

## THE GOVERNMENT PHARMACEUTICAL ORGANIZATION

#### **RAW MATERIAL SPECIFICATION**

Title: Metoclopramide HCl (For injection dosage form) BP (Item No.

Spec. No.

: SP-AK30-M28/1

41032400)

Reference(s): BP 2016 p. II-278-279

Rev. No.

: 05

Other Requirements : GPO Specification

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#### **BP 2016**

Test Items	Specification
Description	White or almost white crystalline powder, odorless.
Solubility	Very soluble in water, freely soluble in alcohol, sparingly soluble in methylene chloride.
Identification	
A. The pH of Solution S	4.5 to 6.0.
B. Infrared absorption	Conforms to IR standard spectrum.
spectrophotometry	
C. Chloride Test	Dilute 1 ml of Solution S to 2 ml of water, Acidify with dilute nitric acid and add 0.4 ml of
	silver nitrate solution R1. Shake and allow to stand. A curdled, white precipitate is formed.
	Suspend the precipitate in 2 ml of water and add 1.5 ml of ammonia. The precipitate dissolves
¥.	easily with the possible exception of a few large particles which dissolve slowly.
Appearance of solution	Solution S is clear and colorless.
Related substances	- Examine in ultraviolet light at 254 nm. Any spot in the chromatogram obtained with <i>Test</i>
	solution (a), apart from the principal spot, is not more intense than the spot in the chromatogram
	obtained with Reference solution (b) (0.5%).
	- Spray with dimethylaminobenzaldehyde solution R1. Any spot in the chromatogram
	obtained with Test solution (a) that has not been visualized in ultraviolet light at 254 nm is not
	more than intense than the spot in the chromatogram obtained with Reference solution (c)
	(0.5%).
Heavy metals	Not more than 20 ppm, Test A.
Water	4.5% to 5.5%.
Sulfated ash	Not more than 0.1%.
Assay	99.0 – 101.0% of C <sub>14</sub> H <sub>22</sub> ClN <sub>3</sub> O <sub>2</sub> ,HCl, calculated with reference to the anhydrous substance.

เอกสารไม่ควบคุม ใช้ในการจัดซื้อ

1	Prepared by:		Reviewed by:		7	Approved by :	Eff. Date
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1	Head of Raw	Head of V	Director of Raw	Director of	Director of Drug	Director of Quality	
	Material	Microbiological	Material Standard	Microbiological	Registration and	Assurance Department	
	Standard Section 1	Analysis Section 1	Division	Analysis Division	Pharmacovigilance	(Active)	
				•	Division	(2/0/19/2)	

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#### **GPO Specification**

Test Items	Specification		
Bacterial endotoxins	Not more than 2.5 USP Endotoxin Units per mg of Metoclopramide.		

#### **Product Information**

Approved source (s)	Refer to current version of Approved Vendor List of Metoclopramide HCl (Injection grade)		
	BP (Item No. 41032400).		
Sampling plan	1. N Plan $(\sqrt{N} + 1)$ : for other tests.		
	2. 100% Identification.		
Testing procedure	Tests to be performed as per current version of WI-AK30-M28/1.		
Storage condition	Store protected from light.		
Retest period	1 year after first testing date.		

เอกสารไม่ควบคุม ใช้ในการจัดซื้อ

Prepared by :	Reviewed by :	^	7 .	Approved by :	Eff. Date
Head of Raw Head of Material Microbiological Standard Section 1 Analysis Section 1	Director of Raw Material Standard Division	Director of Microbiological Analysis Division	Director of Drug Registration and Pharmacovigilance Division	Director of Quality Assurance Department (Active)	30/11/20

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#### History of changes

Rev. No.	Description	Effective date
03	ข้างอิง spec. เป็น BP 2011	28/02/12
04	Update spec. เป็น BP 2016 เนื่องจากเอกสารมีอายุมากกว่า 3 ปี จึงต้องทบทวน โดยเนื้อหาของ BP 2011 และ BP 2016	24/05/16
	เหมือนกัน อ้างอิง CR No. AN80- 59092	
05	เอกสารมีอายุครบ 3 ปี จึงต้องทบทวน อ้างอิงตาม BP 2016 เพื่อให้สอดคล้องกับ Metoclopramide injection ซึ่งขึ้น	30/11/20
	ทะเบียนตาม Monograph BP 2016	

เอกสารไม่ควบคุม ใช้ในการจัดซื้อ

Prepare	d by
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Director of Raw Material Standard Division

Director of Microbiological Analysis Division

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Director of Drug Registration and Pharmacovigilance Division

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Eff. Date 30/11/20

Assurance Department (Active)