United States Food and Drug Administration

Center for Drug Evaluation and Research

10903 New Hampshire Ave, Silver Spring, MD 20993, United States of America

CDERExportCertificateProgram@fda.hhs.gov - Telephone (301) 796-4950

Certificate of a Pharmaceutical Product - Active Pharmaceutical Ingredient (API)

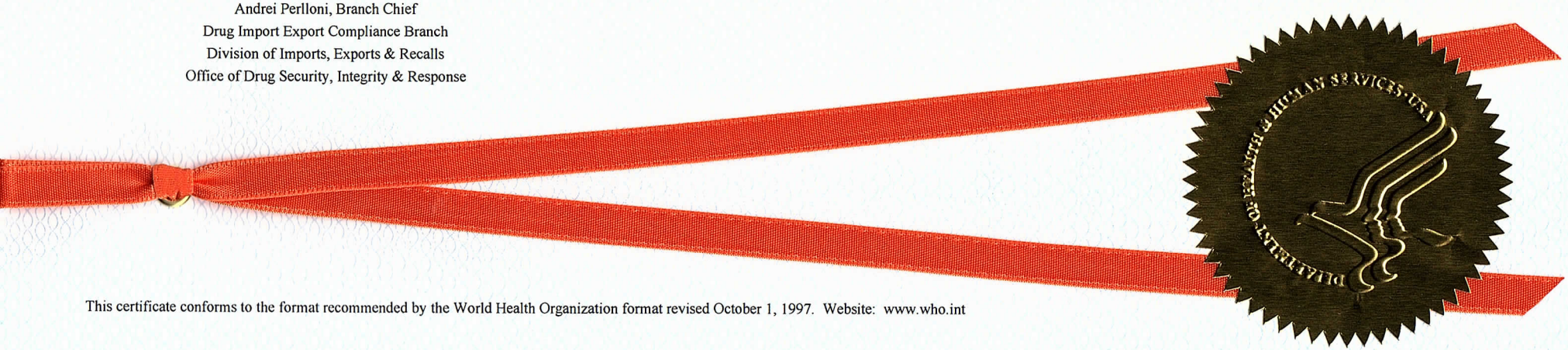
Certificate Number: **WPWN-62TV** Certificate Issue Date: **September 22, 2017** Certificate Expiration Date: **September 21, 2019**

Importing Country: **ITALY** Exporting Country: **UNITED STATES of AMERICA**

1.	Drug Trade Name, International or National non-proprietary name (as applicable) & dosage form: **ZINC OXIDE**
1.1	Active Ingredient(s) and amount(s) per unit dose (complete quantitative composition is preferred): **See Attachments**
1.2	Is this product licensed to be placed on the market for use in the exporting country? **No**
1.3	Is this product actually on the market in the exporting country? **Yes**
2.B.1	Applicant for certificate name & address: **Zinc Oxide LLC, 600 Printwood Drive, Dickson, TN 37055 United States of America**
2.B.2	Status of Applicant: **Manufacturer**
2.B.2.1	Manufacturer name & address: **Zinc Oxide LLC, 600 Printwood Drive, Dickson, TN 37055 United States of America**
2.B.3	Why is marketing authorization lacking? **Not Applicable**
2.B.4	Remarks:**The firm proposes to export the active pharmaceutical ingredients (API) listed above, which when properly labeled with statement "Caution: For further manufacturing, processing or repacking", may be freely marketed in the United States of America at this time.**
3.	Does the certifying authority arrange for periodic inspection of the manufacturing plant in which the dosage form is produced? **Yes**
3.1	Periodicity of routine inspections (years): **Pursuant to section 510(h)(3) of the Federal Food, Drug & Cosmetic Act, Inspections will occur in accordance with a risk-based schedule**
3.2	Has the manufacture of this type of dosage form been inspected? **Yes**
3.3	Do the facilities and operations conform to GMPs as recommended by the WHO? (GMPs including 21 Code of Federal Regulations parts 210, 211, or ICH Q7A): **Yes, at time of inspection, site complies with FDA cGMP**
3.4	Does the information submitted by the applicant satisfy the certifying authority on all aspects of the manufacture of the product undertaken by another party? **Yes**

Andrei Perlloni

Andrei Perlloni, Branch Chief
Drug Import Export Compliance Branch
Division of Imports, Exports & Recalls
Office of Drug Security, Integrity & Response



This certificate conforms to the format recommended by the World Health Organization format revised October 1, 1997. Website: www.who.int