SELVOK PHARMACEUTICAL CO.

147 , G.I.D.C. , Antalia , Bilimora – 396 325

QUALITY CONTROL DEPARTMENT

CERTIFICATE OF ANALYSIS FOR FINISHED PRODUCT
LIGNOCAINE HYDROCHLORIDE GEL IP 2% w/v

Name of Product : **Numbon 2% Gel**	A.R.No. : UFP/26/SG/485
Specification No. : SPC/FPS/STE.GEL/IP/01	Protocol No. : SPC/FPP/STE.GEL/IP/01

Batch No.	:	MA 015	Mfg. Date	:	07/2026
Batch size	:	302 Lits	Exp. Date	:	06/2028
Sampled Quantity	:	21 x 30 gm	Testing Date	:	22/07/2026
Received Date	:	22/07/2026	Release Date	:	05/08/2026

Result of Analysis As Per Specification

(Under the Drugs & Cosmetics Act 1940 and the Drugs & Cosmetics Rules 1945 thereunder)

Sr.No.	Test Parameters	Results	Specifications
1.0	Description	Clear,colourless viscous gel	Clear, Colourless viscous gel.
2.0	Identification	# by IR – Complies Bluish green ppts is produced Yellow colour is produced	The residue complies with the following tests. A. Determine by infrared absorption spectrophotometry. B. a bluish green precipitate is produced. C. a yellow colour is produced
3.0	Uniformity of weight	30.0323 gm	Not Less than Labelled amount 30.0 gm
4.0	pH	6.692	Limit : 6.0 to 7.0
5.0	Weight per Millilitre	1.0135 gm/ml	Limit : 1.0000 gm/ ml to 1.1000 gm/ml at 25° C
6.0	2,6-Dimethylaniline	Less than 400 ppm	Not more than 400 ppm
7.0	Assay (by Titration)	Estimated amount : 2.0095 % w/v (100.48 %)	Not less than 95.0% and Not more than 105.0 % of the Labelled amount of Lignocaine Hydrochloride IP equivalent to Lignocaine Hydrochloride Anhydrous 2.000% w/v
8.0	Sterility	Complies	By membrane filtration method No evidence of Microbial growth is found in 14 days

Tests complies with
Parishil Laboratories COA No : PLPL/PH/260727042 Date :28/07/2026

Opinion :In the opinion of the undersigned, the sample referred to above is OF STANDARD QUALITY / ~~NOT OF STANDARD QUALITY~~ as defined in the Act and the rules made there under for reason given Below. The sample COMPLIES / ~~DOES NOT COMPLY~~ with the prescribed above standard as per I.P./~~B.P. / U.S.P. / IN HOUSE~~ SPECIFICATION.

Prepered by :
05/08/2026
(QC Chemist)

Checked by :
05/08/2026
(QC Head)

Approved by :
05/08/2026
(QA Head)

SELVOK PHARMACEUTICAL CO.

147 , G.I.D.C. , Antalia , Bilimora – 396 325

QUALITY CONTROL DEPARTMENT

CERTIFICATE OF ANALYSIS FOR FINISHED PRODUCT
LIGNOCAINE HYDROCHLORIDE GEL IP 2% w/v

Name of Product : **Numbon 2% Gel**	A.R.No. : UFP/26/SG/486
Specification No. : SPC/FPS/STE.GEL/IP/01	Protocol No. : SPC/FPP/STE.GEL/IP/01

Batch No.	:	MA 016	Mfg. Date	:	07/2026
Batch size	:	302 Lits	Exp. Date	:	06/2028
Sampled Quantity	:	21 x 30 gm	Testing Date	:	22/07/2026
Received Date	:	22/07/2026	Release Date	:	05/08/2026

Result of Analysis As Per Specification

(Under the Drugs & Cosmetics Act 1940 and the Drugs & Cosmetics Rules 1945 thereunder)

Sr.No.	Test Parameters	Results	Specifications
1.0	Description	Clear,colourless viscous gel	Clear, Colourless viscous gel.
2.0	Identification	# by IR – Complies Bluish green ppts is produced Yellow colour is produced	The residue complies with the following tests, A. Determine by infrared absorption spectrophotometry. B. a bluish green precipitate is produced. C. a yellow colour is produced
3.0	Uniformity of weight	30.0319 gm	Not Less than Labelled amount 30.0 gm
4.0	pH	6.778	Limit : 6.0 to 7.0
5.0	Weight per Millilitre	1.0131 gm/ml	Limit : 1.0000 gm/ ml to 1.1000 gm/ml at 25° C
6.0	2,6-Dimethylaniline	Less than 400 ppm	Not more than 400 ppm
7.0	Assay (by Titration)	Estimated amount : 2.0362 % w/v (101.81 %)	Not less than 95.0% and Not more than 105.0 % of the Labelled amount of Lignocaine Hydrochloride IP equivalent to Lignocaine Hydrochloride Anhydrous 2.000% w/v
8.0	Sterility	Complies	By membrane filtration method No evidence of Microbial growth is found in 14 days

Tests complies with
Parishil Laboratories COA No : PLPL/PH/260727043 Date :27/07/2026

Opinion :In the opinion of the undersigned, the sample referred to above is OF STANDARD QUALITY / ~~NOT OF STANDARD QUALITY~~ as defined in the Act and the rules made there under for reason given Below. The sample COMPLIES / ~~DOES NOT COMPLY~~ with the prescribed above standard as per I.P./~~B.P. / U.S.P. / IN HOUSE~~ SPECIFICATION.

Prepered by : 05/08/2026
(QC Chemist)

Checked by : 05/08/2026
(QC Head)

Approved by : 05/08/2026
(QA Head)