### **Features**

- Medical grade DC/DC converter
- 250VAC working, 2MOPP
- Unregulated **Converter**
- 5.2kVDC isolation for 250VAC working voltage
- -40°C up to +90°C operating temperature
- 3rd Ed. safety and 4th. Ed. EMC

### Description

The REM1 complements the existing board-mount REM3, REM6 and REM10 series by offering a 1W medical grade unregulated DC/DC converter in a more compact SIP7 package. The REM1 features reinforced 5.2kVDC/1 minute isolation and 2MOPP/250VAC working voltage. The REM1 is available with 3.3, 5, 12, 15 or 24V inputs and offers 3.3, 5 or 12V outputs with up to 85% efficiency. The operating temperature range is -40°C up to +90°C. The converter is Class B EMC and 60601-1-2 (4th Ed.) medical EMC certified using a simple external LC filter. The converters are fully certified to CB, IEC/EN and ANSI/AAMI 60601 third edition safety standards, RoHS2+ (10/10) and REACH and come with a 5 year warranty.

#### **Selection Guide** Part nom. Input Output Output Efficiency Max. Capacitive typ. (1) Number Voltage Voltage Current Load (2) [VDC] [VDC] [mA] [µF] [%] REM1-3.33.3S 3.3 3.3 303 78 2200 REM1-3.305S 3.3 5 2200 200 81 REM1-3.312S 3.3 12 84 85 470 REM1-053.3S 5 3.3 303 79 2200 5 REM1-0505S 5 200 80 2200 5 12 REM1-0512S 84 84 470 3.3 303 2200 REM1-123.3S 12 78 12 5 200 81 REM1-1205S 2200 REM1-153.3S 15 3.3 303 77 2200 15 5 REM1-1505S 200 81 2200 REM1-243.3S 24 3.3 303 76 2200 REM1-2405S 24 5 200 80 2200



### REM1





IEC/EN60601-1 certified ANSI/AAMI ES60601-1 certified EN62368-1 certified IEC/EN60601-1-2 EN55011 **CB** Report

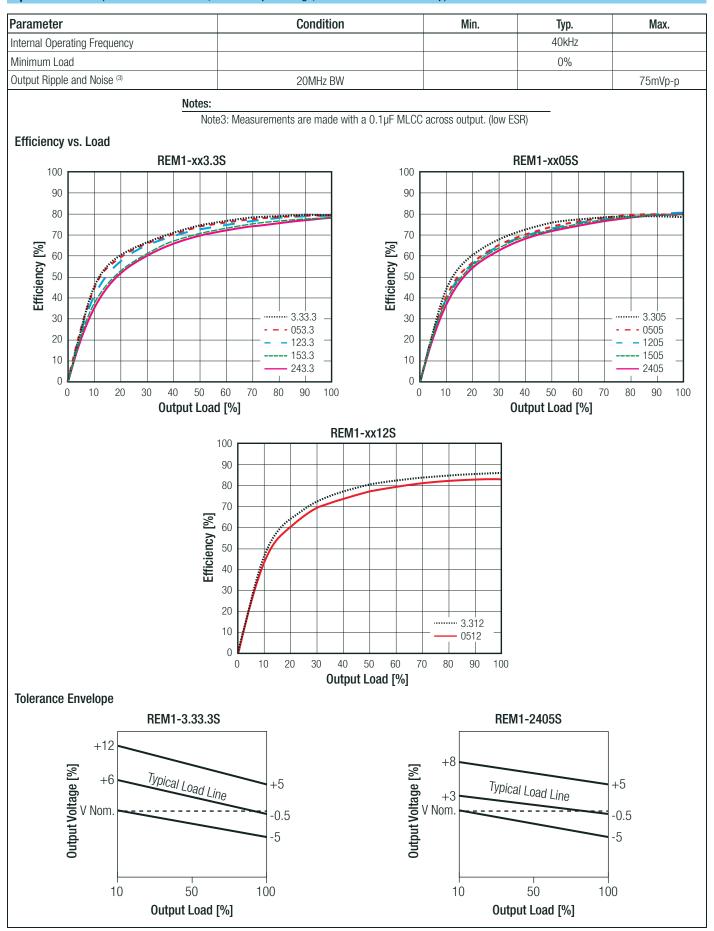
#### Notes:

Note1: Efficiency tested by nominal input and full load at +25°C ambient Note2: Max Cap Load tested by nominal input and full resisitive load

BASIC CHARACTERISTICS				
Parameter	Condition	Min.	Тур.	Max.
Internal Input Filter				capacito
Input Voltage Range			±10%	
Maximum Reverse Voltage				OVDC
Quiescent Current	3.3VDC			40mA
	5VDC			25mA
	nom. Vin= 12VDC			12mA
	15VDc			10mA
	24VDC			7mA

## REM1 Series

#### Specifications (measured @ Ta= 25°C, nominal input voltage, full load and after warm-up)

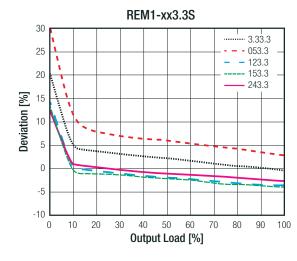


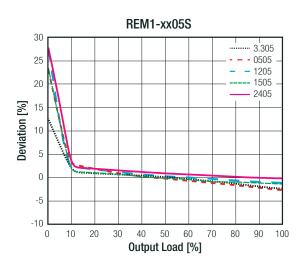
# REM1 Series

#### Specifications (measured @ Ta= 25°C, nominal input voltage, full load and after warm-up)

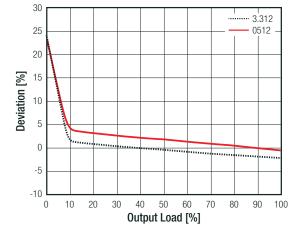
REGULATIONS			
Parameter	Cond	dition	Value
Output Accuracy			±5% max.
Line Regulation	low line to hig	h line, full load	±1.2% typ. @ 1% of Vin
Load Regulation	10% to 100% load	3.3Vout and 5Vout 12Vout	8% typ. / 12% max. 5% typ. / 8% max.

#### Deviation vs. Load





#### REM1-xx12S



Parameter		Туре	Value
raiametei		Туре	
Isolation Voltage (4)	I/P to O/P	tested for 1 minute	5.2kVDC
	1/1 10 0/1		4kVAC
Isolation Resistance			10GΩ min.
Isolation Capacitance			25pF typ.
Insulation Grade			reinforced
Means of Protection	250VAC working voltage		2MOPP
Medical Device Classification			built-in power supply
Creepage and Clearance			≥8mm

Note4: For repeat Hi-Pot testing, reduce the time and/or the test voltage

Note5: Refer to local safety regulations if input over-current protection is also required. Recommended fuse: T1A slow blow type

## REM1 Series

#### Specifications (measured @ Ta= 25°C, nominal input voltage, full load and after warm-up)

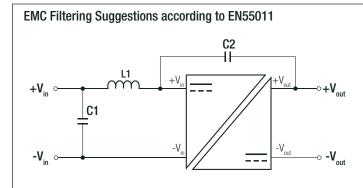
ENVIRONMENTAL		
Parameter	Condition	Value
Operating Temperature Range	without derating (see graph)	-40°C °C to +85°C
Maximum Case Temperature		105°C
Temperature Coefficient		0.02%/K typ.
Operating Altitude	according to IEC/EN60601-1 accroding to IEC/EN62368-1	3000m 5000m
Operating Humidity	non-condensing	5% - 95% RH max.
Pollution Degree		PD2
MTBF	according to MIL-HDBK-217F, G.B. +25°C +85°C	18200 x 10 <sup>3</sup> hours 7500 x 10 <sup>3</sup> hours
Vibration		according to MIL-STD-202G standard
Derating Graph (@ chamber and natural convection 0.1m/s)	100 100 100 100 100 100 100 100	
	Ambient Temperature [°C]	

SAFETY AND CERTIFICATIONS		
Certificate Type (Safety)	Report / File Number	Standard
Audio/video, information and communication technology equipment. Safety requirements (CB Scheme)	L0339m31-A-L	EN62368-1:2014
Medical Electric Equipment, General Requirements for Saftey and Essential Performance	E314885-D1000-1/A0/C0-UL	ANSI/AAMI ES60601-1:2005/®2012 + A1:2012 + C1:2009/®2012 + A2:2010/®2012 CSA C22.2 No. 60601-1:14, 3rd Edition, 2014
Medical Electric Equipment, General Requirements for Safety and Essential Performance	E314885-D1000-1/A0/C0-CB	IEC60601-1:2005 +AM1:2012 EN60601-1:2006 + A12:2014
EAC	RU-AT.49.09571	TP TC 004/2011 TP TC 004/2011
RoHS2+		RoHS-2011/65/EU + AM-2015/863
EMC Compliance	Condition	Standard / Criterion
Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility		IEC60601-1-2:2014 EN60601-1-2:2015
Industrial, scientific and medical equipment - Radio frequency disturbance characteristics - Limits and methods of measurement	with external filter	EN55011, 2009+A1:2010, Class B
ESD Electrostatic discharge immunity test	Air: ±15kV; Contact: ±8kV	IEC61000-4-2:2008, Criteria A
Radiated, radio-frequency, electromagnetic field immunity test	3V/m	IEC61000-4-3:2006+A2:2010, Criteria A
Fast Transient and Burst Immunity	DC Port: ±2kV	IEC61000-4-4:2012, Criteria A
Surge Immunity	DC Port: ±1kV	IEC61000-4-5:2014, Criteria B
Immunity to conducted disturbances, induced by radio-frequency fields	DC Port: 6V	IEC61000-4-6:2013, Criteria A
Power Magnetic Field Immunity	50Hz, 30A/m	IEC61000-4-8:2009, Criteria A
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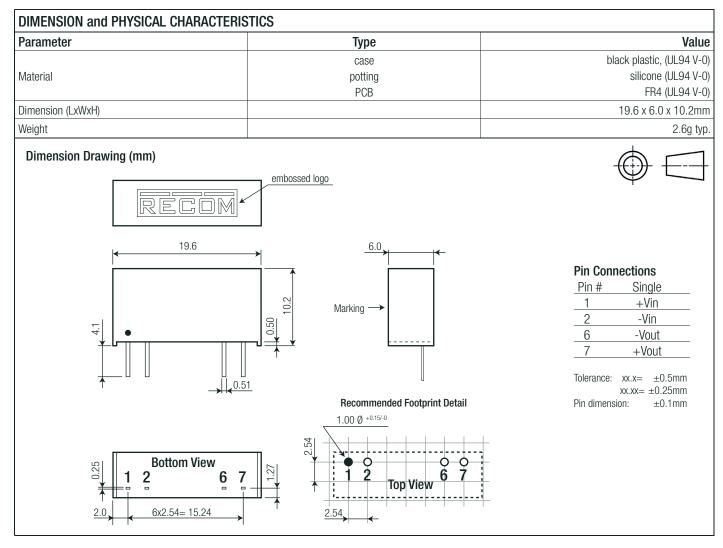
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# REM1 Series

#### Specifications (measured @ Ta= 25°C, nominal input voltage, full load and after warm-up)



Component Lis	t Class B		
Input Voltage	C1	C2	L1
3.3VDC			
5VDC	4 7.5	470pF/6kVDC	22µH Choke
12VDC	4.7µF		
15VDC			
24VDC	2.2µF	-	47µH Choke



PACKAGING INFORMATION			
Parameter	Туре	Value	
Packaging Dimension (LxWxH)	tube	520.0 x 16.0 x 9.3mm	
Packaging Quantity		25pcs	
Storage Temperature Range		-55°C to +125°C	
Storage Humidtiy	non-condensing	95% RH max.	

The product information and specifications may be subject to changes even without prior written notice. The product has been designed for various applications; its suitability lies in the responsibility of each customer. The products are not authorized for use in safety-critical applications without RECOM's explicit written consent. A safety-critical application is an application where a failure may reasonably be expected to endanger or cause loss of life, inflict bodily harm or damage property. The applicant shall indemnify and hold harmless RECOM, its affiliated companies and its representatives against any damage claims in connection with the unauthorized use of RECOM products in such safety-critical applications.