INNOVATION LINE
LOW PRICES, HIGH STANDARDS!

CERTIFICATION OF COMPLIANCE

GENERAL CERTIFICATION OF CONFORMITY

1. **Description Of Product Covered in Certificate:**

   #Sunscreen Bottles

2. **Compliance Testing Certification:**
   - **A. Total Lead and Cadmium (CA Prop 65) -** PASS
   - **B. Total Phthalates (CA Prop 65)-** PASS
   - **C. Not Intended for Children Under 12yrs**

3. **Importer or Manufacturer Certifying Compliance**
   - **A. Name:** Innovation Line
   - **B. Address:** California, USA

4. **Certifying Agent Contact Information**
   - **A. Name:** Compliance Manager
   - **B. Address:** Los Angeles, California, USA
   - **C. Telephone Number:** (310) 398- 8116

5. **Origin and Date of Manufacture**
   - **A. Location:** China
   - **B. Date (Month/Year):** APRIL 2022

6. **Location and Date of Product Testing:**
   - **A. Location:** Hangzhou, China
   - **B. Date (Month/Year)**: MAY 2022

7. **Third Party Testing Agent:**
   - **A. Name:** QIMA
   - **B. Address**:
     5/F A2 Building, No 1213 Huoju South. Rd., Puyan Street. Binjiang District, Hangzhou, China

This certificate is issued pursuant by the Importer of Record for the Consumer Product Safety Improvement Act of 2008, Section 14, where required. This certificate confirms that the tests reported above have been completed for the product by a third-party laboratory under the reasonable program of testing and that the product complies with rules, bans, standards or regulations applicable to the product as listed above. This certificate accompanies the product and is also available from the importer/domestic manufacturer referenced above. The information set forth herein is presented in good faith and is believed to be accurate and correct as of the date of issue and is subject to change without further notice.

Certificate of Registration

This certifies that:

Innovation Specialties, Inc dba. Innovation Line

Is registered with the U.S. Food & Drug Administration, Center for Drug Evaluation and Research, Office of Drug Registration and Listing, as a Drug Establishment pursuant to the Code of Federal Regulations 21 CFR 207.

FDA Drug Establishment #1000300091/Labeler code #76138/ NDC#76138-220-02 for item #5298

ISI - FDA Registrations:
Current

Eddie Blau

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