

## ADMINISTRATIVE BODIES



### \*ADVISORY:

- Drugs Technical Advisory Board (D.T.A.B).
- Drugs Consultative Committee(DCC).

### \*ANALYTICAL:

- The Central Drug Laboratory (CDL).
- Drugs Control Laboratories In State.
- Government Analyst & Power.

### \*EXECUTIVE:

- Licensing Authorities.
- Customs Collections.
- Drug Inspector.

## DRUG TECHNICAL ADVISORY BOARD

**FUNCTION:-**The Central Govt. Appoint the DTAB to advice the central & state govt. on technical matter arising out of the administration of this act.

### CONSTITUTION OF DTAB:

#### EX-officio Members:

- (i) The Director General of Health Services, who shall be Chairman;
- (ii)The Drugs Controller, India, ex officio;
- (iii)The Director of the Central Drugs Laboratory, Calcutta.
- (iv)The Director of the Central Research Institute, Kasauli.
- (v) The Director of Indian Veterinary Research Institute, Izatnagar.
- (vi) The President of Medical Council of India.
- (vii) The President of the Pharmacy Council of India.
- (viii) The Director of Central Drug Research Institute, Lucknow.



## Elected Members:

- (i) One person, to be elected by the Executive Committee of the Medical Council of India, from among teachers in medicine or therapeutics on the staff of an Indian university or a college affiliated thereto;
- (ii) One person to be nominated by the Central Government from the pharmaceutical industry;
- (iii) One pharmacologist to be elected by the Governing Body of the Indian Council of Medical Research;
- (iv) One person to be elected by the Central Council of the Indian Medical Association;
- (v) One person to be elected by the Council of the Indian Pharmaceutical Association;
- (vi) Two persons holding the appointment of Government Analyst under this Act, to be nominated by the Central Government.

**Nominated Members:**(i)The persons among the persons who are incharge of the drugs control in the state.

(ii)One person from the pharmaceutical Industry.

(iii)Two Government Analyst.

## DRUG CONSULTATIVE COMITTE (DCC):- Under section.7



### Functions:-

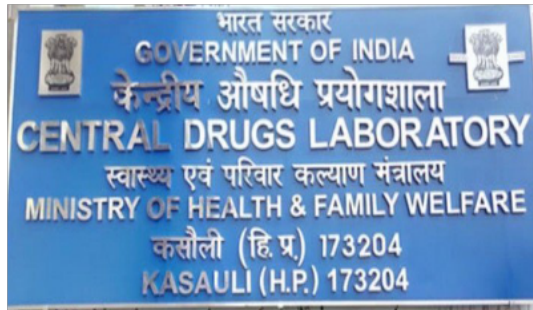
- (i)The advisory committee constitute by central government to adive central & state government and also DTAB.
- (ii)On matter to secure uniformity throughout india in administration of this act.

**CONSTITUTION:-** It consist of,

- 1.Two representatives of central government nominated by central government and,
- 2.One representative of each state government nominated by the concerened government.



## Central Drug Laboratory(CDL)-



(a)customs collectors (under sub-sec.2) or

(b)courts (under sub sec. 4 of sec.25)

### Government Analyst & Power:-

A)a graduate in Pharmaceutical Chemistry with a minimum of 5 years of experience after graduation in testing or analysis of drugs and pharmaceutical in laboratory under the control of Government Analyst

or

(b)a post-graduate in Medicine/Pharmacy/ India with Analysis of Drugs and Pharmaceutical as a

special

subject and at least 3 years of experience (in all cases) in testing



or analysis of drugs and pharmaceuticals under the control of

Government Analyst/ Head of approved analytical institution.

### Drug Inspector-

The central or state government by notification in the officialgazette appoint inspector having prescribed under Section 21

#### QUALIFICATIONS OF DI


-  Person must have a **degree** in Pharmacy/Pharmaceutical Chemistry/Medicine with specialization in Clinical Pharmacology/Microbiology from a recognized Indian University.
-  For inspection of manufacture of substances in Schedule C
  - 1) 18 months experience in **manufacture/ testing** of at least one of the substance specified in schedule C
  - 2) Gained experience of NLT 3 yrs **in inspection of firms** manufacturing any of the substances of Schedule C during their tenure as services as DI



## POWER OF DRUG INSPECTOR:-

1. Inspect:
2. Take samples of any drug or cosmetics:
3. Search
4. Enter and search
5. stop and search
6. give order
7. examine
8. exercise

**DUTIES OF INSPECTORS**



**Inspect for premises licensed for sale**

- **Inspect** NLT twice an year all establishments licensed for sale of drugs within the area assigned to him
- **Procure and send for tests or analysis**, if he has reason to think that the drugs are sold in contravention of provisions of Acts or Rules.
- To **investigate any complaints** made to him in writing & to institute prosecutions in respect to the breaches of the act.
- To **maintain all records** of inspections made & actions taken by him including taking of samples and seizure of stocks & to submit copies of such records to the CA.
- Make **enquiries** such inspections as may be necessary to detect sale of drugs in contravention to the Act.
- When so authorized by State Governments to **detain imported packages** which he has reason to suspect to contain drugs whose import is prohibited.

## Licensing authorities-

Central government appoints an authority called "licensing authority"

to issue licence for import of drugs. each state government

appoints licensing authorities to issue licence for manufacture

, distribution and sale of drugs and cosmetics for a specified areas

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