

CERTIFICATE of ANALYSIS

11- α -HYDROXIPROGESTERONE

BATCH No.: C0373/0 18020

MAN DATE: Oct/18

RE-TEST DATE: Oct/23

TEST	SPECIFICATIONS	RESULTS
1. Description	White to almost white crystalline powder	Complies
2. Identification		
2.1. IR	According to reference standard	Complies
2.2. HPLC	According to reference standard	Complies
3. Loss on drying	$\leq 1.0 \%$	0.1 %
4. Specific optical rotation	+175.0° to +182.0° (c=1; CHCl ₃ ; 25 °C)	+178.6°
5. Residue on ignition	$\leq 0.10\%$	< 0.10 %
6. Heavy Metals	≤ 10 ppm	< 10 ppm
7. Molybdenum	≤ 25 ppm	< 25 ppm
8. HPLC chromatographic purity	I0373-01 $\leq 0.15 \%$ I0373-02 $\leq 0.15 \%$ I0373-03 $\leq 0.15 \%$ I0373-04 $\leq 0.15 \%$ I0373-05 $\leq 0.15 \%$ Unknown $\leq 0.10 \%$ Total $\leq 1.0 \%$	< 0.05 % < 0.05 % < 0.05 % < 0.05 % < 0.05 % < 0.05 % 0.00 %
9. HPLC Assay (dried basis)	98.0 – 102.0 %	100.3 %
10. Residual solvents	Methanol < 3000 ppm Ethyl Acetate < 5000 ppm Tetrahydrofuran < 720 ppm Methyl-THF < 5000 ppm	< LOQ* 1488 ppm < LOQ* < LOQ*

*LOQ: Limit of quantitation



 Complex Science. Expert Solutions.
 Crystal Pharma, S.A.U.
 Parque Tecn. de Boecillo, Parcela 105
 47151 BOECILLO (Valladolid) - SPAIN
 Tel.: 34 923 54 60 72 | www.amriglobal.com
 22/11/18

Quality Assurance Department