GLAMOX LUXO

Declaration of Conformity

(Directive 93/42/EEC as amended by directive 2007/47/EE)

Manufacturer:	Glamox ASA
Adress:	Industrifeltet 2260 Kirkenær Norway
Product:	Table standing luminaire for fluorescent lamp
Product name:	**LFM Medical**

We declare under sole responsibility that above listed products confirms with the standards listed.

Reference	*Date of issue*	*Name*
IEC-60601-1	1988	General requirements (lighting)
IEC 60601-2-41	2000	Particular requirements for the safety of surgical luminaires and luminaires for diagnosis
EN 60601-1-2	2001	Electromagnetic compability – Requirments and tests.

The CE mark was affixed to the product:99.

Place and date: Kirkenær, 27 Aug. 2012

Name and signature of
autorized persons:

Lars Andersson

Lars Andersson Wehlin
Factory Manager

Doc.No: PME-305-0003 Rev:0

CERTIFICATE

No. P06206023

Order No. 55446

Applicant	Luxo Norge AS P.O.Box 54 2261 KIRKENÆR NORWAY
Manufacturer	Luxo Norge AS P.O.Box 54 2261 KIRKENÆR NORWAY
Factory	Luxo Norge AS P.O.Box 54 2261 KIRKENÆR NORWAY
Group	Fluorescent lamp luminaries, for medical use
Model/type	LFM Medical
Data	95VA 230VAC 50Hz
Other specification	Cl. I Type BF
The above product is certified according to the following standard(s)	Safety std.: EN 60601-1:1990 + A1:1993 + A2:1995 + A13:1996 + Corrigendum (June 1995), including possible listed national conditions/deviations for Norway. EMC Std: EN 60601-1-2:2001
Validity	This certificate documents conformity with the standards shown, and also applies as license for use of Nemko's name and certification mark. The certificate and license is valid as long as the applicable conditions are complied with, and provided that any changes to the product are notified to Nemko for acceptance prior to implementation. New standards or amendments to the standards may imply that the product design must be updated and/or that re-testing and re-certification is necessary.
Additional Information:	The above certified equipment complies with current technical statutory requirements in Norway regarding safety and Electromagnetic Compatibility (EMC). It should be noted that by marketing of equipment subject to official registration in Norway, the party responsible for the marketing shall be registered by the Norwegian Directorate for Product- and Electrical Safety.

Date of issue 10 March 2006

Hanne Yndestad

signature

Hanne Yndestad

Certification Department

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Enterprise number:
NO 974404532

IEC IECEE CB SCHEME

Ref. Certif. No.

NO37610

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

Product Produit	Fluorescent lamp luminaries, for medical use
Name and address of the applicant Nom et adresse du demandeur	Luxo Norge AS P.O.Box 54 2261 KIRKENÆR NORWAY
Name and address of the manufacturer Nom et adresse du fabricant	Luxo Norge AS P.O.Box 54 2261 KIRKENÆR NORWAY
Name and address of the factory Nom et adresse de l'usine Note *When more than one factory, please report on page 2* Note *Lorsque il y plus d'une usine, veuillez utiliser la 2ème page*	Luxo Norge AS P.O.Box 54 2261 KIRKENÆR NORWAY ☐ Additional Information on page 2
Ratings and principal characteristics Valeurs nominales et caractéristiques principales	95VA 230VAC 50Hz, Cl. I Type B applied part
Trademark (if any) Marque de fabrique (si elle existe)	LUXO
Model / Type Ref. Ref. De type	LFM Medical
Additional information (if necessary may also be reported on page 2) Les informations complémentaires (si nécessaire,, peuvent être indiqués sur la 2ème page	☐ Additional Information on page 2
	PUBLICATION
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 + A1:1991 + A2:1995
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	55446

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

Nemko

P.O. BOX 73, BLINDERN
N-0314 OSLO, NORWAY

Hanne Yndestad

Date: 2006-03-10

Signature: Hanne Yndestad
Certification Department