

SPECIFICATION

Dried Sodium Carbonate

Japanese Pharmacopoeia

REQUIREMENT	SPECIFICATION
* Description (JP)	White crystals or crystalline powder
Identification (JP)	to pass test
Purity (JP)	-
(1) Clarity and color of solution (JP)	to pass test
(2) Chloride (JP)	not more than 0.071%
Loss on drying (JP)	not more than 2.0%
Assay (after drying) (JP)	not less than 99.0%
* Bacterial endotoxins	less than 10EU/g
Identification (USP-NF)	to pass test
Assay (dried basis) (USP-NF)	99.5~100.5%
Specific tests (USP-NF)	-
Water determination (USP-NF)	NMT 0.5%
* Appearance (Ph.Eur.)	White or almost white, slightly granular powder
Identification (Ph.Eur.)	to pass test
Appearance of solution (Ph.Eur.)	to pass test
Alkali hydroxides and bicarbonates (Ph.Eur.)	to pass test
Chlorides (Ph.Eur.)	maximum 125ppm
Sulfates (Ph.Eur.)	maximum 250ppm
Iron (Ph.Eur.)	maximum 50ppm
Loss on drying (Ph.Eur.)	maximum 1.0%
Assay (dried substance) (Ph.Eur.)	99.5~100.5%

Japanese Pharmacopoeia (Dried Sodium Carbonate)(* : Additional test performed by Wako) United States Pharmacopoeia - National Formulary (Sodium Carbonate) European Pharmacopoeia (Sodium Carbonate)(* : Additional test performed by Wako)