

AMRI India Private Limited, Plot No. G-1/1, 1/2, MIDC Area, Waluj, Aurangabad-431 136, India t:+91-240-2554006 / 2564456 www.amriglobal.com

QUALITY	CONTROL DEPARTMENT		
CERTIFICATE OF ANALYSIS			
Type: Dispatch Page: 1 of 2			
Product Name: ATENOLOL	Batch Number: AM20114040		
Compendia: EP	Quantity: 225 Kg.		
Mfg. Date: MAY -2014	Retest Date: APR-2019		
Date of Sampling: 26/05/2014	Date of Analysis: 27/05/2014		
Date of Release: 10/06/2014	A.R Number: FPA/14/044		
Stange Candidian D. 11 1	1 1: 1		

Storage Condition: Preserve in well-closed, light resistant container. Store at 25°C, excursions permitted between 15°C to 30°C

Sr.No.	Test	Results	Specification	
1.	Appearance	Almost white powder	A white or almost white powder	
2.	Solubility	Complies	Soluble in anhydrous ethanol, sparingly soluble in water, slightly soluble in methylene chloride.	
3.	Identification A) By Melting Point	153.4 °C	First Identification C Second Identification A, B, D. 152°C to 155°C	
	B) By UV	1.16	1.15 to 1.20	
	C) By IR	Complies	Should be comparable with Atenolol working standard.	
	D) By TLC	Complies	Should be comparable with Atenolol working standard.	
4.	Appearance of solution	1.0% w/v solution in water is clear & transparent	1.0%w/v solution in water should be clear & not more intensely colored than degree 6 of the range of reference solutions of the most appropriate color.	
5.	Optical Rotations	0°	$+0.10^{\circ}$ to -0.10°	
6.	Chloride	Less than 0.1%	NMT 0.1%	

Prepared By /Date	Checked By / Date	Reviewed By / Date	Approved By / Date
14106/2014	Jacoby 14/06/1014	14/06/2019	All oblight
Quality Control	Quality Control	Quality Assurance	Quality Assurance



AMRI India Private Limited, Plot No. G-1/1, 1/2, MIDC Area, Waluj, Aurangabad-431 136, India t: \pm 91-240-2554006 / 2564456 www.amriglobal.com

QUALITY CONTROL DEPARTMENT CERTIFICATE OF ANALYSIS			
Product Name: ATENOLOL	Batch Number: AM20114040		
Compendia: EP	Quantity: 225 Kg.		
Mfg. Date: MAY -2014	Retest Date: APR-2019		
Date of Sampling: 26/05/2014	Date of Analysis: 27/05/2014		
Date of Release: 10/06/2014	A.R Number: FPA/14/044		

Sr.No.	Test	Results	Specification
7.	Related Substances By HPLC		
	Impurity B	0.16 %	NMT 0.20 %
	Impurity F	BDRL	NMT 0.15 %
	Impurity G	BDRL	NMT 0.15 %
	Impurity I	0.08 %	NMT 0.15 %
	Unspecified Impurity	0.06 %	NMT 0.10 %
	Total Impurities	0.36 %	NMT 0.5 %
8.	Sulphated Ash	0.06 %	NMT 0.1 %
9.	Loss on drying	0.25 % w/w	NMT 0.5 % w/w
10.	Assay (ODB) by Potentiometer	99.62 % w/w	99.0% to 101.0% w/w
11.	Foreign Particle	No black Particles observed	NMT 5 black particles observed on
		on filter paper	filter paper
12.	*Additional Tests		
12.1	Bulk Density		
	1) Untapped	0.24 gm/mL	Informative
	2) Tapped (50 strokes)	0.31 gm/mL	
12.2	Particle Size	100% sample passes through 40 mesh	100% passing through 40 mesh.

*Additional test determined as per in-house requirement.

• BDRL – Below Disregard Limit (Disregard Limit is 0.05% as per EP 7.0)

• 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 (No: R1-CEP 1999-122-Rev 04) of the monographs of the European Pharmacopoeia.

Prepared By /Date	Checked By / Date	Reviewed By / Date	Approved By / Date
14/06/2014	Jacob 1410611014	14/06/2014	Shipping
Quality Control	Quality Control	Quality Assurance	Quality Assurance