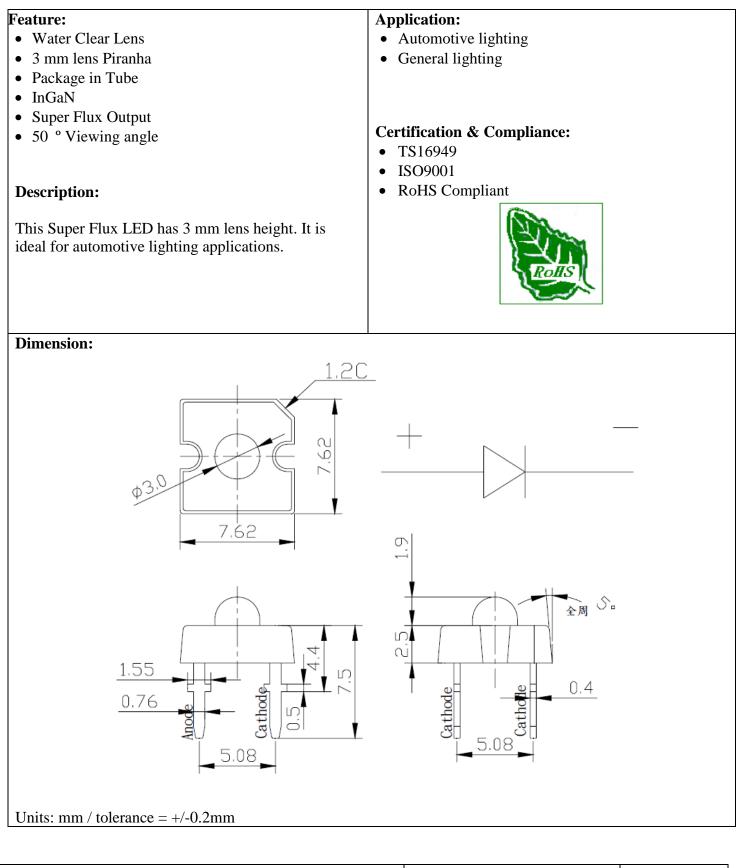
QBPP350C-IB



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#### **Electrical / Optical Characteristic** (T<sub>A</sub>=25 °C)

Product	Color	l <sub>⊧</sub> (mA)	V <sub>F</sub>	(V)		λ <sub>D</sub> (nm)		Φ <sub>V</sub> (m	ılm)
FIOUUCI	COIOI	IF(IIIA)	Тур.	max	Min.	Тур.	Max.	Min	Тур.
QBPP350C-IB	Blue	20	3.2	4.0	462		473	740	1300

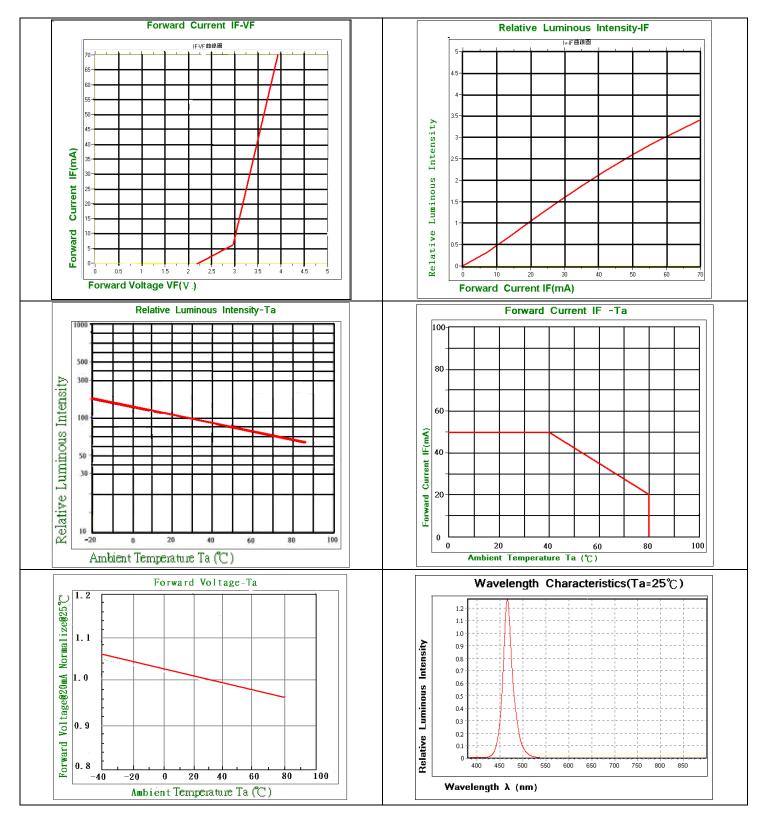
#### **Absolute Maximum Rating**

Material	P <sub>d</sub> (mW)	I <sub>F</sub> (mA)	I <sub>FP</sub> (mA)*	V <sub>R</sub> (V)	T <sub>OP</sub> (⁰C)	T <sub>ST</sub> (⁰C)	T <sub>SOL</sub> (⁰C)**	ESD (V)
InGaN	200	50	100	5	-30 to +80	-40 to +100	260	6000

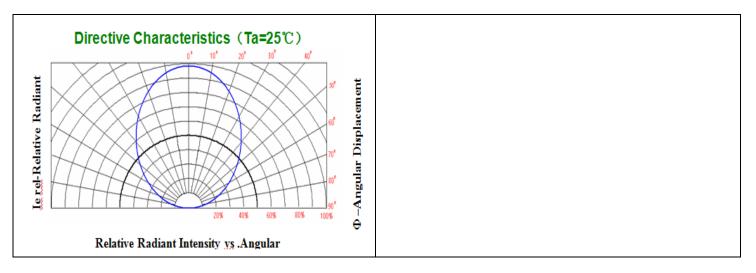
\*Duty 1/10 @0.1ms Pulse Width \*\* IR Reflow for no more than 5 sec @ 260 °C

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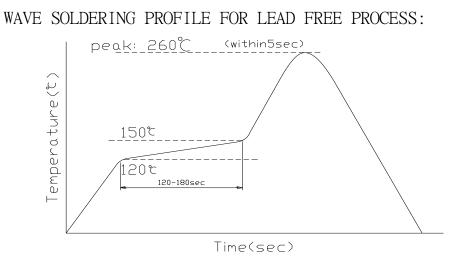
#### Characteristic Curves For InGaN:



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#### **Solder Profile & Footprint:**



Packing: TBD

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### Labeling:

	Þ	QT-Brig	htek	<b>K</b>
∎ Par	t No:			
Cue	stomer	P/N:		
<u>lten</u>	n:			
<u>Q'ty</u>	/:			
$\nabla f$ :		-	_	
<u>Iv:</u>				
<u>VVI:</u>				
Dat	e:	Mada in C	) le i le e	
		Made in C	, nina	

### Ordering Information:

Part #	Orderable Part #	Spec Range	Quantity per Tube
QBPP350C-IB	QBPP350C-IB	Φ v = 1300 mlm typ. @ I <sub>F</sub> =20mA λ <sub>D</sub> =462-473nm	твр

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#### **Revision History:**

Description:	Revision #	Revision Date
New Release of QBPP350C-IB	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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