

**BUPROPION HYDROCHLORIDE USP**

CAS N. 31677-93-7

CODE 1019

BATCH B220036

MAN. DATE 06/2022

RETEST DATE 06/2028

Description : White powder

TEST FOR	SPECIFICATIONS			RESULTS
Identification				
· A) IR	positive			Complies
· B) HPLC	positive			Complies
· C) Chlorides	positive			Complies
Water (KF)		≤ 0.5	%	0.03
Assay (HPLC, on anhydrous basis)	≥ 98.0	≤ 102.0	%	100.5
Impurities (HPLC) - Limit of m-chlorobenzoic Acid				
· m-chlorobenzoic acid		≤ 0.10	%	<0.02 (LOQ)
	(USP spec.: NMT 0.2%)			
Impurities (HPLC) - Organic Impurities				
· RRT~0.38 (Deschloro bupropion)		≤ 0.1	%	<0.05 (LOQ)
	(USP spec.: NMT 0.5%)			
· RRT~0.58 (Bupropion dione derivative)		≤ 0.1	%	<0.05 (LOQ)
	(USP spec.: NMT 0.2%)			
· RRT~0.71 (o-Bupropion)		≤ 0.1	%	<0.05 (LOQ)
· RRT~0.78 (Chloropropiophenone)		≤ 0.1	%	<0.05 (LOQ)
· RRT~0.92 (USP Rel. Comp. A)		≤ 0.1	%	<0.05 (LOQ)
	(USP spec.: NMT 0.2%)			
· RRT~1.14 (USP Rel. Comp. B)		≤ 0.1	%	<0.05 (LOQ)
	(USP spec.: NMT 0.2%)			
· RRT~2.30 (4-Chlorobupropion)		≤ 0.1	%	<0.05 (LOQ)
	(USP spec.: NMT 0.2%)			
· RRT~2.74 (5-Chlorobupropion)		≤ 0.1	%	<0.05 (LOQ)
	(USP spec.: NMT 0.2%)			
· any other impurity (highest)		≤ 0.10	%	<0.05 (LOQ)
	(USP spec.: NMT 0.1%)			
Total impurities (Limit of m-chlorobenzoic Acid + Organic Impurities)		≤ 0.5	%	<0.05 (LOQ)
	(USP spec.: NMT 1.0%)			
Additional specifications				
2-Bromo-3'-chloropropiophenone (Bromochloropropiophenone)	NMT 3 ppm			Complies
Residual solvents (GC)				
· sec-butyl alcohol		≤ 300	ppm	<100 (LOQ)
· methanol		≤ 1000	ppm	<60 (LOQ)
Particle size (sieving)				
· more than 250 µm		≤ 10	%	1.2
Bulk density	≥ 0.40	≤ 0.60	g/ml	0.46
Tapped density (1250 taps)	≥ 0.60	≤ 0.86	g/ml	0.69

OBSERVATIONS:

- Other residual solvents are unlikely to be present
- The impurity 2-Bromo-3'-chloropropiophenone (Bromochloropropiophenone) is evaluated with the USP HPLC method slightly modified

Copies of this CoA are not valid

APPROVED

on Sep 26, 2022

QUALITY CONTROL

Dr. S. Patroni