## **SPECIFICATION**

## Atropine Sulfate Hydrate

for the Japanese Pharmacopoeia Crude Drugs Test (for Assay and Thin-Layer Chroma

|     | REQUIREMENT                                 | SPECIFICATION                                  |
|-----|---|--|
|     | Description (JP method)                     | Colorless crystals or white crystalline powder |
|     | Identification (1) (JP method)              | to pass test                                   |
|     | Identification (2) (JP method)              | to pass test                                   |
|     | Identification (3) (JP method)              | to pass test                                   |
|     | Identification (4) (JP method)              | to pass test                                   |
|     | Purity (JP method)                          | -  |
| (1) | Clarity and color of solution (JP method)   | to pass test                                   |
| (2) | Acidity (JP method)                         | to pass test                                   |
| (3) | Related substances (1) (JP method)          | to pass test                                   |
| (4) | Related substances (2) (JP method)          | to pass test                                   |
| (5) | Hyoscyamine (JP method)                     | to pass test                                   |
| (6) | Readily carbonizable substances (JP method) | to pass test                                   |
|     | Loss on drying (JP method)                  | max.4.0%                                       |
|     | Residue on ignition (JP method)             | max.0.1%                                       |
|     | Assay (after drying) (JP method)            | min.99.0%                                      |
|     | Assay (HPLC)                                | min.99.0%                                      |

(1 / 1) established on 2011/07/19