

THE GOVERNMENT PHARMACEUTICAL ORGANIZATION

RAW MATERIAL SPECIFICATION

Title: Disodium Hydrogen Phosphate Extra Pure USP (Item No. 41031760)

Spec. No.

: SP-AK30-S66/1

(Dibasic Sodium Phosphate (dried form))

Reference(s): USP 43 p. 4087 - 4088

Rev. No.

: 03

Other Requirements: GPO specification

Page

: 1/2

USP 43

Test Items	Specification		
Description	White powder, odorless.		
Solubility	Freely soluble in water, insoluble in alcohol.		
Identification			
A. Sodium Test	A dense white precipitate is formed.		
B. Phosphate Test	With silver nitrate TS, neutral solutions of orthophosphates yield a yellow precipitate that is		
	soluble in 2 N nitric acid and in 6 N ammonium hydroxide.		
Loss on drying	Not more than 5.0%, for the dried form.		
Insoluble substances	Not more than 20 mg (0.4%).		
Chloride	Not more than 0.06%.		
Sulfate	Not more than 0.2%.		
Arsenic	Not more than 16 ppm, Method I.		
Assay	98.0% to 100.5% of Na ₂ HPO ₄ , calculated on the dried basis.		

GPO specification

Test Items		Specification	
Heavy metals	Not more than 20 ppm.		

Prepared by:	1
Surannee, 12/10/20	ز
Head of Raw Material	1
Standard Section 1	1

Reviewed by:
Turmin
Ludword
Allola

Director of Raw Material

Standard Division

Director of Drug Registration and Pharmacovigilance Division

Approved by:

(function /14/0/20

Director of Quality Assurance
Department (Active)

Eff. Date

30/11/20

THE GOVERNMENT PHARMACEUTICAL ORGANIZATION

RAW MATERIAL SPECIFICATION

Title: Disodium Hydrogen Phosphate Extra Pure USP (Item No. 41031760)

Spec. No.

: SP-AK30-S66/1

(Dibasic Sodium Phosphate (dried form))

Reference(s): USP 43 p. 4087 - 4088

Rev. No.

: 03

Other Requirements: GPO specification

Page

: 2/2

Product Information

Approved source (s)	Refer to current version of Approved Vendor List of Disodium Hydrogen Phosphate Extra Pure
	USP (Dibasic Sodium Phosphate (dried)) (Item No. 41031760).
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-S66/1.
Storage condition	Preserved in tight container. Label it to indicate whether it is dried or is the monohydrate, the
	dihydrate, the heptahydrate or the dodecahydrate.
Retest period	1 year after first testing date.
í	

History of changes

Rev. No.	Description	Effective date
01	ประกาศใช้ครั้งแรก อ้างอิง USP 34	17/10/11
02	Update spec. เป็น USP 39 โดยเนื้อหาของ USP 34 และ USP 39 เหมือนกัน	24/05/16
03	เอกสารมีอายุครบ 3 ปี จึงต้องทบทวน โดย update spec. เป็น USP 43 โดยเนื้อหาของ USP 39 และ USP 43 เหมือนกัน	30/11/20

Prepared by:	Reviewed by:	1 -	Approved by:	Eff. Date
Surannee, 12/10/20	Luirent 12/10/20	14/10/20	Yunapon 11/10/20	30/11/20
Head of Raw Material	Director of Raw Material	Director of Drug Registration and	Director of Quality Assurance	
Standard Section 1	Standard Division	Pharmacovigilance Division	Department (Acting)	