CUI DEVICES

MODEL: CPE-2883 | DESCRIPTION: PIEZO BUZZER INDICATOR

FEATURES

- through-hole pins
- 12 Vdc rating
- 83 db
- 3 kHz rated frequency



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SPECIFICATIONS

parameter	conditions/description	min	typ	max	units
rated voltage			12		Vdc
operating voltage		3		28	Vdc
current consumption	at rated voltage			8	mA
rated frequency		2,500	3,000	3,500	Hz
sound pressure level	at 30 cm, rated voltage	83			dB
dimensions	Ø28.6 x 7.5				mm
weight				3.8	g
material	ABS UL94 1/16" HB (black)				
terminal	pin type (gold plating)				
operating temperature		-30		85	°C
storage temperature		-40		95	°C
RoHS	yes				

Notes: 1. All specifications measured at 5~35°C, humidity at 45~85%, under 86~106kPa pressure, unless otherwise noted.

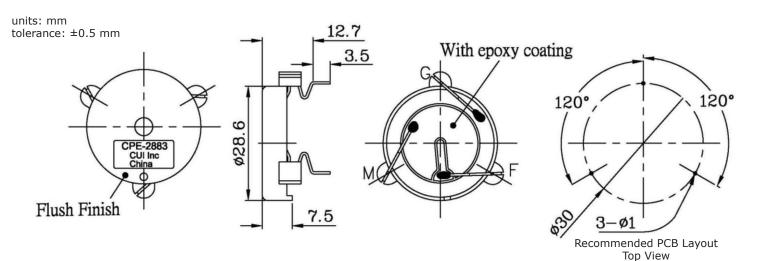
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SOLDERABILITY

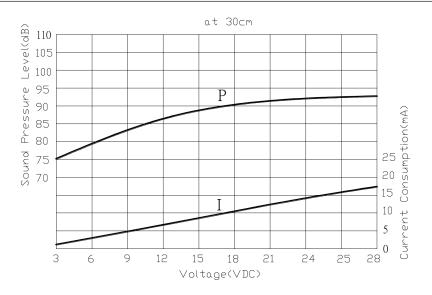
parameter	conditions/description	min	typ	max	units
hand soldering ²	for maximum 3 seconds	320	330	340	°C

Notes: 2. Not recommended for wave soldering

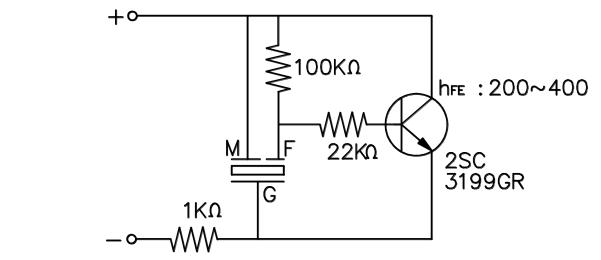
MECHANICAL DRAWING

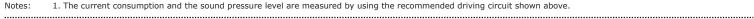


PERFORMANCE CURVES



DRIVING CIRCUIT

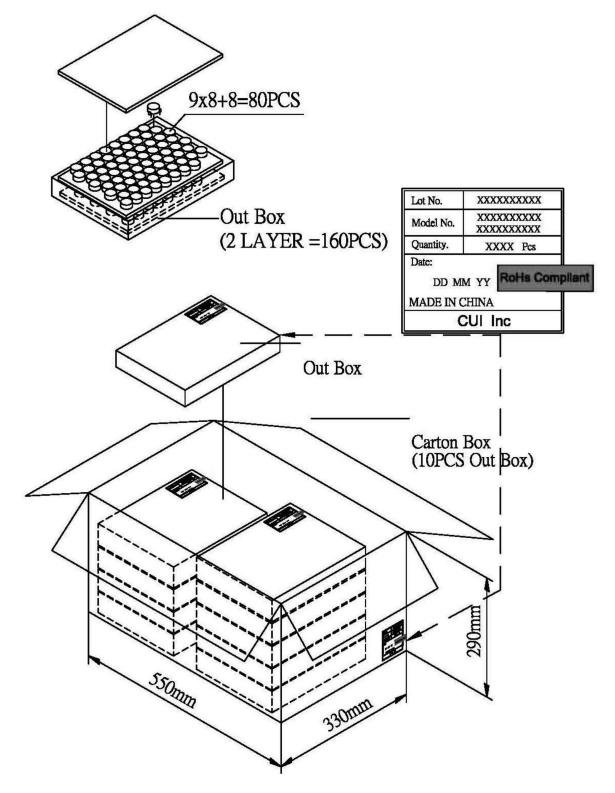




PACKAGING

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Internal Box Size: 310 x 248 x 49 mm Carton Size: 550 x 330 x 290 mm QTY: 1,600 pcs (160 x 10)



REVISION HISTORY

rev.	description	date
1.0	initial release	06/01/2010
1.01	brand update	10/04/2019

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.

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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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