QTB

QT-Brightek Lamp Series

8mm Round LED

Part No.: QBL12XX30C

XX: Color Code

Table of Contents:

Introduction

Feature:	**Application:**
• Clear lens	• General purpose indicator application
• Package in bulk pack	• Electronic signs and electronics board
• 8mm round lamp	
• AllnGaP technology	
• Viewing angle: 30° typ.	**Certification & Compliance:**
	• TS16949
Description:	• ISO9001
	• RoHS Compliant
These bright 8mm round type lamps are suitable for all indicator applications such as electronic signs and electronics board indicator	

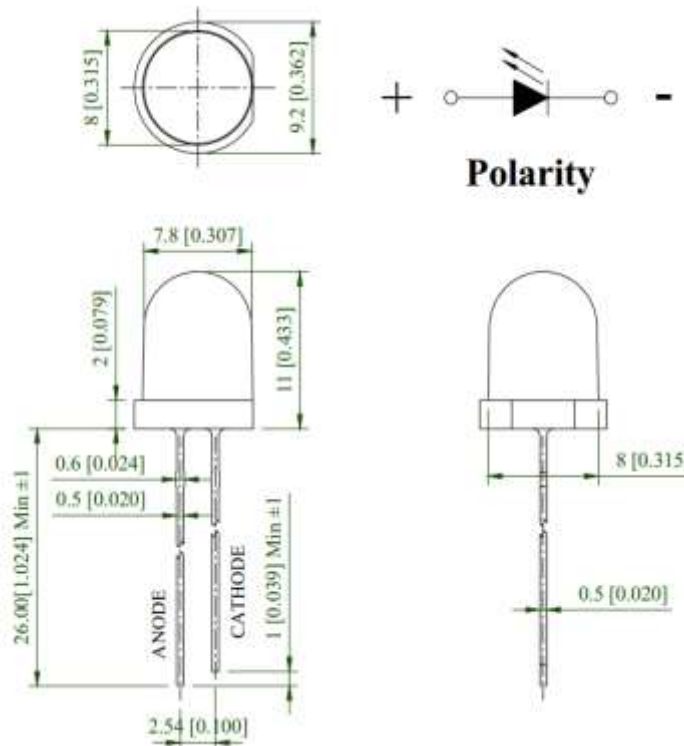**Dimension:**

Polarity

Units: mm / tolerance = +/-0.25mm unless otherwise specified

Electrical / Optical Characteristic (Ta=25°C)

Product	Color	I_F (mA)	V_F (V)		λ_D (nm)	I_V (mcd)	
			Typ.	Max.	Typ.	Min.	Typ.
QBL12R30C	Red	20	2.0	2.4	624	1300	2200
QBL12Y30C	Yellow	20	2.0	2.4	590	780	1300

Absolute Maximum Rating

Material	P_d (mW)	I_F (mA)	I_{FP} (mA)*	V_R (V)	T_{OP} (^{o}C)	T_{ST} (^{o}C)
AlInGaP	60	25	100	5	-40 ~ +85	-40 ~ +100

*Duty Factor=10%, Frequency=1kHz

Characteristic Curves

AlInGaP

Forward Current & Forward Voltage (Ta=25℃)

Luminous Intensity & Ambient Temperature

Forward Current Derating Curve

Luminous Intensity & Forward Current (Ta=25℃, f=1KHz, Duty=1/10)

Relative Intensity vs. Wavelength

Radiation Diagram (Ta=25℃)
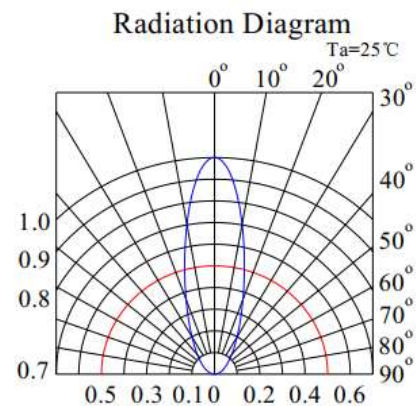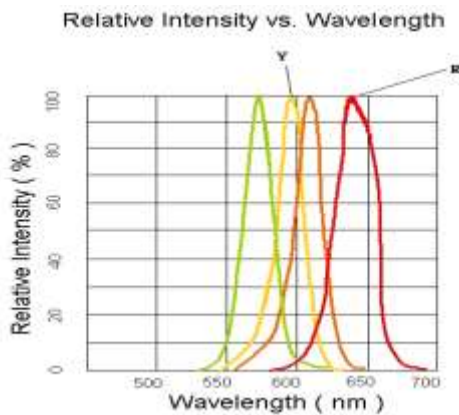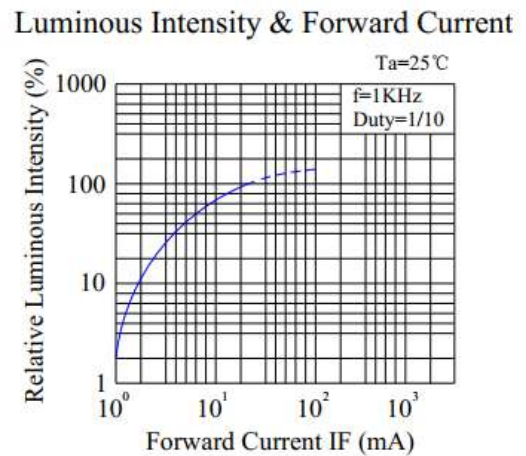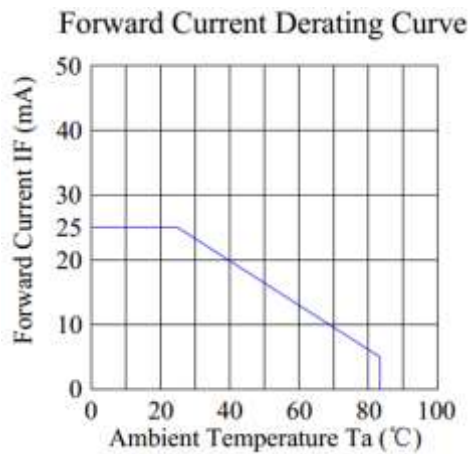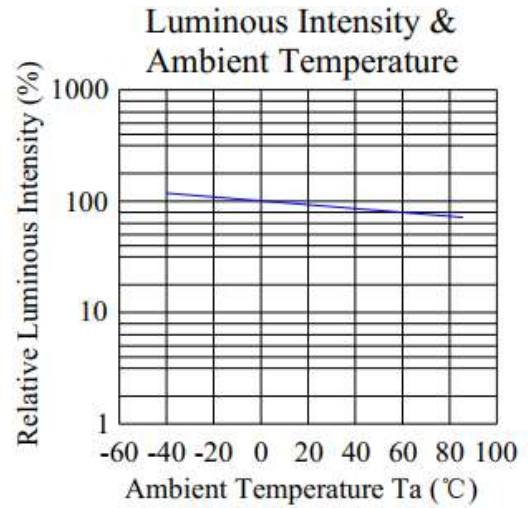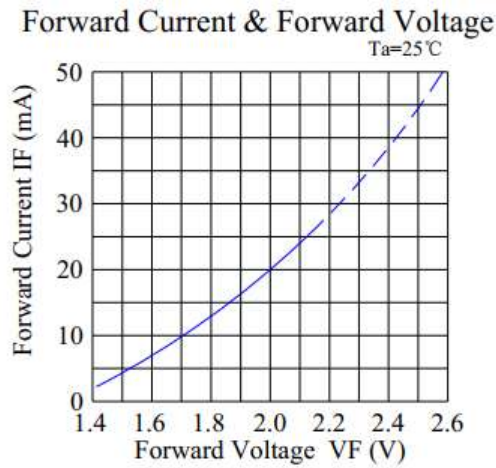
Ordering Information

Part #	Orderable Part #	Spec Range	Quantity per bag
QBL12R30C	QBL12R30C	Iv=2200mcd typ. @ 20mA, λ_D=624nm typ.	400pcs
QBL12Y30C	QBL12Y30C	Iv=1300mcd typ. @ 20mA, λ_D=590nm typ.	400pcs

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Revision History

Description:	Revision #	Revision Date
New Release of QBL12XX30C	V1.0	08/19/2019

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1. Life support devices or systems are devices or systems which, (a) are intended for surgical implant into the body, or (b) support or sustain life, and (c) whose failure to perform when properly used in accordance with instructions for use provided in the labeling, can be reasonably expected to result in a significant injury of the user.

2. A critical component in 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.