



MYLAN LABORATORIES LIMITED
F-4 & F-12, Malegaon MIDC, Sinnar,
Nashik - 422 113, Maharashtra, India.

CERTIFICATE OF ANALYSIS

Name of the Product: MyHEP ALL (Sofosbuvir and Velpatasvir Film-coated Tablets 400 mg / 100 mg)

Batch No.	3112907	A.R. No.	MLNFP20002195
Mfg. Date	Mar-2020	Market	Emerging
Exp. Date	Feb-2022	Batch Size	2,50,000 Tablets
Ref. Spec No.	FPSSFV195R-06	Date of Analysis	Mar 20, 2020

S. No.	TEST	SPECIFICATION	RESULT
1	Description	Light green to green colored, modified capsule shaped biconvex beveled edge film coated tablet debossed with M on one side and SFV on the other side.	Light green colored, modified capsule shaped biconvex beveled edge film coated tablet debossed with M on one side and SFV on the other side.
2	Identification		
2.1	A. By HPLC	The retention time of the Sofosbuvir and Velpatasvir peaks in the chromatogram of the test preparation should corresponds to that in the chromatogram of the standard preparation as obtained in the test "Assay (By HPLC)".	The retention time of the Sofosbuvir and Velpatasvir peaks in the chromatogram of the test preparation corresponds to that in the chromatogram of the standard preparation as obtained in the test "Assay (By HPLC)".
2.2	B. By HPLC (With PDA detector)	Compare the spectrums of the Sofosbuvir and Velpatasvir peaks from the test solution chromatogram with that of the standard solution chromatogram from the PDA data. The test spectrum should match with that of standard spectrum due to Sofosbuvir and Velpatasvir peaks.	Compared the spectrums of the Sofosbuvir and Velpatasvir peaks from the test solution chromatogram with that of the standard solution chromatogram from the PDA data. The test spectrum matches with that of standard spectrum due to Sofosbuvir and Velpatasvir peaks
3	Dissolution (By HPLC)		
3.1	Sofosbuvir 400 mg	Complies with Ph. Eur. General Chapter 2.9.3 Not less than 80% (Q = 75%) of the labeled amount of Sofosbuvir, C ₂₂ H ₂₉ FN ₃ O ₉ P should be dissolved in 30 minutes.	(Tab-1) : 96 % (Tab-2) : 97 % (Tab-3) : 97 % (Tab-4) : 97 % (Tab-5) : 96 % (Tab-6) : 97 % Min : 96 % Max : 97 % Avg : 97 % RSD:0.5%
3.2	Velpatasvir 100 mg	Complies with Ph. Eur. General Chapter 2.9.3 Not less than 80% (Q = 75%) of the labeled amount of Velpatasvir, C ₄₉ H ₅₄ N ₈ O ₈ should be dissolved in 30 minutes.	(Tab-1) : 97 % (Tab-2) : 98 % (Tab-3) : 98 % (Tab-4) : 97 % (Tab-5) : 96 % (Tab-6) : 97 %

Remarks: APPROVED (Sample Conforms to above Specification)

Prepared By : Sandip.Lachake	Approved By : Rajendra.Adavkar
Prepared On : Mar 21 2020 11:41AM	Approved On : Mar 21 2020 11:51AM
Printed by: Sandip.Lachake	Printed on: Mar 21 2020 11:55AM
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CNo: C9000002663

Plot No. 564/A/22, Road No. 92, Jubilee Hills, Hyderabad - 500 034, Telangana, India

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			Min : 96 % Max : 98 % Avg : 97 % RSD:0.6%
4	Uniformity of dosage units (By Content Uniformity)		
4.1	Sofosbuvir 400 mg	Complies with Ph. Eur. General Chapter 2.9.40 The Acceptance Value (AV) should be not more than 15.0.	Sample-1 : 99.7 % Sample-2 : 99.6 % Sample-3 : 100.2 % Sample-4 : 99.9 % Sample-5 : 100.4 % Sample-6 : 100.1 % Sample-7 : 98.9 % Sample-8 : 99.9 % Sample-9 : 101.3 % Sample-10 : 100.0 % Average : 100.0 % AV:1.5
4.2	Velpatasvir 100 mg	Complies with Ph. Eur. General Chapter 2.9.40 The Acceptance Value (AV) should be not more than 15.0.	Sample-1 : 100.5 % Sample-2 : 100.2 % Sample-3 : 100.7 % Sample-4 : 100.6 % Sample-5 : 100.7 % Sample-6 : 101.0 % Sample-7 : 99.7 % Sample-8 : 100.9 % Sample-9 : 101.9 % Sample-10 : 101.0 % Average : 100.7 % AV:1.3
5	Assay (By HPLC)		
5.1	Sofosbuvir 400 mg	Not less than 360.00 mg and not more than 440.00 mg of Sofosbuvir, $C_{22}H_{29}FN$ 3O_9P (90.0% w/w – 110.0% w/w of labeled amount of Sofosbuvir)	In mg : 400.55 mg In %:100.1% w/w
5.2	Velpatasvir 100 mg	Not less than 90.00 mg and not more	

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S. No.	TEST	SPECIFICATION	RESULT
		than 110.00 mg of Velpatasvir, C $^{49}_{15}\text{H}$ $^{54}_{14}\text{N}$ $^{16}_8\text{O}$ (90.0% w/w – 110.0% w/w of labeled amount of Velpatasvir)	In mg : 100.52 mg In %:100.5% w/w
6	Related Substances (By HPLC)		
6.1	A.Velpatasvir impurities:		
6.1.1	a.Amine free base	Not more than 1.0% w/w	Not Detected
6.1.2	b.Lactone impurity	Not more than 1.0% w/w	Below LOQ
6.1.3	c. Any unknown impurity	Not more than 0.5% w/w	0.049 % w/w
6.2	B.Sofosbuvir impurities:		
6.2.1	a.Fluoro uridine impurity	Not more than 0.5% w/w	Not Detected
6.2.2	b. Any unknown impurity	Not more than 0.5% w/w	0.010 % w/w
6.3	C.Total impurities (A + B)	Not more than 3.5% w/w	0.297 % w/w
7	Residual solvents (By GC)		
7.1	Ethanol	Not more than 5000 ppm	369 ppm
8	Water (By KF)	Not more than 5.5% w/w	2.79 % w/w

Storage: Do not store above 30 °C, store in original container.

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