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JP2007277194A - O/w type emulsified preparation containing ...

Japanese Pharmaceutical Codex 2020/09/06 REQUIREMENT SPECIFICATION * Description White crystals or crystalline powder Identification to pass test Optical rotation [α]D20 +24.8□+25.3° pH

6.7□7.2 Purity - (1) Clarity and color of solution to pass test (2) Chloride max.0.041% (3) Sulfate max.0.028% (4) Ammonium max.0.02% (5) Heavy metals max ...

SPECIFICATION - labchem-wako.fujifilm.com

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Pharmaceutical Administration and Regulations in Japan

pharmaceutical manufacturers, and the transfer of manufacturing or import approvals in 1983, and those related to promotion of R&D of orphan drugs and priority reviews for such drugs in 1993. In 2002, the Pharmaceutical Affairs Law (Law No. 96 dated July 31, 2002) was revised based on demands for

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Japanese Pharmaceutical Excipients 2004

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