

date 11/15/2011

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MODEL: CVS-1708 | DESCRIPTION: SPEAKER

FEATURES

- · plastic frame
- · mylar cone
- · plastic case





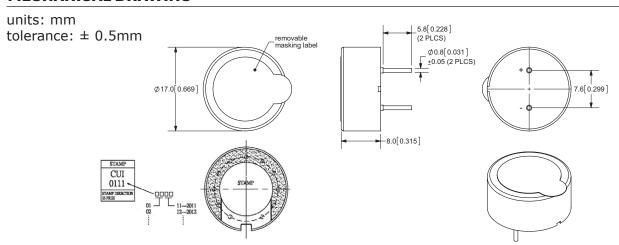
SPECIFICATIONS

parameter	conditions/description	min	typ	max	units
diameter			17		mm
depth			8		mm
input power	max. power: IEC-60268-5, filter 60 s on / 120 s off, 10 cycles at room temp		0.2	0.5	W
impedance	at 2 kHz, 1 V	6.8	8	9.2	Ω
resonant frequency	at 1 V	1,200	1,500	1,800	Hz
sound pressure level	0.2 W, 10 cm ave. at 2.0, 2.5, 3.0, 4.0 kHz 1 W, 1 m ave. at 2.0, 2.5, 3.0, 4.0 kHz	84 71	87 74	90 77	dB dB
response				20,000	Hz
distortion	at 2 kHz, 0.2 W			10	%
buzz, rattle, etc.	must be normal at sine wave 1.26 V				
magnet size	ø6.5 x 1.5 mm				
operating temperature		-20		55	°C
weight			2.0		g
material	PBT +20% glass (black)				
washable	yes				
RoHS	yes		·		

SOLDERABILITY

parameter	conditions/description
hand soldering	350 ±5°C for 5 seconds

MECHANICAL DRAWING



2K

20K

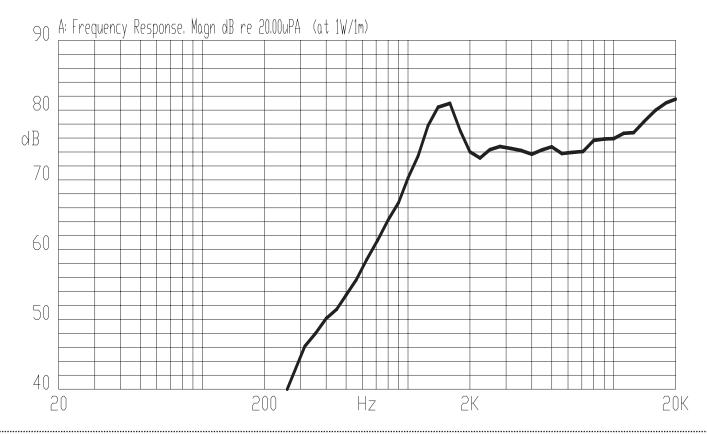
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CUI Inc | MODEL: CVS-1708 | DESCRIPTION: SPEAKER

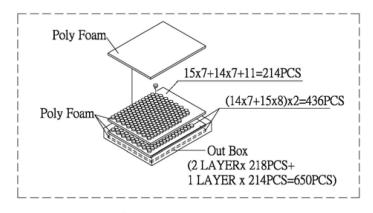
AF Frequency Response. Magn dB re 2000uPA (at 0.2W/10cm) 90 AB 80 70 60

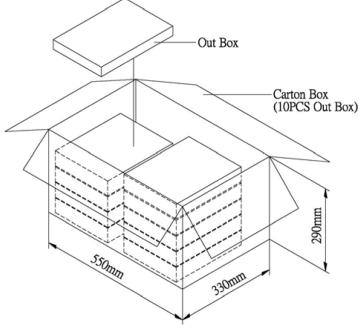
Hz

200



PACKAGING





Out Box 310mmx248mmx49mm 1x650PCS=650PCS 550mmx330mmx290mm Carton Box 650PCSx10=6,500PCS

REVISION HISTORY

rev.	description	date
1.0	initial release	11/15/2011

The revision history provided is for informational purposes only and is believed to be accurate.



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CUI offers a one (1) year limited warranty. Complete warranty information is listed on our website.

CUI reserves the right to make changes to the product at any time without notice. Information provided by CUI is believed to be accurate and reliable. However, no responsibility is assumed by CUI for its use, nor for any infringements of patents or other rights of third parties which may result from its use.

CUI products are not authorized or warranted for use as critical components in equipment that requires an extremely high level of reliability. A critical component is any component of a life support device or system whose failure to perform can be reasonably expected to cause the failure of the life support device or system, or to affect its safety or effectiveness.