



# *Declaration of Conformity*

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

Manufacturer: Glamox ASA

Address: Industriefeltet  
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.

<i>Reference</i>	<i>Date of issue</i>	<i>Name</i>
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  
authorized persons:

A handwritten signature in black ink that reads "Lars Andersson Wehlin".

Lars Andersson Wehlin  
Factory Manager

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

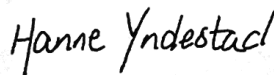
**Validity**

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



signature

Hanne Yndestad  
Certification Department

**IEC****IECEE  
CB  
SCHEME**

Ref. Certif. No.

**NO37610**IEC SYSTEM FOR MUTUAL RECOGNITION OF TEST  
CERTIFICATES FOR ELECTRICAL EQUIPMENT (IECEE)  
CB SCHEMESYSTEME 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 demandeurLuxo Norge AS  
P.O.Box 54  
2261 KIRKENÆR  
NORWAYName and address of the manufacturer  
Nom et adresse du fabricantLuxo Norge AS  
P.O.Box 54  
2261 KIRKENÆR  
NORWAYName and address of the factory  
Nom et adresse de l'usineLuxo Norge AS  
P.O.Box 54  
2261 KIRKENÆR  
NORWAYNote: When more than one factory, please report on page 2  
Note: Lorsque il y a plus d'une usine, veuillez utiliser la 2<sup>ème</sup> page Additional Information on page 2Ratings 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 2<sup>ème</sup> 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**N Nemko**P.O. BOX 73, BLINDERN  
N-0314 OSLO, NORWAY*Hanne Yndestad*

Date: 2006-03-10

Signature: Hanne Yndestad  
Certification Department