



BUPROPION HYDROCHLORIDE USP

CAS N. 31677-93-7

CODE 1019

BATCH 0414S050

MAN. DATE 04/2014

RETEST DATE 01/2020

Description : White powder

TEST FOR	SPECIFICATIONS	RESULTS
Identification		
· IR	positive	Complies
· HPLC	positive	Complies
· Chlorides	positive	Complies
Water (KF)	≤ 0.5 %	0.04
Assay (HPLC, on anhydrous basis)	≥ 98.0 ≤ 102.0 %	100.3
Impurities (HPLC) - Limit of m-chlorobenzoic Acid		
· m-chlorobenzoic acid	≤ 0.2 %	Non quantifiable
Impurities (HPLC) - Organic Impurities		
· RRT~0.38 (Deschloro bupropion)	≤ 0.1 % (USP spec.: NMT 0.5%)	0.02
· RRT~0.58 (Bupropion dione derivative)	≤ 0.2 %	Non quantifiable
· RRT~0.71 (o-Bupropion)	≤ 0.1 %	Non quantifiable
· RRT~0.78 (Chloropropiophenone)	≤ 0.1 %	Non quantifiable
· RRT~0.92 (USP Rel. Comp. A)	≤ 0.1 % (USP spec.: NMT 0.2%)	Non quantifiable
· RRT~1.14 (USP Rel. Comp. B)	≤ 0.1 % (USP spec.: NMT 0.2%)	0.04
· RRT~1.63 (Bromochloropropionphenone)	≤ 0.1 %	Non quantifiable
· RRT~2.30 (4-Chlorobupropion)	≤ 0.1 % (USP spec.: NMT 0.2%)	0.01
· RRT~2.74 (5-Chlorobupropion)	≤ 0.1 % (USP spec.: NMT 0.2%)	Non quantifiable
· any other impurity	NMT 0.10% (USP spec.: NMT 0.1%)	Complies
Total impurities (Limit of m-chlorobenzoic Acid + Organic Impurities)	≤ 1.0 %	0.07
<u>Additional specifications</u>		
Residual solvents (GC)		
· sec-butyl alcohol	≤ 300 ppm	43
Particle size (sieving)		
· more than 250 µm	≤ 10 %	3.3
Bulk density	≥ 0.40 ≤ 0.60 g/ml	0.52
Tapped density (1250 taps)	≥ 0.60 ≤ 0.86 g/ml	0.73

OBSERVATIONS:

- Other residual solvents are unlikely to be present

Copies of this CoA are not valid

APPROVED

on Apr 24, 2014

QUALITY CONTROL

Dr. R. Poli

Apr 28, 2014