are have saved...



140kg of carbon emissions savings from diverting used devices into remanufacturing vs sending them for incineration.



700 kg carbon emissions savings when these remanufactured surgical devices are reused.

Are remanufactured surgical devices on a HSE Procurement Framework?

Yes, remanufactured surgical devices are available to purchase from a HSE Procurement Framework (DPS 202164:1 DPS Qualification for Electro surgery Equipment and Neurology Products).

How does a Surgical Consultant use remanufactured devices?

Set up a used surgical device collection first, followed by purchasing of remanufactured devices.





- 2. Hospital theatre staff are trained in advance of collection.
- 3. Hospital theatre staff dispose of pre-cleaned used surgical devices in a dedicated collection box, including the device reference label.

4. Devices are collected by the remanufacturing company and exported to Germany for remanufacturing.

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Here taking climate action

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