

# THE GOVERNMENT PHARMACEUTICAL ORGANIZATION

### RAW MATERIAL SPECIFICATION

Title: Benzyl Alcohol BP (For injection dosage form)

(Item No. 41030520)

Spec. No. :

: SP-AK30-B002

(

Reference(s): BP 2019 p.I-282 to I-284

Rev. No.

: 06

Other Requirements: -

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#### BP 2019

Tested Items	Specifications		
Description	Clear, colourless, oily liquid.		
Solubility	Soluble in water, miscible with ethanol (96%) and with fatty and essential oils.		
Relative density	1.043 to 1.049.		
Identification - Infrared absorption spectrophotometry	Conforms to FT-IR standard spectrum.		
Appearance of solution	The solution is clear and colourless.		
Acidity	Not more than 1 ml of 0.1 M sodium hydroxide is required to change the colour of the indicator to pink.		
Refractive index	1.538 to 1.541.		
Peroxide value	Not more than 5.		
Related substances	Impurity A	Not more than 0.05%.	
	Impurity B	Not more than 0.10%.	
	Sum of other peaks with a relative retention less than that of benzyl alcohol	Not more than 0.02%.	
	Sum of peaks with a relative retention greater than that of benzyl alcohol	Not more than 0.2%.	
Residue on evaporation	Not more than 0.05%.		
Assay	98.0% to 100.5% of C <sub>7</sub> H <sub>8</sub> O.		

เอกสารไม่ควบคุม ใช้ในการจัดซื้อ

Prepared by:	Reviewed by:		Approved by :	DO Date
Summe , 13/04/19	Tarme	125/04/19	Walshan	Eff. Date 01/06/19
Head of Raw Material	Director of Raw Material		Director of Quality Assurance	01/06/19
Standard Section 1	Standard Division	_	Department (Active)	



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### **Product Information**

Approved source (s)	Refer to current version of Approved Vendor List of Benzyl Alcohol BP (For injection dosa form) (Item No. 41030520)	
Sampling plan	<ol> <li>N Plan (√N + 1) : for other tests.</li> <li>100% Identification.</li> </ol>	
Testing procedure	Tests to be performed as per current version of WI-AK30-B002.	
Storage condition	To be stored in an airtight container, under nitrogen, protected from light and at a temperature between 2 °C and 8 °C.	
Retest period	1 year after first testing date.	

#### History of changes

Rev. No.	Description	Effective date	
05	Update spec. เป็น BP 2014 โดยเนื้อหาของ BP 2009 และ BP 2014 เหมือนกัน	29/05/15	
06	Update spec. เป็น BP 2019 ตามประกาศกระทรวงสาธารณสุข เรื่องระบุตำรา พ.ศ. 2561 โดยให้ใช้ตำรายาฉบับ USP 39/ BP 2016 ขึ้นไป นอกจากนี้เนื่องจากเอกสารมีอายุครบ 3 ปี จึงต้องทบทวน โดยเนื้อหาของ BP 2014 และ	01/06/19	
	BP 2019 เหมือนกัน		

เอกสารไม่ควบคุม ใช้ในการจัดซื้อ

Prepared by:	Reviewed by:	8	Approved by:	Eff. Date
GUYANIYA / LO (4   19 Head of Raw Material Standard Section 1	Director of Raw Material Standard Division	Director of Regulatory Compliance and Documentation Division	Director of Quality Assurance Department (Aching)	01/06/19