

CERTIFICATE OF ANALYSIS

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|----------------------|--|-----------------------------------|---------------------|---------------------|
| Product Name: | Ubiquinol Daily Super Strength Liquid Vege Capsules | | | |
| Product #: | PY086 | Manufactured for: Purity Products | | |
| Lot#: | 211445 | | | |
| Date Manufactured: | 6/2021 | | | |
| Product Appearance: | #0 Green/Green vegetarian capsule with a yellow to orange fill. Result: Passed | | | |
| Weight Variation: | Theoretical Weight: 590 mg | Specification: 531-649 mg | Result: 600.43 mg | Method: Current USP |
| Disintegration Time: | Specification: NMT 30 minutes | Result: 12 mins. | Method: Current USP | |
| Reference: | B1386p172 | | | |

DIETARY INGREDIENTS

| Ingredient Name | LC/2caps | Result | % of LC | Spec | Method |
|----------------------------------|-----------|-----------|---------|----------|--------|
| Ubiquinol (Kaneka QH Ubiquinol®) | 200.00 mg | 200.00 mg | 100.00 | NLT 100% | ** |

OTHER INGREDIENTS

Rice Bran Oil, gelatin (capsule), Antioxidant Stabilizer (d-alpha tocopheryl acetate, Rosemary Oil, Ascorbyl Palmitate), Chlorophyll.

HEAVY METALS

| Heavy Metal | Specification | Result/2caps | Method |
|-------------|------------------------------|--------------|--------|
| Lead | ≤ 2.75 µg/maximum daily dose | 0.018 mcg | ICP-MS |
| Arsenic | ≤ 10 µg/maximum daily dose | ND | ICP-MS |
| Cadmium | ≤ 4.1 µg/maximum daily dose | 0.007 mcg | ICP-MS |

MICROBIOLOGY

| Micro Study#MB0017066 | Specification | Result | Method |
|------------------------|----------------|----------|-------------|
| Total Plate Count: | < 10,000 CFU/g | 10 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 |

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| Prepared By: <i>Rebecca Ruan</i> | Date: 6/8/2021 |
| Reviewed By: <i>Raividya Thacker</i> | Date: 6/8/2021 |
| Approved By: <i>Santosh Kumar</i> | Date: 6/8/2021 |

** - 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.