Additional Resources: Product Page | 3D Model

# **CUI** DEVICES

**date** 03/23/2020

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MODEL: GF0661M | DESCRIPTION: SPEAKER

#### **FEATURES**

- 66 mm
- round frame
- 0.5 W
- 8 Ω
- ferrite magnet
- mylar cone





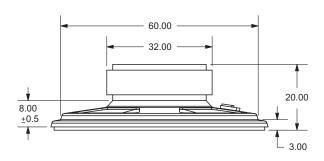
#### **SPECIFICATIONS**

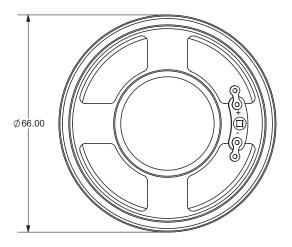
parameter	conditions/description	min	typ	max	units
input power			0.5	1.0	W
impedance	at 0.8 kHz, 1.0 V	6.8	8	9.2	Ω
resonant frequency (Fo)	at 1.0 V	344	430	516	Hz
frequency response		Fo		10,000	Hz
sound pressure level	at 0.5 W, 50 cm, avg at 1.0, 1.4, 1.7, 2.0 kHz	85	88	91	dB
buzz, rattle, etc.	must be normal at sine wave			2.0	V
dimensions	Ø66 x 20				mm
magnet	ferrite				
cone material	mylar				
terminal	solder eyelets				
weight			57.0		g
operating temperature		-25		60	°C
hand soldering	for maximum 3 seconds	360	380	400	°C
RoHS	yes				

### **MECHANICAL DRAWING**

units: mm

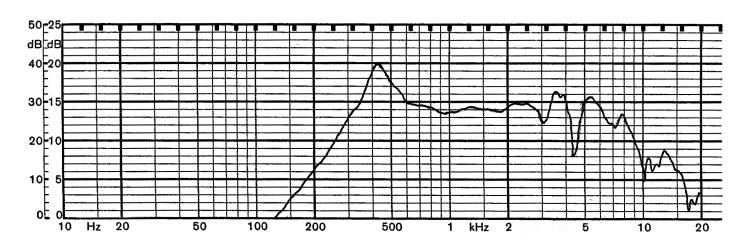
tolerance: ±0.8 mm





## **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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CUI Devices | MODEL: GF0661M | DESCRIPTION: SPEAKER date 03/23/2020 | page 3 of 3

#### **REVISION HISTORY**

rev.	description	date
1.0	initial release	09/27/2006
1.01	brand update	03/23/2020

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.

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