

ECG Electrodes

Instructions for use

These instructions for use are intended for the following Ambu ECG electrodes: BR, BRS, NEOX, NF, 40554, 40556, 40713

Warnings

- Surface electrodes should be applied only to clean, intact skin. Do not use on neonates, infants, or on patients with lesions, infected, or inflamed areas.
- Do not use on patients with a pacemaker or other cardiac device as cause contamination, leading to infections.
- After removal of electrodes from the protective liner, the protective liner should be disposed of in an appropriate waste bin.
- Except for 40713: Remove the electrode from the patient when performing MRI scans as this may cause skin burn on the patient.
- After removal of electrodes from the protective liner, the protective liner should be disposed of in an appropriate waste bin.
- Except for 40713: Remove the electrode from the patient when performing MRI scans as this may cause skin burn on the patient.

1.3. Potential Adverse Events

Potential adverse events in relation to the use of surface electrodes are skin reactions such as skin burn, skin redness, skin itching and small blisters.

2.0. Explanation of Symbols Used

Symbol for the Ambu ECG electrode	Indication	Symbol for the Ambu ECG electrode	Indication
	Medical Device		Solid gel
	Magnetic Resonance Conditional. Static magnetic field of 1.5 Tesla and 3 Tesla, with maximum spatial gradient of 25.300 G/cm or 253 T/m.		Cloth
	Radionuclide		Radiolucent
	Maximum whole body absorbed dose (WBA) specific absorption rate (SAR) of 2 W/kg (max. temp. increase 1.1°C (1.5 Tesla) and 2.3°C (3 Tesla)) or 4 W/kg (max. temp. increase 1.5°C (1.5 Tesla) and 4.6°C (3 Tesla)) for 15 minutes of scanning.		Country of manufacturer
	Electrical Safety Type BF Applied Part		Electrical Safety Type BF Applied Part

3.0. Directions for Use

- Washed, clean application site with a mild soap and water and dry thoroughly.
- The correct placement of electrodes is typically indicated by the manufacturer's or local protocol/procedure.
- Remove the electrode from the protective liner. Peel the sharp corners, and care should be taken when removing electrodes from the protective liner. Peel the electrode from the protective liner and place it on the skin. Dispose of the protective liner immediately after removal of the electrode.
- When using electrodes with connectors or pre-attached wires should be replaced if they are not stuck firmly to the skin.
- Get rid of cable as not removing paper towels.
- Electrodes in an opened pouch may dry out, therefore, do not open the electrode pouch.
- The electrodes are disposable and must be disposed of according to local hospital procedures.

3.1. Product Specific Directions for Use

NEOX removal: Gently roll the 4 leads of the electrode towards the center. Lift up the electrode with your finger between the adhesive and the patient's skin. It is recommended to leave the electrodes on until they fall off.

4.0. Duration of Application (time in hours)

Electrode	System	Color	Material	Neonatal	Indication	Duration of application
BR	Solid Gel	Ag/AgCl	Neonatal			48
BRS	Solid Gel	Ag/AgCl	Neonatal			24
NEOX	Solid Gel	Silicone	Neonatal			72
NF	Solid Gel	Silicone	Neonatal			48
40554	Solid Gel	Ag/AgCl	Neonatal			24
40556	Solid Gel	Ag/AgCl	Neonatal			24
40713	Solid Gel	Ag/AgCl	Pediatric			48

Kasutusjuhend EKG elektroodid

1.0. Kasutusotused

Elektroodid asetatakse keha pinnale, et edastada elektrilisi signaale keha pinnaalt südamiselle, mis jälgimisandmesse, mis kasutatakse elektrokardiogrammi eelvõtkoridogrammi.

1.1. Vastandüstused

1.2. Hoiatused ja ettevaatusabinõud

- Nahaelektroodide tüübi kindlaksmääritamiseks, kasutage kaitseliini, mis on ette nähtud selleks, et vältida naha ärritust, infektsiooni või põletikku.
- Ärge kasutage elektroode, millel on nähtavaid kahjustusi, mis võivad põhjustada kontakti katkestusi või teha elektroodid ebasobivaks kasutamiseks.
- Kui vahelduvvoolu, eriti voolu, mis on ette nähtud selleks, et vältida naha ärritust, infektsiooni või põletikku.
- USA föderaalsete tootmis- ja tervishooldusametite (FDA) soovitusel ei tohiks elektroode kasutada alla kolme süstemaatilise, mis ulatub ECG 60001-1 niivõrd.

1.3. Võimalikud kõrvaltoimed

Kui teie aadame kasutamise ajal teie nahal tekivad kõrvaltoimed (naha punetus, naha sügelus ja väikesed vaha), peate elektroode kohe eemaldama ja võtma ühendust oma arstiga.

1.4. Üldised märkused

Kui teie aadame kasutamise ajal teie nahal tekivad kõrvaltoimed (naha punetus, naha sügelus ja väikesed vaha), peate elektroode kohe eemaldama ja võtma ühendust oma arstiga.

2.0. Kasutatud sümbolite selgitused

Symbol for the Ambu ECG electrode	Indication	Symbol for the Ambu ECG electrode	Indication
	Medical Device		Tahke geel
	MRI Compatible. Static magnetic field of 1.5 Tesla and 3 Tesla, with maximum spatial gradient of 25.300 G/cm or 253 T/m.		Rii
	Radionuclide		Radionuclaan
	Maximum whole body absorbed dose (WBA) specific absorption rate (SAR) of 2 W/kg (max. temp. increase 1.1°C (1.5 Tesla) and 2.3°C (3 Tesla)) or 4 W/kg (max. temp. increase 1.5°C (1.5 Tesla) and 4.6°C (3 Tesla)) for 15 minutes of scanning.		Tootja riik
	Electrical Safety Type BF Applied Part		Elektrilise turvata tüüpi BF rakendatud osa

3.0. Kasutusjuhised

- Väljastatud puhastage ja kuivatage hoolikalt enne elektroodide kasutamist. Ärge kasutage elektroode, millel on nähtavaid kahjustusi, mis võivad põhjustada kontakti katkestusi või teha elektroode ebasobivaks kasutamiseks.
- Plastiktahtide eemaldamiseks kasutage kaitseliini, mis on ette nähtud selleks, et vältida naha ärritust, infektsiooni või põletikku.
- Kui vahelduvvoolu, eriti voolu, mis on ette nähtud selleks, et vältida naha ärritust, infektsiooni või põletikku.
- USA föderaalsete tootmis- ja tervishooldusametite (FDA) soovitusel ei tohiks elektroode kasutada alla kolme süstemaatilise, mis ulatub ECG 60001-1 niivõrd.

3.1. Tootespetsiifilised kasutusjuhised

NEOX eemaldamine: Kergelt rullige 4 juhtviiba elektroodi nähale ja suruge kergelt selle servadele. Eemaldage lüüdi 2 suruga kergelt selle servade külalt 4 juhtviiba. Eemaldage lüüdi 2 suruga kergelt selle servade külalt 4 juhtviiba.

4.0. Kasutamiseaeg (vältus tundides)

Electrode	System	Color	Material	Neonatal	Indication	Duration of application
BR	Solid Gel	Ag/AgCl	Neonatal			48
BRS	Solid Gel	Ag/AgCl	Neonatal			24
NEOX	Solid Gel	Silicone	Neonatal			72
NF	Solid Gel	Silicone	Neonatal			48
40554	Solid Gel	Ag/AgCl	Neonatal			24
40556	Solid Gel	Ag/AgCl	Neonatal			24
40713	Solid Gel	Ag/AgCl	Pediatric			48

Инструкции за употреба

1.0. Предназначение

- Прикрепяне на електрода на кожата на пациента, за да предаде електрокардиален сигнал от повърхността на тялото до устройството, което обработва сигнала и генерира електрокардиограма или векторкардиограма.
- Предупреждения и предпазни мерки
- След отстраняване на защитния покривач на електрода, защитният покривач трябва да се изхвърли в съответно предвиден контейнер за отпадъци.
- След като е извършено изпитане, електрода трябва да се използва само с водеща кабелна система съгласно с МРТ.

1.3. Потенциални неблагоприятни събития

Potential adverse events in relation to the use of surface electrodes are skin reactions such as skin burn, skin redness, skin itching and small blisters.

2.0. Обяснение на използваните символи

Symbol for the Ambu ECG electrode	Indication	Symbol for the Ambu ECG electrode	Indication
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3.0. Directions for Use

- Washed, clean application site with a mild soap and water and dry thoroughly.
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1.1. Противопоказания

1.2. Предупреждения и предпазни мерки

- След отстраняване на защитния покривач на електрода, защитният покривач трябва да се изхвърли в съответно предвиден контейнер за отпадъци.
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