

# CERTIFICATE OF ANALYSIS

|                      |  |  |                     |
|----------------------|--|--|---------------------|
| Product Name:        | <b>Clinical Curcumin™ Capsules</b>   |  |                     |
| Product #:           | <b>PY097</b>   | Manufactured for: Purity Products                      |                     |
| Lot#:                | T21014   |  |                     |
| Date Manufactured:   | 8/2021   |  |                     |
| Product Appearance:  | #00 clear/clear oblong capsule with orangish-brown powder fill. Result: Passed |  |                     |
| Weight Variation:    | Theoretical Weight: 869.95 mg<br>Result: 923.71 mg                             | Specification: 782.96-956.94 mg<br>Method: Current USP |                     |
| Disintegration Time: | Specification: NMT 30 mins.  | Result: 8 mins.  | Method: Current USP |
| Reference:           | B6p214   |  |                     |

## DIETARY INGREDIENTS

| Ingredient Name  | LC/3caps   | Result     | % of LC | Method |
|--|------------|------------|---------|--------|
| Meriva® Enovita® & Green Tea Phytosome™ Blend [Providing 1000mg Meriva® Turmeric (Curcuma longa rhizome), 150 mg of Enovita® Grapeseed (Vitis vinifera L.) and 150 mg Green Tea Phytosome™ Camellia sinensis leaves] | 1300.00 mg | 1300.00 mg | 100.00  | **     |

## OTHER INGREDIENTS

Rice flour, gelatin (capsule), turmeric powder, calcium palmitate, silicon dioxide, and sunflower lecithin.

## HEAVY METALS

| Heavy Metal | Specification                | Result/3caps | Method |
|-------------|------------------------------|--------------|--------|
| Lead:       | ≤ 2.75 µg/maximum daily dose | 0.156 mcg    | ICP-MS |
| Arsenic:    | ≤ 10 µg/maximum daily dose   | 0.159 mcg    | ICP-MS |
| Cadmium:    | ≤ 4.1 µg/maximum daily dose  | 0.049 mcg    | ICP-MS |

## MICROBIOLOGY

| Micro Study#MB0019557  | Specification  | Result    | Method      |
|------------------------|----------------|-----------|-------------|
| Total Plate Count:     | < 10,000 CFU/g | 50 cfu/g  | Current USP |
| Yeast & Mold Count:    | < 1,000 CFU/g  | <10 cfu/g | Current USP |
| Escherichia Coli:      | Negative       | ND        | Current USP |
| Salmonella:            | Negative       | ND        | Current USP |
| Staphylococcus Aureus: | Negative       | ND        | Current USP |

|                                     |                |
|-------------------------------------|----------------|
| Prepared By: <i>Rebecca Rivera</i>  | Date: 9/8/2021 |
| Reviewed By: <i>Rajivdya Thakur</i> | Date: 9/8/2021 |
| Approved By: <i>Santoshkumar</i>    | Date: 9/8/2021 |

\*\* - In Accordance with 21 CFR 111.75(d) (1) the following finished dietary ingredients product specifications are Exempt from direct finished batch testing requirements as set forth in paragraph 21 CFR 111.75(c) (1). Also Confirmed by proper raw material identification, verified by production process controls and QA batch record Review and approval to ensure finished product meets all approved MMR in-process specifications. Ref: SOP No. QC-3.