By Speed Post

No RTI/ 360/142-15/15
Office of Commissioner
Food & Drug Administration,
Maharashtra State, 341,
Bandra-Kurla Complex.
Bandra (East). Mumbai-51.
Date:- 25/8/2015

To

Prashant Reddy T.
Advocate
C/o Advocate Harsh Paraskar.
Lex one Partners E-19.LGF.
Jungpura Extension.
New Delhi- 110014

Subject: Drugs and Cosmetics Act 1940 & Rules 1945.

Ref: - Your application dated 20.07.2015 received by this office from the on 23/07/2015

Sir,

With reference to above, this is to inform you that I am forwarding herewith information furnished by Assistant Commissioner, Desk-10 of this office.

In case you wish to prefer an appeal the same can be made within 30 days to the Appellate Authority. The details of the Appellate Authority are mentioned below.

Appellate Authority,
Shri O. S. Sadhwani
Joint Commissioner (H. Q),
Food & Drugs Admn
S. No 341, Bandra Kurla Complex,
Bandra (E), Mumbai 400051

Yours,
(Dr. Rakesh Tirpude)
Assistant Commissioner
And Information Officer, (H.Q)
Food & Drugs Admn. (M. S)

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Tipani | Desk - 10

Sub :- Application under Right to Information Act 2005
Application of Prashant Reddy, Advocate, New Delhi, dated – 20.07.2015

Ref :- Your Tipani No. Mau A/360/14215/15, dt. 14.08.2015

This is to inform you that, as per above said application information is as follows –

Sr.No	Information Sought	Particulars of information
	Does the Controller follow any specific rules or guidelines to recall a drug that is detected as being of Not of Standard quality. Please provide the applicant with a copy of such rules of guidelines.	The Food & Drug Administration, M.S. recall Not of Standard quality drug as per provision laid down in Drug & cosmetics Rules 1945.
2	What is the procedure followed by the Controller while deciding appropriate legal action when a sample is detected to be of Not of Standard quality. Does the Controller initiate criminal prosecution in all cases or is suspension of licenses enough. The PIO is requested to please provide the applicant with a copy of procedure/rules to be followed while deciding appropriate legal action in such cases.	In case of Not of Standard quality drug the Food & Drug Administration, M.S. follows the DCGI, New Delhi guidelines and also follows the guideline prepared by this Administration.

Kindly inform the applicant accordingly.

(J.B.Mantri)
For Joint Commissioner,
Food & Drugs Administration,
Maharashtra State
Mumbai – 51
Out No. - kemaa A- 10/806-15/10
Date - 25/8/15

To,
Assistant Commissioner(H.Q.) &
Public Information Officer, (Desk -15)
Food & Drugs Administration,
Maharashtra State

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