THE NORTH WEST FRONTIER PROVINCE DRUGS RULES, 1982

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S.O.H(H)TECH/DHC-1/79, dated 31-05-1982. In exercise of the powers conferred by Section 14 of the Drugs Act, 1976 (XXXI of 1976), the Government of North West Frontier Province is pleased to make the following Drugs Rules, namely:-

Part I

Preliminary

1. Short title and commencement. (1) These rules may be called the North West Frontier Province Drugs Rules, 1982.

(2) They shall come into force at once.

2. Definitions. In these rules, unless there is anything repugnant in the subject or context:-

(a) "Act" means the Drugs Act, 1976 (XXXI of 1976);

(b) "Analyst" means an Analyst appoint by Government under the Act;

(c) "Board" means the Quality Control Board for the North West Frontier Province set up under Section 11;

(d) "Form" means a form specified in Schedule "A";

(e) "Government" means the Government of North West Frontier Province;

(f) "Inspector" means an Inspector appointment by Government under the Act;

(g) "Licensing Authority" means an authority specified in Rule 12;

(h) "Narcotics" means the drug specified in Schedule B;

(i) "Pharmacy" means a shop, store or place where drugs are compounded or prepared on prescriptions;

(j) "Schedule" means a schedule to these rules; and

(k) "Section" means section of the Act;

Part II

Appointment and function of Enforcement Staff

3. Procedure in case of prosecution. (1) An inspector and an Analyst shall submit monthly returns in Form 1 and Form 2, respectively, to the Board and a summary on the overall situation of quality control in the

area under their respective jurisdiction and the Board shall maintain such information in a manner so as to monitor, the quality of all the drugs sold and to keep watch on the performance of all manufactures.

(2) The Board shall, as far as possible, meet at least once in a month and review situation of the quality control of drugs on the whole including consideration of any specified point arising during the period on the working of the various firms, drugs testing laboratories and inspectors.

(3) The Board shall, examine carefully the cases referred to it by any Inspector under the Act, and provide an opportunity of hearing to the accused to explain this position before directing the Inspector to prosecute the accused.

(4) Before referring any cases to the Drug Court, the Board shall ascertain the names of the directors, partners and employees of the company, corporation, firm or institution who the *prima facie* responsible for the Commission of the offence under the Act, or the rules made thereunder and allow an Inspector to institute prosecution only against such persons.

(5) Where a drug is found to be substandard or adulterated the Board, before referring the case to the Drug Court, on the request of the complainant or the accused may cause a sample of the drug to be tested and analysed and provide an opportunity to the accused to explain his position in view of the contents of the report of the test:

Provided that where the retesting is ordered by the Board under this rule, the test result shall be final.

4. Qualifications etc. of Inspector and Analyst. (1) No person shall be appointed as an Inspector unless he possess a degree in Pharmacy from a Pakistan University or any other institution recognised for this purpose by the Pharmacy Council of Pakistan and has at least one year's experience of the manufacture, sale, testing or analysis of drugs or in the drugs Control Administration or in a hospital, or pharmacy.

(2) No person shall be appointed as an analyst unless he possess a degree in pharmacy from a Pakistani University or any other institution recognised for this purpose by the Pharmacy Council of Pakistan and has at least five year's experience in the manufacture, testing or analysis of drugs or in the Drugs Control Administration:

Provided that if a person of the requisite qualification is not available, a person possessing a degree in medicine or Master's degree in Pharmaceutical Chemistry, Micro Biology or pharmacology with five years experience in testing of drugs and medicines in public health laboratories may be appointed:

Provided further that the provision of this rule shall not apply to the Inspectors and Analyst who were appointed as such on regular basis before the coming into force of these rules.

(3) Government may, by notification in the official Gazette, appoint a person possessing a degree in Pharmacy, medicine and Master's degree in pharmaceutical Chemistry or Microbiology or Pharmacology as an *ex-officio* Inspector from amongst its officers working in the Drug Administration or in any other recognized pharmacy or medical institution, who otherwise does not fulfil the qualifications laid down in sub-rule (1):

Provided that the *ex-officio* Inspector shall be appointed for the purpose of,--

(i) Conducting inspection of any premises wherein any drug is sold or is stocked or exhibited for sale or distribution;

(ii) Conducting Inspection of storage arrangements and relevant records and registers in such premises; and

(iii) taking samples of any drug which is being sold or is stocked or exhibited for sale or is being distributed.

(4) Government may, by notification in the official Gazette, for the exercise of such powers as may be specified in the notification appoint as *ex-officio* Analyst any person who holds a degree in pharmacy or Medical or Master's Degree in Pharmaceutical Chemistry or Microbiology or Pharmacology and is engaged in testing and analysis in a Government Testing Laboratory or in a Chemical Examiner's Laboratory or is working in a pharmaceutical or medical educational or Research Institution.

5. Duties of Inspectors. Subject to the instruction of the licensing authority, it shall be the duty of an Inspector,--

(a) to inspect not less than twice a year all establishments of drugs licensed for sale and once year all establishment licensed for manufacture of drugs within the area assigned to him, and to keep record of such inspections;

(b) to satisfy himself that the conditions of the licenses are being observed;

(c) to take and send for testor analysis, sample of any drugs where there is reason to suspect that the drug is being manufactured or sold stocked or exhibited for sale in contravention of any of the provisions of the Act;

(d) to investigate any complaint in writing which may be made to him and furnish the report in respect thereof to the licensing authority;

(e) to institute prosecution in respect of contravention of the Act and the rules;

(f) to maintain record of all inspections made and actions taken by him in the performance of his duties, including the taking of

samples and the seizure of stocks, and submit reports of such record as may be required by the licensing authority; and

(g) to make such enquiries and inspections as may be necessary to stop manufacture and sale of drugs in contravention of the Act and these Rules.

6. Duties of Analyst. An analyst shall cause to be analysed or tested such complex of drugs as may be sent to him under the Act, and shall furnish report of the results of tested and analysis in Form 3 in accordance with the Act and these rules.

(2) As analyst shall cause to be tested and analysed such samples of drugs as may be sent to him in writing from a department of Government or any other public institution and shall furnish report of the result of test and analysis to the Department of the public institution concerned.

(3) An analyst shall forward monthly reports giving results of samples tested and analysed during the period under report with a view to their publication at the description of the Federal Government and furnish such other information as may be required by that Government.

7. Prohibition of disclosure of information. Except of the purpose of official business or when required by Court of law and inspection of Analyst shall not disclose to any person any information acquired by him in the course of his official duties.

8. Forms of order not to be dispose of stock. An order in writing by an Inspector under clause (i) of sub-section (1) of Section 18 requiring a person not to dispose of any stock in his possession shall be in Form 4.

9. Forms of institution of purpose of taking samples. (1) Where an Inspector takes a sample of drug under clause (c) of sub-section (1) of Section 18 for the purpose of test or analysis, he shall intimate such purpose in writing in Form 5 to the person from whom he takes it and where he seizes stock of a drug of other material under clause (f) of Section 18 the receipt for such drug and material shall be in Form 6.

(2) The Inspector shall send a portion of the sample or the container to the analyst for test or analyses under clause (i) of sub-section (3) of Section 19 through a memorandum in Form 7.

(3) In case the sample is delivered to the analyst by an indirect means such as post, a copy of the memorandum, a specimen impression of the seal or mark used to seal the packet together with the specimen impression of the person from whom the sample is drawn shall be sent to the analyst separately by registered post or by hand.

10. Procedure on receipt of samples from Inspection. On receipt of a package from all Inspector containing a sample for test and

analysts, the analyst shall compare the seals of the package with specimen impression received separately and shall note the condition of the seal on the package and after the test or analysis has been completed, he shall forthwith supply the analysis with protocols under the Act.

11. Fee for test and analysis of drugs. The fee for test and analysis of drugs in respect of samples sent by persons other than an Inspector or a Government Institution shall be determined by the analyst or the person incharge of the Government laboratory in accordance with the fees specified in Schedule 'C'.

Part III

Sale of Drugs

12. Licensing Authority. (1) The Secretary to Government, Health Department shall be the licensing authority for the purpose of these rules.

(2) The licensing authority may, by order in writing, authorise any person under his control to sign the licenses and to exercise such other powers and in respect of such areas as may be specified in the order.

13. Type of licenses to sell drugs. The licenses under these rules shall be of the following types, namely,--

(i) Licence to sell drugs by way of retail sale;

(ii) Licence to sell drugs by way of wholesale;

(iii) Licence to sell narcotics; and

(iv) Licence to sell drugs in a pharmacy.

14. Application for licence to all drugs and fees therefore. Application for the grant or renewal of a licence referred to in rule 13 shall be made in Form 8 to the licensing authority.

[1][(2) An application under sub-rule (1) shall be accompanied by the following fee:--

(i) in the case of Provincial Headquarter, one thousand rupees for fresh licenses and seven hundred fifty rupees for renewal;

(ii) in the case of divisional/district headquarter, seven hundred fifty rupees for fresh licenses and five hundred rupees for renewal; and

(iii) in other cases, five hundred rupees for fresh licences and three hundred rupees for renewal.]

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1. Substituted by Not. No. SO (Drugs) Health/3-5/92 (Vol. I), dated 25.11.1997. Before substitution read as:
(2) An application under sub-rule (I) shall be accompanied by a fee two hundred rupees in case of a fresh licence and one hundred rupees of a renewal.

(3) A fee of fifty rupees shall be paid for any change of proprietor or qualified persons or duplicate copy of the licence if the original is defaced, damaged or lost, and such copy of the licence shall bear the words "duplicate copy".

15. Forms of licence to sell drugs. (1) A licence to sell, store, exhibit for sale or distribute drugs by way of retail sale shall be issued in Form 9.

(2) A licence to sell, store, exhibit for sale or distribute drugs by way of wholesale shall be issued in Form 10.

(3) A licence to sell, store, exhibit for sale or distribute narcotics shall be issued in Form 11.

(4) A licence to sell drugs in a pharmacy shall be in Form 12.

16. Sale at more than one place. If drugs are sold, stored, exhibited for sale or distributed at more than one place, a separate licence shall be required in respect of each such place.

17. Duration of licence. A licence issued under these rules shall, unless sooner suspended or cancelled, remain in force for two years from the date of issue or with the disposal of the application for renewal of such licence whichever is later. An application of renewal of all licence shall be made within one month of the expiry thereof:

Provided that an application for renewal of a licence may be entertained by the licensing authority if such application is made within one month after the expiry of the licence and the licensing authority is satisfied that the application could not be made earlier for reasons beyond the control of the licencee.

(2) An application for renewal of licence shall be disposed within three months of the receipt of such application.

18. Pre-condition of the issue of licence. (1) The licensing authority shall not issue,--

(a) Licence in Form 9 and Form 12 unless;

(i) the premises have proper and adequate facilities for storage of drugs and for their protection from direct sunlight, dust or dirt including refrigeration facilities where necessary for preserving the properties of the drugs to which the licence applies;

(ii) the premises are clean and in hygienic and tidy condition; and

(iii) in the case of a pharmacy, the requirement laid down in Schedule "F" are complied with;

(b) Licence in Form 10 unless the applicant is an indentor, importer, manufacturer or distributor of a manufacturer drug and fulfils the conditions laid down is sub-clauses.

(i) and (ii) of clause (a) and

(c) Licence in Form 11 unless--

(i) the applicant possesses a licence in Form 9 or Form 10 or Form 11; and

(ii) the applicant has never been convicted of any offence under the Act.

(2) The sale of drug shall be supervised,--

(a) Under licence in Form 9 or Form 11 by a person--

(i) Who is registered under Section 24(1) (a) and (b) of the Pharmacy Act, 1967 (XI of 1967); or

(ii) Who was approved as qualified person for grant of drug sales licence under the West Pakistan Drug Rules, 1958; or

(iii) Who was on the 19th day of June, 1972 qualified for registration under Section 24(1)(b) of Pharmacy Act, 1967 (XI of 1967); or

(iv) Who has before the commencement of these rules passed the examination of compounder or dispenser and has complete two years period of apprenticeship under Section 24(1)(c) of the Pharmacy Act, 1967.

(b) Under licence in Form 10 by a person--

(i) Who fulfils the conditions laid down in clause (a), or

(ii) Who has been a student or apprentice in pharmacy under clause (iii) of sub-section (2) of Section 25 of the Pharmacy Act, 1967 (XI of 1967):

Provided that this provision shall be applicable after 2 years of the commencement of these rules;

(c) Under licence in Form 12 by a person who is registered as pharmacist under Section 24(1)(a) of Pharmacy Act, 1967 (XI of 1967) or by a person who is registered under Section 24(1)(b) of Pharmacy Act, 1967 (XI of 1967) and possess at least 3 years experience in compounding.

19. Condition of licence. (1) Licences of Form 9, 11 and 12 shall be issued subject to the conditions stated therein and to the following general conditions, namely,--

(a) The supply by way of retail sale of any drug shall be recorded suitably and such records, bills or conditions shall be preserved for a period of at least three years from the date of such sale;

(b) Drug specified in Schedule B and D and preparations containing such drugs shall not be sold by retails except on and in accordance with the prescription of a registered medical practitioner:

Provided that no such prescription shall be required for sale of these drugs to a registered medical practitioner, hospital, dispensary or any other institution approved by an order of the licensing authority for such sale:

(c) The sale of any drug specified in Schedule B and D by way of retail sale shall be recorded at the time of supply in a register specially maintained for the purpose and the serial No. of the entry in the register shall be entered in the prescription and the following particulars shall be entered in the register, namely.

(i) Serial No.

(ii) Date of sale.

(iii) Name of the prescriber.

(iv) Name of the patient/purchaser.

(v) Name of the drug.

(vi) Name of manufacturer.

(vii) Quantity.

(viii) Batch No.

(ix) Signature of the qualified person:

Provided that if the drug specified in Schedule (D) is sold on a prescription on which the drug has been sold on a previous occasion, it shall be sufficient if the entry in the register includes Serial No. the date of sale, the quantity sold, and sufficient references to any entry in the register recording the dispensing of the drug on a previous occasion.

(2) For the purpose of this rule, a prescription shall,--

(a) be in writing and signed by the person giving it with his usual signature and be dated by him;

(b) specify the name and address of the person for whose treatment it is given; and

(c) indicate the total quantities of drugs to be supplied and doses to be taken.

(3) All invoices and bills of purchase of drugs shall be reserved for a period of at least three years.

(4) Records shall be maintained of all purchases and sales of drugs by way of wholesale and such records shall be preserved for three years and shall include the following particulars, namely,--

(a) the date of purchase and sale;

(b) the name and address of the concern from which purchased and the concerns to which sold;

(c) the names of the drugs, their batch number, their dates of expiry, where applicable and the quantities; and

(d) the name of the manufacturer.

(5) Except as otherwise provided in these rules, all registers and records maintained under these rules shall be preserved for a period of not less than three years from the date of last entry.

(6) The licencee shall produce for inspection by an Inspector on demand all registers and records maintained under these rules, and shall supply to the Inspector such information as he may require.

(7) Substances specified in Schedule 'E' and falling under the list of poison and those specified in Schedule 'B' shall be stored in the retail shop,--

(a) in a part of the premises to which customers do not have access; or

(b) in an almiarh or cupboard or drawer locked, and reserved solely for the storage of such drugs.

(8) Substance falling under the list of poisons in Schedule 'E' shall be stored in containers, impervious to the poison and sufficiently stout to prevent leakage arising from the ordinary risk of handling and transport.

(9) A substance falling in the list of poison under Schedule 'E' when compounded and dispensed, shall be labelled with the word "Poison".

20. Cancellation and suspension of licences. (1) The Licensing Authority may, on the report of an Inspector or on its own motion, after giving the licensee an opportunity to show cause, by an order in writing stating the reasons therefore cancel a licence issued under these rules or suspend it for such period as it thinks fit, either wholly or in respect of some of the drugs to which it relates, if in its opinion, the licencee has failed to comply with any of the condition of the licence or with any of the provisions of the Act or these rules when the offence is of serious nature.

(2) A licensee whose licence has been cancelled or suspended may, appeal to the Appellate Board within sixty days of the date of such order.

SCHEDULE A

[See Rule 2(e)]

FORM 1

[See Rule 4(1)]

MONTHLY PROPERTY FROM INSPECTOR FOR THE MONTH OF SUMMARY OF INSPECTIONS

Place Inspected Manufactures.

No. of Firms Inspected.

No. of Firms found violating law-specify main offences.

No. of samples drawn, if any.

Remarks.

Shops Chemists and Druggists.

Others Please Specify.

DETAILS OF VIOLATIONS IN RESPECT OF DRUGS.

Reports of samples of drugs not in compliance with law

Name of Drugs

Regn. No. and Manufacturer's Name

Batch No.

Place of taking sample.

Date of despatch sample and name of Laboratory.

Date of receipt of test report with nature of results.

Action taken including details of seizure and sale restriction.

Copy of inspection report of the Pharmaceutical Manufacturing Unit should be supplied along with comments about overall situation of quality control.

FORM 2

[See Rule 4(1)]

DRUG TESTING LABORATORY

PROGRESS REPORT FOR THE MONTH .....

Number of sample in the beginning of month

Samples received during the month

Total

Tested

New

Old

Total

Samples upto standard with percentage.

Samples below standard.

Details of samples pending for more than 2 months.

Remarks/Reasons.

Spurious.....

Substandard.....

Adulterated.....

Counterfeit.....

Others.....

Total .....

DETAILS OF DRUG FOUND IN CONTRAVENTION OF LAW

DURING THE MONTH OF .....

S. No.

Name and Registration No. of drug

Batch No.

Manufactured by

Test Report No. date and nature of contravention

FORM 3

[See Rule 6]

CERTIFICATE OF TEST OR ANALYSIS BY THE DRUGS TESTING LABORATORY/GOVERNMENT ANALYST

1. Certified that the samples, bearing number ..... purporting to be a sample of ..... received on ..... with Memorandum No. .....

dated ..... from ..... has been tested/analysed and that the result of such test/analysis is as stated below.

2. The condition of the seals on the packet on receipt was as follows.

3. In the opinion of the undersigned the sample is not/is adulterated/sub-standard/misbranded/spurious as defined in the Drugs Act, 1976 for the reasons given below:

Director, Drugs Testing-Laboratory or other authorized officer/Government Analyst

Details of results of test or analysis, (with protocols of tests applied)

Director, Drugs Testing Laboratory or other authorized officer/Government Analyst

FORM 4

[See Rule 8]

ORDER UNDER SECTION 18(1)(f) OF THE DRUGS ACT, 1976 REQUIRING A PERSON NOT TO DISPOSE OF STOCK IN HIS POSSESSION

Whereas I have reason to believe that stock of drugs in your possession detailed below contravenes the provisions of Section ..... of the Drugs Act, 1976.

Now, therefore I hereby direct you not to dispose of the said stock for a period of ..... days from this date.

Date .....Inspector

Details of stock of drugs

Date .....Inspector

FORM 5

[See Rule 9(1)]

Intimation of purpose to person from whom sample is taken to

I have this day taken from the premises of ..... situated at ..... samples of the drugs specified below for the purpose of test/anyalysis details of samples drawn.

Name of drug

Name of Manufacturer

Registration No.

Batch No.

Quantity

Bill No.

Value

Dated ..... Inspector .....

FORM 6

[See Rule 9(1)]

Receipt for stock of drug and other material seized under Section 18(1)(f) of the Drugs Act, 1976.

The stock of drug detailed below has this day been seized by me under the provisions of clause (f) of sub-section (1) of Section 18 of the Drugs Act, 1976 from the premises of .....

Situated at .....

Dated .....

Inspector.....

Details of Drugs, other material and articles of drugs seized

Dated ..... Inspector .....

FORM 7

[See Rule 9(2)]

MEMORANDUM TO ANALYST

Serial No. of Memorandum.....

Form

To

The Analyst.

The portion of sample/container described below is sent herewith for test or analysis under the provisions of clause (i) of sub-section (3) of Section 19 of the Drugs Act, 1976.

The sample is of the drug ..... and purport to contain

.....

The portion of sample has been marked by me with the following marks:

Dated ..... Inspector .....

FORM 8

[See Rule 14(1)]

APPLICATION FOR A LICENCE TO SELL STOCK AND EXHIBIT FOR SALE AND DISTRIBUTE DRUGS

I/We ..... of .....

.....

..... hereby apply for a licence to sell

1. Drugs by way of retail sale.
2. Drugs by way of wholesale.
3. Narcotic and other drugs.
4. Drugs in Pharmacy.

on the premises situated at .....

.....

2. The sale of drugs will be under the personal supervision of.....

.....

(Name)..... (Qualification) .....

(Name)..... (Qualification) .....

3. I/We/am/are submitting herewith the following documents evidence of being importer/indenter/manufacturers, agent/distributor, etc.

Treasury Challan(s) for Rs. .....

Date.....

Delete whichever is not applicable.

Signature

Name and Permanent Home Address.

FORM 9

[See Rule 15(1)]

LICENCE TO SELL, STOCK AND EXHIBIT FOR SALE AND DISTRIBUTE DRUG BY WAY OF RETAIL SALE

..... is hereby licensed to sell stock and exhibit for sale and distribute drugs by way of retail sale on the premises situated at..... subject to the condition specified below and to be provisions of the Drugs Act, 1976 and the rules made thereunder.

2. This licence will be in force for a period of two years from the date given below.

.....

3. Name(s) of qualified person(s).

1. _____

2. _____

4. Address of down/godowns where drugs shall be stored.

Date ..... LICENSING AUTHORITY

CONDITIONS OF LICENCE

1. This licence shall be displayed in a prominent place in part of the premises open to the public.

2. The licensee shall comply with the provisions of the Drugs Act, 1976, and the rules made thereunder for the time being in force.

3. The licensee shall report forthwith the Licensing Authority any change in the qualified staff incharge.

4. No drug requiring special storage conditions of temperature and humidity shall be stored or sold unless the precautions necessary for preserving the properties of the contents have been observed throughout the period during which it has been in possession of the licence.

FORM 10

[See Rule 15(2)]

LICENSE TO SELL, STOCK AND EXHIBIT FOR SALE AND DISTRIBUTE DRUGS BY WAY OF WHOLESALE

..... is hereby licensed to sell, stock and exhibit for sale and distribute drugs by way of wholesale on the premises situated at.....

.....
subject to the conditions specified below and to the provision of the Drugs Act, 1976 and the rules made thereunder.

2. The licence will be in force for two years from the date given below.

3. Name(s) of qualified person(s).

LICENSING AUTHORITY

1 .....

2 .....

CONDITIONS OF THE LICENCE

1. This licence shall be displayed in a prominent place in parts of the premises open to the public.

2. The licensee shall comply with the provisions of Drugs Act, 1976 and the rules made thereunder for the time being in force.

3. The licensee shall report forthwith to the Licensing Authority any change in the qualified staff incharge.

4. No drug requiring special storage conditions of temperature and humidity shall be sold unless the precautions necessary for preserving the properties of the contents have been observed throughout the period during which it has been in possession of the licensee.

Address of the godown/godowns where the drugs are stocked should also be given.

FORM 11

[See Rule 13(3)]

LICENCE TO SELL NARCOTIC AND OTHER DRUGS SPECIFIED IN SCHEDULE B

..... holder of Licence No. ..... on Form 9/10 is licensed to self/stock narcotic and other drugs specified in Schedule B on the premises situated at ..... subject to the conditions specified below and to the provisions of the Drugs Act, 1976 and rules made thereunder.

2. This licence will be in force for two years from the date given below or till the validity of licence in Form 9/10.

3. Name(s) of qualified person(s).

Date .....

LICENSING AUTHORITY

CONDITIONS OF THE LICENCE

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.

2. The licensee shall report forthwith to the licensing authority any charge in qualified staff incharge.

3. No drug to which this licence applies shall be sold unless the precaution necessary for preserving the properties of the contents have been observed throughout the period during which it has been in possession of the licensee.

Address of the godown/godowns where the drugs are stocked should also be given.

FORM 12

[See Rule 15(4)]

Licence to sell drugs in Pharmacy ..... hereby licensed to compound or prepare on prescription the drugs and distribute drugs by way of retail sale on the premises situated at ..... subject to the conditions specified below and to the provisions of the Drugs Act, 1976 and the rule made thereunder.

2. This licence will be in force for a period of two years from the date given below.

3. Name(s) of qualified person(s).

(1) .....

(2) .....

4. Addresses of godown/godowns where drugs shall be stored.

Date ..... LICENSING AUTHORITY

CONDITIONS OF THE LICENCE

1. This licence shall be displayed in a prominent place in part of the premises open to the public.

2. The licensee shall comply with the provisions of the Drugs Act, 1976, and the rules made thereunder for the time being in force.

3. The licensee shall report forthwith to the Licensing Authority any change in the qualified staff incharge.

4. No drug requiring special conditions of temperature and humidity shall be stored or sold unless the precautions necessary for preserving the properties of the conventions have been observed throughout the period during which it has been in possession of the licensee.

SCHEDULE 'B'

[See Rule 2(h) and 19(1)(b) and (c)]

NARCOTICS

Acetorphine	Dipipanone
Acelylmethadol	Dorotebano
Allyiprodine	Econino
Alphacelyiemethadol	Ethylmothylhiambutone
Alphamethadol	Etonitazene
Alfantil	Etorphine
Alphaprodine	Etoxeridne
Aplrazolan	Fantayl
Atileridine	Furethidine
Benzethidin	Heroin
Benzylmorpine	Hydrocodone
Betacoylethadol	Hydromorphinol
Betaprodine	Hydromorphone
Beitramide	Hydroxyperthidine
Bronozapan	Isomethadone
Bupremorphine	Katobemidone
Cannabis	Lerazopam