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Rauwolfia Serpentina Tablets

» Rauwolfia Serpentina Tablets contain an amount of reserpine-rescinnamine group alkaloids, calculated as reserpine, equivalent to not less than 0.15 percent and not more than 0.20 percent of the labeled amount of powdered rauwolfia serpentina.

Packaging and storage—Preserve in tight, light-resistant containers.

USP REFERENCE STANDARDS (11).—
[USP Rauwolfia Serpentina RS](#)
[USP Reserpine RS](#)

Identification—The powdered rauwolfia serpentina in the Tablets conforms to the requirements for *Ground Rauwolfia Serpentina root* under *Botanic characteristics* and meets the requirements of the test for *Chemical identification* under [Rauwolfia Serpentina](#).

MICROBIAL ENUMERATION TESTS (61), and **TESTS FOR SPECIFIED MICROORGANISMS** (62).—Tablets meet the requirements of the test for absence of *Salmonella* species.

DISINTEGRATION (701): 1 hour, with disks using simulated gastric fluid TS, without enzyme.

UNIFORMITY OF DOSAGE UNITS (905): meet the requirements.

Assay—Weigh and finely powder not less than 20 Tablets. Weigh accurately a portion of this powder, equivalent to 2.5 g of powdered rauwolfia serpentina, and proceed as directed for *Procedure* in the Assay under [Rauwolfia Serpentina](#).

Auxiliary Information - Please [check for your question in the FAQs](#) before contacting USP.

Topic/Question	Contact	Expert Committee
RAUWOLFIA SERPENTINA TABLETS	Nam-Cheol Kim Scientific Liaison	BDSHM2020 Botanical Dietary Supplements and Herbal Medicines
REFERENCE STANDARD SUPPORT	RS Technical Services RSTECH@usp.org	BDSHM2020 Botanical Dietary Supplements and Herbal Medicines

Chromatographic Database Information: [Chromatographic Database](#)

Most Recently Appeared In:
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