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CERTIFICATE OF ANALYSIS SO0335_25496501/1

Product di-Sodium hydrogen phosphate anhydrous, Pharmpur®, Ph Eur, BP, USP Batch 25496501
Quality release date 27/01/2025
SO0335 Expiry date 01/2032

Analysis	Batch value	Specifications
Assay USP (Acidimetric, referred to dried sample)	98,50 %	98,0 - 100,5
Assay EP (Acidimetric, referred to dried sample)	100,25 %	98,0 - 101,0
Identification sodium	passes test	passes test
Identification phosphate	passes test	passes test
Identification A (EP)	passes test	passes test
Appearance of solution	clear and colourless	clear and colourless
Insoluble in water	0,0040 %	≤ 0,4
Chlorides (CI)	40 ppm	≤ 200
Sulfates (SO4)	< 50 ppm	≤ 500
Sodium dihydrogenphosphate (NaH2PO4)	0,023 %	≤ 2,5
Arsenic (As) (EP)	< 1 ppm	≤ 2
Arsenic (As) (USP)	< 1 ppm	≤ 16
Iron (Fe)	< 20 ppm	≤ 20
Reducing substances	passes test	passes test
Loss on drying (105 °C, 4h)	0,17 %	≤ 1,0
Loss on drying (130 °C)	0,21 %	≤ 5,0

This product is intended to be suitable for the manufacture of pharmaceuticals, medical devices, cosmetics and food products, but is entirely under the user's responsibility.

