



Revision 4050-00

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SODIUM CROMOGLICATE

Batch N°:	4050/06/16	ANALYSIS CERTIFICATE N°	161042	Date:	27/04/2016
Kg:	208.00	Formula:	C ₂₃ H ₁₄ Na ₂ O ₁₁	M.W.:	512.3
Man. date:	11/03/2016	Retest date:	Mar/2021	Complies with:	Ph.Eur. BP USP JP
Solubility:	soluble in water; practically insoluble in ethanol (96 per cent) and in chloroform				

TESTS	SPECIFICATIONS	RESULTS
DESCRIPTION	white or almost white hygroscopic crystalline powder	complies
IDENTIFICATION	(1): IR spectrum (2): UV spectrum (3): sodium reaction	complies complies complies
ACIDITY OR ALKALINITY	(1): complies to Ph. Eur. test (2): complies to USP test	complies complies
APPEARANCE OF SOLUTION	a 2.0% solution in water is not more intensely coloured than <i>reference solution BY₅</i>	complies
CLARITY OF SOLUTION	a 2.0% solution in water is not more opalescent than <i>reference suspension II</i>	complies
COLOUR OF SOLUTION	Abs at 440 nm not more than 0.05 (5.0% solution)	0.02
WATER	not more than 10.0%	7.4%
LOSS ON DRYING	not more than 10.0%	7.0%
OXALATE	not more than 0.35%	complies
HEAVY METALS	not more than 20 ppm	complies
This material has been prepared following the current Good Manufacturing Practice (cGMP).		

Q.C. Manager

Q.A. Manager

Qualified Person

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Batch N°: 4050/06/16 ANALYSIS CERTIFICATE N° 161042 Date: 27/04/2016
 Kg: 208.00 Formula: $C_{23}H_{14}Na_2O_{11}$ M.W.: 512.3
 Man. date: 11/03/2016 Retest date: Mar/2021 Complies with: Ph.Eur. BP USP JP

Solubility: soluble in water; practically insoluble in ethanol (96 per cent) and in chloroform

TESTS	SPECIFICATIONS	RESULTS
RELATED SUBSTANCES (TLC)	each impurity not more than 0.5%	no impurities detected
RELATED SUBSTANCES (HPLC)	(1): impurity 1 not more than 0.10% (2): impurity 2 not more than 0.10% (3): impurity 3 not more than 0.10% (4): any other impurity not more than 0.10%	< 0.01% < 0.01% < 0.01% RRT 0.96: 0.02% RRT 1.14: 0.06% RRT 1.17: 0.03% RRT 1.20: 0.05% RRT 1.23: 0.01% RRT 1.28: 0.01% RRT 1.44: 0.01%
	(5): sum of all impurities not more than 0.30%	0.19%
ASSAY	99.0% to 101.0% with reference to the dried substance (pot.) 98.0% to 101.0% calculated on the anhydrous basis (UV)	99.5% 99.5%
RESIDUAL SOLVENTS	(1): methanol not more than 200 ppm (GC) (2): dimethylformamide not more than 100 ppm (HPLC)	< LOQ 11 ppm

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Q.C. Manager

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Qualified Person

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28/04/2016

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