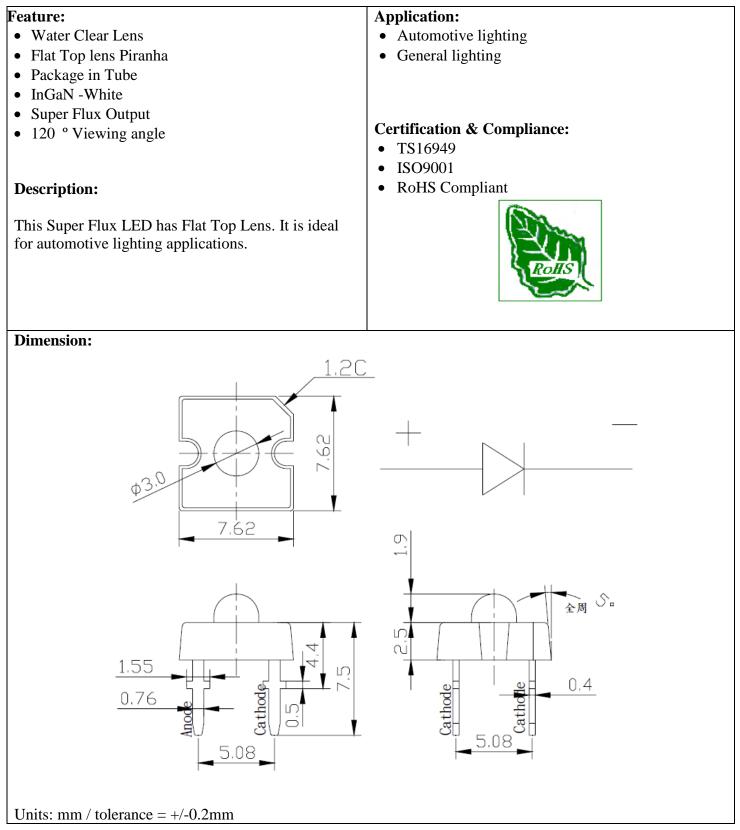
QT BRIGHTEK QBPP0120C-IW



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Electrical / Optical Characteristic (T_A=25 °C)

Droduct	Color	I(mA)	V _F	(V)		λ_{D} (nm)		Φ _V (r	nlm)
Product	Color	l _F (mA)	Тур.	max	Min.	Тур.	Max.	Min	Тур.
QBPP0120C-IW	White	20	3.2	4.0		X=0.271 Y=0.263		2120	4400

Absolute Maximum Rating

QT BRIGHTEK

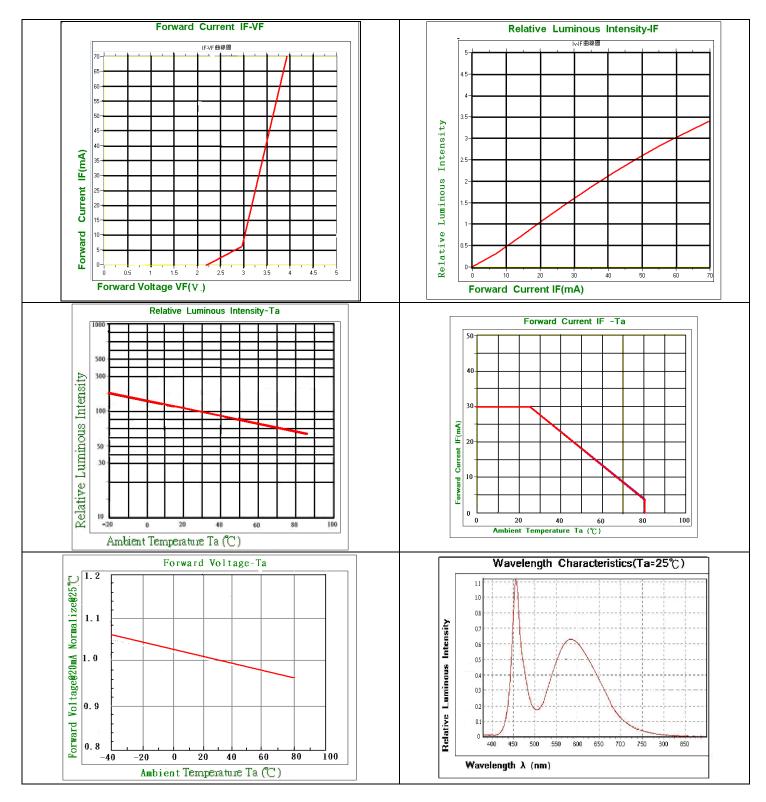
Material	P _d (mW)	I _F (mA)	I _{FP} (mA)*	V _R (V)	T _{OP} (⁰C)	T _{ST} (⁰C)	T _{SOL} (°C)**
InGaN	120	30	100	5	-30 to +80	-40 to +100	260

*Duty 1/10 @0.1ms Pulse Width ** IR Reflow for no more than 5 sec @ 260 °C

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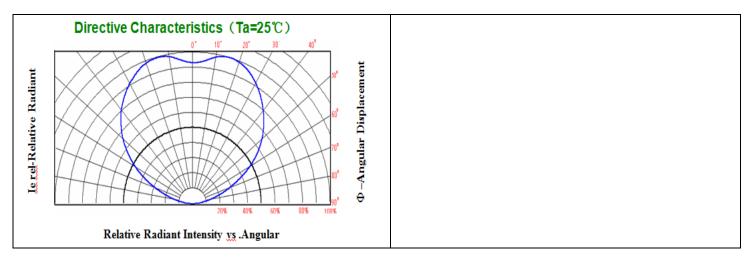
Characteristic Curves For InGaN:

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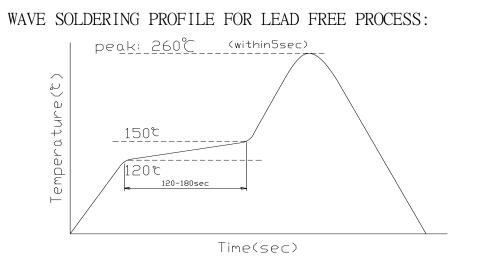


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Solder Profile & Footprint:



Packing: TBD

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Labeling:

N	QT-Brightek	K
Part No		
<u>Custom</u>	er P/N:	
ltem:		
Q'ty:		
Vf:		
lv:		
WI:		
Date:		
	Made in China	

Ordering Information:

Part #	Orderable Part #	Spec Range	Quantity per Tube
QBPP0120C-IW	QBPP0120C-IW	Φ v = 4400 mlm typ. @ I _F =20mA (X,Y)=(0.335,0.355)	TBD

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Revision History:

Description:	Revision #	Revision Date
New Release of QBPP0120C-IW	V1.0	06/25/2010

Disclaimer

QT-BRIGHTEK reserves the right to make changes without further notice to any products herein to improve reliability, function or design. QT-BRIGHTEK does not assume any liability arising out of the application or use of any product or circuit described herein; neither does it convey any license under its patent rights, nor the rights of others.

Life Support Policy

QT-BRIGHTEK's products are not authorized for use as critical components in life support devices or systems without the express written approval of QT-BRIGHTEK. As used herein:

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.

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