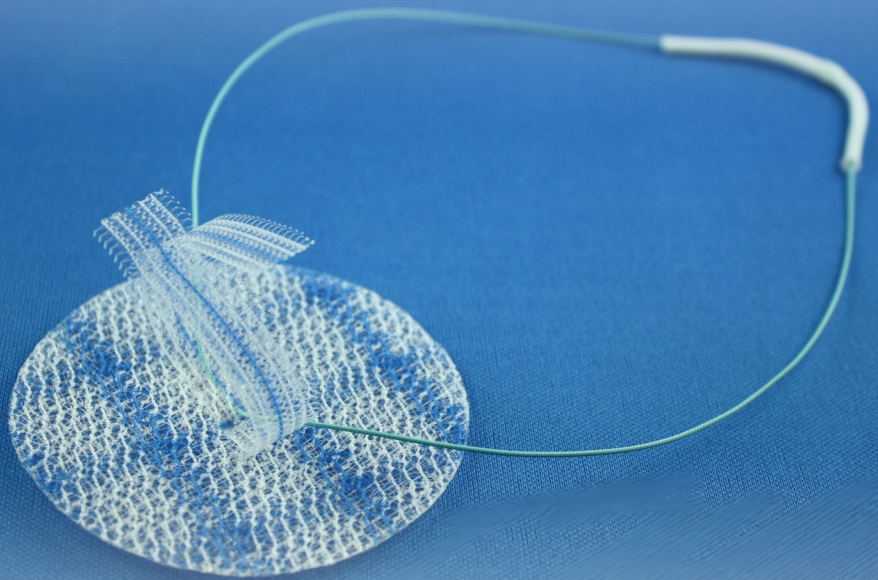


2P UCMC[®]



Umbilical ClearMesh Composite

*Treatment of abdominal wall defects
umbilical and epigastric hernias
small incisional hernias and
hernias caused by trocars*

dipromed
Dipro Medical Devices

2P-UCMC® Umbilical ClearMesh Composite - Protesi composita ombelicale non riassorbibile

2P-UCMC® è una protesi chirurgica, trasparente, leggera, ultra sottile, non riassorbibile, in polipropilene, composta da una rete in monofilamento macroporosa e da un film trasparente per minimizzare le aderenze.

Indicazioni:

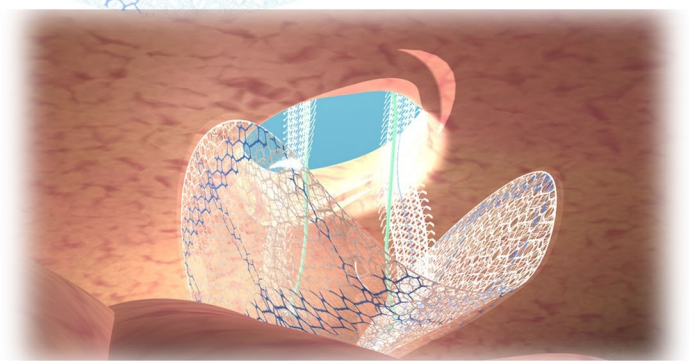
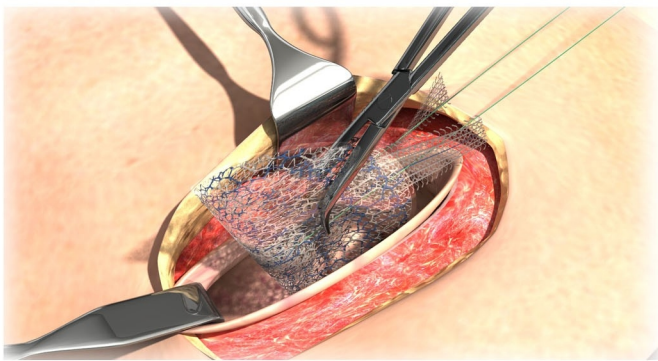
Trattamento dei difetti della parete addominale in particolare ernie ombelicali, epigastriche, piccoli laparoceli ed ernie da trocar.

2P-UCMC® Umbilical ClearMesh Composite - Umbilical composite nonabsorbable prosthesis

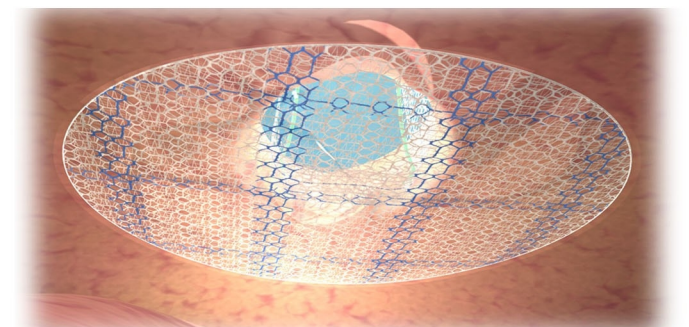
2P-UCMC® is a transparent, lightweight, ultra-thin, nonabsorbable surgical prosthesis in polypropylene, made by a monofilament macroporous mesh and a transparent film in order to minimize the adhesion formations.

Indications:

Treatment of abdominal wall defects, such as umbilical and epigastric hernias, small incisional hernias and hernias caused by trocars.



Codice - Code	Misura - Size	Pezzi/Cf. - Units/Box
UCMCI-045	Ø 4,5 cm	2
UCMCI-05	Ø 5 cm	
UCMCI-064	Ø 6,4 cm	
UCMCI-07	Ø 7 cm	
UCMCI-08	Ø 8 cm	



Possibili complicazioni includono sieroma, adesioni, ematomi, infiammazione, fistole ed estrusione, infezione, reazioni allergiche e recidive o difetti dei tessuti molli.
Possible complications include seroma, adhesion, hematomas, inflammation, extrusion, fistula formation, infection, allergic reaction and recurrence of the hernia or soft tissue defect.

DIPRO MEDICAL DEVICES S.r.l è fondata su un efficace sistema di gestione per la qualità, in grado di assicurare il mantenimento delle prestazioni dei propri prodotti e migliorare continuamente la qualità.

Il sistema qualità per la progettazione, produzione e distribuzione è conforme alla norma UNI CEI EN ISO 13485:2012. I dispositivi recano marcatura CE rilasciata ai sensi dell' Allegato II della Direttiva Europea 93/42 CEE.

DIPRO MEDICAL DEVICES S.r.l is established on an effective quality control system that assures a constant performance maintenance on the entire range of our products, as well as the constant endeavour of quality improvement.

The quality assurance system for the phases of planning, production and distribution is in correspondence with the regulation UNI CEI EN ISO 13485:2012. Our devices bear the CE seal of approval, issued under Enclosure II of European Directive 93/42 CEE.



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