



INSTRUCTIONS FOR USE ECG Gel: ContiGel ECG&EEG&DEFI

Conti Gel is a very good echo-transmission gel for ECG/EEG diagnostics and for defibrillator's use. Conti Gel is designed to reduce the resistance between skin and electrodes Conti Gel is watersoluble, odourless, easily removable. Conti Gel doesn't stain, does not grease. It does not contain any active principles that might damage the electrodes.

INSTRUCTIONS FOR USE

- Gel is ready to use
- Clean and degrease the patient skin.
- Apply the desired amount of gel
- After the procedure remove all residues of the gel with paper towel or gauze moistened with water.
- Stop using the gel for hypersensitive patients, if skin reactions occur.

WARNINGS

- External use only.
- The product is not sterile.
- Do not apply gel on skin affected by abrasions, erythema or wounds.
- Close the bottle after use.
- May cause irritations or reddenings with allergic persons.
- Contact with eyes may cause irritation.
- Analyst with sensitive skin should wear protective gloves during the use.
- Do not use the gel after the "use by" date on the bottle.
- If the package contains a labeled, empty refillable bottle, only the product with the name on the label should be filled

INGREDIENTS: aqua, carbomer, sodium hydroxide, sodium chloride, EDTA, phenoxyethanol, ethylhexylglycerin, E 129

PACKAGING

Carton with 50 bottles of 260g
 Carton with 24 bottles, of 500g
 Carton with 10 bottles of 1000g
 Carton with 4 cubitainers of 5000g
 Carton with 4 canisters of 5000g













DISPOSAL CONSIDERATION: Product, waste: Chemicals must be disposed of in compliance with the respective national regulations. Packaging must be disposed in compliance with the country-specific.

STORAGE: Temperature: +10°C ÷ +30°C - Relative humidity: 30% ÷ 75%. Keep container firmly closed, away from direct sunlight and sources of UV radiation.

EXPIRY:36 months from production date. Use only three month after opening.

NOTICE: any adverse event that has occurred in relation to the device should be reported to the manufacturer and any serious incident (MDR Article 2 (65)) should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established;

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	Manufacturer		Temperature limit		Batch-Code		Attention
	User Manual		polyethylene packaging		Use by date		Do not reuse
	no steril		away from UV		Medical device		

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