AC Filter



		See below: Approvals and Co	ompliances		
Description - Line filter in standard and medical version - 1-stage - standard attenuation		Characteristics - Designed for standard applications - Protection against interference voltage from the mains Possible interferences generated in the equipment are strongly attenu- ated - Universal line filter for standard applications - Suitable for use in equipment according to IEC/UL 62368-1			
		Other versions or - Version with wire c - Version with varisto			
		References	now applications the type EMAR NEO		
		We recommend for new applications the type FMAB NEO Weblinks pdf data sheet, html datasheet, General Product Information, Approvals, Distributor-Stock-Check, Detailed request for product, Microsite			
Technical Data					
Ratings IEC	1 - 10A @ Ta 40 °C / 250 VAC; 50 Hz	Line Filter	Standard and Medical Version, IEC		
Ratings UL/CSA	1 - 10A @ Ta 40 °C / 125 VAC; 60 Hz		60939, UL 1283, CSA C22.2 no. 8		
Leakage Current	standard < 0.25 mA (250 V / 60 Hz)		Technical Details		
	medical (M5) < 5 μ A resp. (M80) < 80	MTBF	> 200'000 h acc. to MIL-HB-217 F		
Dielectric Strength	μΑ (250 V / 60 Hz) 1.7 kVDC between L-N				
Dielectric Strength	2.7 kVDC between L/N-PE				
	Test voltage (2 sec)				
Allowable Operation Tempe- rature	-25 °C to 85 °C				
Climatic Category	25/100/21 acc. to IEC 60068-1				

 Climatic Category
 25/100/21 acc. to IEC 60088-1

 Protection Class
 Suitable for appliances with protection class I acc. to IEC 61140

 Terminal
 Quick connect terminals 6.3 x 0.8 mm

 Material
 Nickel plated steel

Approvals and Compliances

Detailed information on product approvals, code requirements, usage instructions and detailed test conditions can be looked up in Details about Approvals

SCHURTER products are designed for use in industrial environments. They have approvals from independent testing bodies according to national and international standards. Products with specific characteristics and requirements such as required in the automotive sector according to IATF 16949, medical technology according to ISO 13485 or in the aerospace industry can be offered exclusively with customer-specific, individual agreements by SCHURTER.

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Approvals

The approval mark is used by the testing authorities to certify compliance with the safety requirements placed on electronic products. Approval Reference Type: FMW2

Approval Logo	Certificates	Certification Body	Description
10	VDE Approvals	VDE	Certificate Number: 40004673
c FL [°] us	UL Approvals	UL	UR File Number: E72928

Product standards

Product standards that are referenced

Organization	Design	Standard	Description				
IEC	Designed according to	IEC 60939	Passive filters for suppressing electromagnetic interference				
(UL)	Designed according to	UL 1283	Electromagnetic interference filters				
CSA Group	Designed according to	CSA C22.2 no. 8	Electromagnetic interference (EMI) filters				

Application standards

Application standards where the product can be used

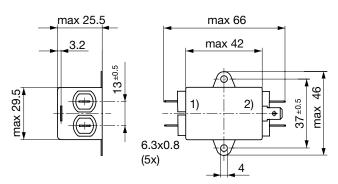
Organization	Design	Standard	Description
IEC.	Suitable for applications acc.	IEC/UL 62368-1	Audio/video, information and communication technology equipment - Part 1: Safety requirements
IEC	Suitable for applications acc.	IEC 60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance

Compliances

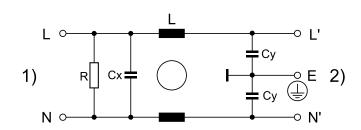
The product complies with following Guide Lines

Identification	Details	Initiator	Description				
CE	CE declaration of conformity	SCHURTER AG	The CE marking declares that the product complies with the applicable requirements laid down in the harmonisation of Community legislation on its affixing in accordance with EU Regulation 765/2008.				
UK CA	UKCA declaration of conformity	SCHURTER AG	The UKCA marking declares that the product complies with the applicable requirements laid down in the British Amendment of Regulation (EC) 765/2008.				
RoHS	RoHS	SCHURTER AG	Directive RoHS 2011/65/EU, Amendment (EU) 2015/863				
©	China RoHS	SCHURTER AG	The law SJ / T 11363-2006 (China RoHS) has been in force since 1 March 2007. It is similar to the EU directive RoHS.				
REACH	REACH	SCHURTER AG	On 1 June 2007, Regulation (EC) No 1907/2006 on the Registration, Evaluation, Authorization and Restriction of Chemicals 1 (abbreviated as "REACH") entered into force.				
T	Medical Equipment	SCHURTER AG	Suitable for use in medical equipment according to IEC/UL 60601-1 (1 MOOP, 1 MOPP)				

Dimension [mm]

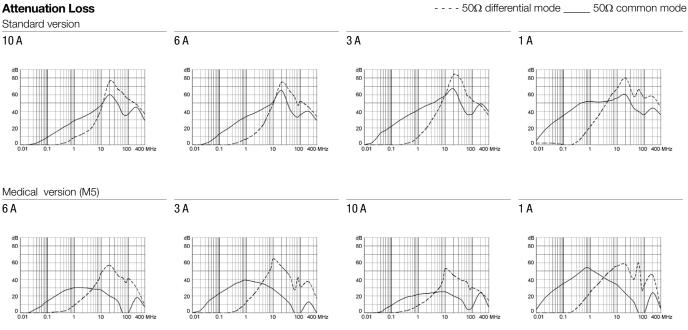


Diagrams



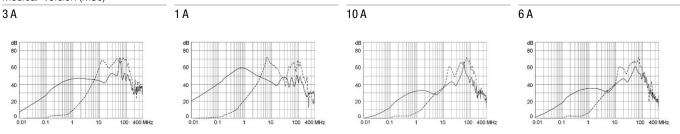
R and Cy according to version, see variants 1) Line, 2) Load

Attenuation Loss



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Medical version (M80)



All Variants

Rated Current [A]	Filter-Type	Leakage Cur- rent [mA]	L [mH]	Cx [nF]	Cy [nF]	R [M Ω]	Weight [g]	Order Number
1	Standard version	0.25	2 x 10	15	2.2	-	55 g	5500.2039
3	Standard version	0.25	2 x 2	15	2.2	-	55 g	5500.2040
6	Standard version	0.25	2 x 0.8	15	2.2	-	55 g	5500.2041
10	Standard version	0.25	2 x 0.4	15	2.2	-	55 g	5500.2042
1	Medical Version (M5)	0.005	2 x 10	15	-	1	55 g	5500.2098
3	Medical Version (M5)	0.005	2 x 2	15	-	1	55 g	5500.2104
6	Medical Version (M5)	0.005	2 x 0.8	15	-	1	55 g	5500.2100
10	Medical Version (M5)	0.005	2 x 0.4	15	-	1	55 g	5500.2102
1	Medical Version (M80)	0.08	2 x 10	15	0.47	1	55 g	5500.2099
3	Medical Version (M80)	0.08	2 x 2	15	0.47	1	55 g	5500.2105
6	Medical Version (M80)	0.08	2 x 0.8	15	0.47	1	55 g	5500.2101
10	Medical Version (M80)	0.08	2 x 0.4	15	0.47	1	55 g	5500.2103

Most Popular.

Availability for all products can be searched real-time:https://www.schurter.com/en/Stock-Check/Stock-Check-SCHURTER

Packaging unit

25 Pcs