LabCorp				P	atientReport
Specimen ID: Control ID:		Acct #	#:	Phone:	Rte: 00
Patient Details DOB: Age(y/m/d): Gender: SSN: Patient ID:	Specimen Detail Date collected: Date received: Date entered: Date reported:	s	Physic Orderir Referri ID: NPI:		0
General Comments & Additional Information Alternate Control Number: Total Volume: Ordered Items	n		ternate Patient ID: sting:		
HCV FibroSure; Venipuncture					
TESTS HCV FibroSure	RESULT	FLAG	UNITS RI	EFERENCE IN	NTERVAL LAB
HCV FibroSURE Results:					
Fibrosis Score	0.62	High		0.00 -	0.21
Fibrosis Stage					
F3-Brid Necroinflammat Activity Sco	lging fibros:	is with m	any septa		
Necromitanual Activity Sec	0.77	High		0.00 -	0.17
Necroinflammat Activity Gra				0.00	0.17
A3-Sever	e activity				
•					
Analysis:	306	II i sh	(17	110	076
Alpha 2-Macroglobulins, Qn Haptoglobin	308 117	High	mg/dL mg/dL	110 - 34 -	
Apolipoprotein A-1	110		mg/dL	101 -	
Bilirubin, Total	0.5		mg/dL	0.0 -	
GGT	38		IU/L	0 -	
ALT (SGPT) P5P	128	High	IU/L	0 -	
<pre>. Interpretations: Quantitative results of a computational algorit marker (0.0-1.0) for 15 necroinflammatory activ Fibrosis Scoring: <0.21 = Stage F0 0.21 - 0.27 = Stage F0 0.27 - 0.31 = Stage F1 0.31 - 0.48 = Stage F1 0.48 - 0.58 = Stage F1 0.48 - 0.58 = Stage F2 0.58 - 0.72 = Stage F3 0.72 - 0.74 = Stage F3 >0.74 = Stage F4</pre>	<pre>chm to provid ver fibrosis vity (METAVI) - No fibros: - F1 - Portal fib - F2 - Bridging : - Bridging : - F4 - Cirrhosis</pre>	de a quan s (METAVI R A0-A3). is prosis fibrosis	titative su: R FO-F4) and with few se	rrogate d for pta	

Date Issued:

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S	a	b	F	Dr	D
		1022-02-			

Patient: Specimen ID: Patient ID: **Control ID:** Date collected: TESTS RESULT FLAG UNITS REFERENCE INTERVAL LAB <0.17 = Grade A0 - No Activity 0.17 - 0.29 = Grade A0 - A10.29 - 0.36 = Grade A1 - Minimal activity 0.36 - 0.52 = Grade A1 - A20.52 - 0.60 =Grade A2 - Moderate activity 0.60 - 0.62 = Grade A2 - A3>0.62 = Grade A3 - Severe activity

Limitations:

DOB:

The negative predictive value of a Fibrotest score <0.31 (absence of clinically significant fibrosis) was 85% when compared to liver biopsy in 1,270 HCV infected patients with a 38% prevalence of significant liver fibrosis (F2, 3 or 4). The positive predictive value of a Fibrotest score >0.48 (F2, 3, 4) was 61% in that same patient cohort. HCV FibroSURE is not recommended in patients with Gilbert Disease, acute hemolysis (e.g. HCV ribavirin therapy mediated hemolysis) acute hepatitis of the liver, extra-hepatic cholestasis, transplant patients, and/or renal insufficiency patients. Any of these clinical situations may lead to inaccurate quantitative predictions of fibrosis and necroinflammatory activity in the liver.

Comment:

This test was developed and its performance characteristics determined by LabCorp. It has not been cleared or approved by the Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. For questions regarding this report please contact customer service at 1-800-788-9223.

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PatientReport