- 32. लेवोफ्लोक्सासीन
- 33. मेरापेनाम
- 34. मीडाजोलाम
- 35. मोक्सीफ्लोक्सासीन
- 36. निट्राजेपाम
- 37. पेंटाजोसीन
- 38. प्रतीफ्लोक्सासीन
- 39. पाइराजीनामाइड
- 40. रिफाब्टीन
- 41. रिफाम्पिसेन
- 42. सोडियम पैरा-अमिनोसालिसाइलेट
- 43. स्पारफ्लोक्सासीन
- 44. थायसीटाजोन
- 45. टारामाडोल
- 46. जोलपीडेम

टिप्पण: पूर्वीक्त ओषि पदार्थों और उनके लवणों को अंतर्विष्ट करने वाली निर्मितियां, जिनमें वे शामिल नहीं है, जो सीमित या बाह्य प्रयोग (आँख, और कान/नाक विनिर्मितियों के सिवाय) के लिए आशयित हैं, जिनमें पूर्वीक्त पदार्थ अंतर्विष्ट है, भी उक्त सूचि के अधीन आती हैं।

> [फा. सं. एक्स-11014/6/2010-डीएफक्यूसी] अरूण के. पण्डा, संयुक्त सचिव

पाद टिप्पण: मूल नियम राजपत्र में अधिसूचना सं एफ.28-10/45-एच(1), तारीख 21 दिसंबर, 1945 द्वारा प्रकार्शित किए गए थे और इनमें अंतिम संशोधन अधिसूचना सं. सा.का.नि. तारीख द्वारा किया गया।

## MINISTRY OF HEALTH AND FAMILY WELFARE (Department of Health and Family Welfare) NOTIFICATION

New Delhi, the 30th August, 2013

G.S.R. 588(E).— Whereas certain draft rules further to amend the Drugs and Cosmetics Rules, 1945, were published, as required by sections 12 and 33 of the Drugs and Cosmetics Act, 1940 (23 of 1940), without consulting the Drugs Technical Advisory Board vide notification of the Government of India, Ministry of Health and Family Welfare (Department of Health and Family Welfare), number G.S.R. 228(E), dated the 20<sup>th</sup> March, 2012, published in the Gazette of India, Extraordinary, Part II, Section 3, Sub-section (i), dated the 20<sup>th</sup> March, 2012, inviting objections and suggestions from all persons likely to be affected thereby before the expiry of a period of forty-five days from the date on which the copies of the Official Gazette containing the said notification were made available to the public;

And whereas the copies of the Gazette in which the said notification was published were made available to the public on the 20<sup>th</sup> March, 2012;

And whereas, the Drugs Technical Advisory Board has been consulted in the matter,

And whereas, objections and suggestions received in respect of the said draft rules have been considered by the Central Government;

Now, therefore, in exercise of the powers conferred by sections 12 and 33 of the Drugs and Cosmetics Act, 1940 (23 of 1940), the Central Government, after consultation with the Drugs Technical Advisory Board, hereby makes the following rules further to amend the Drugs and Cosmetics Rules, 1945, namely:-

- 1. (1) These rules may be called the Drugs and Cosmetics (Fourth Amendment) Rules, 2013.
  - (2) They shall come into force after six months of their publication in the Official Gazette.
- 2. In the Drugs and Cosmetics Rules, 1945 (hereinafter referred to as said rules),-
  - (i) in rule 65,-
    - (a) in condition (3), in clause (1),-
      - (A) in sub-clause (f) and in item (ii) of the third proviso to sub-clause (g), for the words and letter 'Schedule H', the words and letters "Schedule H and Schedule H1" shall respectively be substituted;
      - (B) after clause (g) and the provisos thereof, the following shall be inserted, namely:-
      - "(h) the supply of a drug specified in Schedule H1 shall be recorded in a separate register at the time of the supply giving the name and address of the prescriber, the name of the patient, the name of the drug and the quantity supplied and such records shall be maintained for three years and be open for inspection.
    - (b) in condition (9), in clauses (a) and (b), for the words and letter 'Schedule H', the words and letters "Schedule H and Schedule H1" shall respectively be substituted;
    - (c) in condition (11), for the words and letter 'Schedule H', the words and letters "Schedule H and Schedule H1" shall be substituted;
    - (d) in condition (11A), for the words and letter 'Schedule H', the words and letters "Schedule H and Schedule H1" shall be substituted;
- (ii).in rule 97, in sub rule (1), after clause (d), the following shall be inserted, namely,-
  - '(e) if it contains a drug substance specified in Schedule H1, the drug formulation shall be labelled with the symbol Rx which shall be in red and conspicuously displayed on the left top corner of the label, and shall also be labelled with the following words in a box with a red border:

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## "SCHEDULE H1 DRUG - WARNING:

- -It is dangerous to take this preparation except in accordance with the medical advice.
- -Not to be sold by retail without the prescription of a Registered Medical Practitioner." ';
- 3. In the said rules, in Schedule H, the following entries shall be omitted, namely:-
  - "1. Alprazolam
  - 2. Cefdinir
  - 3. Cefepime Hydrochloride
  - 4. Cefetamet Pivoxil
  - 5. Cefpirome
  - 6. Cefpodoxime Poxetil
  - 7. Ceftazidime Pentahydrate
  - 8. Ceftizoxime Sodium
  - 9. Chlordiazepoxide
  - 10. Clofazimine
  - 11. Codeine
  - 12. Diazepam
  - 13. Diphenoxylate and its salts
  - 14. Ethambutol Hydrochloride
  - 15. Ethionamide
  - 16. Levofloxacin
  - 17. Meropenam
  - 18. Midazolam
  - 19. Moxifloxacin
  - 20. Nitrazepam
  - 21. Pentazocine
  - 22. Pyrazinamide
  - 23. Sparfloxacin
  - 24. Thiacetazone
  - 25. Tramadol hydrochloride
  - 26. Zolpidem";

 In the said rules, after Schedule H, the following Schedule shall be inserted, namely:-

## "Schedule H1

(See rules 65 and 97)

- 1. Alprazolam
- 2. Balofloxacin
- 3. Buprenorphine
- 4. Capreomycin
- 5. Cefdinir
- 6. Cefditoren
- 7. Cefepime
- 8. Cefetamet
- 9. Cefixime
- 10. Cefoperazone
- 11. Cefotaxime
- 12. Cefpirome
- 13. Cefpodoxime
- 14. Ceftazidime
- 15. Ceftibuten
- 16. Ceftizoxime
- 17. Ceftriaxone
- 18. Chlordiazepoxide
- 19. Clofazimine
- 20. Codeine
- 21. Cycloserine
- 22. Diazepam
- 23. Diphenoxylate
- 24. Doripenem
- 25. Ertapenem
- 26. Ethambutol Hydrochloride
- 27. Ethionamide
- 28. Feropenem
- 29. Gemifloxacin

- 30. Imipenem
- 31. Isoniazid
- 32. Levofloxacin
- 33. Meropenem
- 34. Midazolam
- 35. Moxifloxacin
- 36. Nitrazepam
- 37. Pentazocine
- 38. Prulifloxacin
- 39. Pyrazinamide
- 40. Rifabutin
- 41. Rifampicin
- 42. Sodium Para-aminosalicylate
- 43. Sparfloxacin
- 44. Thiacetazone
- 45. Tramadol
- 46. Zolpidem

**Note.-** Preparations containing the above drug substances and their salts excluding those intended for topical or external use (except ophthalmic and ear or nose preparations) containing above substances are also covered by this Schedule."

[F. No. X-11014/6/2010-DFQC]
ARUN K. PANDA, Jt. Secy.

Foot note: The principal rules were published in the Official Gazette vide notification No. F.28-10/45-H (1), dated 21<sup>st</sup> December, 1945 and last amended vide notification number G.S.R. 72 (E), dated the 8<sup>th</sup> February, 2013.