

Guidelines, Procedure, Checklist & Fee

Retail Drug License (ONDLS)

Health & Medical Education (State Drug Controller)

Government of J&K

Name of Service

Issuance of **Retail Drug License** through “Online National Drug Licensing System” (ONDLS)

Name of Department

Health & Medical Education Department (Office of State Drug Controller)

Policy/Government Order

Drugs & Cosmetic Act, 1940 & Rules there under.

Introduction to Online National Drug Licensing System (ONDLS)

The National Drug Authority of the Country viz CDSCO has designed & developed an Online Module/Portal in collaboration with Centre for Development of Advanced Computing, Noida. The said portal is extended to all types of Regulatory Clearances in terms of Licensing / Certifications for Manufacturing, Blood Centres & Sales.

Guidance for Applicant Stakeholders

The module is based on the principle of self – registration on the portal by Firm / Blood Centre & Technical Person.

Accessibility to the ONDLS module: -

- The individual / firm intending to register on the portal may logon at <https://statedrugs.gov.in>
- On the Homepage of the Module at top right corner, click on Sing on /up.
- Enter Mobile No. & OPT will be sent on Mobile Number. Enter OTP with 60 seconds.
- Enter email ID. OTP will be sent on email ID. Enter OTP within 60 seconds.
- New Page will open up which provides below tiles: Firm: Technical Person : Firm & Technical Person
- The Firm / Individual has to select one tile as per requirement. The selection of tiles is simplified as below:
 - ✓ a) If user is a Firm or Corporate, Tile “Firm” is required to be selected & proceed accordingly.
 - ✓ b) If user is an individual & intends to register as a Technical Person, Tile “Technical person” is required to be selected & proceed accordingly.
 - ✓ c) If Firm & Technical Person is one entity i.e Proprietor &

Registered Pharmacist is one person only, Tile "Firm & Technical Person" is required to be selected & proceed accordingly. In case Proprietor & Registered Pharmacist are two different persons (Technical Person on arrangement basis) , then firm & technical person are to be registered on the portal by selecting relevant tile respectively.

Submission of Application by Firm / Uploading of Details by Technical Person:

- A person who has registered himself / herself as technical person (Registered Pharmacist) has to update below details along with uploading of relevant documents.

- A) Add Academic Qualification
- B) Add Professional Experience
- C) Add Technical Registration.

After completing the details, System generated Technical Registration Number is allotted to Technical Person which is available in the profile option.

Application Submission by Firm

- The firm after successful Registration may hire technical person (Registered Pharmacist) through add technical member tile by using Technical registration Number to be provided by Technical Person. After hiring , the technical person will be available in Hired Technical Member.
- The user has to click on Fresh Application & update basic details about the premises, add technical person details from Hired Technical Member Tile & selection of intended License Forms.
- Proceed to checklist of documents related to type of License Form(s) selected
- Upload documents against the checklist & proceed.
- Fill Payment details form.
- Download Legal Form viz Form 19 in case of Retailer / Wholesaler
- Sign the legal form & proceed to upload.
- Submit the application & message will be displayed that your application has been submitted successfully. Kindly note your File No. for future correspondence.

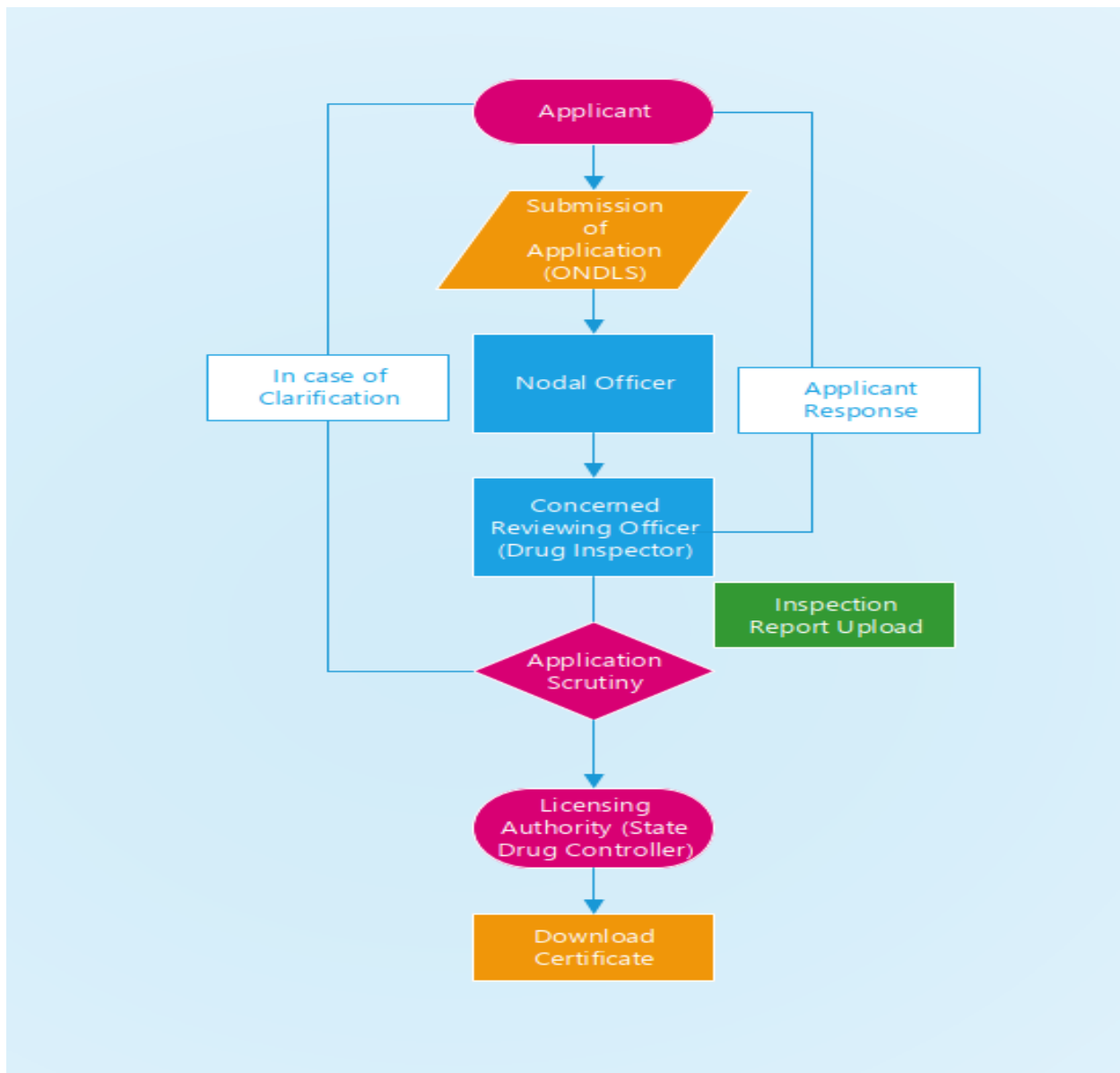
Note: The module provides option of hiring of technical person by the firm. The Technical person has to share his system generated Technical Registration Number with the firm. The firm searches this number in the Hiring of Technical Staff option. After completion of offer & acceptance by & between firm & technical person, the hiring process is complete & the services of such hired person can be utilised as Competent Technical Staff on Licenses if fulfilling eligibility criteria for same. However, in case of registration as "Firm & Technical person" auto-hiring & switch role option as Firm or Technical Person are provided.

Guidance for Official Stakeholder (Department Workflow)

The working principle for official users is based on Mobile Number & Email ID based Registration of Officials on the portal, Creation of Application Type Zones, & Mapping of registered Officials with Created Zones. Three tier hierarchy system is customized for UT of J&K.

- a) Reviewing Officer (Drug Inspector)
 - b) Nodal Officer
 - c) Licensing Authority
- The applications submitted by the stakeholders are received on the dashboard of Nodal Officer.
 - Nodal Officer assigns the application to concerned Reviewing Officer.
 - Reviewing Officer after scrutiny of uploaded documents & uploading of Inspection Report forwards the application to Licensing Authority with specific recommendation.
 - Based on recommendations, Licensing Authority shall dispose off the application. The approved application is downloaded , digitally / physically signed & uploaded along with Cover Letter.
 - The applicant can download the final certificate from his /her user Id credentials.

- Department Level Processing of Application



1. Procedure, Checklist & Fee of Grant of Retail Sale Drug License

The step wise procedure for grant of retail sale drug Licence/s is given as below:

Step 1: Application for grant of a licence [to sell, stock, exhibit or offer for sale or distribute] drugs, other than those included in Schedule X, [shall be made in Form-19 accompanied by “**a fee of rupees three thousand for Licenses on Form 20 & 21**” is made in Form-19 [Rule 59 (2)] to the Licensing Authority. (Online through ONDLS)

The statutory / supportive requirements to the Form-19 is as:

1. Valid Registration Certificate issued by President J&K Pharmacy Council & named to be as Registered Pharmacist on the Licence/s.
2. Constitution of the Firm / Photoproforma (Proprietorship / Partnership) & mentioning the full particulars of the Registered Pharmacist.
3. Documentary evidence in respect of the ownership or occupation on rental or other basis of the premises
4. Site Plan of the Premises (Minimum carpet area 10 square meters)
5. Undertaking /Affidavit/s in respect of Proprietorship / Partnership.
6. Agreement deed in case of hiring of Qualified person.

Step 2: The documents complete in all respects as per step 1 is forwarded to concerned area inspector by the Licensing Authority for on spot inspection of the premises.

Step 3: The concerned area inspector after inspection of the premises along with Inspection report submits the above documents / file along with inspection report to the Licensing Authority with the recommendation as: Licence/s may / may not be granted in favor of the applicant along with reasons.

Step 4: The Licensing Authority after being satisfied with the recommendations of concerned area inspector for grant of licence issues the Licence/s in favor of the applicant on Form 20 / 21

Eligibility Criteria (Minimum Carpet Area)

- For Retail sale - 10 square meter
- For Retail - Wholesale together - 15 square meter

Note: For more clarification on checklist documents, Please visit homepage of <https://statedrugs.gov.in> & browse “Service” tab at the top of the page & download checklist of required documents available against each type of application form.

Online Mandate Order for ONDLS

OFFICE OF THE STATE DRUGS CONTROLLER
DRUGS & FOOD CONTROL ORGANIZATION
PATOLI MANGOTRIAN J&K (JAMMU)

Regd. Post / Email / WAP

Sub: Adoption of "Online National Drug Licensing System" for Manufacturing & Sales Licensing in the UT of J&K – Regarding.

ORDER No. DFO of 119, 2021

Dated : 02-12-2021

Whereas , XLN Online Module for Licensing System (Sales) was adopted by this UT on 13.08.2018 in terms of Notification bearing No. HD/Drugs/69/2017 , dt: 13.08.2018 ;

Whereas , Apex Drug Authority of the Country viz Central Drugs Standard Control Organization , New Delhi (**herein referred to as CDSCO**) in collaboration with Centre for Development of Advanced Computing , Noida (**herein referred to as CDAC**) has developed a PAN India Portal under the banner "**Online National Drug Licensing System (ONDLS)**" for all states which shall extend to all regulatory clearances related to Manufacturing ; Blood Centers , Retail Sale & Wholesale under the provisions of Chapter IV of Drugs & Cosmetics Act, 1940 ;

Whereas , in order to ensure smooth transition , XLN Module was shut down vide this office Order No. DFO 105 of 2021 dt : 10.11.2021 ;

Whereas , the procedural trial / validation of ONDLS Module has been completed by the CDAC Authorities ;

Whereas , the said portal / module bearing URL as <https://statedrugs.gov.in> has been launched on 30th November, 2021 ;

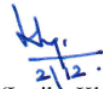
Whereas , under **XLN Module** , Wholesale Licensing (Fresh) were disposed off as per "**Guidance Manual for issuance of Drug Sale Licenses on Form 20B & 21B**" issued vide this office Communication No. DFO/D-845/7418-20 , dt: 04.02.2019 ;

Whereas , **XLN Module** & Newly adopted Module (**ONDLS**) are different in operation design & therefore necessary amendment is required in the existing Guidance Manual ;

Now , therefore in light of the contents explained herein above , all the statutory authorities appointed by the Government of J&K to carryout statutory duties under the provisions of the Drugs & Cosmetics Act, 1940 & Drugs Rules 1945 thereunder are hereby ordered to adopt Online Module viz **“Online National Drug Licensing System” (ONDLS)** with immediate effect for regulatory clearances under the said Act & Rules.

No. DFO/D-936/ 4193-4242

Dt: 02-12-2021


2/12.
(Lotika Khajuria)
State Drugs Controller
Drugs & Food Control Organization
J&K (Jammu)

Copy To :

1. The Additional Chief Secretary to Government , H&ME Department, Civil Secretariat (J&K) for favor of information.
2. Drugs Controller General (I) , CDSCO (HQ) , FDA Bhawan , Kotla Road , New Delhi for favor of information.
3. Commissioner, FDA (J&K) –Drugs & Food Control Organization J&K for favor of information.
4. Jt. Director Planning (H&ME) Department , Civil Secretariat , J&K for information. Process of integration of Portal with UT specific SWC System has been already taken up with concerned quarters for necessary action at an earliest convenience.
5. SIO-NIC (J&K) for information.
6. **Dy. Controller –Drugs & Food Control Organization Jammu / Kashmir for information & necessary action with the request to ensure implementation of revised guidelines in letter & spirit. The revised Guidance Manual is attached herewith as Annexure A.**
7. **All Assistant Controller Drugs / All Licensing Authorities for necessary compliance.**
8. Assistant Controller Drugs (I), CDSCO Sub Zone Jammu for information.
9. Mr. Rahul Gautam –CDAC , Noida for information.
10. **All Drug Control Officers (Enforcement) for strict compliance.**
11. **Mr. Surinder C/o M/s Saytechnologies , Jammu with the request to upload the document on our official website by today only.**

Weblink for the Notification: - [ONDLS.pdf \(dfcojk.org\)](https://dfcojk.org/ONDLS.pdf)

Notified Timelines Under Jammu & Kashmir Public Service Guarantee Act



GOVERNMENT OF JAMMU AND KASHMIR
GENERAL ADMINISTRATION DEPARTMENT
(Public Services Management Cell)
Civil Secretariat, Jammu/Srinagar

NOTIFICATION

Srinagar, the 13th September, 2013

SRO 400 - In exercise of the powers conferred by section 4 read with section 8 of the Jammu and Kashmir Public Services Guarantee Act, 2011 (Act No. IX of 2011), the Government hereby specify the following services and the designated officers/appellate authorities as under:-

I. Health and Medical Education Department

Issuance of Drug Licenses under Drugs and Cosmetics Act-1940/Rules

S. No.	Name of the Service	Designated Officer (the officer who will provide the service)	Stipulated time frame for providing the service	First Appellate Authority	Second Appellate Authority
01	License to sell, stock or exhibit (or offer) for sale, or distribute drugs by retails (Allopathic /Homeopathic drugs) on various forms viz. Form 20, Form 21 and Form 20F (Rule 64), Form 20A and Form 21A (Rule 62B), Form 20 C (Rule 67F)	Assistant Controller Drugs of the concerned District.	1 Month	Deputy Controller Drugs and Food Control Organization of the respective Division	Controller, Drugs and Food Control Organization J&K, Jammu/ Srinagar
02	License to sell, stock or exhibit (or offer) for sale, or distribute drugs by wholesale (Allopathic/Homeopathic drugs) on various forms viz. Form 20 B, Form 21B and Form 20G (Rule 64), Form 20BB and Form 21BB (Rule 62D), Form 20D (Rule 67F).	Deputy Controller Drugs and Food Control Organization of the respective Division	1 Month	Controller, Drugs and Food Control Organization J&K, Jammu/ Srinagar	Administrative Secretary to Government, Health & Medical Education Department
03	License to manufacture For sale or for distribution of drugs/cosmetics/ Ayurvedic (including Siddha or Unani drugs) on various forms viz. Form 25 and Form 25F (Rule 71), Form 25A (Rule 71 B), Form 25 B (Rule 71A), Form 25C (Rule 85E), Form 25D (Rule 157), Form 25E (Rule 154A), Form 26 (Rule 73), Form 26A (Rule 76A), Form 26 B (Rule 76), Form 29 (Rule 69), Form 32 (Rule 139), Form 32 A (Rule 142 B).	Controller, Drugs and Food Control Organization J&K, Jammu/ Srinagar	1 Month	Divisional Commissioner, Jammu/ Kashmir	Administrative Secretary to Government, Health & Medical Education Department