



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 \pm 5.0 % | 898.0 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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 sustained 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 | 14/06/2025 |
| Name: | Raynuree | Manish Mishra | Manish Mishra |
| Designation: | Officer Q.C. | Officer | Officer |



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 peak 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 \pm 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 | 25/06/2025 | 25/06/2025 |
| Name: | Harpreet | Manish Mishra | Manish Mishra |
| Designation: | Officer in | Officer | Officer |



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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 \pm 5% | 300.1 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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/USP/In house specification.

| Particulars | Prepared By | Checked By | Approved By |
|--------------|-------------|--------------|-------------|
| Sign & Date | 19/02/2025 | 19/02/2025 | 19/02/2025 |
| Name: | Rajhans | Mukul Mishra | Sh. Office |
| Designation: | Officer | Sh. Office | Officer |

ZEE/QC/072/02-00



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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 with reference solution. | Complies |
| 3 | Average Weight of Tablet | 120.0 mg \pm 7.5 % | 121.6 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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. Manager |
|--------------|--------------|----------------|------------------|
| Sign & Date | L 26/09/2024 | MKS 26/09/2024 | 26/09/2024 |
| Name: | Rajhans | Mukul Mishra | Tarun Deen Singh |
| Designation: | Officer | Officer | Officer |

ZEE/QC/072/02-00



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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 \pm 5.0 % | 397.5 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 5%. | Complies |
| 5 | Disintegration Test (A) | a) Do not disintegrate in 0.1 M hydrochloric acid in 2 hours. | Complies |
| 6 | Disintegration Test (B) | b) Disintegrate 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 | <i>24/07/2025</i> | <i>24/07/2025</i> | <i>24/07/2025</i> |
| Name: | <i>Manoj Kumar</i> | <i>Manoj Kumar</i> | <i>Manoj Kumar</i> |
| Designation: | <i>QA Officer</i> | <i>QA Officer</i> | <i>QA Officer</i> |





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CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

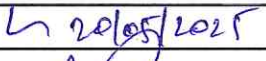

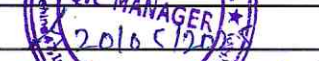
(The Drugs & Cosmetics Act 1940 at the under form 39)

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 \pm 7.5 % | 217.2 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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: | Prayanshu | Mukesh Tansky | Thakur Deep Singh |
| Designation: | Officer Q.C. | SS. officer | QC |





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CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

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

Work Order No. : 252610784

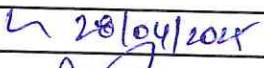
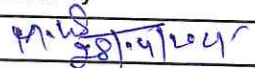
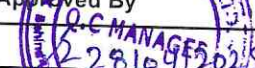
Sample Rec No. : 252611751

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 \pm 5.0% | 326.0 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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 IP/BP/USP/In house specification. Quality with respect to above tests as per

| Particulars | Prepared By | Checked By | Approved By |
|--------------|---|--|---|
| Sign & Date |  28/04/2025 |  28/04/2025 |  28/04/2025 |
| Name: | Deepak Kumar | Manish Mishra | Manish Mishra |
| Designation: | Officer in Charge | Officer in Charge | Officer in Charge |





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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 \pm 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 | 6/09/2025 | 6/09/2025 | 6/09/2025 |
| Name: | Prashant | Prashant | Prashant |
| Designation: | Officer | Officer | Officer |





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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 \pm 10 % | 77.3mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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 | <i>[Signature]</i> 16/06/2025 | <i>[Signature]</i> 16/06/2025 | <i>[Signature]</i> 16/06/2025 |
| Name: | Rayhunnas | Amrisha Prasad | Thakur An Singh |
| Designation: | officer a/c | SR officer | QRM |



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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 \pm 5% | 1031.0 mg |
| 4 | Uniformity of Weight | Average weight of tablet \pm 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/USP/In house specification.

| | | | |
|--------------|-------------|---------------|--------------|
| Particulars | Prepared By | Checked By | Approved By |
| Sign & Date | 13/03/2024 | 13/03/2024 | 13/03/2024 |
| Name: | Rayluces | Mukesh Mishra | Thakur Singh |
| Designation: | Officer QC | Officer | OCM |