

Mouth ulcers and oral lesions are inflammatory forms which occur on the oral mucosa. These erythematous lesions can be painful and often make it difficult to chew or swallow, compromising the quality of life of the patient.

Indications:

Indicated for the treatment of aphthae, aphthous ulcers, irritations and mouth lesions due to oral surgery, braces and ill-fitting dentures. Iglu Ultra Clear gel works by forming a protective barrier over the mouth ulcer, adhering to the inside of the mouth providing protection, pain relief and promoting healing. Iglu Ultra Clear gel is alcohol free.

Instructions for use:

Apply 3-4 times a day directly on the lesion using the applicator. Avoid eating and drinking for at least 1 hour after application. Ready to use. Iglu Ultra Clear gel is safe if ingested. It can be used in adults and children.

Contraindications, warnings and precautions:

Do not use if the package is opened or damaged or after the expiry date. Do not swallow the product.

Do not exceed recommended dosage. Keep out of the sight and reach of children. No data is available for use in pregnancy and breast-feeding. No data related to concomitant use of drugs is available, consult the doctor. Avoid contact with eyes; in case of contact rinse with plenty of water. The expiry date refers to the product unopened and properly stored. Close the tube after use.

Store between 5°C and 30°C, away from direct heat.

In the event of problems during treatment interrupt the use and consult a doctor.

In case of any serious incident occurring in relation to Iglu Ultra Clear gel, it should be reported to the manufacturer and to local competent authority via: United Kingdom: Device Yellow Card Scheme, website: www.mhra.gov.uk/yellowcard.

Ingredients: Aqua, PVP (12%), xanthan gum (1%), taurine, zinc gluconate, PEG-40 hydrogenated castor oil, sodium saccharin, aroma.

Do not use Iglu Ultra Clear gel if you have a known allergy to any of the ingredients.



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UK Responsible Person: Diomed Developments Ltd, SG4 7QR, UK.



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'Iglu' is a Registered Trademark.

8 ml tube

LOT and Expiry Date are imprinted on the front and back of the tube.



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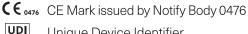
LOT

Batch number



Expiry date







Unique Device Identifier



Medical Device



See leaflet before using the product



Manufacturer

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