

SCHEDULE	FOR
A	application for the licenses, issue and renewal of licenses, for sending memoranda under the act
B	Rate of fee for test or analysis by the Central Drugs Laboratory or the Government Analyst.
2nd schedule	Standard to be complied with by
C	List of biological and other special products whose import, sale, import, distribution and manufacture are governed by special provisions.
C1	List of other special products whose import, sale, import, distribution and manufacture are governed by special provisions.
D	List of drugs exempted from the provisions of import of drugs.
E1	List of poisonous substances under the Ayurvedic , Sidha and Unani systems of medicine.
F&F1	provisions applicable to the production, testing, storage, packing, and labeling of biological and other special products
F2	Standard for surgical dressings
F3	Standards for sterilized umbilical tapes.
FF	Standards of ophthalmic preparations .
G	List of substances that are to be used only under medical supervision and which are to be labeled accordingly.
H	List of prescription drugs which are sold by retail and only on prescription of registered Medical Practitioner.
J	Diseases or ailments which a drug may not claim to prevent are cure.
K	Drugs exempted from certain provisions relating to the manufacture of drugs.
M	GMP requirements of factory premises, plants and equipments.
M1	Requirements of factory premises for the manufacture of homoeopathic preparations.
M2	Requirements of factory premises for the manufacture of cosmetic preparations.
N	List of minimum equipment for efficient running of pharmacy
O	Standards for disinfectant fluids .
P	Life period of drugs.
Q	List of coal tar colors permitted to be used in cosmetics.
R	Standards for mechanical contraceptives.
S	Standards for cosmetics.
T	Requirements of factory premises and hygienic conditions for ayurvedic and Unani drugs.
U	Particulars to be shown in manufacturing, raw material and analytical records of cosmetics.
V	Standards for plant and proprietary medicines .
W	List of drugs which are to be marketed under generic names only.
X	List of drugs whose import, manufacture and sale and labeling and packaging are governed by special provisions .
Y	Requirements and guidelines on clinical trials for import and manufacture of new drugs.