
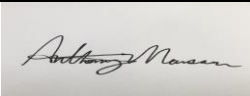





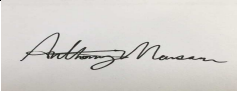


Test Report issued under the responsibility of:



IEC 60601-1 Medical electrical equipment Part 1: General requirements for basic safety and essential performance	
Report Reference No.....:	4786488108-5
Date of issue	2014-08-14 (Amendment 2: 2017-06-07)
Total number of pages.....:	144
CB Testing Laboratory.....:	UL Fremont
Address	47173 Benicia Street, Fremont, CA 94538 USA
Applicant's name.....:	XP Power LLC
Address	15641 Red Hill Ave., Suite 100 Tustin, CA 92780 USA
Test specification:	
Standard	IEC 60601-1: 2005 + CORR. 1:2006 + CORR. 2:2007 + AM1:2012 (or IEC 60601-1: 2012 reprint)
Test procedure.....:	CB Scheme
Non-standard test method.....:	N/A
Test Report Form No.....:	IEC60601_1J
Test Report Form Originator	UL(US)
Master TRF.....:	2014-07
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General disclaimer: The test results presented in this report relate only to the object tested. This report shall not be reproduced, except in full, without the written approval of the Issuing CB testing laboratory. The authenticity of this Test Report and its contents can be verified by contacting the NCB, responsible for this Test Report.	

Test item description	Component Switching Power Supply	
Trade Mark		
Manufacturer	XP Power LLC 15641 Red Hill Ave., Suite 100, Tustin, CA 92780 USA	
Model/Type reference.....	AHM100PSXXYY-ZZ (where XX is any number between 12-48 designating output voltage and YY can be blank or "C2", -ZZ can be "-A", "-6", "-6A", "8", "-8A", or blank), AHM100PS24 XD0112A	
Ratings.....	Input Rated: 100-240 Vac, 50/60 Hz, 1.2 A Output Rated: See Model Differences for details.	
Testing procedure and testing location:		
<input checked="" type="checkbox"/>	CB Testing Laboratory:	
Testing location/ address		UL Fremont 47173 Benicia Street, Fremont, CA 94538 USA
<input type="checkbox"/>	Associated CB Testing Laboratory:	
Testing location/ address		
Tested by (name + signature).....		Anthony Moussa/Project Handler 
Approved by (name + signature)		Haydee Gonzalez/Project Reviewer 
<input type="checkbox"/>	Testing procedure: TMP/CTF Stage 1:	
Testing location/ address		
Tested by (name + signature).....		
Approved by (name + signature)		
<input type="checkbox"/>	Testing procedure: WMT/CTF Stage 2:	
Testing location/ address		
Tested by (name + signature).....		
Witnessed by (name + signature)		
Approved by (name + signature)		
<input checked="" type="checkbox"/>	Testing procedure:	
<input type="checkbox"/>	SMT/CTF Stage 3 or 4:	
Testing location/ address		XP Power LLC, 15641 Red Hill Ave., Suite 100 Tustin, CA 92780 USA

Tested by (name + signature).....	Ron Nabong/Tester	
Witnessed by (name + signature)		
Approved by (name + signature)	Haydee Gonzalez/Project Reviewer	
Supervised by (name + signature).....	Anthony Moussa/Project handler	

List of Attachments (including a total number of pages in each attachment):

Enclosures (4 pages)

National Differences (9 pages) - See original report

Test Report IEC 60601-1-6 (5 pages) - See original report

Test Report IEC 62366 (16 pages) - See original report

Summary of testing: Unless otherwise indicated, all original tests were conducted at XP POWER LLC, 15641 Red Hill Ave., Suite 100, Tustin, CA 92780 USA

All testing conducted under the Applicant's IEC 60601-1, 3rd Ed investigation issued under CBTR No. 11CA41872, CBTC No. US-18097-UL was consider to cover the requirements of IEC 60601-1, Edition 3 with Am. 1

Tests performed (name of test and test clause):

Testing location:

Power Input Test (4.11)

Durability of Markings (7.1.3)

Voltage or Charge Limitation (8.4.3)

Working Voltage Measurement (8.5.4)

Dielectric Voltage Withstand (8.8.3)

Ball Pressure (8.8.4.1)

Temperature Test (11)

Abnormal Operation and Single Fault Conditions (13)

Push (Rigidity) (15.3.2)

Ball Impact (15.3.3)

Drop Impact (15.3.4)

Mold Stress Relief (15.3.6)

Transformer Overload and Short-Circuit Tests (15.5.1)

Leakage Current Test (8.7)

Ingress (11.6.5), Dielectric Voltage Withstand (8.8.3), Earth Leakage (8.7)

Humidity Preconditioning Treatment (5.7)

UL Camas

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UL Camas

UL Camas

UL Camas

UL Camas

UL Camas

UL CamasUL Fremont

**XP Power 15641 Red Hill
Ave., Suite 100 Tustin, CA
92780 USA**

RISK MANAGEMENT FILE Review (4.2)

Summary of compliance with National Differences

List of countries addressed: Austria, Canada, Republic of Korea, Sweden, UK, USA

☒ The product fulfils the requirements of IEC 60601-1, Edition 3 with Am 1

Copy of marking plate

The artwork below may be only a draft. The use of certification marks on a product must be authorized by the respective NCBs that own these marks.

Multiple files attached to Enclosure - Marking Plate ID 13-01 and 13-02

Labels provided are considered representative of the entire series.

GENERAL INFORMATION	
Test item particulars (see also Clause 6):	
Classification of installation and use	External Transportable
Device type (component/sub-assembly/ equipment/ system):	Component
Intended use (Including type of patient, application location) :	Component switching power supply
Mode of operation	Continuous
Supply connection	Appliance coupler
Accessories and detachable parts included.....	None
Other options include	None
Testing	
Date of receipt of test item(s)	2011-07-06, 2010-03-22, 2010-02-03, 2010-02-02, 2017-04-06, 2017-05-11
Dates tests performed	2011-10-04 to 2011-10-18, 2010-06-23, 2010-03-08 to 2010-03-23, 2010-02-02 to 2010-06-08, 2017-04-06, 2017-04-13, 2017-05-24
Possible test case verdicts:	
- test case does not apply to the test object	N/A
- test object does meet the requirement.....	Pass (P)
- test object was not evaluated for the requirement	N/E (collateral standards only)
- test object does not meet the requirement.....	Fail (F)
Abbreviations used in the report:	
- normal condition	N.C.
- single fault condition.....	S.F.C.
- means of Operator protection	MOOP
- means of Patient protection	MOPP
General remarks:	
<p>Before starting to use the TRF please read carefully the 4 instructions pages at the end of the report on how to complete the new version "J" of TRF for IEC for 60601-1 3rd edition with Amendment 1.</p> <p>"(See Attachment #)" refers to additional information appended to the report. "(See appended table)" refers to a table appended to the report. The tests results presented in this report relate only to the object tested. This report shall not be reproduced except in full without the written approval of the testing laboratory. List of test equipment must be kept on file and available for review. Additional test data and/or information provided in the attachments to this report.</p> <p>Throughout this report a <input type="checkbox"/> comma / <input checked="" type="checkbox"/> point is used as the decimal separator.</p>	

Manufacturer's Declaration per sub-clause 4.2.5 of IEC60061-1

The application for obtaining a CB Test Certificate includes more than one factory location and a declaration from the Manufacturer stating that the sample(s) submitted for evaluation is (are) representative of the products from each factory has been provided..... :

☒ Yes☐ Not applicable

When differences exist; they shall be identified in the General product information section.

Name and address of factory (ies)..... :

XP POWER LLC
990 BENECIA AVE
SUNNYVALE CA 94085
UNITED STATES

XP POWER (KUNSHAN) LTD
230 BIN JIANG NAN RD
ZHANGPU TOWN
KUNSHAN
JIANGSU 215300 CHINA

General product information:

Products covered are external power supplies intended to be used with Medical Electrical Equipment. Units may be either Class I or Class II. Double insulated symbol is provided on Class II units.

Earthing symbol may only be provided for Class I power supplies.

Model Differences:

All models in the Model AHM100PSXXYY-ZZ series are identical with exception to the Mains Transformer, T2, and minor secondary components that allow for different output voltage ratings per the output voltage range noted below. See Table below for Model Ratings at 40°C:

Model AHM100PS12: 10.1-13.5 Vdc, 8.33 A max. (100W max.)
 Model AHM100PS15: 13.5-17.0 Vdc, 6.67 A max. (100W max.)
 Model AHM100PS19: 17.1-21.0 Vdc, 5.26 A max. (100W max.)
 Model AHM100PS24: 21.0-26.0 Vdc, 4.16 A max. (100W max.)
 Model AHM100PS28: 26.1-31 Vdc, 3.57 A max. (100W max.)
 Model AHM100PS33: 31.1-33 Vdc, 3.03 A max. (100W max.)
 Model AHM100PS36: 33.1-42 Vdc, 2.78 A max. (100W max.)
 Model AHM100PS48: 42.1-54.0 Vdc, 2.08 A max. (100W max.)

See Enclosure - Miscellaneous for de-rated output values for higher ambient.

Models may have an additional YY identifier which can be blank or "C2" to designate a Class II configuration.

Models may have an additional -ZZ identifier which can be "-A", "-6", "-6A", "-8", "-8A", or blank to designate the type of input connector:

blank = C14 style input connector (Class I construction) or C18 input connector (Class II construction);
 "A" = C14 style input connector with optional IEC cable retention;
 "6" = C6 style input connector (Class I or Class II construction);
 "6A" = C6 style input connector with optional IEC cable retention;
 "8" = C8 style input connector (Class I or Class II construction)
 "8A" = C8 style input connector with optional IEC cable retention.

Model AHM100PS24 XD0112A is identical to Model AHM100PS24 with exception to the addition of an alternate input connector.

Model AHM100PS24, with additional A or B in the Part Number are suitable for non-continuous operation and have the following rated output and duty cycle. Refer to the Enclosure 13-03 and 13-04 for marking labels:

- Model AHM100PS24 with A in Part Number: 21.0-26.0 Vdc, 6.05Apk @ 24% duty(on time <1.3s), Average Power 95W
- Model AHM100PS24 with B in Part Number: 21.0-26.0 Vdc, 7Apk, 13% Duty, (<0.3s)

Additional Information:

The schematics are kept on file at the CBTL and can be provided by the manufacturer upon request by NCB's/CBTL's.

When submitting this Test Report to other Certification Body, the manufacturer is responsible for providing any additional information that the Body may need in order to issue its Mark, including testing for compliance with the applicable collateral standards.

The nameplate markings provided are considered representative of the entire series.

Multiple Location Manufacturer Codes:

"K" XP Power, Jiangsu, China

"FS" XP Power, Sunnyvale, CA

No additional testing was deemed necessary to evaluate the models covered under this Report to IEC 60601-1, Edition 3 with Am.1 based on previously conducted testing and the review of product technical documentation including photos, schematics, wiring diagrams, etc. conducted under separate CB Scheme investigation to IEC 60601-1, 3rd ed issued under CBTR No. 11CA41872, CBTC No. US-18097-UL.

Amendment 1:

Report was amended to add non-continuous Models AHM100PS24, with additional A or B in the Part Number and to evaluate the models under this report to the collateral standard IEC 60601-1-6. No tests were deemed necessary. Address for XP Power LLC was updated from Suite 150, 1241 E Dyer Road Santa Ana, CA 92705 USA to 15641 Red Hill Ave., Suite 100, Tustin, CA 92780 USA

Amendment 2:

This report was amended to update the IPXX rating of all models to IP22. Representative testing was conducted on models AHM100PS24 and AHM100PS12-1. Humidity testing was conducted at XP Power LLC, 15641 Red Hill Avenue Suite 100, Tustin CA, while Earth leakage and dielectric test were conducted at UL Fremont 47173 Benicia Street, Fremont, CA 94538 USA. The Collateral report to IEC 60601-1-6 was not affected by this change. This report is meant to be read in conjunction with CBTR Ref. No: 4786488408-5, CB Test Certificate Ref. No. US-23769-A1-UL.

When submitting this Test Report to other Certification Body, the manufacturer is responsible for providing any additional information that the Body may need in order to issue its Mark, including testing for compliance with the applicable collateral standards.

Technical Considerations:

- The product was investigated to the following additional standards:: ANSI/AAMI ES60601-1 (2005 + C1:09 + A2:10)+AM1 (2012) (Medical Electrical Equipment - Part 1: General Requirements for Basic Safety and Essential Performance) (includes Deviations for United States), CAN/CSA-C22.2 No. 60601-1 (2008) +AM1 (2014) (Medical Electrical Equipment - Part 1: General Requirements for Basic Safety and Essential Performance) (includes National Differences for Canada), EN 60601-1:2006 / A1:2013 / A12:2014 (Medical electrical equipment Part 1: General requirements for basic safety and essential performance), IEC 60601-1, Edition 3 with Am. 1 (2012), IEC 60601-1-6 (General requirements for safety - Collateral Standard: Usability)
- The product was not investigated to the following standards or clauses:: Biocompatibility (ISO 10993-1), Clause 14, Programmable Electronic Systems, Electromagnetic Compatibility (IEC 60601-1-2)
- The degree of protection against harmful ingress of water is: IPX2
- The mode of operation is:: Continuous
- The product is suitable for use in the presence of a flammable anesthetics mixture with air or oxygen or with nitrous oxide:: No
- Scope of Power Supply evaluation defers the following clauses to the be determined as part of the end product: Clause 7.5 (Safety Signs), Clause 7.9 (Accompanying Documents), Clause 9 (ME Hazard), Clause 10 (Radiation), Clause 14 (PEMS), Clause 16 (ME Systems)
- Scope of Power supply evaluation excludes the following:
 - Patient applied parts clauses: 4.6, 7.2.10, 8.3, 8.5.2, 8.5.5, 8.7.4.7-8.7.4.9, 8.9.1.15
 - Battery related clauses: 7.3.3, 15.4.3
 - Hand Control related clauses: 8.10.4
 - Oxygen related clauses: 11.2.2
 - The following fluids related clauses: 11.6.2, 11.6.3, 11.6.4, and 11.6.6
 - Sterilization clause: 11.6.7
 - Biocompatibility Clause: 11.7 (ISO 10993)
 - Motor related clauses: 13.2.13.3, 13.4
 - Heating Elements related clause: 13.2

Flammable Anaesthetic Mixtures Protection: Annex G

- The product is evaluated only to the following hazards: Casualty, Fire, Shock
- Software is relied upon for meeting safety requirements related to mechanical, fire and shock: No
- Unit also complied with spacing requirements of UL60601-1 (1st), CSA C22.2 No. 60601-1 (2nd), and IEC 60601-1 (2nd) for Basic for 250 Vac from Primary to Ground, Double/Reinforced for 250Vac from Primary to Secondary, and Supplementary for 250 Vac from Secondary to Earth.

Risk Controls/ Engineering Condition of Acceptability

- The component shall be provided in compliance with the Marking (clause 7) and Separation (clause 8) requirements of the end use application.
- The power supply was evaluated for use in 40°C ambient at Full Rated Output and 60% of the Rated Output in 60°C ambient. (See De-rating Curve, Enclosure 7-01 for details)
- Repeating leakage current testing should be considered in the end product application.
- This power supply was evaluated with Two MOPP between Primary and Secondary; One MOPP primary and Earth/Secondary Reference Conductor; and One MOPP between Secondary and Earth/Secondary Reference Conductor.
- This power supply has been evaluated as a continuous operation, ordinary equipment and has not been evaluated for use in the presence of a flammable anesthetic mixture with air, oxygen, or nitrous oxide. The output circuits have not been evaluated for direct patient connection (Type B, BF or CF).
- The end product should ensure that the requirements related to accompanying documents, clause 7.9, are met.
- The available voltage for the secondary outputs does not exceed 25 Vac or 60 Vdc, under normal and single fault conditions.
- The output connectors are not acceptable for field connections; they are only intended for connection to mating connectors of the end-use machine.
- The Electric Strength Test conducted on this power supply was based upon a maximum working voltage of: Primary-Earthed Dead Metal (Class I units): 430 Vpk, 240 Vrms; Primary-SEC: 430 Vpk, 240 Vrms.
- The following magnetic devices (e.g. transformers or inductor) are provided with an OBJY2 insulation system with the indicated rating greater than Class A (105°C): L1-L4 and T2 are Class B (130°C).
- Cleaning test to be considered as part of end product evaluation.
- The need for Marking Durability and Marking Legibility Testing to be considered as part of the end product installation.
- Power cord suitable for the application to be provided as part of the end product evaluation.