



Ref. Certif. No.
SE-56495

IEC SYSTEM FOR MUTUAL RECOGNITION OF TEST CERTIFICATES FOR ELECTRICAL EQUIPMENT (IECEE) CB SCHEME

SYSTEME CEI D'ACCEPTATION MUTUELLE DE CERTIFICATS D'ESSAIS DES EQUIPEMENTS ELECTRIQUES (IECEE) METHODE OC

CB TEST CERTIFICATE

CERTIFICAT D'ESSAI OC

Product Produit	Fetal heart monitor
Name and address of the applicant Nom et adresse du demandeur	Neoventa Medical AB Ågatan 32 SE-431 35 Mölndal, SWEDEN
Name and address of the manufacturer Nom et adresse du fabricant	Same as applicant
Name and address of the factory Nom et adresse de l'usine	Same as applicant
Ratings and principal characteristics Valeurs nominales et caractéristiques principales	100-240VAC, 50/60Hz (IEC/EN 60601-1), 100-120VAC, 50/60Hz (UL 60601-1), 90VA, Class I, Type: B, BF, CF (for additional information see page 2), IPX1
Trademark (if any) Marque de fabrique (si elle existe)	STAN [®]
Model / Type Ref. Ref. De type	STAN S31
Additional information (if necessary) Les informations complémentaires (si nécessaire)	See page 2
A sample of the product was tested and found to be in conformity with Un échantillon de ce produit a été essayé et a été considéré conforme à la	IEC 60601-1:1988 and A1+A2 IEC 60601-1-8:2003 IEC 60601-2-30:1999 IEC 60601-2-49:2001
As shown in the Test Report Ref. No. which forms part of this Certificate Comme indiqué dans le Rapport d'essais numéro de référence qui constitue partie de ce Certificat	807713 and 800413

This CB Test Certificate is issued by the National Certification Body
Ce Certificat d'essai OC est établi par l'Organisme National de Certification

Intertek Semko AB
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Signature:
Mikael Goffhé

Date: 3 December 2008

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Additional information (if necessary)
Information complémentaire (si nécessaire)

Type B on US1/US2 ultrasound transducer and event marker. Type BF on NIBP cuff and SpO2 sensor. Type CF on FECG and TOCO/IUP transducer.

The product is also in conformity with EN 60601-1:1990 with A1 + A2.

The product is also in conformity with EN 60601-1-8:2004 with A1.

The product is also in conformity with EN 60601-2-30:2000.

The product is also in conformity with EN 60601-2-49:2001.

The product is also in conformity with EN ISO 9919:2005.

Biocompatibility according to EN ISO 10993 has not been evaluated.

Software development according to IEC/EN 60601-1-4 has not been evaluated.

Date: 3 December 2008

Signature: 

TB