Z-28020/256/2015-DC
Directorate General of Health Services
Office of DCG (I)
(RTI CELL)

FDA Bhawan, New Delhi.
Dated: 22/5/15

To,

Sh. Prashant Reddy T.
C/o Advocate Harsh Parashar,
Lex One Partners, E-19,
LGF, Jungpura Extention, New Delhi – 110 014

Sub: Information under RTI Act, 2005 – Regarding.

Sir,

Please refer to your letter no. Nil dated 15-04-2015 received in this office on dated 16-04-2015 regarding information under RTI Act, 2005.

The point wise information is placed as below:

POINT NO. 1, 2 & 3

The responsibility of regulating manufacturing, Sale & Distribution of drugs in the Country is primarily the concern of State Licensing Authority appointed by respective State Governments. Being the licensing Authorities, State Licensing Authorities are empowered to issue Safety Alert or direct the manufacturing to recall the drug declared as Not of Standard Quality.

Yours Faithfully,

(Sunil Kulshrestha)
Central Public Information Officer

Address of first appellate authority is as below:
The Drugs Controller General (India)
FDA Bhawan, Opp. Mata Sundri College,
Kotla Road, New Delhi – 110002.