

HETERO LABS LIMITED (UNIT-II)

(Formulations Division)

CERTIFICATE OF ANALYSIS

Product Name: LEDI	FOS (Ledipasvir 90mg and	Sofosbuvir 400mg Table	ets)
Product Code	4013077	A.R. No.	H5FP19002056
Specification ID	FPS/B-3007107-1-02	Batch No.	31172446
Mfg. Date	05/2019	Batch Size	1.0 Lac.
Exp. Date	04/2021	Date Of Release	27-05-2019

S. No.	TEST	RESULT	SPECIFICATION
1	Description	Brown coloured, capsule	Brown coloured, capsule
		shaped, bevel biconvex film	shaped, bevel biconvex film
		coated tablets debossed	coated tablets debossed
		with 'H' on one side and	with 'H' on one side and
		'L18' on other side.	'L18' on other side.
2	Identification (By HPLC)	The retention time of the	The retention time of the
		major peak in the	major peak in the
	•	chromatogram of the sample	chromatogram of the sample
		solution corresponds to that	solution should correspond
		in the chromatogram of the	to that in the chromatogram
		standard solution, as	of the standard solution, as
		obtained in the assay.	obtained in the assay.
3	Average weight	1029.57 mg	1025.00mg ± 3.0%
			(994.25mg to 1055.75mg)
4	Uniformity of weight		± 5% of Average weight
		Highest : 1.16 %	
		Lowest::-0.93 %	
5	Water content (By KF)	3.40 %w/w	Not more than 5.0%w/w
6	Uniformity of Content (By		Not less than 85.0% and Not
	HPLC) Content of	Min.: 98.3 %	more than 115.0% of
	Ledipasvir	Max. : 99.4 %	average content.
	·	Average::99.0 %	_
7	Dissolution (By HPLC)		
7.1	Ledipasvir		Not less than 75 % (D) of
		Tablet 1-: 97.7 %	labeled amount of Ledipasvir
		Tablet 2- : 95.6 %	should dissolve in 30
		Tablet 3- : 93.7 %	minutes.
		Tablet 4- : 93.9 %	
		Tablet 5- : 97.1 %	
		Tablet 6- : 97.1 %	
		Average::95.8 %	
7.2	Sofosbuvir		Not less than 75 % (D) of
		Tablet 1-: 86.9 %	labeled amount of
		Tablet 2- : 85.0 %	Sofosbuvir should dissolve

Remarks: APP	ROVED (Sample Conforms to a	bove Specification)		
Checked By	MANU.RAJ	Approved By	D.S.N.Reddy	
Date	27-05-2019 17:26	Date	27-05-2019 17:31	
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CERTIFICATE OF ANALYSIS

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		Tablet 3- : 96.7 %	in 30 minutes.
		Tablet 4- : 97.0 %	
		Tablet 5-: 84.7 %	
		Tablet 6- : 84.7 %	
		Average::89.1 %	
8	Related Substances (By HPLC)		
8.1	Sofosbuvir Related compound -01	0.00 %	Not more than 0.50%
8.2	Ledipasvir Related compound -04	0.08 %	Not more than 1.0%
8.3	Max. single Unknown Impurity	0.18 %	Not more than 0.50%
8.4	Total Impurities	0.44 %	Not more than 2.0%
9	Assay (By HPLC) Each film coated tablet contains		
9.1	Ledipasvir (C49H54F2N8O6), in mg	91.33 mg	Not less than 85.5mg and Not more than 94.5mg
9.2	(%) Labeled amount	101.5 %	Not less than 95.0 and Not more than 105.0
9.3	Sofosbuvir (C22H29 FN3O9P) in mg	397.34 mg	Not less than 380.0mg and Not more than 420.0mg
9.4	(%) Labeled amount	99.3 %	Not less than 95.0 and Not more than 105.0

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