



PHARMA DEVILS

QUALITY CONTROL DEPARTMENT

ANALYTICAL SPECIFICATION FOR POVIDONE IP

Manufacturer Name:		Supplier Name:
Manufacturer's Batch No.:		Manufacturer's Batch Size:
Manufacturing Date:		Expiry Date:
Quantity Received:		Test Quantity:
Document No.:	Effective Date:	Change Control No.: N/A
Control No./ A.R. No.:		Retest Date:

Reference : IP 1996 Page No 610-611

Description : White or yellowish white powder or flakes; odourless or almost odourless; hygroscopic free from foreign particles.
Complies/Does Not Comply

Solubility : Freely soluble in distilled water, in *chloroform*, and in *ethanol (95%)* (1 in 1-10); practically insoluble in *ether* (1 in more than 10,000).
Complies/Does Not Comply

SNo.	Test	Reference	Result	Specification	Remark
1.	Identification (B,C,D or A,D) A. IR B. Precipitation reaction C. Colour reaction D. Colour reaction	IP		Note : Routinely Identification test B,C,D are carried out A. Infra Red absorption spectrum is concordant with that of reference standard of povidone. B. Orange yellow precipitate is formed. C. A pink colour is produced. D. A red colour is produced.	



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S.No.	Test	Reference	Result	Specification	Remark
2.	Clarity and colour of solution	IP		Solution A is clear and not more intensely coloured than reference solution BS6 or BYS6.	
3.	Heavy Metals	IP		Not More Than 10 ppm	
4.	Aldehydes as C ₂ H ₄ O	IP		Not More than 0.2 %	
5.	Vinylpyrrolidone	IP		Not More Than 0.2 %	
6.	Sulphated ash	IP		Not More Than 0.1%	
7.	Loss on drying	IP		Not More Than 5.0%	
8.	K-value	IP		Not less than 27.0 and not more than 32.1 %.	
9.	Nitrogen	IP		Not less than 11.5 and not more than 12.8 %.	

Raw Data Reference:

Analyst Name:

Analyst Name:

Analyst Name:

Analyst Hard Book No. :

Page No. :

Analyst Hard Book No. :

Page No. :

Analyst Hard Book No. :

Page No. :

Results : Sample Conforms / Does Not Conform to Specification

Analyzed By : _____

Date : _____

Checked By : _____

Date : _____

Approved By : _____

Date : _____