

## Hydrocortisone acetate

<b>CODE ARTICLE:</b> 93917	<b>BATCH:</b> 0111786	<b>Active pharmaceutical ingredient</b>
<b>TEST DATE:</b> 09/06/2022	<b>TEST:</b> QC-00243597	
<b>EXPIRY DATE:</b> 30/04/2027	<b>FABRICATION DATE:</b> 08/05/2022	<b>MR:</b> 404,50
<b>CAS:</b> [50-03-3]	<b>FORMULA:</b> C23H32O6	
<b>SYNONYMS / BOTANICAL NAME:</b> Acetato de cortisol; 21-Acetato de hidrocortisona; Acetilhidrocortisona; Hidrocortisona acetilada		
<b>STORAGE:</b> Store in tightly closed containers in a cool, dry place.		

TEST	RESULTS	Units	MIN	MAX	METHOD
APPEARANCE	POWDER				PH.EUR.*
COLOUR	WHITE				PH.EUR.*
<b>SOLUBILITY</b>					
SOLUBLE	SLIGHTLY SOLUBLE IN ANHYDROUS ETHANOL				PH.EUR.*
SOLUBLE	SLIGHTLY SOLUBLE IN DICHLOROMETHANE				PH.EUR.*
INSOLUBLE	PRACTICALLY INSOLUBLE IN WATER				PH.EUR.*
<b>IDENTIFICATION</b>					
IR IDENTIFICATION	CONFORM				PH.EUR.*
IDENTIFICATION (HPLC)	CONFORM				PH.EUR.*
SPECIFIC ROTATION	160 DEGREES		158	167	PH.EUR.*
<b>RELATED SUBSTANCES</b>					
TOTAL IMPURITIES	0,6% (CONFORME) -			<= 1.5% (<15xArea RS b)	PH.EUR.*
UNSPECIFIED IMPURITIES	<DL (0,05%) (PASS) -			<= 0.10% (<area RS b)	PH.EUR.*
IMPURITY A	0,3%(PASS) -			<= 0.5% (<5xArea RS b)	PH.EUR.*
IMPURITY B	<DL (0,05%) (PASS) -			<= 0.3% (<3xArea RS b)	PH.EUR.*
IMPURITY C	0,1% (PASS) -			<= 0.6% (<6xArea RS b)	PH.EUR.*
IMPURITY D	<DL (0,05%) (PASS) -			<= 0.3% (<3xArea RS b)	PH.EUR.*
IMPURITY E	0,1% (PASS) -			<= 0.3% (<3xArea RS b)	PH.EUR.*
IMPURITY G	0,12% (PASS) -			<= 0.15% (<1.5xarea rs b)	PH.EUR.*
LOSS ON DRYING	0,1 %		-	0,5	PH.EUR.*
ASSAY (DRY SUBSTANCE)	101,4 %		97	102	PH.EUR.*
<b>RESIDUAL SOLVENTS</b>					
ACETONE	873 ppm		-	3.000	Manufacture's Standard
PYRIDINE	N.D. ppm			<=200	Manufacture's Standard
DMF	381 MG/KG		-	880	Manufacture's Standard
SAMPLING	COLLECT				Manufacture's Standard

The data expressed in this certificate of analysis is facilitated by our supplier and/or obtained in our control laboratory, in no case being exempted of the controls demanded by each sector. Tests marked with (\*) are verified in Guinama.

Manufacturer: 405601 H. L. PHARMACEUTICAL CO.,LTD. (China) Manufacturer Batch: K06B20220501 Manufacturer original CoA available under request

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TEST	RESULTS	Units	MIN	MAX	METHOD
CORRECT BATCH	YES				Manufacture's Standard
SPECIFICATION	PH.EUR 10				Manufacture's Standard
Risk Assessment Evaluation for Nitrosamines	CONFORM				
Elemental Impurities ICHQ3D	CONFORM				

*This product has been handled in a non-sterile room, for batches suitable for sterile use, consult availability.*



**Silvia Sancho-Tello Ripoll**  
Technical Director.

**This product has been manipulated in a NON-sterile clean room, for batches suitable for sterile use, consult availability.**

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