

MET LABORATORIES, INC. CLASSIFICATION RECORD



The applicant named below has been authorized by MET Laboratories, Inc. to represent the product(s) listed in this record as "MET Classified" and to mark this/these product(s) according to the terms and conditions of the MET Mark Utilization Agreement, MET Listing Reports, and the applicable marking agreements. Only the product(s) bearing the MET Mark and under a follow-up service are considered to be included in the MET Classification program.

FILE NUMBER: E212295

APPROVAL DATE: June 19, 2006

REVISED: April 4, 2007

PRODUCT	MODEL(S)	ELECTRICAL RATINGS
Medical Beds	1030018XXX, 1030017XXX, 1030019XXX, 1030100XXX, & 1030101XXX	115V, 60Hz 230watt
	1040026XXX, 1040027XXX, 1040028XXX, 1040120XXX, 1040121XXX, 1040023XXX, 1040024XXX, 1040025XXX, 1040029XXX, 1040030XXX, 1040031XXX, 1040220XXX, 1040221XXX, 1040032XXX, 1040033XXX, 1040034XXX, 1040035XXX, 1040036XXX, 1040037XXX, 1040038XXX, 1040039XXX & 1040040XXX	120V, 50/60Hz 4.0A

STANDARD NUMBER	STANDARD TITLE	EDITION
UL60601-1	Medical Electrical Equipment	1st
CSA C22.2 No. 601-1	Medical Electrical Equipment	M90-2001

MET LABORATORIES, INC. requires that any and all changes proposed in the previously identified product(s), that affects the information contained in the above referenced listing report, must be submitted to MET for evaluation prior to implementation to assure continued MET Classification status.

The above identified product(s) has/have been submitted by the applicant:

APPLICANT:

NOA Medical Industries Inc.
801 Terry Lane
Washington, MO 63090

The covered products shall be subjected to quarterly follow-up inspections to ensure that the Classified product(s) are identical to the representative product sample evaluated by MET LABORATORIES, INC. and that all manufacturer's responsibilities are being fulfilled as specified in the MANUFACTURING RESPONSIBILITY section of the Certification report.

CLASSIFICATION CONDITIONS: *

- The products were not evaluated to comply with clause 52.1 (Programmable Electronic Systems) and clause 48 (Biocompatibility).
- This device has been classified by the manufacturer as a medical bed (medical device). This device has no patient connections and is not known to connect to patient equipment.
- The physiological effects of the bed's use have not been investigated by MET Laboratories Inc.
- Any accessories requested or referenced for use with this medical bed will need to comply with the Standards noted above to maintain level of Safety anticipated for Medical Devices.

Rick Cooper
Director of Laboratory Operations,
Safety Laboratory



*MET Laboratories, Inc. is accredited by OSHA and the Standards Council of Canada.
The Nation's First Nationally Recognized Testing Laboratory*

