GULF PHARMACEUTICAL INDUSTRIES

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Julphar

Quality Control Department

شردة النليج للصناعات الدوانية

رأس الخيمة - ص.ب. ٩٩٧ تليفون : ٢٤٦١٤٦١ -٩٧١٧ - فاكس : ٢٤٦٢٤٦٢ -٩٧١٧ الامارات العربية المتحدة

WRS M9-6 1554ed On 19/4/16

<u>Certificate of Analysis</u> <u>Secondary Reference Standard</u>

Name: Mometasone Furoate Material No: 10000490 Batch No: 0000028385/IV Insp. Lot No: 10000026857

Test	Specification	Result
Description	White or almost white powder, melts at about 220°C with decomposition, free from extraneous substances like black particles.	Conform
Identification	To pass tests under analytical method.	Conform
Loss on drying	NMT 0.50%	0.44%
Related Substances	Impurities A,B,C,D,E,F,G,H,I for each impurity: NMT 0.30%	0.06%
	Total impurities: NMT 0.60%	0.06%
Assay	NLT 97.0% and NMT 103.0% of C ₂₇ H ₃₀ Cl ₂ O ₆ , calculated on the dried basis.	99.04% (as is basis) 99.48% (on dried basis)

Standardization date: 10/03/2016

Standardized by: Sathya

Exp. date: 03/2017

Weight per vial: 500mg

Checked by: Pensee Mohamed, B. S. Pharm

Quality Control Deputy Manager

Head of Quality Control Department Sheikha Khamis

