1-817 NG 2

## SQUARE FORMULATIONS LIMITED TANGLADESH



## Certificate of Analysis (COA) for Working Standard

Ref. Log Book ; LB/QC/0120/14, Page No : 49-53	Storage Condition : Below 25 ° C
Batch no of Reference standard : 5.0	Date of storage: 05 Aug'15
Reference standard: Fluoxetine Hydrochloride EP CRS	No. of Container: Two
Method of Analysis: HPLC	Ref. Spec. No : RMSP/QC/003/02
Date of Analysis: 05 Aug'15	Date of Expiry (as std.): 04 Aug'16
Supplier: Cadila Pharmaceuticals Ltd	Expiry date of materials : 31 Mar'19
Manuf. : Cadila Pharmaceuticals Ltd	Date of Mfg : 01 Apr'14
Maferial: FLUOXETINE HYDROCHLORIDE ВР	Batch No. : 1000022028

Results	Specification	stsaT
White crystalline powder.	White or almost white, crystalline powder.	1. Appearance
Complies (Test A and Test B)	A. The infrared absorption spectrum of the sample must be concordant with the standard spectrum.  B. Chloride test	2, Identification
%611.0	%03.0 TM <i>V</i>	3. Water Content
Impurity A: ND, Impurity B: ND, Impurity B: ND, Impurity C: ND, Any Unknown impurity: 0.06%; Total impurity: 0.16%.	: 8,A Viruqml,%31.0 TMM ;) yinuqml sətiruqmi bəhləqqanU,%32.0 TMM TMM : sətiruqmi latoT,%01.0 TMM 0.50%	4. Related substances
(8AO)%49,66 (ai 1i aA)%23.66	no) % 0.201 TMM and UMT 102.0 % (on anhydrous basis)	ұвеед. <u>.</u> д

Sr. Executive, QA	Sr. Executive, QC	Sxecutive, QC
Musabbir Humayun	duddeM risetnuM	Md. Hanifur Rahman
1.120m) Dr	Markery	S1770 22 D
Approved by:	Срескед ру:)	Prepared by: