

THE GOVERNMENT PHARMACEUTICAL ORGANIZATION

RAW MATERIAL SPECIFICATION

Title: Talcum (325 Mesh) USP (Item No. 41011490)

Spec. No.

: SP-AK30-T001

Reference(s): USP 43 p. 4207-4209

Rev. No.

: 07

Other Requirements: GPO Specification

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USP 43

Test Items	Specification		
Description	Very fine, white or grayish-white, crystalline powder, odorless and free from grittiness.		
Identification			
A. Infrared absorption	The IR spectrum of a potassium bromide dispersion of it exhibits maxima at 3677 ± 2 cm ⁻¹ ,		
	at $1018 \pm 2 \text{ cm}^{-1}$, and $669 \pm 2 \text{ cm}^{-1}$.		
B. Chemical test	A white, crystalline precipitate of magnesium ammonium phosphate is formed.		
C. Chemical test	Within a short time, a white ring is rapidly formed around the drop of water.		
Acidity or alkalinity	Not more than 0.4 ml of 0.01 N HCl is required to change the color of the bromothymol blue		
	indicator.		
	Not more than 0.3 ml of 0.01 N NaOH is req	uired to change the color of the phenolphthalein	
,	indicator to pink.		
Loss on ignition	Not more than 7.0%.		
Water-soluble substances	ater-soluble substances The filtrate is neutral to litmus paper, and the weight of the residue is not more than 5 r		
	(0.1%).		
Limit of iron	Not more than 0.25%.		
Limit of lead	Not more than 10 ppm.		
Limit of calcium	Not more than 0.9%.		
Limit of aluminum	Not more than 2.0%.		
Absence of asbestos.	Absence.		
Content of magnesium	Between 17.0% and 19.5% of magnesium.		
Microbial enumeration tests and	Total aerobic microbial count	: Not more than 1000 cfu/g.	
Tests for specified microorganisms	Total combined molds and yeasts count	: Not more than 100 cfu /g.	

Prepared by: Chavalum Thungton / 27/03/12 Janjangashal /29/03/22	Reviewed by: Sywannee, 29/03/22	Pyry ,31/03/22	31/03/22	Approved by: (Actim)	Eff. Date 09/05/22
Head of Raw Head of	Director of Raw	Director of	Director of Regulatory	Director of Quality	07/03/22
Material Microbiological	Material Standard	Microbiological	Strategy Division	Assurance Department	
Standard Section 1 Analysis Section 2	Division	Analysis Division	3	,	



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GPO Specification

Test Items	Spe	Specification		
Fineness	Pass through sieve No. 325	: Not less than 60.0%.		
Labeling	The label states, where applicable, that the	The label states, where applicable, that the substance is suitable for oral or topical		
	administration. The certificate of analysis states the absence of asbestos. It also indicates			
	which method specified in the test for Abse	which method specified in the test for Absence of Asbestos was used for analysis.		

Product Information

Approved source (s)	Refer to current version of Approved Vendor List of Talcum (325 Mesh) USP (Item No. 41011490).
Sampling plan	 N Plan (√N + 1): for other tests. 100% Identification.
Testing procedure	Tests to be performed as per current version of WI-AK30-T001.
Storage condition	Preserve in well closed containers. No storage requirements specified.
Retest period	1 year after first testing date.

Prepared by:	-
Thungton / 20/03/12	J
Head of Raw	
Material	
Standard Section 1	

thavalum minampuni/29/03/22 Head of Microbiological Analysis Section 2 Reviewed by:

SUMANNEL 12.03 22

Director of Raw

Material Standard

Division

Director of
Microbiological
Analysis Division

Director of Regulatory Strategy Division Approved by: (Active)
Towns
Limberty 0104121
Director of Quality
Assurance Department

Eff. Date 09/05/22



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History of changes

Description	Effective date
Update spec. เป็น USP 36	15/05/14
Update spec. เป็น USP 39 เนื่องจากเอกสารมีอายุครบ 3 ปีจึงต้องทบทวน โดยเนื้อหาของ USP 36 และ USP 39 เนื้อหา	20/06/17
เหมือนกัน	
Update spec. เป็น USP 43 เนื่องจากเอกสารมีอายุครบ 3 ปีจึงต้องทบทวน โดยเนื้อหาของ USP 43 และ USP 39 เนื้อหา	09/05/22
เหมือนกัน	
	Update spec. เป็น USP 39 เนื่องจากเอกสารมีอายุครบ 3 ปีจึงต้องทบทวน โดยเนื้อหาของ USP 36 และ USP 39 เนื้อหา เหมือนกัน Update spec. เป็น USP 43 เนื่องจากเอกสารมีอายุครบ 3 ปีจึงต้องทบทวน โดยเนื้อหาของ USP 43 และ USP 39 เนื้อหา

Prepared by:
(Aching)
Thurston / 27/03/12
Head of Raw
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Standard Section 1

Chavaleum Head of Director of Raw Microbiological Analysis Section 2

Reviewed by: Material Standard

Division

Director of Microbiological Analysis Division

(1) 31/03/21 Director of Regulatory Strategy Division

Approved by: (Action) turni , 01/14/22 Director of Quality Assurance Department

Eff. Date 09/05/22