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N.B.: The Japanese Pharmacopoeia Drugs are to be tested according to the provisions given in the pertinent monographs, General Notices, General Rules for Crude Drugs, General Rules for Preparations, and General Tests for their conformity to the Japanese Pharmacopoeia. See the General Notices 5.1.4.

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The Japanese Pharmacopoeia, Seventeenth Edition (JP17)

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Japanese Pharmacopoeia(JP) - Present and Future

Pharmaceuticals and Medical Devices Agency (PMDA) 2 Office of Standards and Guidelines Development Division of Pharmacopoeia and Standards for Drugs

Japanese Pharmacopoeia(JP) - Present and Future

PMRJ produces and distributes the Japanese Pharmacopoeia Reference Standards prescribed in the Japanese Pharmacopoeia, "General Tests 9.01 Reference standards, (1) The reference standards which are prepared by those who have been registered to prepare them by the Minister of Health, Labour and Welfare, according to the Ministerial ordinance established by the Minister separately."

Japanese Pharmacopoeia Reference Standards - PMRJ

A pharmacopoeia, pharmacopeia, or pharmacopoea, in its modern sense, is a legally binding collection, prepared by a national or regional authority, of standards and quality specifications for medicines used in that country or region.

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1. Name of pharmacopoeia - WHO

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