

SPECIFICATION

Propranolol Hydrochloride

for the Japanese Pharmacopoeia General Tests (for Determination)

REQUIREMENT	SPECIFICATION
Description (JP method)	White crystalline powder
Identification (JP method)	to pass test
pH (JP method)	5.0~6.0
Melting point (JP method)	163~166°C
Purity (JP method)	-
(1) Clarity and color of solution (JP method)	to pass test
(2) Heavy metals (JP method)	max.20ppm
(3) Related substances (JP method)	to pass test
Loss on drying (JP method)	max.0.5%
Residue on ignition (JP method)	max.0.1%
Assay (after drying) (JP method)	min.99.5%