

COVER PAGE FOR TEST REPORT

Product Category:	Power Supplies, Medical and Dental
Product Category CCN:	QQHM2, QQHM8
Test Procedure:	Component Recognition
Product:	AC-DC Adaptor
Model/Type Reference:	<p>BP(a)030(b)(c)(e)(f) and (a)ENB1030(b)(c)(d)(e)(f)</p> <p>(a) can be A to Z for family related designs. (b) can be S for single output in model BP(a)010 series and (b) can be A to Z for design revision changes in model (a)ENB1010 series. (c) can be 05, 06, 07, 09, 12, 15, 16, 18, 24 and 48 for output voltage. (d) can be can be 00 thru 99 for standards output cord options ("d)" is not provided in model BP(a)010series). (e) can be F or N or Q or B or H or G or M or C for input plug type. F-Class I appliance inlet type: IEC60320-C14 Q-Class II appliance inlet type: IEC60320-C18 N-Class II appliance inlet type: IEC60320-C8 B or C-Class I & Class II direct-plug-in for North America, China, Japan and Argentina (Changeable Direct-plug-in type is only used for Class II) H-Class I & Class II direct-plug-in for Australia (AS/NZS 3112) G-Class I & Class II direct-plug-in for British (BS 1364) M-Class I & Class II direct-plug-in for European (CEE /16)] & Korea. (f) can be 00 thru 99 for customer options.</p> <p>PENB1030(a)(b)(c)(d)(e)</p> <p>(a) can be A to Z for family related designs. (b) can be output voltages, may be 48. (c) can be can be 00 thru 99 for standards output cord options (d) can be F or N or Q or B or H or G or M or C for input plug type. F-Class I appliance inlet type: IEC60320-C14 Q-Class II appliance inlet type: IEC60320-C18 N-Class II appliance inlet type: IEC60320-C8 B or C-Class I & Class II direct-plug-in for North America, China, Japan and Argentina (Changeable Direct-plug-in type is only used for Class II) H-Class I & Class II direct-plug-in for Australia (AS/NZS 3112) G-Class I & Class II direct-plug-in for British (BS 1364) M-Class I & Class II direct-plug-in for European (CEE /16)] & Korea. (e) can be 00 thru 99 or AA to ZZ for customer options.</p>
Rating(s):	<p>Rated Input; 100-240Vac, 50/60Hz, 1.0 A</p> <p>Rated Output;</p> <p>5Vdc, 4.0A or 7.5Vdc, 3.0A or 9Vdc, 3.0A or 12Vdc, 2.5A or 14Vdc, 2.1A or 15Vdc, 2.0A or 18Vdc, 1.67A or 24Vdc, 1.33A or 48Vdc, 0.4A or 48Vdc, 0.67A (Rated output voltage is designated according to the model name designation system).</p>
Standards:	<p>UL 60601-1, 1st Edition, 2006-04-26 (Medical Electrical Equipment, Part 1: General Requirements for Safety)</p> <p>CAN/CSA-C22.2 No. 601.1-M90, 2005 (Medical Electrical Equipment - Part 1: General Requirements for Safety)</p>

Applicant Name and Address:	BRIDGEPOWER CORP 964 GOSAEK-DONG GWONSEON-GU SUWON-SI GYEONGGI-DO 441-813 KOREA
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This Report includes the following parts, in addition to this cover page:

1. Specific Inspection Criteria
2. Specific Technical Criteria
3. Clause Verdicts
4. Critical Components
5. Test Results
6. National Differences
7. Enclosures

This is to certify that representative samples of the products covered by this Test Report have been investigated in accordance with the above referenced Standards. The products have been found to comply with the requirements covering the category and the products are judged to be eligible for Follow-Up Service under the indicated Test Procedure. The manufacturer is authorized to use the UL Mark on such products which comply with this Test Report and any other applicable requirements of Underwriters Laboratories Inc. ('UL') in accordance with the Follow-Up Service Agreement. Only those products which properly bear the UL Mark are considered as being covered by UL's Follow-Up Service under the indicated Test Procedure.

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