ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : **Document No. : 252619066**

Ref. STS No.	4514	**Product Code**	FG03889
Product Name	TAB. TENLIZEM-M 500 SR		
Generic Name	TENELIGLIPTIN 20MG & METFORMIN HYDROCHLORIDE 500MG (SUSTAINED RELEASE) TABLETS		
Batch No.	425-1115	**Date Of Sampling**	07/06/2025
Batch Size	240000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	05/2025	**Exp. Date**	04/2028
A. R. No.	ZL25FG04301	**Date Of Release**	14/06/2025

S NO.	Tests	Specification	Results
1	Description	One side light yellow and other side white colour, elongated, biconvex ,Sustained Released tablets, scored on one side.	Complies
2	Identification	Should be positive for Metformin Hydrochloride and Teneligliptin.	Complies
3	Average Weight of Tablet	900.0 mg ± 5.0 %	898.0 mg
4	Uniformity of Weight	Average weight of tablet ± 5.0 %.	-0.44% to + 0.44%
5	Friability Test	NMT 1.0 %	0.20 %
6	Dissolution A (Metformin)	NLT 20.0% and NMT 60.0% in 1 Hours.	47.3% to 48.7 %
7	Dissolution B (Metformin)	NLT 40.0% and NMT 90.0 % in 3 Hours.	67.5% to 69.3 %
8	Dissolution C (Metformin)	Q NLT 70 % in 10 Hours. At 100 RPM of labeled amount of Metformin HCl Dissolve 1000 ml Phosphate Buffer pH-6.8.	97.3% to 99.9 %
9	Dissolution Teneligliptin	Q NLT 70 % of the stated amount of Teneligliptin	83% to 98 %
10	Assay Each uncoated bilayer tablet contains: Teneligliptin Hydrochloride Hydrate I.P. Eq. to Teneligliptin 20mg	NLT 90.0 % and NMT 110.0 % of the labeled amount	96.1 %
11	Metformin Hydrochloride I.P. 500 mg (in sustaned release from)	NLT 90.0 % and NMT 110.0 % of the labeled amount	97.8 %

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP/USP~~/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	14/06/2025	14/06/2025	
Name:			
Designation:	Officer QC	Sr. Officer	QC MANAGER

ZEE LABORATORIES LTD. Baddi (H.P.) QC MANAGER 14/06/2025

ZEE/QC/072/02-00

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252615158 **Sample Rec No. :** 252617434 **QC No. :** 252620857

Ref. STS No.	2541	**Product Code**	FG03267
Product Name	TAB. VOGLITER-0.3		
Generic Name	VOGLIBOSE TABLETS I.P. 0.3MG		
Batch No.	425-1271	**Date Of Sampling**	23/06/2025
Batch Size	200000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	06/2025	**Exp. Date**	05/2028
A. R. No.	ZL25FG04779	**Date Of Report**	25/06/2025

S NO.	Tests	Specification	Results
1	Description	Light yellow colour, rounded, biconvex, uncoated tablets, both side are smooth.	Complies
2	Identification By hplc	In the Assay,the principal pak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with reference solution.	Complies
3	Average Weight of Tablet	120.0 mg ± 7.5%	122.3 mg
4	Uniformity of Weight	Not more than two of the individual weights deviate from the Average weight by more than the 7.5% and none deviates by more than 15%.	Complies
5	Disintegration Test	Not more than 15 minutes	05:41 Min
6	Friability Test	NMT 1.0 % w/w	0.23%
7	Uniformity of content	Not less than 85.0% and Not more than 115.0% of the Average content.	Complies
8	Composition: Assay Each Uncoated tablet contains : Voglibose I.P. 0.3 mg	NLT 90.0 % and NMT 110.0 % of the labeled amount of Voglibose.	99.3 %

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP/USP~~/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	25/06/2025	25.6.2025	25/06/2025
Name:	Manpreet	Mukesh Kashyap	Jaideep Singh
Designation:	Officer QC	Sr. officer	QCM

ZEE LABORATORIES LTD. QC MANAGER

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. :

Document No. : 242550143

Ref. STS No.	325	**Product Code**	FG05557
Product Name	TAB. TELMIZEM AMH		
Generic Name	TELMISARTAN 40MG, AMLODIPINE 5MG, HYDROCHLORTHIAZIDE 12.5MG TABLETS		
Batch No.	425-280	**Date Of Sampling**	17/02/2025
Batch Size	300000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	01/2025	**Exp. Date**	12/2027
A. R. No.	ZL25FG01331	**Date Of Release**	19/02/2025

S NO.	Tests	Specification	Results
1	Description	White colour rounded film coated tablet scored on one side.	Complies
2	Identification	Should be positive for Telmisartan, Amlodipine & Hydrochlorothiazide.	Complies
3	Average Weight of Tablet	300.0 mg ± 5%	300.1 mg
4	Uniformity of Weight	Average weight of tablet ± 5%.	Complies
5	Disintegration Test	NMT 30 minutes	03:14min
6	Assay Each film coated tablet contains Amlodipine Besylate I.P Eq to Amlodipine 5 mg	NLT 90.0% and NMT 110.0% of the labeled amount	103.2%
7	Telmisartan I.P 40 mg	NLT 90.0% and NMT 110.0% of the labeled amount	98.6%
8	Hydrochlorothiazide I.P 12.5 mg	NLT 90.0% and NMT 110.0% of the labeled amount	101.8%

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per ~~IP/BP/U~~SP/In ~~h~~ouse specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	19/02/2025	19/02/2025	QC MANAGER 21/02/2025
Name:	Raghuveer	Mukesh Mishra	[illegible]
Designation:	Officer QC	Sr. Officer	QCM

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : **Document No. : 242531350**

Ref. STS No.	216	**Product Code**	FG6377
Product Name	TAB. MISS ME		
Generic Name	Tadalafil Tablets I.P		
Batch No.	Z24-763	**Date Of Sampling**	21/09/2024
Batch Size	81000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	09/2024	**Exp. Date**	08/2027
A. R. No.	ZL24FG07451	**Date Of Release**	26/09/2024

S NO.	Tests	Specification	Results
1	Description	A light pink colour, rounded, film coated tablets and both sides are smooth.	Complies
2	Identification By hplc	In the Assay, the principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained wih reference solution.	Complies
3	Average Weight of Tablet	120.0 mg ± 7.5 %	121.6 mg
4	Uniformity of Weight	Average weight of tablet ± 7.5%.	Complies
5	Disintegration Test	Not more than 30 minutes	02:27 Min
6	Dissolution (%)	Q.Not less than 40 % of the labeled amount of Tadalafil in 10 min .	78.21% - 94.85 %
7	Dissolution (%)	Q. Not less than 80 % of the labeled amount of Tadalafil in 30 min.	95.60% - 106.95 %
8	Uniformity of Content	NLT 85% TO 115% of the Average content.	92.37% - 102.08 %
9	Related substance	The area of secondary peak 0.2%.The sum of the areas of all the secondary peaks 0.3%.	Complies
10	Assay Each Film coated tablet contains Tadalafil I.P. 10 mg	NLT 90.0% and NMT 110.0% of the labeled amount of Tadalafil.	96.09 %

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP/USP~~/In house specification.

Particulars	Prepared By	Checked By	Q.A. Approved By
Sign & Date	26/09/2024	26/09/2024	26/09/2024
Name:	Rajkumari	Mukesh Mishra	Thakur Bhim Singh
Designation:	Officer Q.C	Sr. Officer	QCM

ZEE LABORATORIES LTD. MANAGER

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252614285 **Sample Rec No. :** 252620590 **QC No. :** 252624990

Ref. STS No.	548	**Product Code**	FG03221
Product Name	TAB. ESZONE-40		
Generic Name	ESOMEPRAZOLE TABLETS I.P. 40MG		
Batch No.	425-1137	**Date Of Sampling**	24/07/2025
Batch Size	120000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	05/2025	**Exp. Date**	04/2028
A. R. No.	ZL25FG05719	**Date Of Report**	24/07/2025

S NO.	Tests	Specification	Results
1	Description	A brown colour, elongated, biconvex, enteric coated tablets both sides are plain	Complies
2	Identification	Should be positive for Esomeprazole Magnesium.	Complies
3	Average Weight of Tablet	407.0 mg ± 5.0 %	397.5 mg
4	Uniformity of Weight	Average weight of tablet ± 5%.	Complies
5	Disintegration Test (A)	a) Do not disentegrate in 0.1 M hydrochloric acid in 2 hours.	Complies
6	Disintegration Test (B)	b) Disentegrate in mixed phosphate buffer pH 6.8 in 60 minutes .	23:25Min
7	Dissolution Test (A)	A) Not more than 10% of the labeled amount of Esomeprazole dissolves in 120 minutes at 75 rpm in 1000ml 0.1 M HCl as dissolution medium.	Complies
8	Dissolution Test (B)	B) Q Not less than 70% of the labeled amount of Esomeprazole dissolves in 60 minutes at 75 rpm in 900 ml Phosphate Buffer pH 6.8 as dissolution medium.	82% to 98%
9	Assay Each enteric coated tablet contains Esomeprazole Magnesium I.P. Eq. to Esomeprazole 40 mg	NLT 90.0 % and NMT 110.0 % of the labeled amount of Esomeprazole	100.0 %

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP~~/USP/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	24/07/2025	24/7/2025	24/07/2025
Name:			
Designation:			

ZEE LABORATORIES LTD. Q.C MANAGER

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252612545 **Sample Rec No. :** 252613903 **QC No. :** 252616206

Ref. STS No.	265	**Product Code**	FG06582
Product Name	TAB. TENLIZEM-20		
Generic Name	TENELIGLIPTIN TABLETS I.P. 20MG		
Batch No.	425-915	**Date Of Sampling**	20/05/2025
Batch Size	150000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	05/2025	**Exp. Date**	04/2028
A. R. No.	ZL25FG03718	**Date Of Report**	20/05/2025

S NO.	Tests	Specification	Results
1	Description	A yellow colour, rounded, biconvex shaped film coated tablets, Scord on one side.	Complies
2	Identification	Should be positive for Teneligliptin Hydrobromide.	Complies
3	Average Weight of Tablet	224.4 mg ± 7.5 %	217.2 mg
4	Uniformity of Weight	Average weight of tablet ± 7.5 %.	Complies
5	Disintegration Test	Not more than 30 minutes	06:41 Min
6	Dissolution Test	Q NLT 70.0 % of the labeled amount of Teneligliptin.	85% to 92%
7	Assay Each film coated tablet contains: Teneligliptin Hydrobromide Hydrate I.P. Eq. to teneligliptin 20mg	NLT 90.0 % and NMT 110.0 % of the labeled amount	96.3%

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP/USP~~/~~In house~~ specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	20/05/2025	20/05/2025	20/05/2025
Name:	Rajkumari	Mukesh Thakur	Thakur Rajesh Singh
Designation:	Officer A/C	Sr. officer	QC Mg

ZEE LABORATORIES LTD. Q.C MANAGER

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252610784 **Sample Rec No. :** 252611751 **QC No. :** 252612802

Ref. STS No.	1254	**Product Code**	FG03281
Product Name	TAB. METOLASH-AM 50		
Generic Name	METOPROLOL SUCCINATE PROLONGED-RELEASE & AMLODIPINE BESILATE TABLETS I.P.		
Batch No.	425-711	**Date Of Sampling**	28/04/2025
Batch Size	100000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	04/2025	**Exp. Date**	03/2028
A. R. No.	ZL25FG03065	**Date Of Report**	28/04/2025

S NO.	Tests	Specification	Results
1	Description	A orange colour, rounded, prolonged tablets, both side are smooth.	Complies
2	Identification	Should be positive for Metoprolol succinate & amlodipine Besilate.	Complies
3	Average Weight of Tablet	320.0 mg ± 5.0%	326.0 mg
4	Uniformity of Weight	Average weight of tablet ± 5.0%	Complies
5	Dissolution Test (A)	NLT 30.0% and NMT 65.0% in 2Hours	42% to 55%
6	Dissolution Test (B)	NLT 60.0% and NMT 95.0% in 6Hours	72% to 86%
7	Dissolution Test (C)	Q Not less than 70% of the labeled amount of dissolves in 10th Hours.	92% to 93%
8	Uniformity of Content.	NLT 85% to 115% of the Average content.	Complies
9	Related substances.	Should be complies.	Complies
10	Assay Each prolonged tablet contains: Amlodipine Besilate I.P. eq. to Amlodipine 5mg	NLT 90.0% and NMT 110.0% of the labeled amount of amlodipine Besilate.	98.4 %
11	Metoprolol Succinate I.P. 47.5 mg Eq to Metoprolol Tartrate 50 mg (Extended Release)	NLT 90.0% and NMT 110.0% of the labeled amount of Metoprolol Succinate	100.3%

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP/USP~~/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	28/04/2025	28/4/2025	28/04/2025
Name:	Raghuveer	Mukesh Mishra	Thakur Dinu Singh
Designation:	Officer Q.C	Sr. Officer	QCM

ZEE LABORATORIES LTD. Q.C MANAGER

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252613390 **Sample Rec No. :** 252615500 **QC No. :** 252618416

Ref. STS No.	2022	**Product Code**	FG05753
Product Name	TAB. TELMIZEM BETA 50		
Generic Name	METOPROLOL SUCCINATE ER 50MG & TELMISARTAN 40MG TABLETS		
Batch No.	425-1015	**Date Of Sampling**	03/06/2025
Batch Size	100000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	05/2025	**Exp. Date**	04/2028
A. R. No.	ZL25FG04202	**Date Of Report**	09/06/2025

S NO.	Tests	Specification	Results
1	Description	A White colour, rounded, film coated tablets, scored on one side.	Complies
2	Identification	Should be positive for Metoprolol Succinate and Telmisartan.	Complies
3	Average Weight of Tablet	440.0 mg ± 5.0%	439.2 mg
4	Uniformity of Weight	NMT two of the individual weights deviate from the Average weight by more than the 5% and none deviates by more than 10%.	Complies
5	Dissolution Test	NLT 30.0% and NMT 65.0% In 2 hours	42% to 55%
6	Dissolution Test	NLT 60.0% and NMT 95.0% In 6 hours	72% to 82%
7	Dissolution Test	Q NLT 70.0% In 10 hours	86 % to 98%
8	Assay Each film coated tablet contains: Metoprolol Succinate I.P Eq to Metoprolol Tartrate (Extended Releas) 50 mg	NLT 90.0% and NMT 110.0% of the labeled amount of Metoprolol Tartrate	99.2%
9	Telmisartan I.P. 40mg	NLT 90.0% and NMT 110.0% of the labeled amount of Telmisartan	102.4%

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~BP~~/~~USP~~/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	09/06/2025	09.6.2025	09/06/2025
Name:	Rajkumari	Mukesh Mishra	Thakur Dev Singh
Designation:	Officer Q.C	Sr. officer	QCM

ZEE LABORATORIES LTD. Q.C MANAGER

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252614314 **Sample Rec No. :** 252616095 **QC No. :** 252619232

Ref. STS No.	265	**Product Code**	FG05049
Product Name	TAB. DEXZEE 0.5		
Generic Name	DEXAMETHASONE TABLETS I.P. 0.5MG		
Batch No.	425-1147	**Date Of Sampling**	10/06/2025
Batch Size	180000.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	05/2025	**Exp. Date**	04/2028
A. R. No.	ZL25FG04378	**Date Of Report**	16/06/2025

S NO.	Tests	Specification	Results
1	Description	White colour, rounded uncoated tablets and Scored on one side.	Complies
2	Identification	Should be positive for dexamethasone.	Complies
3	Average weight of Tablet	80.0 mg ± 10 %	77.3mg
4	Uniformity of Weight	Average weight of tablet ± 10 %	Complies
5	Disintegration Test	Not more than 15 minutes	03:45 Min
6	Friability Test	Not more than 1.0 % w/w	0.15%
7	Uniformity of content	NLT 85.0% and NMT 115.0%	Complies
8	Assay: Each uncoated tablet contains: dexamethasone I.P. Eq. to dexamethasone 0.5 mg	NLT 90.0 % and NMT 110.0 % of the labeled amount	98.5%

Remarks ~~The~~ above sample complies/~~does not~~ comply with prescribed standards of Quality with respect to above tests as per IP/~~BP/USP~~/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	16/06/2025	16/06/2025	16/06/2025
Name:	Raghuveer	Mukesh Mishra	Thakur ... Singh
Designation:	officer QC	Sr. officer	QCM

QC MANAGER (stamp: ZEE LABORATORIES LTD.)

ZEE LABORATORIES LTD

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 232430507 **Sample Rec No. :** 232439256 **QC No. :** 232450332

Ref. STS No.	265	**Product Code**	FG06140
Product Name	TAB. HEPADEM-400		
Generic Name	Ademetionine Tablets 400mg		
Batch No.	424-421	**Date Of Sampling**	01/03/2024
Batch Size	29500.000 Tab	**Sample Quantity**	60.000 Tab
Mfg. Date	02/2024	**Exp. Date**	01/2027
A. R. No.	ZL24FG01910	**Date Of Report**	13/03/2024

S NO.	Tests	Specification	Results
1	Description	A white colour, elongated, biconvex shaped uncoated tablet. One side plain and other side having mid dividing line.	Complies
2	Identification	Should be positive for Ademetionine	Complies
3	Average Weight of Table	1010.0 mg ± 5%	1031.0 mg
4	Uniformity of Weight	Average weight of tablet ± 5.0%.	Complies
5	Disintegration test	Not more than 15 minutes	09:31 Min
6	Assay Each enteric coated tablet contains: Ademetionine eq to Ademetionine Ion 400mg	NLT 90.0% and NMT 110.0% of the labeled amount	107.01 %

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per ~~IP/BP/U~~SP/In house specification.

Particulars	Prepared By	Checked By	Approved By
Sign & Date	13/03/2024	13/3/2024	13/03/2024
Name:	Rajkumar	Mukesh Mishra	Thakur ... Singh
Designation:	Officer QC	Sr. officer	QCM

QC MANAGER