

P
1-10P₁₋₁₀
INTAS PHARMACEUTICALS LTD.P₁₋₁₀
INTAS

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**QUALITY CONTROL DEPARTMENT
CERTIFICATE OF ANALYSIS**

Name of working standard	: PARACETAMOL		
Evaluated with EP Batch No.	: 4	Date of preparation	: 10/03/2016
Working standard No.	: S PARA 161	Valid Up to	: 10/03/2017
Raw material Batch No.	: 1447/15-16	Page	: 1 of 1
Raw material A. R. No.	: 101603994		
Storage condition	: Keep container tightly closed and store below 25°C in amber colour vial.		
Reference : Ph.Eur & Inhouse specification.			
SR. NO.	TEST	RESULT	LIMITS
01.	Characters	White, crystalline powder.	White or almost white, crystalline powder.
02.	Identification	By IR: The IR spectrum of the substance being examined is concordant with the reference spectrum of Paracetamol EP reference standard.	By IR: The IR spectrum of the substance being examined should be concordant with the reference spectrum of Paracetamol EP reference standard.
03.	Loss on drying	0.3%.	Not more than 0.5%.
04.	Related substances A) Impurity J B) Impurity K C) Impurity F D) Any other impurity E) Total of other impurities	A) Not detected. B) 4 ppm. C) Not detected. D) 0.01%. E) 0.01%.	A) Not more than 10 ppm. B) Not more than 50 ppm. C) Not more than 0.05%. D) Not more than 0.05%. E) Not more than 0.1%.
05.	Assay	99.6%. 99.3% as such.	99.0% to 101.0% calculated on dried basis. Record.

Remarks : The above material is compared with EP standard for above tests and is suitable to use as a working standard

PREPARED BY :
DATE :

9/10/03/16

CHECKED BY :
DATE :

10/03/16

APPROVED BY :
DATE :

10/03/16

