



ABSOLUTE LENZUOLINO

MEDICAL ROLL 100% CELL. MT 150 H 50 1X6

Lenzuol. Carta

C61250



Technical Details

Raw Material	100% PURE CELLULOSE	Absorption by immers.%(+/- 10)	480	Nr.Sheets	395
Nr.Plies	2	Absorption Klemm mm/10(+/-5)	100	Roll Length (mt)	150,00
Grammage per ply (gr/mq)	16,5	Longit.Dry Tract. Res.N/m(+/-50)	470	Roll Diameter	16,00
Colour	WHITE	Transv.Dry Tract. Res.N/m(+/-50)	235	Core	RIGID
Whiteness Point	85	Longit.Wet Tract. Res.N/m(+/-50)	164	Core Diameter (mm)	39
Whitening Method	ECF	Transv.Wet Tract. Res.N/m(+/-50)	82	Rolls per Pack	1
WS Treatment	YES	Sheet Height (cm)	50,00	Packs per Carton	6
Embossing	MICRO EMBOSSED	Sheet Length (cm)	38,00	EAN Pack	8022650612503
Ply Bond veli	PUNTA PUNTA SYSTEM	Perforation	YES	EAN Carton	18022650612500



Logistic Details

Pack Dimensions (cm HxLxD)	--	Packaging Material	POLYETHYLENE	Cartons per Pallet	20
Carton Dimensions (cm HxLxD)	50,90x48,00x32,30	Carton Material	CARTON	Cartons per Layer	5
Pallet Dimensions (cm HxLxD)	218,60x120,00x80,30	Pallet Certification	EPAL	Layers per pallet	4
Carton Volume (mc)	0,0789	Pallet Construction	--	Demipallet	NO



This product meets EU GPP criteria, therefore it can be offered in public tenders organized by contracting authorities.



The technical data indicated above is generated by testing performed on articles belonging to our standard spectrum and are provided only for informational purposes. Additional information may be supplied upon request. All dimensions are subject to tolerance as by law enacted.

Montecarlo, 30/01/2026

DECLARATION OF CONFORMITY CE

Mr. Andrea Bernacchi, as C.E.O. of INDUSTRIE CELTEX S.p.A., with registered office in Via Traversa del del Marginone, 21/23- 55015 Montecarlo (LU), **SRN: IT-MF-000027968**, manufacturer of the Medical Device "LENZUOLINO MEDICO" (Disposable medical roll), **Basic UDI-DI: 8022650C60000ZN**, in the variant:

C61250 MEDICAL ROLL 100% CELL. MT 150 H 50 1X6

meant to be used to protect beds, stretchers, chairs or similar equipment during their use for medical activities,

DECLARES

Under its own responsibility, that the Medical Device referred to in object meets all the general requirements on safety and performance, applicable, of Annex I of Regulation EU 745/2017 on Medical Devices.

To this end, the company as represented above, warrants and declares the following:

- that the device in object complies with the applicable provisions of Regulation EU 2017/745 of European parliament and council;
- that the Device in object is sold in a NON STERILE form:
- that the Device in object belongs to Class I according to Regulation EU 745/2017, being a non-invasive device according to Rule 1 and not being applicable the other rules defined in ANNEX VIII, Chapter III, of that Regulation;
- that the Device complies with the general requirements on safety and performance and provisions of EU 745/2017 as per the current version of the Technical File;
- that the Device is manufactured in accordance with the Quality System that meets the requirements of Annex IX of the above mentioned Regulation.

The manufacturer also declares:

- to have established and maintain an appropriate procedure to ensure the after-sales surveillance required by Regulations EU 745/2017;
- to keep the technical documentation relating to the Medical Device at the disposal of the competent Authorities at their premises.

Applicable harmonised European standards

The list is provided in the relevant technical file Annex 3 to section 1.



INSTRUCTIONS FOR USE

Product meant to cover the examination couch or other dry surface where the patient may lie during a medical examination or outpatient treatment.

Ensures patient's contact with a clean, non-irritating surface and avoids the need for frequent equipment cleaning procedures.

Do not use in contact with exposed wounds or injured skin. Do not use in a sterile environment (clean room); contact with sterile equipment does not guarantee that the equipment remains sterile.

Do not reuse. In case of accidental contact with potentially harmful fluids, dispose it properly. The medical device is not sterile.

No sterilization or cleaning is provided for the medical device by the user.

The device is for single use only after which it has to be disposed according to local regulations. This medical device does not contain any toxic or dangerous materials.

INDUSTRIE CELTEX S.p.A.
C.E.O.
Andrea Bernacchi



Montecarlo, 30/01/2026

DECLARATION OF CONFORMITY
DERMATOLOGICALLY TESTED

We hereby declare that the following product:

C61250 MEDICAL ROLL 100% CELL. MT 150 H 50 1X6

passed a dermatological test on humans. The supervision of these tests and the evaluations were carried out by medical staff.

No toxic/irritative intolerance reactions carried out in the test and therefore the material type can be declared as dermatologically tested.

*INDUSTRIE CELTEX S.p.A.
QEHS Director
Manolo Papini*

Montecarlo, 30/01/2026

EU ECOLABEL DECLARATION

We hereby declare that the following product:

C61250 MEDICAL ROLL 100% CELL. MT 150 H 50 1X6

*complies with EU ECOLABEL certification.
Additionally, we will inform you if any changes regarding the above take place.*

License EU ECOLABEL: IT/004/010

INDUSTRIE CELTEX S.p.A.
QEHS Director
Manolo Papini



Montecarlo, 30/01/2026

PEFC DECLARATION

We hereby declare that the following product:

C61250 MEDICAL ROLL 100% CELL. MT 150 H 50 1X6

*is certified as PEFC product.
Additionally, we will inform you if any changes regarding the above take place.*

Logo-License PEFC: PEFC/18-32-24

*INDUSTRIE CELTEX S.p.A.
PEFC Responsible
Manolo Papini*

