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Monthly Newsletter

For circulation among DCOIWA members only

In collaboration with



The Health Master

Published on 1st day of every month

## Landmark Seminar by FDA Haryana on Spurious drugs and NDPS Misuse

### FDA Haryana

The northern belt of India is set to benefit tremendously with the intensified action against **spurious drugs** and illicit **narcotic drugs and psychotropic substances (NDPS)**. The Food and Drugs Administration Haryana (FDA Haryana) recently organized a high-level, cross-boundary, **interstate coordination seminar** in Chandigarh an unprecedented event for any state government.

Not just any run-of-the-mill gathering, this was a high-level, **milestone meeting** that included drug regulators from seven states and important enforcement officers from the CID and Police Departments.

The assembly is a pivotal move toward drafting a cohesive regional enforcement plan to combat cross-border drug trafficking.

### Why is Inter-State Coordination Essential?

The core objective of the seminar was to take **mutual coordination** and rapid **intelligence** sharing to the next level



among adjoining jurisdictions that need to be on high alert.

Drug traffickers of **counterfeit medicines** and diverted NDPS drugs that have transformed into dangerous **intoxicantuse** don't recognize state boundaries.

According to Lalit Kumar Goel, Haryana's State Drug Controller (SDC), this high-level gathering was all possible from the strategic foresight of the leaders of FDA Haryana who

*(Continued on page 17)*



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**EDITORIAL****Rakesh Dahiya****Editor-in-Chief  
DCOIWA Newsletter****A Milestone Edition: Steering the Ship of Public Health Together**

Welcome to the **22<sup>nd</sup> Edition** of the DCOIWA Newsletter, published on December 1st, 2025! This edition feels like a true milestone, not just for the number itself, but for the sheer volume of critical updates and inspiring stories packed within these pages.

It's a testament to the fact that while our work—ensuring the safety and quality of medicines—is often challenging, it is also incredibly dynamic and impactful.

**Our Collective Voice in Action**

We begin by acknowledging the proactive steps taken by our association. The **Representation to the DCGI regarding the amendment of the Drugs Inspector Recruitment Rules** is a vital effort.

It shows that DCOIWA isn't just a platform for discussion; it's a powerful agent for positive change.

By advocating for improved rules, we are essentially investing in the **future quality** and professionalism of our entire regulatory body.

Similarly, the **President's Note** highlighting the analysis by the **Nagaland State Testing**

Laboratory reinforces a crucial message: **Standardization and rigorous quality control** must be a nationwide priority. Our strength lies in a robust, interconnected system where every state contributes to the national goal of public health protection.

**Shining a Light on Excellence and Experience**

This month, we are thrilled to feature an exclusive article by **Rakesh Dahiya, FDA Haryana**, on the **Procedure to obtain a license for a Blood Centre**.

This detailed guidance isn't just procedural; it's a deep dive into an area of regulation where lives literally hang in the balance.

Knowledge sharing like this is the bedrock of a competent and efficient regulatory body.

The news from **FDA Haryana** about their **Landmark Seminar on Spurious Drugs and NDPS Misuse** and the subsequent **20 years Imprisonment for selling NDPS in Haryana** sends a clear, powerful signal.

Our enforcement efforts are becoming stronger and the penalties are serving as serious deterrents.

This is the **frontline of drug control**, and our officers are delivering results.

We also celebrate the well-deserved **Promotions of FDA officers Karan Singh and Rakesh Dahiya**.

*(Continued on page 4)*



*(Continued from page 3)*

Seeing our peers advance is a reminder that dedication and hard work do not go unnoticed. Conversely, we extend our heartfelt best wishes and gratitude to the retiring FDA officers in **Maharashtra**, whose years of service have laid the foundation for the protections we enjoy today. Their wisdom and experience are invaluable legacies.

## **Innovations and Avenues for Growth**

The regulatory landscape is moving fast! The new mandate for a **QR Code at every Medical Store for ADR reporting** is a fantastic leap into modern, patient-centric surveillance.

This simplifies the process for consumers and allows for quicker identification of potential drug risks.

Furthermore, the **CDSCO granting powers to Joint Drug Controllers for key approvals** is a smart move towards **decentralizing** and speeding up decision-making.

This efficiency will surely make our work more streamlined. The invitations to the **74th IPC 2025** and the **Odisha Pharma Summit 2025** are golden opportunities for DCOIWA members to stay ahead of the curve, network, and bring back cutting-edge knowledge.

## **More Than Just Rules: Knowledge and Camaraderie**

Beyond the headlines, the wealth of information in this edition—from **Key notes on Revised**

**Schedule M** to the detailed **NSQ drugs lists** and the comprehensive **FAQs**—equips every member with the necessary tools for their daily work.

And finally, we must mention the importance of connection.

**Honoring Sh. Rajeev Singh Raghuvanshi, DCGI, with DCOIWA Membership** and the inclusion of **FDA memorable moments** and a much-needed **Laughter Dose** reminds us that this association is a **family**.

We support each other, we learn from each other, and we celebrate together.

As we close out the year, let us take pride in the crucial, often unsung role we play in safeguarding the nation's health. Please read the whole newsletter, absorb the insights, and continue to serve with dedication.

We eagerly await your feedback and participation for the next edition!





Representation to DCGI



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(Regd.No. 634 of 2022)

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Ref. No.: DCOIWA/NP/2025-26/101

Date: 26th Nov.2025

To
Shri.Rajiv Singh Raghuvanshi jee,
The Drugs Controller General of India
Central Drugs Standard Control Organisation
New Delhi
emial: dci@nic.in

Subject: Request for Amendment of the Recruitment Rules for the Post of Drugs Inspector to Include Pharm.D Qualification

Respected Sir,

I respectfully submit the following for your kind consideration with reference to the Ministry of Health and Family Welfare Notification G.S.R. 832(E) dated 11th November, 2025, issued under the proviso to Article 309 of the Constitution. This notification supersedes the Central Drugs Standard Control Organisation (Drugs Inspectors) Recruitment Rules, 2010 and prescribes the revised method of recruitment for the post of Drugs Inspector (419 posts) under the Directorate General of Health Services, Ministry of Health and Family Welfare. (Annexure-1 enclosed)

As per the currently prescribed educational qualifications for direct recruitment, the notification specifies:

Bachelor's degree in Pharmacy or Pharmaceutical Sciences or Medicine with specialisation in Clinical Pharmacology or Microbiology from a recognised University or Institute.

In this regard, I humbly request that the qualification Doctor of Pharmacy (Pharm.D) be included among the eligible qualifications for the post of Drugs Inspector.

Justification for Inclusion of Pharm.D

1. Statutory Recognition of Pharm.D

The Pharm.D Regulations, 2008 were notified in The Gazette of India, No. 19, Part III, Section 4, dated 10th May 2008, under Section 10 of the Pharmacy Act, 1948.

These regulations were approved by the Government of India vide letter No.

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V.13013/1/2007-PMS dated 13th March 2008, establishing the Pharm.D as a fully recognised and government-approved qualification.

2. Curriculum and Training Aligned to the Role of Drugs Inspector

The Pharm.D programme includes:
Five years of structured academic study
One year of internship/residency with exposure to pharmacy practice, regulatory affairs, drug safety, pharmacology, and pharmacovigilance

The training equips graduates with hands-on experience essential for drug inspection, quality assurance, and regulatory functions.

3. Parity with State Recruitment Rules

Several State Drug Control Departments have already included Pharm.D as an eligible qualification for Drugs Inspector positions, such as:

- Odisha - OPSC Advertisement No. 8 of 2022-23 (Annexure-2)
Telangana - TSPSC Advertisement No. 21/2022 dated 08/12/2022 (Annexure-3)
Maharashtra - As reported in The Times of India (Annexure-4)

This reflects an emerging national consensus on the suitability of Pharm.D graduates in regulatory roles.

4. Relevance to Drug Safety, Quality Control & Clinical Knowledge

Pharm.D graduates are trained extensively in:

- Clinical Pharmacology
Drug Interactions & Safety Monitoring
Rational Use of Medicines
Regulatory & Quality Standards

These competencies directly support the technical and clinical responsibilities outlined under the Drugs and Cosmetics Act and Rules.

Prayer

In view of the above, we earnestly request that the Pharm.D qualification be formally added to the Recruitment Rules for the post of Drugs Inspector under CDSCO. This inclusion would:

Handwritten signature

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- Ensure fairness, uniformity, and equitable opportunities for pharmacy professionals
Strengthen India's drug regulatory framework
Enhance public health safeguards through clinical and regulatory expertise
We sincerely request your favourable and urgent consideration of this matter.

Thanking you,

Yours faithfully,

[Handwritten signature]

(G. Koteswar Rao)
National President
(DCOIWA)

Copy forward to:

- 1. Shri Jaya Prakash Nadda, Hon'ble Union Health Minister, Minister of Health and Family Welfare, Room No. 402D, Nirman Bhavan, New Delhi - 110 011
Email: india-hfm@gov.in
2. Smt. Punya Shila Srivastava, Union Health Secretary, Minister of Health and Family Welfare, Room No. 156A, Nirman Bhavan, New Delhi - 110 011
Email: secyhfww@nic.in
3. To registrar, pharmacy council of India, New Delhi.
Email: registrar@pci.nic.in



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## Press Note: Nagaland

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**Mr. Tapan Choudhry**  
Cell: 9230610226 - West Bangal

Date: 26 November 2025

Place: Kohima, Nagaland

Lt. No.: DCOIWA/NV26/2025/01

### Press Note

The analysts of the Nagaland State Drug Testing Laboratory, Kohima, have successfully completed an intensive week-long training program held from 21st to 26th November 2025. The program was organised by Drugs Control Officers India Welfare Association (DCOIWA) in coordination with Industry Subject Matter Expert (SME) Mr. K. Sridhar, Former Director, Sipra Labs, Hyderabad.

This initiative was aimed at strengthening the laboratory's technical competence and fostering a robust culture of regulatory compliance.

During the training, participants were oriented on the fundamental principles and regulatory requirements of Quality Management Systems as defined under Schedule L1 of the Drugs and Cosmetics Act, WHO Good Practices for Pharmaceutical Quality Control Laboratories (Annex 1, WHO TRS 957), and ISO/IEC 17025.

The modules covered Good Analytical Practices (GAP), laboratory safety, and structured compliance management using the 6M framework—Man, Machine, Mother Nature, Method, Material, and Measurement.

Analysts also underwent hands-on scientific and operational training on key analytical instruments and techniques integral to the laboratory's routine operations. These included UV-Visible Spectrophotometry, HPLC, Polarimetry, Analytical Balances, Refractometry, FTIR, Water Purification Systems, Auto-Titrators, and Dissolution Apparatus. Special focus was placed on the importance of calibration and the instrument-specific calibration procedures currently practiced in the laboratory.

As part of the laboratory's continuous capacity-building efforts, DCOIWA along with the SME will continue to conduct weekly virtual sessions covering quality systems, instrumentation functionality, laboratory safety, and regulatory expectations in pharmaceutical analysis.



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## Press Note: Nagaland

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The overarching goal of this initiative is to support the Nagaland State Drug Testing Laboratory in its journey towards achieving ISO/IEC 17025 (NABL) accreditation, and to position it as one of the country's leading drug testing facilities. The program also aims to strengthen analytical excellence through practical skill-building and structured performance assessments, ensuring that all analysts consistently operate at the proficiency levels required in modern pharmaceutical testing laboratories.

DCOIWA extends heartfelt thanks to Shri Ethungbemo Ezung President, DCOIWA Nagaland chapter and Shri Anoop Khinchi, IAS, Commissioner and Secretary, Health & Family Welfare, Government of Nagaland, for inviting us and providing this valuable opportunity.

Regards,

**G. Koteswar Rao**  
National President,



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# Invitation to DCOIWA for 74th IPC 2025



## 74<sup>th</sup> Indian Pharmaceutical Congress, 2025 Bengaluru

Hosted By: Association of Pharmaceutical Teachers of India  
Organised By: Indian Pharmaceutical Congress Association  
Organised at: Bangalore International Exhibition Centre (BIEC)



Theme: AI & Technology in Pharma: Educate, Innovate and Transform

Nov 27, 2025

To  
**Mr. Koteswar Rao**  
President, Drug Control Officers Welfare Association (DCOWA)

**Subject:** Invitation to grace the Inaugural Function of the 74<sup>th</sup> Indian Pharmaceutical Congress (IPC)

Respected Sir,

Greetings from the Organising Committee of the 74<sup>th</sup> Indian Pharmaceutical Congress (IPC) 2025.

It gives us immense pleasure to extend our cordial invitation to Your Honour to grace the **Inaugural Function of the 74<sup>th</sup> Indian Pharmaceutical Congress** is scheduled to be held on **19/12/2025** at 10am at **Bangalore International Exhibition Centre, Madanayakahalli, Bengaluru** (BIEC).

The Indian Pharmaceutical Congress is landmark annual event that brings together leading scientists, uniting researchers, academicians, industry leaders, and policymakers from across the country and abroad to discuss advances, challenges, and opportunities in the pharmaceutical sector. This prestigious event is organized by the Indian Pharmaceutical Congress Association (IPCA) and will be hosted by the Association of Pharmaceutical Teachers of India (APTI).

Your august presence and address will be a source of great inspiration to the delegates and students attending the conference and will add immense value to the occasion.

We would be deeply honoured by your acceptance of this invitation and look forward to welcoming you to the 74th Indian Pharmaceutical Congress.

With regards,



Dr. Milind J  
Umekar  
President, IPCA,  
74<sup>th</sup> IPC



Mr. Harish  
Kumar Jain  
LOC Chairman  
74<sup>th</sup> IPC



Dr. Deependra  
Singh  
Organising  
Secretary  
74<sup>th</sup> IPC



Dr. Raman  
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**22**

**DCOIWA News**

December 2025

# 74th IPC 2025 19-21 December 2025 Bengaluru

19-21 DECEMBER 2025

# 74<sup>th</sup> INDIAN PHARMACEUTICAL CONGRESS



# IPC



**B E N G A L U R U**

**22****Invitation to Odisha Pharma Summit 2025**

Hemant Sharma, IAS  
Additional Chief Secretary

**ODISHA**  
NEW OPPORTUNITIES**1**

D.O. No. 11729/ACS.,  
IND-HI2-INVPRO-0008-2025  
Dated: 26/11/2025

Dear Sir,

**Sub:** Invitation to Odisha Pharma Summit 2025

The Government of Odisha is organising **Odisha Pharma Summit 2025**, a flagship initiative to catalyse growth and investment in the pharmaceutical and healthcare sector, in collaboration with leading industry stakeholders.

The **Odisha Pharma Summit 2025**, with **FICCI** as the National Industry Partner, is a first-of-its-kind stakeholder interaction aimed at shaping the next phase of growth in Pharma & Healthcare sector in Odisha. The event will be chaired by **Shri Mohan Charan Majhi, Hon'ble Chief Minister of Odisha**.

The summit is being convened with the intent to engage key stakeholders across the value chain, manufacturers, investors, skilling partners, and pharma associations, to present Odisha as a new emerging destination in the pharmaceutical manufacturing and healthcare sector.

The details of Odisha Pharma Summit 2025 are as per below:

**Date: Tuesday, 16<sup>th</sup> December 2025**

**Time: 10:00 AM onwards**

**Venue: Plumeria Hall, Taj Vivanta, Bhubaneswar, Odisha**

The Government of Odisha is committed to establishing a robust ecosystem for the Pharmaceutical and Medical Devices manufacturing sector by developing integrated industrial infrastructure and enabling seamless regulatory facilitation. In addition, concerted efforts are underway to strengthen workforce skilling and capacity building through dedicated training programs aligned with industry needs. Simultaneously, the State is reviewing fiscal and non-fiscal incentives to attract investment and ensure a globally competitive manufacturing environment, positioning the region as a preferred destination for healthcare, pharma and related sectors.

**Industries Department, Government of Odisha**  
Kharavela Bhavan, Bhubaneswar-751001, Odisha, INDIA  
☎ +91-674-2536640, 2390253  
✉ indsec.or@od.gov.in 🌐 www.investodisha.org



# Invitation to Odisha Pharma Summit 2025

1

We would be honored to invite you, your leadership team and your member companies to the Summit. We believe this interaction will provide your members with an excellent opportunity to directly engage with the State leadership. We look forward to your presence in this important dialogue.

For any queries or coordination support, you may please contact **Mr. Deepak Singh, Nodal Officer, IPICOL** (deepaksingh@investodisha.org, +91 9692210100), **Mr. Arpan Sanyal, IPA** (arpan.sanyal@pwc.com; +918052210009) or **Mr. B. K. Nayak, FICCI** (bk.nayak@ficci.com; +91 99457 90735).

A line of confirmation is requested from your end. Alternatively, associated members of your esteemed organization may register their interest for the Odisha Pharma Summit 2025 via the QR code below. We look forward to welcoming you to **Odisha Pharma Summit 2025** and to partner with you in shaping a robust and future-ready pharma ecosystem.

Yours sincerely,

(Hemant Sharma)

QR code for registration:



Link: [bit.ly/3XftNYz](https://bit.ly/3XftNYz)

**Odisha Pharma Summit 2025  
to be held on 16th December 2025  
at Bhubaneswar, Odisha**

**Procedure to obtain license for Blood Centre (Blood Bank)****1****Rakesh Dahiya**

**Editor-in-Chief,  
DCOIWA News  
Organising Secretary,  
DCOIWA  
Asstt. State Drugs Controller,  
FDA Haryana**

**How to obtain manufacturing license for Blood Centre / Bank****Documents required for obtaining Blood Centre**

For obtaining manufacturing license for Blood Centre, the list of documents required is provided below. Download the pdf file and prepare the documents accordingly.

**[Documents-required-for-obtaining-Blood-Centre](#)****Procedure for obtaining license**

Procedure for obtaining manufacturing license for Blood Centre. Download the pdf file for more detail and prepare the documents accordingly.

**[Procedure-for-obtaining-Blood-Centre](#)****Form**

Download the pdf file and prepare the Form accordingly and submit the required fee.

**Form-27C**

Download the below pdf files for various Forms which suits your requirement

**[Form-27C](#)****Download area requirement**

Click below link to download the

**requirement for the manufacturing of Drugs, Cosmetics, Homoeopathic and Blood Centre****[Area requirement for manufacturing](#)**

Submit your application (Online and / or hard copy) to DCGI and State Drugs Controller / Licensing Authority of your area after completing all the required documents.

Note: Requirements of some of documents and procedure for submission of application may vary from State to State

**License conditions**

Conditions of licenses is to be maintained after obtaining the required Blood Centre. Download the pdf file for ready reference.

**[License-conditions-for-Blood-Centre](#)****License renewal fee: Procedure**

Procedure for submission of license renewal fee. Download the pdf file for more detail and prepare the documents accordingly.

**[Procedure-for-License-renewal-fee](#)****Requirements for running a Blood Centre****Part X-B**

Requirements for the collection, storage, processing and distribution of Whole Human Blood, Human Blood Components by Blood Banks / centres and manufacture of blood products are mentioned in "part X B of drugs and cosmetics act 1940 and rules framed thereunder. For more details click below:

**[Requirements-for-running-a-Blood-Bank-Part-X-B](#)****Part XII-B**

*(Continued on page 15)*

## Procedure to obtain license for Blood Centre (Blood Bank)

**2**

Requirements for the functioning and operation of a blood Centre and / or for preparation of blood components are mentioned in "part XII B of drugs and cosmetics act 1940 and rules framed thereunder. For more details click below:

[Requirements-for-running-a-Blood-Centre – Part XII B](#)

### Bulk transfer of Blood and Blood Components

According to the notification GSR No. 328 (E) dt 03-04-2017, bulk transfers of whole human blood and blood components are permitted to other blood centers.

National Blood Transfusion Council in its meeting of 05.08.2015 has also issued some guidelines and conditions for the Bulk transfer of blood and its components from one blood centre to another blood centre.

#### Procedure

The formats for the request and issue of a bulk transfer of blood and its components are as follows:

**Request Form for Recipient Blood Centre: Click the below link to download the Form**

[Request-Form-for-Recipient-Blood-centre](#)

**Issue Form for Supplier Blood Centre: Click the below link to download the Form**

[Issue-Form-for-Supplier-Blood-centre](#)

#### Conditions

**Bulk transfer of blood and blood components amongst licensed blood centres in the country would henceforth**



**be allowed under the following conditions:**

- Bulk transfers of whole human blood and blood components can be done between licensed blood centres.
- Bulk transfers of whole human blood and blood components can be done among Blood centres in the state or across the state borders.
- All transfers shall be done at the recommended temperature and as per prescribed storage conditions for whole human blood and blood components.
- The recipient blood banks shall not further transfer units obtained from another blood bank except to another blood storage centre or a patient
- The supplier blood centre shall be responsible for compliance thereof.
- The recipient blood centre should have the capacity to hold the units requested, at an appropriate temperature until the time of utilization.
- Broad-based donor consent should be incorporated in the standard donor form to

*(Continued on page 16)*

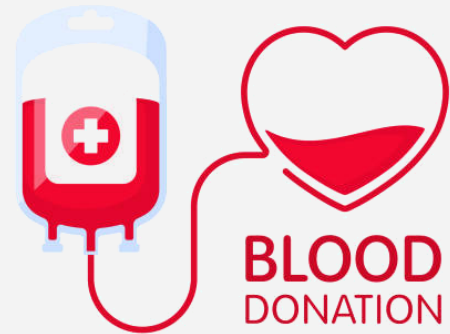
**Procedure to obtain license for Blood Centre (Blood Bank)****3**

ensure that the donor agrees to his blood unit being utilized beyond the blood centre where it is donated.

- The supplier blood centre can levy the prescribed processing charges on the patient or recipient, or the recipient blood centre as per NBTC norms.
- The recipient blood centre can levy only processing charging for compatibility testing (cross-matching), in addition to charges levied by the supplier blood centre, from the patient/recipient for such transferred units.
- Records of traceability shall be retained throughout the process.
- The supplier blood centre would be responsible for all the complications except those related to compatibility testing, which will be the responsibility of the recipient blood bank.
- The recipient blood bank shall report and evaluate all the adverse transfusion reactions, including those happening due to blood that has been transferred from the supplier blood bank.
- The documents accompanying the transfer shall include a TTI testing report and a record of transport at the appropriate temperature.
- Blood centres would be informed regarding bulk transfers to SBTC and, in the case of interstate bulk transfers, to NBTC.

**Note:** Also follow the guidelines or instructions issued by the concerned State Authorities where the Supplier or Recipient Blood centre is situated.

**List of emergency Equipment / Items / Drugs to be kept in the**

**Blood Centre / Bank:**

- (i) Oxygen cylinder with mask, gauge and pressure regulator.
- (ii) 5 per cent Glucose or Normal Saline.
- (iii) Disposable sterile syringes and needles of various sizes.
- (iv) Disposable sterile I.V. infusion sets.
- (v) Ampoules of Adrenaline, Noradrenaline, Mephentin, Betamethasone or Dexamethasone, Metoclorpropamide injections.
- (vi) Aspirin.

**List of SOPs used in Blood Centres**

List of SOPs used in Blood Centres, click the link below to download:

[List of SOPs used in Blood Centres](#)

**Compiled by:**

[Rakesh Dahiya](#)

**Asstt. State Drugs Controller**

[FDA Haryana](#)

[Click here for more articles of the Author](#)

## Landmark Seminar by FDA Haryana ..... continue

*(Continued from page 1)*

had communicated to form a formal interstate coordination committee to connect all northern states.

The idea is to make the entire belt an inhospitable region for traffickers.

### Setting the Tone: Serious Business

The event was flagged off by Sudhir Rajpal, Additional Chief Secretary of Health, Government of Haryana, who acknowledged this knowledge-sharing workshop as the 'need of the hour'.

- **The Cross-Border Challenge:** According to him, the issue of **spurious/counterfeit drugs** and NDPS is a menacing challenge that must be tackled by all neighboring states as one.
- **Data Sharing Is Critical:** He emphasized the need for substantial **data sharing between states** for effective monitoring and movement tracking.

**Protect India's Reputation: He implored all parties involved with the utmost level of integrity and accountability to protect India's global reputation as a quality-oriented pharmaceutical supplier.**

### Leaders Weigh In from FDA Haryana's Perspective



Dr. Manoj Kumar, Commissioner of FDA Haryana, welcomed all dignitaries, mentioning how FDA Haryana has successfully intervened multiple times in **NDPS cases** where officials were on board during prosecution.



**He was followed by SDC Lalit Kumar Goel, who raised concerns over three critical factors:**

- The illegal trafficking, abuse and diversion of controlled substances.
- The detection of **Not-of-Standard-Quality (NSQ) drugs**, which ultimately translates to patient safety being put at risk and public confidence diminished.

Both of these occurrences require a **holistic, multi-stakeholder approach** and seamless access to formal and informal resources to eliminate the crime nexus.

### Enforcement Perspective – National Database

The enforcement perspective was provided by Dr. Keshav Kumar, former Joint Director of CBI and DGP of Gujarat, who advised attendees with real-life experiences while commending FDA Haryana for this initiative.

He updated attendees about the Indian Pharmaceutical Alliance's intention to develop a national database of violators engaged in **spurious drugs** and **NDPS matters**.

This would benefit enforcement efforts massively for any major investigations across

*(Continued on page 18)*

## Landmark Seminar by FDA Haryana ..... continue

*(Continued from page 17)*

India. He also suggested that **forensics** play an important role in exposing exceedingly elaborate **spurious drug rackets**.

### Maintaining the Global Pharmacy Status

O. S. Sadhwani, the former Joint Commissioner-cum-State Drugs Commissioner of Maharashtra, spearheaded the session dedicated to **NDPS drugs**.

He lamented the current status of NDPS drugs after several recent incidents related to the trafficking of abused substances resulting in **NSQ drugs** causing pediatric fatalities in numbers.

He continued to delve into India's position as the leading '**global pharmacy**' without compromise and called upon all drug regulators and manufacturers to seriously **prioritize quality medicine above all else** to uphold reputation standards set nowhere else.

The seminar concluded after a fruitful

discussion on action plans for **spurious drugs** and **NDPS drugs** led by Manish Kapoor, Drugs Controller of Himachal Pradesh.

**This successful seminar was attended by over 100 officers from 7 neighboring states:**

- Jammu and Kashmir
- Himachal Pradesh
- Delhi
- Uttarakhand
- Punjab
- Chandigarh

The nodal officer was Karan Singh Godara; formal thanks were rendered by Parjinder Singh.

SDC Lalit Goel announced that a report would be generated to compile steps taken and proposed policy recommendations to send to state and central governments for immediate implementation.

**Source:** [The Health Master](#)



# FDA Haryana in news

## One-day Educational Seminar at Chandigarh

FDA Haryana successfully organized a one-day Educational Seminar at JW Marriott, Chandigarh, with active participation from officers of FDA, CID & Narcotics Departments of Haryana, Punjab, HP, Uttarakhand, Chandigarh, Delhi & J&K to strengthen coordination in regulating NDPS and combating spurious drugs.



# FDA Haryana in news

## One-day Educational Seminar at Chandigarh



## QR Code mandatory at every Medical Store for ADR reporting

### QR Code

India has officially entered a new chapter of Drug Safety Regulation. The **Central Drugs Standard Control Organization (CDSCO)** has issued a transformative order for citizens to ensure access to any side effects one may experience from taking drugs by means of displaying [QR code](#) at every medical store.

This is all a part of India's national **Pharmacovigilance Programme of India (PvPI)**; thus, any changes in patient safety will now travel even quicker from patient to the agency.

### But why is this Drug Safety Regulation making waves?

For many years, reporting an **Adverse Drug Reaction (ADR)** has not been the easiest option for the non-drug professional. But now, it should be easier.

According to a CDSCO circular dated November 20, 2025, with immediate effect **all retail/wholesale pharmacy stores** across India will now be obligated to prominently place:

A dedicated PvPI **QR Code**.

- The PvPI **Toll-Free Number: 1800-180-3024**.

Effective from the 16th Working Group Meeting of PvPI on 18 June 2025, this new piece of information will add to goals of improved patient safety and Pharmacovigilance.

By placing this in front of patients where drugs are being dispensed, this now gives power to all pharmacies as reporting agencies.

This is significant as never before have retail pharmacies been recognized physical



locations to report ADRs.

This will greatly benefit future goals of **Indian Pharmacovigilance** and **drug oversight** to come.

Thus, the global citizenry and any healthcare professional should feel empowered by ADRMS data that this mechanism of reporting is finally feasible and functional!

### Where Will You See This Change?

In your local or wholesale pharmacy! Effective immediately, the Licencing Authorities of States and Union Territories will ensure compliance within all pharmacies.

**This is what you need to expect from newly posted signs:**

#### The PvPI QR Code:

This is to be published in such a way that it is easily accessible by the average Indian so that they can access the indigenous PvPI Adverse Drug Reaction Monitoring System (ADRMS). The main priority is that any adverse events or suspected events from drugs taken can be instantly reported online for investigation. This

*(Continued on page 22)*

## QR Code mandatory at every Medical Store ..... continue

*(Continued from page 21)*

is India's basis for **ADR reporting**.

### **The PvPI Toll-Free Number (1800-180-3024):**

As an alternative option if someone would prefer calling in a suspected ADR, this alternative but equally beneficial method gives citizens an empowering ability to remain engaged with patient safety monitoring after they suspected a side effect.

This aspect of the system will be managed by the National Coordination Centre – the Indian Pharmacopoeia Commission – which is also the PvPI secretariat and a WHO Collaborating Centre for Pharmacovigilance.

### **How You Can Best Assist in Public Health Safeguarding**

The only way a report ever gets processed as intended is if everyday citizens report an adverse event, as this provides useful information that can lead to practical recommendations or even conclusions as to

whether the drug should be available in the first place.

This is known as **public health protection**.

Thus, whether it's your responsibility to report via **QR code** or toll-free number at your pharmacy is irrelevant to your health; either way you should let someone know if you suspect an ADR from your use of a particular drug.

Additionally, if your pharmacy fails to comply and does not show this required information, inform them that the CDSCO requests "wide dissemination among all the license holders and monitor strict compliance".

This is a new and exciting chapter in **India's drug safety** existence and the reality has been made overwhelmingly clear by government authorities about timely reporting of any medicine side effects as soon as you get them!

**Source:** [The Health Master](#)



**IPC Initiative to Boost Quality Standards for Blood and Blood Components****Blood and Blood Components**

In a transformational move to safeguard patients and uphold the integrity of the nation's healthcare system, the [Indian Pharmacopoeia Commission \(IPC\)](#) undertakes a series of extensive training and online sensitization programs for officials and staff working at [blood centres](#) across the country, ensuring compliance with mandatory, urgent new quality standards for [Blood and Blood Components](#) for transfusion safety.

**Blood and Blood Components Qualify as a Drug**

The underlying rationale is simple: because blood and its separated components (plasma or platelets, for example) come from a human donor, legally, it falls under the definition of a 'drug' as per the prevailing **Drugs Rules, 1945**, which, in turn, means that it must qualify for the same stringent, mandatory, enforceable quality criteria as any other life-saving drug.

[Download: GSR-166 \(E\) dated 11-03-20 regarding – Blood Bank / Blood Centre, Qualification of Doctor and technical staff etc.](#)

**New Standards to Reduce Transfusion Risks**

Since safe blood transfusion practices are paramount, the **IPC** has taken the lead in developing several **general requirements and monographs** or comprehensive "quality parameters" for **Blood and Blood**

**Components.**

These new standards exist with the ultimate goal of reducing the likelihood of transfusion-transmitted infections (TTIs) mandatory accountability means that only safe **Blood and Blood Components** will be administered once each unit/component passes safety screening.

Moreover, this concentrated effort increases **regulatory compliance** in **transfusion medicine**, meaning that all **blood banks / centres** will have the criteria needed for practical implementation by way of comprehensive yet feasible quality control operations issued under national regulations and in line with **international standards**.

**Training Webinar: November 7th**

The national sensitization campaign will kick off with a **National Webinar on November 7, 2025**, which will include presenters from some of the most trusted speakers in the field, from **IPC** subject matter experts to significant officials from drug regulatory agencies and transfusion medicine professionals across **IPC** regulations.

*(Continued on page 24)*

## IPC Initiative to Boost Quality Standards ..... continue

*(Continued from page 23)*

This major meeting will center around **Gazette Notification – GSR 166(E): Drugs and Cosmetic (Second Amendment) Rules, 2020**, published on March 11, 2020, which, for the first time, legally integrates the new quality standards for **Blood and Blood Components** into the **Indian Pharmacopoeia (IP)**.

### What Will Be Discussed?

#### Key topics will cover:

- **Integrated Disclosure:** Analysis of quality standards for **blood and blood components** as per the IP.
- **Best Practices:** Insights from transfusion medicine experts regarding compliance requirements.

**Questions: Technical and compliance-related questions from blood centre personnel will be fielded in practical sessions.**

### Who Will Benefit?

All personnel engaged with safe blood supply

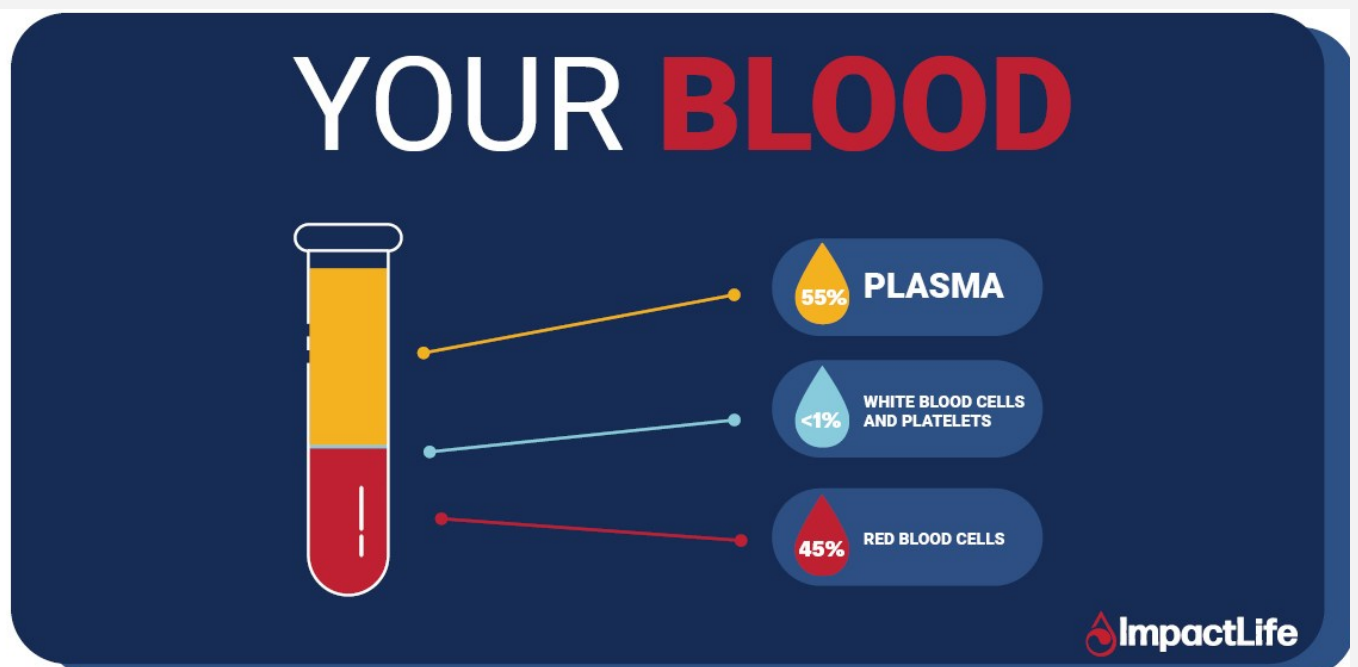
transfusion from this comprehensive training will receive practical clarifications to offer relevant assessments to primary levels of responsibility.

#### Benefitted personnel include:

- **Blood Bank/Blood Centre Professionals:** Those who collect, draw, test, and house various types of blood.
- **Regulatory Officers:** Those charged with assessment and compliance.
- **Institutions for Blood Irradiation:** Those who operate facilities that irradiate blood components.
- **Researchers:** Those engaged in transfusion medicine studies.

According to an official statement, “The sessions are invaluable to ensure that every **blood transfusion** performed in India is **safe, efficient, and effectively quality-assured as per regulations**, which ultimately strengthens the public’s trust in the healthcare delivery system.”

Source: [The Health Master](#)



## CDSCO grants powers to Joint Drug Controllers for key approvals

The [Central Drugs Standard Control Organisation \(CDSCO\)](#), the national drug regulatory agency of India has made a vital change in policies with an intent to improve its performance and response times dramatically.

The official delegation of substantial licensing authority powers to **Joint Drug Controllers (India)** comes at a crucial time to **expedite approvals for drug import licenses** and facilitate faster obligations for important **serious adverse events (SAE)**.

Given that all such crucial considerations need to be managed at present by the [Drugs Controller General of India \(DCGI\)](#), this is a final shift in the longstanding, highly centralized operations reigning up until now.

### Which powers have been delegated?

According to the official Office Order from the **DCGI's** front, specific powers have been transferred and will have a significant impact.

This is intended to alleviate the burden from the office of the top regulator and empower the **Joint Drug Controllers** for timely receipt and response times relative to the pharmaceutical industry.

**The specific powers transferred for regulatory empowerment include:**

- **Import License for Drugs for Human Use:** Approvals via India for much-needed drugs on an international scale.
- **License for Blood Stem Cell Products:** A necessary approval given how complicated and relatively new these products can be.

**Approvals of New Sites for Clinical Trials:** Approvals for sites where necessary [Clinical Trials](#) can occur.

### Decisions for Serious Adverse Events

In addition to these approvals, there's one part of the process which was previously regulated under extra cautionary circumstances that will now benefit as well: **Serious Adverse Events (SAE)**.

These events are unfortunate situations that arise during clinical trials or even **Bioavailability/Bioequivalence (BA-**



**BE)** studies.

They are life-threatening incidents (death) or life-changing incidents (disability).

Thus, the new mandate transfers powers to assess and address **SAE** compensations over time with timely recommendations relative to existing provisions.

This means that approval for compensation payments will now be more quickly addressed as previously they took too long.

The more promptly SAE are addressed, the better for ethical compliance for clinical trial participants. (**High CPC Keyword: SAE Compensation Payout**)

### Ease of Doing Business

According to industry experts, the delegation is a major benefit to a complicated ecosystem of the entire **pharmaceutical industry**, with a senior drug controller stating that it makes sense how all players will benefit from such changes.

In addition, the **CDSCO** has improved its accountability as a result, which complements the aims of the Indian Government as a whole.

By facilitating easier connections with major decision makers, it provides a more reputable stance on the **"Ease of Doing Business"** campaign in India and thus makes international investments much more attractive.

**Download:** [DCGI dt 30-10-2025 Delegation of Power to CDSCO Officers](#)

**Source:** [The Health Master](#)



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# FDA Haryana

**सिरसा में मेडिकल नशा तस्करों को 20 साल की सजा: बिना लाइसेंस बेच रहे थे, 1.75 करोड़ की खेप मिली थी**



जानकारी के अनुसार, मार्च 2018 में एसपी नरेंद्र बिजारणिया के नेतृत्व में सिरसा पुलिस ने सीटीएम वेदप्रकाश बेनीवाल, बीडीपीओ रवि कुमार व स्वास्थ्य विभाग के अधिकारियों को साथ लेकर अनाज मंडी स्थित गोदाम पर रेड मारी थी। सीनियर ड्रग आफिसर एनके गोयल, ड्रग आफिसर सुरेश चौधरी ने दवाओं की जांच की थी।

Source: Suresh Chaudhary, DCO, FDA Haryana

Click any where to read  
all the DCOIWA Newsletters



E-Newsletter

# DCOIWA News

Drugs Control Officers (I) Welfare Association (Regd)

Send your news and articles to:  
[dcoiwanewsletter@gmail.com](mailto:dcoiwanewsletter@gmail.com)





# Tamil Nadu

On 28/11/2025, A meeting was held at Guidance, Tamil Nadu and Hon'ble Industries Minister of Tamil Nadu, Dr. T R B. Rajaa has awarded appreciation to the Tamil Nadu Drugs Control Administration for successful implementation of Business Reform Action Plan, 2024.





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# Best Wishes On Promotion



## Haryana Regular Promotion

The DCOIWA family extends its best wishes for the future endeavours of the officers who have recently assumed their position following their well-deserved promotion.



**Sh. Rakesh Dahiya**  
promoted as Asstt.  
State Drugs Controller,  
FDA Haryana



**Sh. Karan Singh  
Godara** as Asstt.  
State Drugs  
Controller, FDA  
Haryana

# Congratulations



# Best Wishes on retirement



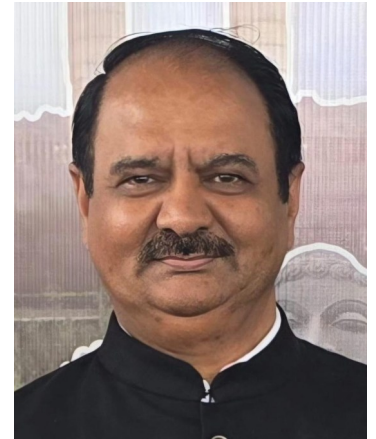
The DCOIWA family extends its best wishes for the future endeavours of the following officer who have recently retired. May God bless them with all happiness and healthy life.



**Mrs. Lorna  
Dorothy Pinto,  
FDA  
Maharashtra**



**Sh. Biswajit  
Mukhopadhyay  
ADDC West  
Bengal**



**Sh. Raj  
Chaudhari  
FDA  
Maharashtra**



*Congratulations*

# Congratulation Message



## *Superannuation*

### **D. R. GAHANE**

Joint Commissioner,  
Controlling & Licensing Authority,  
FDA Maharashtra

Date: 28th November 2025  
Place: Mumbai

On behalf of the Drugs Control Officers India Welfare Association (DCOIWA), I extend my deepest respect and warmest wishes to Shri **D. R. Gahane**, Joint Commissioner, Controlling & Licensing Authority, FDA Maharashtra; Vice President, DCOIWA; Board Member, DCOI Welfare Trust; and the esteemed recipient of the DCOIWA Lifetime Achievement Award 2024–25, on the occasion of his superannuation today.

Shri Gahane's long and distinguished career stands as an inspiring example of dedication, integrity, and commitment to strengthening the drug regulatory system in our country. His visionary leadership, unwavering work ethic, and tireless contribution towards ensuring the availability of safe and quality medicines have left an indelible mark on the profession and the community we serve.

As a pillar of DCOIWA, his guidance, wisdom and constant encouragement have shaped many initiatives and motivated officers across India. His remarkable service to the FDA Maharashtra and to the nation will always be remembered with gratitude and pride.

On this special day, the entire DCOIWA fraternity joins me in wishing Shri Gahane a healthy, peaceful, and fulfilling retired life. May this new chapter bring him joy, relaxation, and cherished moments with his family.



**G. Koteswar Rao**

National President, DCOIWA



**Baldev Choudhary**

National General Secretary



# Congratulation Message



## DRUGS CONTROL OFFICERS (I) WELFARE ASSOCIATION

(Regd.No. 634 of 2022)

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**Mrs. Kanchan Sinha**

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**Mr. Pooran Chand**

Cell: 9760366100 - Uttar Pradesh

**Mrs. Dipika Chauhan**

Cell: 7990351251 - Gujarat

**Mr. Chandan Kumar Giri**

Cell: 8895478543 - Odisha

**Mr. Amit Kumar Bansal**

Cell: 9198463261 - Uttar Pradesh

**Mr. Biswajit Talukdar**

Cell: 7086084833 - Assam

#### Treasurer:

**Dr. Parmanand Verma**

Cell: 9977177574 - Chhattisgarh

#### EC Members:

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Cell: 9443959680 - Puducherry

**Mr. Venkatesh Sinari**

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**Mr. Irap T. D. Toi**

Cell: 9233323204 - Meghalaya

**Mr. Ibom Ete**

Cell: 8414819858 - Arunachal Pradesh

**Mr. Neeraj Kumar**

Cell: 9458900286 - Uttarakhand

**Dr. Mrinal Kanti Sarkar**

Cell: 8974822360 - Tripura

**Mr. Tapan Choudhry**

Cell: 9230610226 - West Bangal



Date: 28 November 2025

Place: Hyderabad

### Congratulations Message

On behalf of the Drugs Control Officers India Welfare Association (DCOIWA), I extend my heartfelt congratulations to Smt. L. D. Pinto, Joint Commissioner (Drugs), Amaravati Division, FDA Maharashtra, on the occasion of her superannuation on 30th November 2025.

Your distinguished career, marked by integrity, dedication, and an unwavering commitment to strengthening the drugs regulatory system, stands as an inspiration to officers across the country. Your leadership in the Amaravati Division and your contribution to ensuring quality, safety, and compliance in the pharmaceutical sector have earned deep respect among your colleagues and stakeholders alike.

As you retire after an illustrious journey of service, we wish you a joyful, healthy, and fulfilling new chapter in life. May this new phase bring you peace, happiness, and many memorable moments with your loved ones.

With warm regards and best wishes,

**G. Koteswar Rao**

National President

Drugs Control Officers India Welfare Association (DCOIWA)



H.Q. 15-21-150/6, New Balaji Nagar,  
Kukatpally, Hyderabad (T.S), INDIA.  
Phone : 8121296397, 8094357800, 9977177574.

# Memorable moments



**Presented association's memento to Mrs Pinto Madam, jt. Commissioner, FDA, Amaravathi, Maharashtra by DCOIWA Maharashtra chapter**



**Mumbai: Retirement function of Shri. D. R. Gahane, Drugs Controller, Maharashtra**

# Memorable movements



**Honouring Sh. Rajeev Singh Raghuvanshi, DCGI with Membership of DCOIWA on 14th November 2025**



**Continuous education services, Week long training to Nagaland drug testing laboratory analysts at Kohima. The trainer Dr. Sridhar is senior most subject expert in analysis, from SIPRA Labs, Hyderabad. Present : Shri. Ethungbemo, DCOIWA Nagaland president and Smt. Imlilila, ADC, Nagaland**



# Honoring Sh. Rajeev Singh Raghuvanshi, DCGI with DCOIWA Membership



## DRUGS CONTROL OFFICERS (I) WELFARE ASSOCIATION

(Regd.No. 634 of 2022)

E-mail: dcoidcwa@gmail.com Website: www.dcoiwa.com

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**Dr. HEMANT G. KOSHIA**  
Cell: 99784 05054 - Gujarat

**President**  
**G. KOTESHWAR RAO**  
Cell: 8121296397 - Telangana

**General Secretary**  
**BALDEV CHOUDHARY**  
Cell: 8094357800 - Rajasthan

- Organizing Secretary:**  
**Mr. Rakesh Dahiya**  
Cell: 99116 00019, Haryana
- Vice-Presidents:**  
**Mr. Lalit Kumar Goel**  
Cell: 70567 02999, Haryana  
**Dr. M. Dhilip Kumar**  
Cell: 9840071715, Tamilnadu  
**Mr. D. R. Gahane**  
Cell: 98928 32289, Maharashtra  
**Mr. V. D. Dobaria**  
Cell: 98790 60666, Gujarat  
**Dr. A. Ramkishan**  
Cell: 63057 50926, CDSCO  
**Dr. Manish Kapoor**  
Cell: -9418081270 - Himachal Pradesh  
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**Dr. Mrinal Kanti Sarkar**  
Cell: 8974822360 - Tripura  
**Mr. Tapan Choudhry**  
Cell: 9230610226 - West Bengal

Date: 15 November 2025  
Place: Hyderabad



Yesterday, I had the privilege of meeting **Shri Rajiv Singh Raghuvanshi**, Drugs Controller General of India, at his office. We held a very fruitful discussion focused on strengthening the regulatory system to ensure the availability of safe and quality medicines to the public.

During the meeting, I requested special emphasis on Cosmetics enforcement and the creation of dedicated Cosmetics Inspector posts in the future. I also placed a request for official recognition of our Association, for which we received a positive and encouraging response.

I briefed him on the forthcoming initiatives and activities planned by the Association. On this occasion, we were honoured to present him with Life Membership of DCOIWA, in the esteemed presence of **Shri N. K. Ahuja**, Former Drugs Controller of Haryana.

On behalf of all DCOIWA members, we express our sincere thanks to the DCGI Sir for his positive response and support.

Regards,

  
**G. KOTESHWAR RAO**  
National President  
Drugs Control Officers India Welfare Association (DCOIWA)



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Kukatpally, Hyderabad (T.S), INDIA.  
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Meeting Sh. Arun Kumar, DI Ranchi in Hospital



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Date: 14 November 2025
Place: New Delhi

Today, along with Shri N. K. Ahuja and Shri Jithen, Drugs Inspector, Delhi, I visited Apollo Indraprastha Hospital, New Delhi, where Shri Arun Kumar, Drugs Inspector, Ranchi, Jharkhand, is undergoing treatment for acute pancreatitis and brain infarct.

We interacted with the treating doctors and were informed that he has just begun responding to the treatment, which is a positive sign.

The officers from Delhi, under the guidance of Chawla ji and Deepak Sharma ji, are showing excellent coordination in ensuring proper treatment and support.

Shri Arun Kumar is unmarried, and his aged parents are unable to attend to him. His elder brother is currently taking care of him at the hospital. As an immediate support measure, a cheque of Rs. 2.0 lakhs has been handed over to him on behalf of the DCOI Welfare Trust.

Further financial assistance and an appeal letter will be released shortly along with the bank details.

Regards
G. Koteswar Rao
National President
Drugs Control Officers India Welfare Association

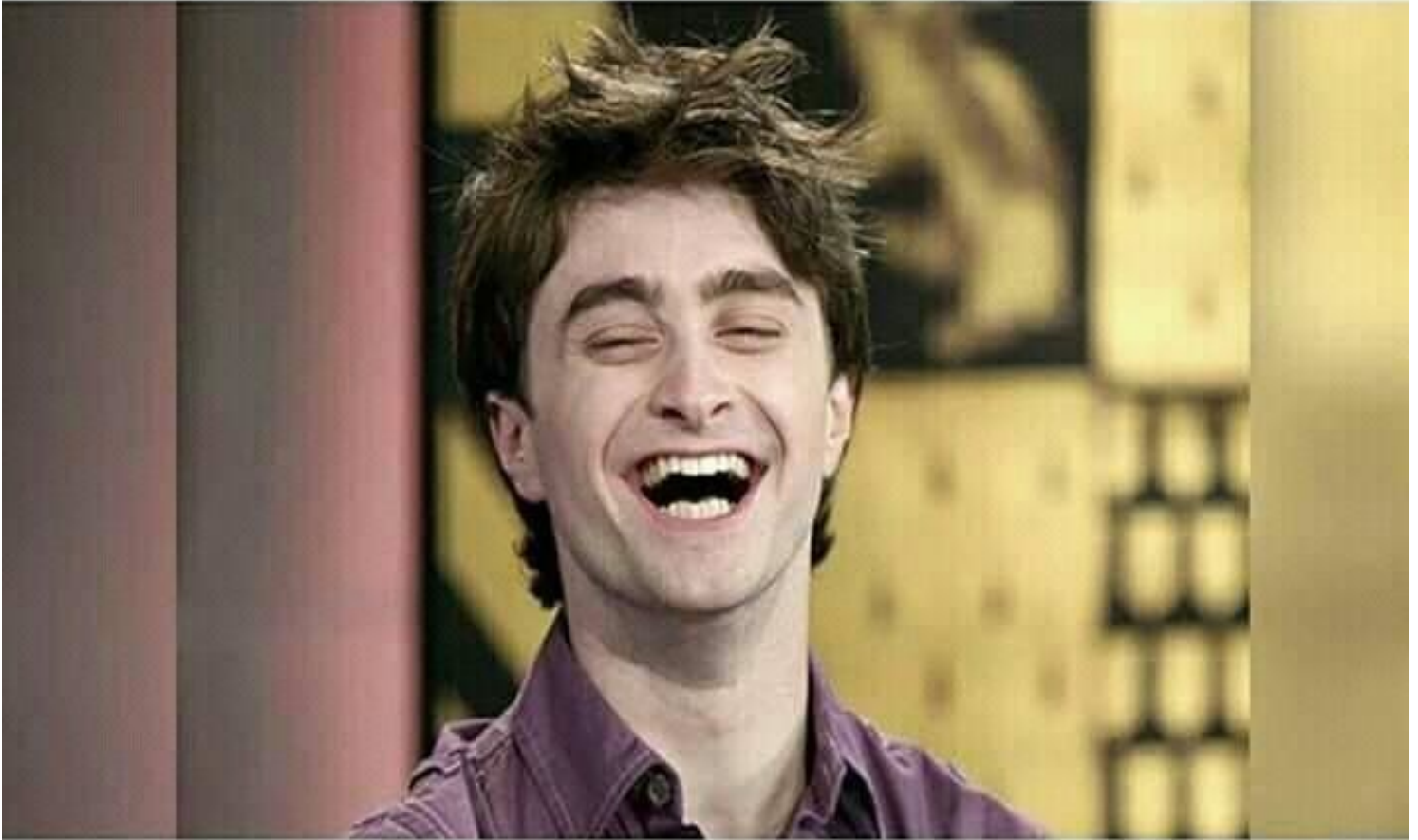
Handwritten signature of G. Koteswar Rao



H.Q. 15-21-150/6, New Balaji Nagar,
Kukatpally, Hyderabad (T.S), INDIA.
Phone : 8121296397, 8094357800,9977177574.

Laughter dose

Laughing is the **best** medicine



But if you're laughing for  
**no reason**, you need  
medicine 😂

Source: [PK Jaggi](#), Co-Editor






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**CDSCO**

**States**

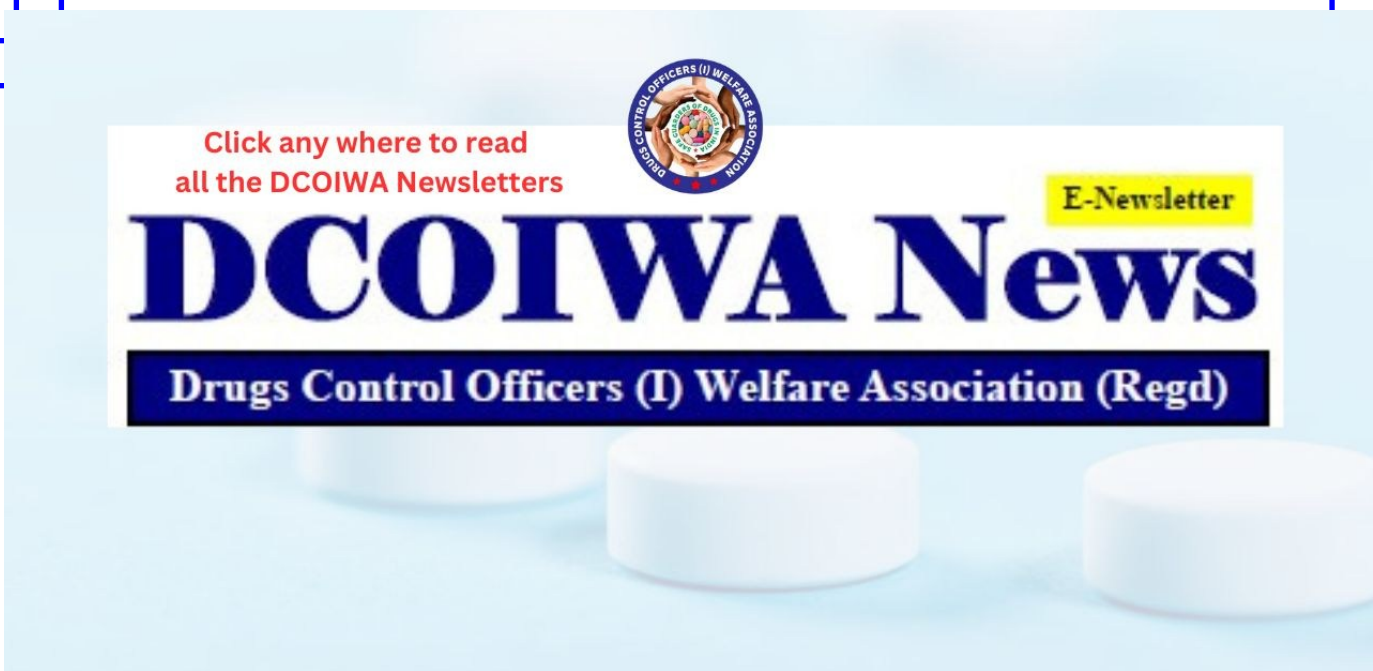
**NSQ List: October 2025**

Click any where to read  
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**DCOIWA News** E-Newsletter

**Drugs Control Officers (I) Welfare Association (Regd)**



Click below links to download:

[NSQ: October 2025 CDSCO](#)

[NSQ: October 2025 STATES](#)

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Source: CDSCO Website



## Sub-Standard Drugs



Send your news and articles to:  
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## Important Links



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# NPPA

AFFORDABLE MEDICINES FOR ALL

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- [NPPA fixed retail price of 28 formulations: November 2025](#)



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**DCOIWA News** E-Newsletter

Drugs Control Officers (I) Welfare Association (Regd)



## Drug alert

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**Important short notes for Industry and Regulators**  
**By Lalit Kr. Goel, FDA Haryana**



**IMPORTANT**  
**Short Notes**  
**for**  
**Industry and Regulators**  
by Lalit Kr. Goel FDA Haryana

**To read click anywhere in the picture**

**Key Notes on Revised Schedule M: Compilation**  
**By Rakesh Dahiya, FDA Haryana**



**The Health Master**  
**Key Notes**  
**Revised**  
**Schedule M**  
**Compilation**  
By Rakesh Dahiya, FDA Haryana

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Appeal to DCOIWA members and Well-Wishers

1



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25 November 2025
Kohima, Nagaland

APPEAL TO DCOIWA MEMBERS & WELL-WISHERS
Medical Status Update – Shri Amit Kumar, Drugs Inspector, Jharkhand

Dear Members and Well-Wishers,

I wish to share the latest medical update of our colleague, Shri Amit Kumar, Drugs Inspector, Jharkhand, who is undergoing treatment for acute pancreatitis with complications at Apollo Indraprastha Hospital, New Delhi. As of today, his condition is as follows:

- 1. Consciousness / Brain Condition: The patient is opening his eyes and showing basic responses, although not fully alert yet. Doctors have stated that brain recovery will take time, but his current status is stable. EEG and ongoing observations will determine further course of treatment.
2. Breathing / Ventilator: He continues to be on ventilator support. A weaning trial is planned in the next 1–2 days, depending on his tolerance. A tracheostomy tube is in place for safer and more comfortable breathing. Oxygen requirement remains stable.
3. Kidneys / Dialysis: Kidney function is gradually improving. Creatinine has reduced to 1.2, which is a very positive indication. SLED dialysis is continuing to support the kidneys.
4. Blood Pressure & Heart: Blood pressure is stable without major medication support. Pulse and circulation are satisfactory.
5. Lungs: No fluid is seen in the lungs presently. Breathing status has improved compared to earlier days.
6. Infection & Other Tests: Infection markers have improved. Fever is under control.

H.Q. 15-21-150/6, New Balaji Nagar,
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## Appeal to DCOIWA members and Well-Wishers

### 2



### DRUGS CONTROL OFFICERS (I) WELFARE ASSOCIATION

(Regd.No. 634 of 2022)

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Antibiotics are being continued as per medical advice.

#### 7. Digestive System:

Patient is passing motion; stool remains watery, which is common during heavy medication and infection recovery.

#### 8. Overall Condition:

Overall condition is stable, though recovery is slow. Noticeable improvement is seen in kidneys and lungs. Brain recovery will require more time and close monitoring. The next 48 hours are crucial for the ventilator weaning trial.

#### Financial Support Appeal

A request has been received from his brother seeking additional financial assistance, as the treatment costs continue to rise significantly.

On behalf of DCOIWA, I earnestly appeal to all our members and well-wishers to contribute a minimum of ₹1,000 towards his medical expenses. Your small support today can make a big difference to a family in distress. Let us come together to help our colleague in this difficult time.

Account Name: **Arun Kumar**  
Bank: Punjab National Bank  
Account Number: **7608000100026534**  
IFSC Code: PUNB0760800  
Branch: HEC, Sector 2, Dhurwa

Google pay number: 8102052494

Regards,

**G. Koteswar Rao**  
National President, DCOIWA



H.Q. 15-21-150/6, New Balaji Nagar,  
Kukatpally, Hyderabad (T.S), INDIA.  
Phone : 8121296397, 8094357800, 9977177574.

# Fee structure: All types of drugs licenses

by  
**Lalit Kr. Goel**  
State Drugs Controller,  
FDA Haryana



- [Drug sale licenses](#)
- [Homeopathic sale licenses](#)
- [Drug manufacturing licenses](#)
- [Homeopathic manufacturing licenses](#)
- [Cosmetics manufacturing licenses](#)
- [Drug repacking](#)
- [Medical devices manufacturing licenses](#)
  
- [LVP](#)
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**structure of all types of drug licenses**

[License fee structure for all licenses](#)

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## FAQs



by  
**Lalit Kr. Goel**  
State Drugs Controller,  
FDA Haryana



**To read all FAQs on various topics, Click the below links**

[\*\*FAQs – on Blood Pressure Monitoring Devices\*\*](#)

[\*\*FAQs – on Alcohol \(in Pharma Industry\)\*\*](#)

[\*\*Blood Centre \(Bank\) – requirements at a glance\*\*](#)

[\*\*FAQs on – Cosmetics Rules 2121\*\*](#)

[\*\*FAQs – on Mouthwash\*\*](#)

[\*\*Fee structure: All types of drugs licenses\*\*](#)

[\*\*FAQs – on Ear Drops\*\*](#)

[\*\*FAQs – on Drug Permission in Brand or Generic Name\*\*](#)

[\*\*FAQs – on Disinfectants \(Series-2\)\*\*](#)

[\*\*Gist of Notification 25th September 2121: Medical Oxygen\*\*](#)

[\*\*Salient features of Supreme Court order dated 28.08.2121\*\*](#)

[\*\*Pharmacopoeial status of Blood and its components\*\*](#)

[\*\*Difference between Sanitizer and Disinfectant\*\*](#)

[\*\*FAQs on Legal Metrology & Blood Bags\*\*](#)

[\*\*FAQs on Sanitizer, N95 Mask & Digital Thermometer\*\*](#)

[\*\*FAQs on Medical Oxygen\*\*](#)

[\*\*FAQs – on Cosmetics \(Series-1\)\*\*](#)

[\*\*FAQs – on Blood Bank \(Series-1\)\*\*](#)

[\*\*FAQs – on Blood Bank \(Series-2\)\*\*](#)

[\*\*FAQs – on Blood Bank / Centre \(Series-3\)\*\*](#)

[\*\*FAQs on Medical Devices Rules, 2117\*\*](#)

[\*\*FAQs about New Drug, Banned drugs etc.\*\*](#)

[\*\*FAQs on Disinfectant \(Series-1\)\*\*](#)

[\*\*FAQs – on Disinfectants \(Series-2\)\*\*](#)

[\*\*FAQs – On ‘Good Night’, ‘All Out’, ‘Hit’ and ‘Harpic’ etc.\*\*](#)

[\*\*FAQs – on Ranitidine tablets and injections in India\*\*](#)

[\*\*FAQs – On Narcotic Drugs, Brand Names of drug \(G.S.R. no. 828 \(E\)\*\*](#)





# Schedules: All types of Cosmetics



## Schedules

Compiled by  
**Rakesh Dahiya**  
FDA Haryana



For Schedules on following topics  
Click below links

1

# Schedules

## Schedules –All types of Cosmetics under Cosmetics Rules 2020

As per Cosmetics Rules 2020, we have provided all the schedules introduced in these rules. Click the below links for more information:

**First Schedule** – Authorisation from manufacturer

[First-Schedule-Cosmetics](#)

**Second Schedule – Part-I** – Information and undertaking required to be furnished by the manufacturer or his authorised importer or distributor or agent with the application form for import registration certificate

[Second-Schedule-Part-I-Cosmetics](#)

**Second Schedule – Part-II** – Information and undertaking required to be furnished by the manufacturer with the application form for grant of manufacturing licence or loan licence

[Second-Schedule-Part-II-Cosmetics](#)

**Third Schedule** – Fee payable for licence, permission and registration certificate

[Third-Schedule-Cosmetics](#)

**Fourth Schedule** – List of categories of cosmetics for import

## [Fourth-Schedule-Cosmetics](#)

**Fifth Schedule** – Fee for test or analysis by the Central cosmetics laboratories or by the state laboratories

[Fifth-Schedule-Cosmetics](#)

**Sixth Schedule** – Undertaking for the import of cosmetics to be submitted by the importer with application form for Import Registration Number

[Sixth-Schedule-Cosmetics](#)

**Seventh Schedule** – (GMP) Good manufacturing practices and requirements of premises, plants and equipment for manufacture of cosmetics

[Seventh-Schedule-Cosmetics](#)

**Eight Schedule** – Particulars to be shown in the manufacturing raw material records

[Eighth-Schedule-Cosmetics](#)

**Ninth Schedule** – Standards for cosmetics

[Ninth-Schedule-Cosmetics](#)

**Tenth Schedule Part-I** – List of colourants allowed for use in cosmetic products as given under IS: 4707 (Part 1) as amended by the Bureau of Indian Standards from time to time.

[Tenth-Schedule-Part-I-](#)

(Continued on page 48)



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## Schedules: All types of Cosmetics



### Schedules

Compiled by  
**Rakesh Dahiya**  
FDA Haryana



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# Schedules

### Cosmetics

**Tenth Schedule Part-II** – List of colours permitted to be used in soaps

### [Tenth-Schedule-Part-II-Cosmetics](#)

**Eleventh Schedule** – (GLP) Good laboratory practices and requirements of premises and equipment

### [Eleventh-Schedule-Cosmetics](#)

**Twelfth Schedule** – Class of cosmetics and Extent and conditions of exemption

### [Twelfth-Schedule-Cosmetics](#)

**Thirteenth Schedule** – Substitutions in the Rules

### [Thirteenth-Schedule-Cosmetics](#)

### Also read

### [Schedules: Drugs](#)

### [Schedules: Medical Devices](#)

### [Schedules: Clinical Trials](#)

### [Cosmetics – Manufacturing](#)

### License

### [Latest Notifications: Cosmetics](#)

### [FAQs on – Cosmetics Rules 2020](#)

### [FAQs – on Cosmetics \(Series-1\)](#)

### [FAQs on Pollution in Drug, Cosmetics & Homeopathic Industries](#)

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### [Rakesh Dahiya](#)

Asstt. State Drugs Controller

### [FDA Haryana](#)

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## Pharmaceuticals

### Important Notifications



भारत का राजपत्र  
The Gazette of India

### Important Notifications

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Rakesh Dahiya  
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[New Drugs](#)

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DCOIWA News

December 2025

# CDSCO Circulars

भारत का राजपत्र  
The Gazette of India

CDSCO Circular

CDSCO

- CDSCO circular dt 20-11-2025 Display of QR Code at retail or wholesale pharmacies across the country to report Adverse Drug Reactions to PvPI
- [CDSCO circular dt 20-11-2025 Display of QR Code at retail or wholesale pharmacies across the country to report Adverse Drug Reactions to PvPI](#)

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E-Newsletter

Drugs Control Officers (I) Welfare Association (Regd)

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**Remembering late shri Dhanpal, DI, Tamil Nadu (2nd Nov 2022)****DRUGS CONTROL OFFICERS (I) WELFARE ASSOCIATION**

(Regd.No. 634 of 2022)

E-mail: [dcoidcwa@gmail.com](mailto:dcoidcwa@gmail.com) Website: [www.dcoiwa.com](http://www.dcoiwa.com)**President****G. KOTESHWAR RAO**

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**General Secretary****BALDEV CHOUDHARY**

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**Treasurer****Dr. PARMANAND VERMA**

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**Vice-Presidents:****Dr. M. Dhillip Kumar**

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**Larap T.D. Toi**

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**Venkatesh Sinari**

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**Tapan Choudhry**

Cell: 9230610226 - West Bangal

**Joint Secretaries:****Chandan Kumar Giri**

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**Pooran Chand**

Cell: 9760366100 - Uttar Pradesh

**Dr. Mrinal Kanti Sarkar**

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**Mrs. Dipika Chauhan**

Cell: 7990351251 - Gujarat

**Organizing Secretary:****Amit Kumar Bansal**

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**EC Members:****Dr. E. Ananda Kirouchenane**

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Kukatpally, Hyderabad (T.S), INDIA.  
Phone : 8121296397, 8094357800, 9977177574.

# Obituary



## DRUGS CONTROL OFFICERS (I) WELFARE ASSOCIATION

(Regd.No. 634 of 2022)

E-mail: [dcoidcwa@gmail.com](mailto:dcoidcwa@gmail.com) Website: [www.dcoiwa.com](http://www.dcoiwa.com)

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Cell: 99784 05054 - Gujarat

### President

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### General Secretary

**BALDEV CHOUDHARY**

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**Dr. M. Dhillip Kumar**

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Cell: 98928 32289, Maharashtra

**Mr. V. D. Dobarla**

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## Shri. M. Venkat Reddy Garu

### CONDOLENCE MESSAGE

The Drugs Control Officers India Welfare Association (DCOIWA) expresses its deepest condolences on the sad demise of Shri M. Venkat Reddy garu, Former Director, Drugs Control Administration of the erstwhile combined Andhra Pradesh, who passed away last night due to ill health.

Shri Venkat Reddy garu was an eminent administrator, a highly respected leader, and a guiding force in strengthening the drug regulatory system. His dedication, integrity, and lifelong commitment to ensuring the availability of safe and quality medicines have left an indelible mark on the profession and the regulatory fraternity.

His passing is an irreparable loss to the Drugs Control community and to all who had the privilege of working with him.

DCOIWA prays for eternal peace to the departed soul and conveys heartfelt sympathies to the bereaved family. May they find strength and comfort during this difficult time.

Om Shanti



**G. Koteswar Rao**

National President, DCOIWA

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# Upcoming Events

## UPCOMING EVENTS

**2025**

### December-2025

#### 74th Indian Pharmaceutical Congress 2025

**Date:** December 19-21, 2025

**Location:** BIEC, Bengaluru, India

**Description:** 74th Indian Pharmaceutical Congress (IPC), being held in the dynamic city of Bengaluru. As President of the Indian Pharmaceutical Congress Association (IPCA), I am deeply honored to lead this prestigious gathering that brings together the best minds from academia, industry, regulatory bodies, and healthcare.

**This year's theme, "AI & Technology in Pharma: Educate, Innovate, Empower,"** captures the transformative journey we are undertaking as a profession. The integration of artificial intelligence and emerging technologies is revolutionizing how we approach drug discovery, patient care, regulatory processes, and pharmacy education. It is our responsibility to ensure that this transformation is inclusive, ethically sound, and focused on public health outcomes.

[Click for more details](#)

### December-2025

#### PharmaTech Expo Bengaluru

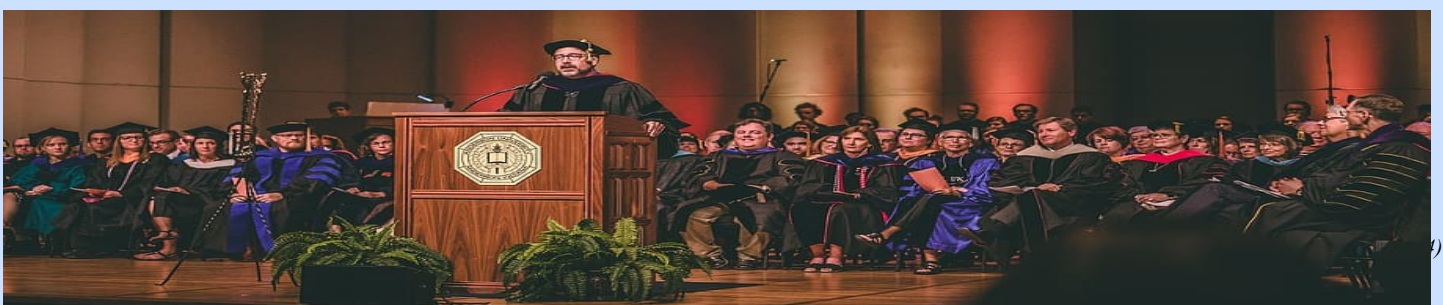
**Date:** December 19-21, 2025

**Location:** Bengaluru, India

**Description:** PharmaTech Expo is one of the largest pharma exhibitions in India and is a place for thousands of people from the business to share their experiences related to products, customers, business, and sales. This pharmaceutical and lab expo brings together people from across the globe to one destination. It is one of the biggest B2B trade shows of the sector that involves people from the healthcare and pharma machinery industries to participate and share innovation related to advanced technologies in the pertinent sector.

It will showcase pharma products, machinery, and technological innovation to buyers from various countries, including India, China, the USA, & Germany, which are major markets for this sector. This event will surely give you a huge platform to establish and enhance your business by meeting active suppliers looking for collaboration with the Indian pharma and healthcare market. Meeting new investors and fellow businessmen from the same fraternity would definitely be a win-win situation for both parties. If you are from the pharmaceuticals and

[Click for more details](#)



# Upcoming Events

## UPCOMING EVENTS

**2026****April-2026**

### PharmaTech Expo Chandigarh

**Date:** April 09-11, 2026  
**Location:** Chandigarh, India

**Description:** PharmaTech Expo is one of the largest pharma exhibitions in India and is a place for thousands of people from the business to share their experiences related to products, customers, business, and sales. This pharmaceutical and lab expo brings together people from across the globe to one destination. It is one of the biggest B2B trade shows of the sector that involves people from the healthcare and pharma machinery industries to participate and share innovation related to advanced technologies in the pertinent sector.

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[Click for more details](#)





## DCOIWA Mission

To unite and organize the working and retired Drugs Control Officers from Indian States, Union Territories and CDSCO, with an object of coordinating their activities for establishment of social justice and regulating the relations of the officers with government agencies. Call : 8121296397, 8094357800,9977177574

- a) To unite and organize the working and retired Drugs Control Officers from Indian States, Union Territories and CDSCO, with an object of coordinating their activities for establishment of social justice and regulating the relations of the officers with government agencies.
- b) To safeguard and promote interest of its members all over the country
- c) To redress the grievances of the members.
- d) To promote a sense of fraternity, feeling of belonging and brotherhood amongst its members.
- e) To cooperate, accept affiliations and federate with the officers associations, federations, and confederations in the country where similar objectives are seen with international bodies.
- f) To achieve professional excellence through better coordination amongst its members.
- g) To offer better services to the public.
- h) To make dedicated efforts for welfare of its members.
- i) To conduct seminars, webinars, social activities, competitions, quiz programs etc. time to time.
- j) To take up any other activity conducive to the betterment in the discharge of their functions effectively and efficiently.

## How to become a member

[Register Online](#)

[Download Form](#)

Dear Members,

As we conclude this edition of our e-newsletter, I would like to express my gratitude to our contributors and readers for their continued support. Your engagement is invaluable, and we appreciate the diverse perspectives that make our community thrive.

We strive to bring you relevant and insightful content, and we welcome any feedback or suggestions you may have for future editions. Our goal is to foster a collaborative space for knowledge-sharing among DCOIWA members, regulators, and pharmacy professionals.

Thank you for being a part of our community. We look forward to bringing you more enriching content in the upcoming editions.

Best regards,

**Rakesh Dahiya**  
Editor-in-Chief  
DCOIWA News



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