



Office of The Commissioner,
Food & Drugs Administration M.S.
Bandra – Kurla Complex,
Bandra (E),
Mumbai – 400 051
Date : 17/06/2017

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

This Certificate conforms to the format recommended by the World Health Organization.
(General instructions and explanatory notes attached).

Certificate No.: **NEW-WHO-GMP/CERT/KD/58604/2017/11/19684**

On the basis of the inspection carried out on 14/03/2017, 15/03/2017 and 20/04/2017, we certify that the site indicated on this Certificate complies with **Good Manufacturing Practices** for the dosage forms, categories and activities listed in Table 1.

1. Name of the Firm : **AARTI DRUGS LTD**
Address : **PLOT NO G-60 MIDC TARAPUR TAL-PALGHAR
DIST THANE 401506 MAHARASHTRA STATE,
INDIA**
2. Licence No. : **KD217 In Form 25**

Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(ies)
1	Active Pharmaceutical Ingredients (Bulk Drugs)	General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones)	Synthesis, Purification, Packing, Labelling, Quality Control, Quality Assurance

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate remains valid until 16 Jun 2019 . It becomes invalid if the activities and / or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Address of certifying authority :
Food & Drug Administration, M.S.
Bandra-kurla Complex,
Bandra (E), Mumbai – 400 051.
Maharashtra, INDIA.
Tel: +91-22-26592363/64
Fax: +91-22-26591959
1RAA6395860420170617

Name of the Authorised person : **O S SADHWANI**

Signature : 
Stamp and Date : **Joint Commissioner (HQ) & Controlling
Authority
Food & Drug Administration, M.S.
Bandra (E), Mumbai.
Maharashtra State, India
Date: 17 Jun 2017**

LIST OF PRODUCT APPROVED UNDER WHO GMP¹

No. of certificate : NEW-WHO-GMP/CERT/KD/58604/2017/11 VALID UP TO :16 Jun 2019
/19684
Name of Manufacturing Firm : AARTI DRUGS LTD
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INDIA
Drug License No : KD217 In Form 25

Sr.No.	Name of the Product	Composition
1	CIPROFLOXACIN HYDROCHLORIDE BP	
2	CIPROFLOXACIN HYDROCHLORIDE EP	
3	CIPROFLOXACIN HYDROCHLORIDE IP	
4	CIPROFLOXACIN HYDROCHLORIDE USP	
5	CLOPIDOGREL BISULPHATE IP	
6	CLOPIDOGREL BISULPHATE USP	
7	CLOPIDOGREL HYDROGENSULFATE BP	
8	CLOPIDOGREL HYDROGENSULFATE EP	
1 2 3		

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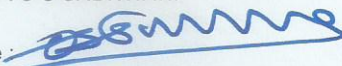
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Sr.No.	Name of the Product	Composition
9	DICLOFENAC DIETHYL AMMONIUM BP	
10	DICLOFENAC POTASSIUM BP	
11	DICLOFENAC POTASSIUM EP	
12	DICLOFENAC POTASSIUM USP	
13	DICLOFENAC SODIUM BP	
14	DICLOFENAC SODIUM EP	
15	DICLOFENAC SODIUM IP	
16	DICLOFENAC SODIUM JP	
1 2 3		

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Sr.No.	Name of the Product	Composition
17	DICLOFENAC SODIUM USP	
18	GATIFLOXACIN SESQUIHYDRATE	
19	METFORMIN HYDROCHLORIDE BP	
20	METFORMIN HYDROCHLORIDE EP	
21	METFORMIN HYDROCHLORIDE IP	
22	METFORMIN HYDROCHLORIDE USP	
23	PIOGLITAZONE HYDROCHLORIDE IP	
24	PIOGLITAZONE HYDROCHLORIDE USP	

1 2 3

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