Kerala State Drugs and Pharmaceuticals Ltd
(A Govt. of Kerala Enterprise)
KALAVOOR, ALAPPUZHA
Phone :0477-2258184 (Extn:207) ; Email :purchase@ksdp.in
Web :http://www.ksdp.co.in

KSDP/PS/EQ/2024-25/450019/T-491 **22-3-2025**

Notice Inviting Email Quotation for the Supply of Raw Materials (LVP/SVP)

Quotations are invited for the supply of undermentioned goods/ services as per the attached specifications on FOR destination basis at our factory site at Kalavoor,Alapuzha, Kerala state

Sl No.	ITEM CODE & DESCRIPTION	UNIT	QUANTITY	EMD	TENDER FEES
1	10300043 Gentamicin sulphate IP	KG	1.000	♦ Not Applicable	♦ Not Applicable
2	10300044 Timolol maleate IP	KG	2.000		
3	10300045 Heparin Sodium IP (Derived from porcine intestinal mucosa)	KG	75.000		
4	10300046 Lignocaine Hydrochloride IP	KG	8.000		

Quotations should be submitted as per the Proforma given below on your letter head.

Sl No.	Name of item	Make	Rate per unit (including freight, if any)	GST %	Offer validity	Remarks, if any

Due Date/Time : 26-Mar-25 / 1.30pm **Opening Date/Time: 26-Mar-25 / 2pm**

Note: Please provide COA along with the quotation.

Offer validity*: Minimum 7 days offer validity from the date of closure of bid submission.

Quotation received with offer validity less than 7 days from the date of closure of bid submission will be entirely rejected.

Please send your lowest offers of the item to our e-mail ksdptender@gmail.com before 1 PM,26/03/2025.

The Quotation should be submitted through a password protected excel sheet.

Please share your Password to our email ksdptender@gmail.com@1.30 PM, 26.03.2025.

TERMS & CONDITIONS

Continued...

1.Payment terms	**:-**	**: 30 days after the receipt of the material along with documents, subject to QC approval.**
2.Mode of payment:-		**: E-Payment.**
3.Delivery Period:-		**Supply should be effected within 15 days on award of PO**
4.Special instructions:-		**Quotation number Should be Mention in the subject line**

HOD - Purchase

	KERALA STATE DRUGS AND PHARMACEUTICALS LTD, KALAVOOR PO, ALAPPUZHA, KERALA-688522
	Raw Material Specification

Name of the Material: **GENTAMICIN SULFATE I.P**			**(LVP/SVP/OPTH)**
SOP No:	KSDP/SOP/SPEC01	Spec. No:	KSDP/R/SPEC01/328
Effective date:	03/02/2025	Revision No:	00
		Item code:	10300043

Sl. No.	TEST	SPECIFICATION
1.	Description	White or almost white, hygroscopic powder.
2.	Solubility	Freely soluble in water, practically insoluble in ethanol (95 per cent).
3.	Identification	A. By Thin-layer chromatography. b. As per IP C. By chemically.
4.	Appearance of Solution	Not more intensely coloured than degree 6 of the appropriate range of reference solution
5.	pH	NLT 3.5 And NMT 5.5
6.	Specific Optical Rotation	NLT 107 And NMT 121(anhydrous substance).
7.	Composition	By Liquid chromatography.
8.	Related Substances*	By Liquid chromatography
9.	Sulfate	32.0 % to 35.0 % (anhydrous substance).
10.	Water*	Maximum 15.0%
11.	Bacterial Endotoxins	NMT 1.67 E/units per mg of gentamicin
12.	Assay	Minimum 590 µg/mg(anhydrous substance).

*In House specification.

The product complies to I.P. 2022 with respect to the above tests.

Prepared by	Checked by	Reviewed By	Approved By	Authorized By	
QA Officer	QC Head	Dy. Manager Production	Production Head	QA Head	MASTER CO
07/02/25	07/02/25	07/02/25	07/02/25	08/02/25	UNCONTROLLED COPY

	KERALA STATE DRUGS AND PHARMACEUTICALS LTD, KALAVOOR PO, ALAPPUZHA, KERALA-688522
	Raw Material Specification

Name of the Material: **LIGNOCAIN HCL I.P**			**(LVP/SVP/OPTH)**
SOP No:	KSDP/SOP/SPEC01	Spec. No:	KSDP/R/SPEC01/330
Effective date:	08/02/2025	Revision No:	00
		Item Code:	10300046

Sl. No.	PARAMETER	SPECIFICATION
1.	Description	A white, crystalline powder.
2.	Solubility	Highly soluble in water, ethanol, and organic solvent.
3.	Identification	A. By IR : To meet the test B. By Gives the chemically reaction. (chloride', a) C. By Gives the chemically reaction. D. By Gives the chemically reaction. E. By It gives reaction (A) of chlorides
4.	Appearance of solution	Water is clear and colorless.
5.	PH	4.0 to 5.5
6.	Heavy metals	2.0 g complies with the limit test for heavy metals
7.	Sulphates.	The remaining portion of the solution to which nothing has been added.
8.	Sulphated ash	Not more than 0.1 per cent.
9.	Water	5.0 to 7.5 per cent, determined
10.	Bacterial endotoxins*	Not more than 1.1 Eu/mg of Lignocaine hydrochloric.
11.	Assay	Not less than 99.0 % and Not more than to 101.0 % of $C_{14}H_{22}N_2O$, HCL. Calculated on dried basis.

*In House specification.

The product complies to I.P. 2022 with respect to the above tests.

Prepared by	Checked by	Reviewed By	Approved By	Authorized By
QA Officer	QC Head	Dy. Manager Production	Production Head	QA Head
07/02/25	07/02/25	07/02/25	07/02/25	08/02/25

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	KERALA STATE DRUGS AND PHARMACEUTICALS LTD, KALAVOOR PO, ALAPPUZHA, KERALA-688522
	Raw Material Specification

Name of the Material: **TIMOLOL MALEATE I.P**			**(LVP/SVP/OPTH)**
SOP No:	KSDP/SOP/SPEC01	Spec. No:	KSDP/R/SPEC01/333
Effective date:	08/02/2025	Revision No:	00
		Item Code:	10300044

Sl. No.	PARAMETER	SPECIFICATION
1.	Description	A white or almost white, crystalline powder or colorless crystals
2.	Solubility	Soluble in water, methanol, and alcohol.
3.	Identification	A. By IR : To meet the test B. By In the test for related substances, the principal peak in i chromatogram. C. By chemically.
4.	Appearance of solution	Not. more intensely coloured than reference solution BS8
5.	pH	3.8 to 4.3
6.	Specific optical rotation	-6.2° to - 5.7°.
7.	Enantiomeric purity	Determine by liquid chromatography.
8.	Related substances	Determine by liquid chromatography
9.	Sulphated ash	Not more than 0. I per cent
10.	Loss on drying	Not more than 0.5 per cent, determined on 1.0 g by drying in an oven 105°C
11.	Assay	Not less than 98.5 % and Not more than to 101.0 % of $C_{13}H_{24}N_4O_3S$, $C_4H_4O_4$. Calculated on dried basis.

*In House specification.

The product complies to I.P. 2022 with respect to the above tests.

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07/02/25	07/02/25	07/02/25	07/02/25	08/02/25

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	KERALA STATE DRUGS AND PHARMACEUTICALS LTD, KALAVOOR PO, ALAPPUZHA, KERALA-688522
	Raw Material Specification

Name of the Material: **HEPARIN SODIUM IP**			**(LVP/SVP/OPTH)**
SOP No:	KSDP/SOP/SPEC01	Spec. No:	KSDP/R/SPEC01/329
Effective date:	08/02/2025	Revision No:	00
		Item Code:	10300045

Sl. No.	TEST	SPECIFICATION
1.	Description	A white or grayish-white powder; moderately hygroscopic.
2.	Solubility	Soluble in water
3.	Identification	
A.	Clotting of freshly shed blood.	It delays the clotting of freshly shed blood.
B.	By Chemical.	White precipitate is formed
C.	By HPLC.	In the test for Over-sulphated chondroitin sulphate, the principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with reference solution (a).
4.	pH	5.0 to 7.5.
5.	Oversulphated chondroitin sulphate	Inject reference solution (b). The test is not valid unless there solution between the peaks due to oversulphated chondroitin sulphate and heparin is not less than 1.5. There tention time of heparin is about 30 minutes 'and of oversulphated chondroitin sulphate is about 50 minutes
		Inject the test solution and reference solution (a). The retentiontime of the principal peak obtained from the test solution corresponds to the peak obtained from reference solution (a).In the chromatogram obtained with the test solution, no peak corresponding to OSCS is observed.
6.	Heavy metal	Maximum 40 ppm

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	KERALA STATE DRUGS AND PHARMACEUTICALS LTD, KALAVOOR PO, ALAPPUZHA, KERALA-688522
	Raw Material Specification

7.	Protein and nucleotidic impurity	About 260 nm (2.4.7) (for nucleotides) and about 280 nm(for proteins) is not more than 0.2 and 0.15 respectively.
8.	Nitrogen	NMT 2.5 per cent, on the dried basis.
9.	Loss on drying	Maximum 8.0%.
10.	Sulfated ash	28.0 TO 41.0
11.	BET*	0.03 EU/Unit
12.	Sterility	Complies with the test for Sterility
13.	ASSAY (As on basis)	NLT 150 IU/ mg

*In House specification.

The product complies to I.P. 2022 with respect to the above tests.

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