

AMRI India Private Limited

Factory / Office : Plot No. G-1/1, 1/2, Near MIDC Water Tank, MIDC Area, Waluj, Aurangabad, India -431 136 t : +91-240-2554006 / 2564456, www.amriglobal.com

		QUALITY CONTR	OL DEP	ARTMENT		
×		CERTIFICATE	OFAN	ALYSIS		
Tym	o. Dignatch for Metanharm		Page: 1			
Type: Dispatch for Metapharmaceutical IND.S.L. Product Name: FUROSEMIDE				Batch Number: AM20214079		
			Quantity: 25 Kg.			
Compendia: EP Mfg. Date: AUG-2014			Retest Date: JUL-2019			
Date of Sampling: 09/09/2014			Date of Analysis: 09/09/2014			
Date of Release: 17/09/2014			A.R Number: FPF/14/112			
Stor	age Condition: - Preserve	e in well-closed, light d between 15°C to 30	resistant °C	container. Store at	25°C, excursions	
Sr. No.	Test	Results		Specification		
1.	Appearance	Almost white crystalline powder		A white or almost white crystalline powder.		
2.	Solubility	Complies		Practically insoluble in water, soluble in Acetone, sparingly soluble in Ethanol (96%), practically insoluble in Methylene chloride. It dissolves in dilute solutions of alkali hydroxide.		
3.	Polymorphism test by DSC	Complies		DSC pattern of sample should be concordant with that of the standard pattern		
§ 4 .	Identification	0.54		0.52 to 0.57		
Fre-	A) By UV B) By IR	Complies		Should be comparable with Furosemide working standard.		
1000	C) By Color Test	Formation of red-violet color		Formation of a re	ed to red-violet color.	
5.	Related Substances					
	Impurity C	BDRL		NMT 0.2%		
¥3	Impurity D	BDRL		NMT 0.15%		
8	Unspecified Impurities	BDRL		NMT 0.10% NMT 0.5 %		
10 1	Total Impurities	BDRL		TNIVI U.3 70		
62	- Identification				ID /D	
Prepared By /Date Check		hecked By / Date	Revi	ewed By / Date	Approved By / Date	
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	Quality Control	Quality Control	Qua	lity Assurance	Quality Assurance	



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QUALITY CONT	ROL DEPARTMENT		
CERTIFICAT	E OF ANALYSIS		
Type: Dispatch for Metapharmaceutical IND.S.L.	Page: 2 of 2		
Product Name: FUROSEMIDE	Batch Number: AM20214079		
Compendia: EP	Quantity: 25 Kg.		
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Sr. No.	Test	Results	Specification			
6.	Chlorides	Less than 200 ppm	NMT 200 ppm			
7.	Sulphates	Less than 300 ppm	NMT 300 ppm			
8.	Heavy Metals	Less than 20 ppm	NMT 20 ppm			
9.	Loss on drying At 105 °C ± 2°C	0.25 % w/w	NMT 0.5% w/w			
10.	Sulphated Ash	0.06 % w/w	NMT 0.1% w/w			
11.	Assay by Potentiometry (ODB)	99.27 % w/w	98.5% to 101.0% w/w			
112.	*Additional Tests					
12.1	Foreign Particle	No black particles observed on filter paper	10gm sample dissolve in 100ml acetone and filtered through 0.45µ filter paper. NMT 5 black particles on filter paper.			
12.2	Residual EDTA (In-house specification)	Less than 200 ppm	NMT 200 ppm			
12.3	Bulk Density 1) Untapped 2) Tapped (By 50 strokes)	0.22 gm/mL 0.31 gm/mL	Informative			
12.4	Particle size (By sieve method)	100% sample passes through 40 mesh	100% passing through 40 mesh			

*Additional tests determined as per in-house requirement.

• BDRL - Below Disregard Limit (Disregard Limit is 0.05% as per EP 8.0)

Remarks:- The product Complies as per EP specification
We certify that the material manufactured in our factory, address - AMRI India Pvt. Ltd. G-1/1,
MIDC, Aurangabad is as per process described in the certificate of suitability. We also certify
that the quality of the material complies according to the certificate of suitability (COS No.: R1-CEP 1999-069-REV 05) of the monographs of the European Pharmacopoeia

Prepared By /Date Checked		Checked By / Date	Reviewed By / Date	Date Approved By / Date	
	Pays 11/10/2014	Jacob "Morrord	11/18/14	Spirotal	
	Quality Control	Quality Control	Quality Assurance	Quality Assurance	