GULF PHARMACEUTICAL INDUSTRIES
Ras Al Khaimah, Post Box No. 997
Tel.: +9717-2461461, Fax: +9717-2462462
United Arab Emirates

Julphar
Quality Control Department

شركة الخليج للصناعات الدوائية
رأس الخيمة - ص.ب. ٩٩٧
تليفون : ٢٤٦١٤٦١-٩٧١٧+ فاكس : ٢٤٦٢٤٦٢-٩٧١٧+
الامارات العربية المتحدة

WRS
P49-1

Certificate of Analysis
Secondary Reference Standard

Name: Paracetamol (Acetaminophen)
Material No: 10000533

Batch No: 0000030446/III
Insp. Lot No: 10000029594

Test	Specification	Result
Description	White, odourless, crystalline powder, having a slightly bitter taste, free from extraneous substances like black particles.	Conform
Identification	To pass tests under analytical method.	Conform
Water Content	NMT 0.50%	0.31%
Free p-aminophenol	NMT 0.005%	0.0004%
Limit of p-chloroacetanilide	NMT 0.001%	0.00%
Assay	NLT 98.0% and NMT 101.0% of $C_8H_9NO_2$, calculated on the anhydrous basis.	99.38% (as is basis) 99.69% (on anhydrous basis)

Standardization date: 18/10/2015

Standardized by: Shaima

Exp. date: 10/2016

Weight per vial: 500mg

Standardized against: Paracetamol (Acetaminophen) USP Reference Standard (Lot No: K0I244)

Checked by: Pensee Mohamed, B. S. Pharm
Quality Control Deputy Manager

QUALITY CONTROL Julphar DEPARTMENT

Head of Quality Control Department
Sheikha Khamis

NATIONAL QUALITY CONTROL LABORATORY
RECEIVED
07 JAN 2016
TEL: 2718963 / 3544528/30
FAX: 2718073
P. O. Box 29126 - 00202, NAIROBI