

GENERAL CERTIFICATE OF CONFORMITY

U.S. Importer or Domestic Manufacturer

(Liquid manufacturer, not bottle manufacturer)

Company Name: Paragon Liquids LLC
Address Line 1: 69-B Mall Drive
Address Line 2:
City/State/Zip: Commack, NY 11725

Individual Maintaining Records of Test Results

Person's Name: Kimberly Sanders
Address Line 1: 217 West Marie Street
Address Line 2:
City/State/Zip: Hicksville, NY 11801
Email Address: ks@serumvape.com
Telephone #: (631) 707-6139

Date and Place Where Product Was Manufactured or Assembled

(For the date[s] when the product was manufactured, provide at least the month and year. For the place of manufacture provide at least the city [or administrative region] and country where the product was manufactured or finally assembled. If the same manufacturer operates more than one location in the same city, provide the street address of the factory.)

Company Name: Paragon Liquids LLC
Address Line 1: 69-B Mall Drive
Address Line 2:
City/State/Zip: Commack, NY 11725
Country: USA
Date of Manufacture: 03/20/2015

Citation to Safety Regulation to which this product is being Certified

16 CFR Part 1700 - Poison Prevention Packaging
(As amended by S. 142 - The Child Nicotine Poisoning Prevention Act of 2015) (CNPPA)
Pub. L. No. 114-116

Product Being Certified

Describe the product(s) covered by this certification in enough detail to match the certificate to each product it covers and no others.

E-Liquid Name: Brand: Dr. Juice LLC
Flavor(s): Funnel Vision
Bottle Type/Size: Flint Glass Boston Round, 60ml
Nicotine Level: 0mg, 3mg, 6mg and 12mg

Date(s) and Place When the Product Was Tested for Compliance

Provide the location(s) of the testing and the date(s) of the test(s) or test report(s) on which certification is being based.

Company Name: Perritt Laboratories
Address Line 1: 145 South Main Street
Address Line 2: P.O. Box 147
City/State/Zip: Heightstown, NJ 08520
Country: U.S.A.
Date of Test(s) or
Test Reports: 11/16/1999

Identification of any third party laboratory on whose testing the certificate depends.

Generally, this section should be labeled "N/A" for a GCC because third party laboratory testing is not a requirement for non-children's products. (It is only a requirement for children's products and must be included in a CPC.) However, if a certifier voluntarily uses test results from a third party laboratory as the basis for issuing its GCC, the law requires that the certifier must then provide the name, full mailing address, and telephone number of the third party laboratory.

Company Name: N/A
Address Line 1:
Address Line 2:
City/State/Zip:
Country:
Telephone #:

NOTES: