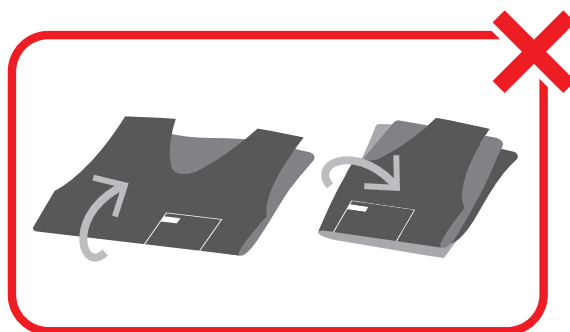
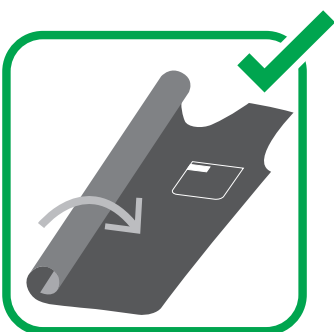
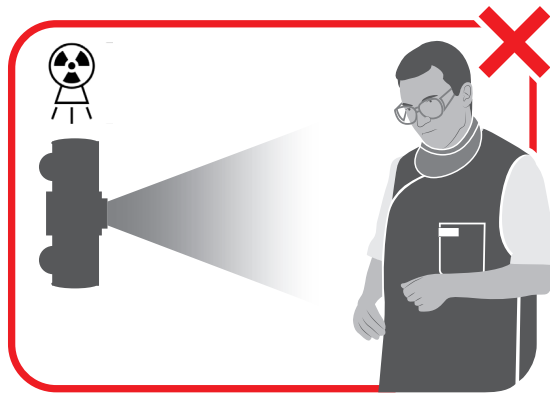
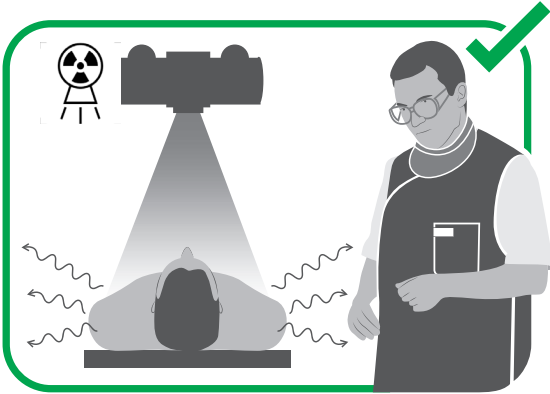


903, 905, 907, 909, 910, 911, 915, 917, 919,
921, 951, 953

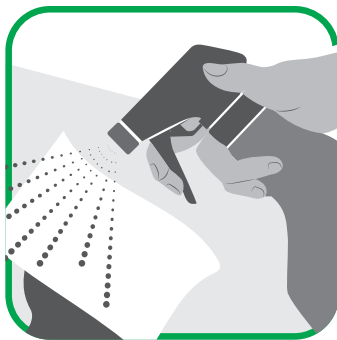
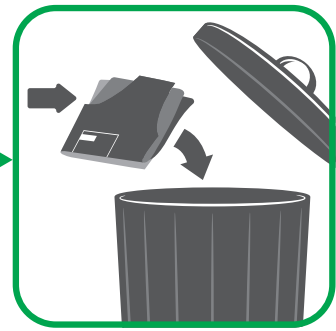
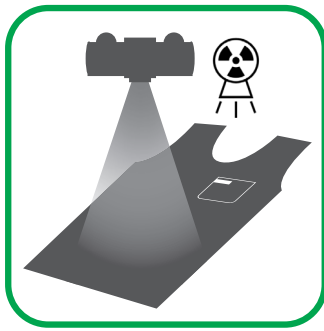


IP002





365



Incidin® OxyFoam S (pH 2.01 - 2.3, 100%)

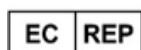


Kenex (Electro-Medical) Limited is certified as meeting the requirements of Regulation (EU) 2016/425 (PPE) for modules B & D related to design and manufacture of x-ray protective clothing. LytaType aprons are in conformity with state-of-the-art harmonised standards BS EN 61331-1:2014 relating to the x-ray attenuating properties of flexible materials used, and to BS EN 61331-3:2014 relating to x-ray protective clothing. The lead equivalent values are tested and certified for use within a 50 to 150 kV range of x-ray tube voltages

Kenex (Electro-Medical) Limited is also certified as meeting the requirements of BS EN ISO 13485:2016 related to medical devices quality management systems. Audited by SGS UK Limited.

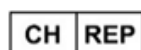
Notified body:
SGS Fimko OY, Takomotie 8, FI-00380 Helsinki, Finland.
Identification number: CE 0598.

UK Approved Body:
SGS (UK) Limited, Rossmore Business Park, Ellesmere Port, Cheshire,
CH65 3EN, UK
Identification number: UKCA 0120



EU authorised representative:

Emergo Europe, Westervoortsedijk 60, 6827 AT Arnhem,
The Netherlands.
Emergo Single Registration Number (SRN): NL-AR-000000116



Swiss authorised representative:

MedEnvoy Switzerland, Gotthardstrasse 28, 6302 Zug, Switzerland.

Kenex Single Registration Number (SRN): GB-MF-000009022



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