# **CUI** DEVICES

### MODEL: CPI-1375IC-80T | DESCRIPTION: PIEZO BUZZER INDICATOR

#### **FEATURES**

- 12 Vdc rated
- 88 dB
- through hole
- piezo
- internally driven
- narrow frequency range
- used in medical & security applications





#### **SPECIFICATIONS**

parameter	conditions/description	min	typ	max	units
rated voltage			12		Vdc
operating voltage		3.5		16	Vdc
current consumption	at rated voltage			7	mA
rated frequency	at rated voltage	3,900	4,000	4,100	Hz
sound pressure level	at 10 cm, rated voltage	88			dB
tone	single				
dimensions	Ø13.8 x 7.5				mm
weight			1.0		g
material	МРРО				
terminal	pins (red coppper with tin plating)				
operating temperature		-30		70	°C
storage temperature		-40		85	°C
washable	yes				
RoHS	yes				

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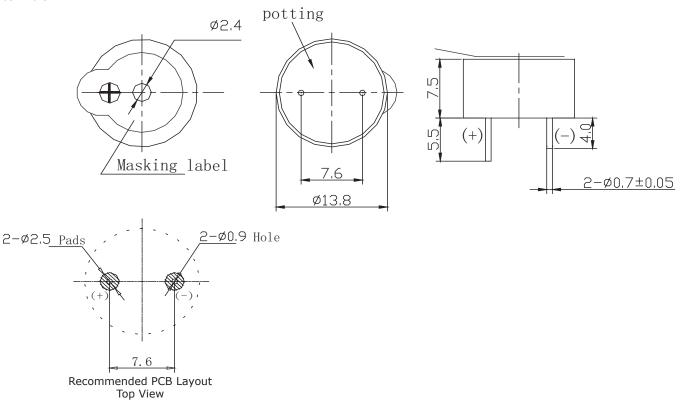
#### **SOLDERABILITY**

parameter	conditions/description	r	min	typ	max	units
wave soldering	see recommended wave soldering profile				250	°C
	250 250 250 250 200 200 200 200	210 240 270				

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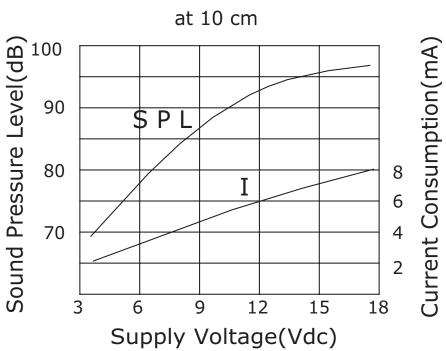
#### **MECHANICAL DRAWING**

units: mm tolerance: ±0.5 mm



#### **PERFORMANCE CURVES**

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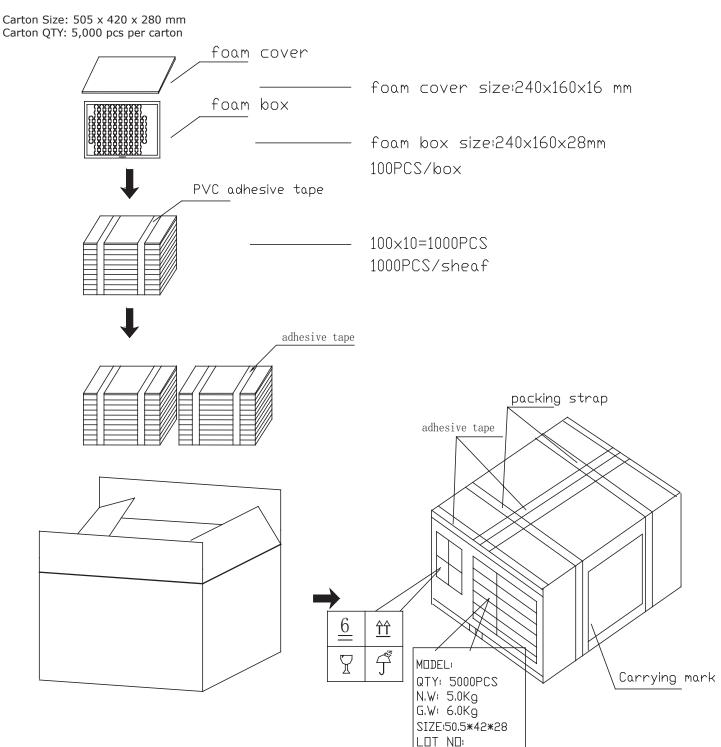


SPL: Voltage vs. Sound Pressure Level I: Voltage vs. Current Consumption

#### PACKAGING

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#### **REVISION HISTORY**

rev.	description	date
1.0	initial release	11/13/2019

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.

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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