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CUI DEVICES

date 02/21/2020

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MODEL: CLS0362MA-1 | DESCRIPTION: SPEAKER

FEATURES

- 36 mm
- round frame
- 0.5 W
- 8 Ω
- neodymium magnet
- mylar cone





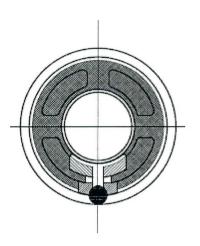
SPECIFICATIONS

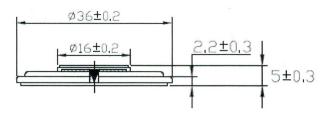
parameter	conditions/description	min	typ	max	units
input power			0.5	1.0	W
impedance	at 1.0 kHz, 1.0 V	6.8	8	9.2	Ω
resonant frequency (Fo)	at 1.0 V	416	520	624	Hz
frequency response		Fo		15,000	Hz
sound pressure level	at 1.0 W, 1 m, avg. at 0.8, 1.0, 1.2, 1.5 kHz	89	92	95	dB
distortion	at 2.0 kHz, 0.5 W			10	%
buzz, rattle, etc.	must be normal at sine wave			2	V
dimensions	Ø36 x 5.0				mm
magnet	Nd-Fe-B				
cone material	mylar				
terminal	solder pads				
weight			8.8		g
operating temperature		-25		60	°C
hand soldering	for maximum 3 seconds	340	350	360	°C
RoHS	yes				

Notes: 1. All specifications measured at 15~35°C, humidity at 25~75%, under 86~106 kPa pressure, unless otherwise noted.

MECHANICAL DRAWING

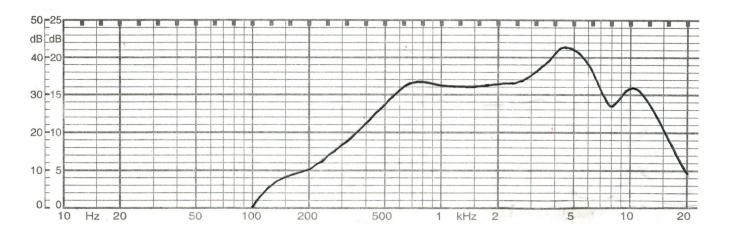
units: mm tolerance: ±0.2 mm





FREQUENCY RESPONSE CURVE

parameter	conditions/description
potentiometer range	50 dB
rectifier	RMS
lower limit frequency	20 Hz
wr. speed	100 mm/sec
zero level	60 dB



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REVISION HISTORY

rev.	description	date
1.0	initial release	07/20/2007
1.01	brand update	02/21/2020

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

CUI DEVICES

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

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

CUI Devices 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.