



HBV Seroconversion Panel Donor No. 71612
Part Number: HBV11058

Donor Demographics: Donor source plasma was collected from a U.S. licensed blood collection facility.
 Donor was un-treated and asymptomatic throughout the collection period.

PANEL MEMBER	DATE OF DRAW	ALT Normal Range (10-45u/L)	ABBOTT HBV REAL TIME CP/ML	ROCHE HBV Monitor Assay (copies/mL)	ROCHE HBV Ampliscreen	GENETIC SYSTEMS HBsAg 2.0 Static	ABBOTT MUREX HBsAg Version 3	ABBOTT PRISM HBsAg
		TEST DATE: 10/8/1999	TEST DATE: UNK	TEST DATE: UNK	TEST DATE: UNK	TEST DATE: UNK	TEST DATE: UNK	TEST DATE: UNK
11058-01	04/11/99	9	NOT DETECTED	NOT TESTED	NEGATIVE	NEGATIVE	0.46	0.72
11058-02	04/14/99	6	NOT DETECTED	NOT TESTED	NEGATIVE	NEGATIVE	0.46	0.33
11058-03	04/18/99	8	NOT DETECTED	NOT TESTED	NEGATIVE	NEGATIVE	0.45	0.35
11058-04	05/01/99	6	<52	NOT TESTED	NEGATIVE	NEGATIVE	0.48	0.49
11058-05	05/19/99	12	1489	6800	POSITIVE	NEGATIVE	2.05	4.02
11058-06	05/22/99	7	4293	16000	POSITIVE	NEGATIVE	4.79	8.81
11058-07	05/29/99	7	30392	120000	POSITIVE	POSITIVE	27.39	59.82
11058-08	06/05/99	9	264700	1100000	POSITIVE	POSITIVE	30.80	211.73

PANEL MEMBER	ORTHO HBsAg Proc B	ABBOTT HBsAg Proc B	ABBOTT HBsAg Proc C	ORTHO VITROS ECi HBsAg	ORTHO VITROS ECi ANTI-HBc TOTAL	ORTHO VITROS ECi HBeAg	ORTHO VITROS ECi ANTI-HBe	ORTHO VITROS ECi ANTI-HBc IgM	ORTHO VITROS ECi ANTI-HBs QUANT
	TEST DATE: UNK	TEST DATE: UNK	TEST DATE: UNK	TEST DATE: 8/30/2018	TEST DATE: 8/30/2018	TEST DATE: 8/30/2018	TEST DATE: 8/30/2018	TEST DATE: 8/30/2018	TEST DATE: 8/30/2018
11058-01	0.097	0.019	0.037	0.07	3.34	0.14	0.18	0.01	0.0
11058-02	0.097	0.019	0.296	0.06	3.39	0.14	NOT TESTED	NOT TESTED	0.0
11058-03	0.097	0.057	0.333	0.10	3.41	0.15	0.15	0.01	0.0
11058-04	0.032	0.075	0.148	0.10	3.19	0.15	0.18	0.01	0.0
11058-05	0.387	1.132	0.741	0.89	3.24	0.14	0.17	0.01	0.0
11058-06	0.938	2.642	1.296	3.21	3.19	0.15	0.18	0.01	0.0
11058-07	11.563	21.132	11.593	24.6	3.17	0.24	0.14	0.01	0.0
11058-08	68.469	>37.736	65.926	218	3.15	2.32	0.15	0.01	0.0

Notes on Data:

1. This panel is for **Research Use Only. Not for use in diagnostic procedures.**
2. Data is generated at ZeptoMetrix™ LLC and by our Laboratory Partners in the United States and Europe.
3. All specimens collected in this longitudinal series are unadulterated 4% sodium citrated plasma samples collected from a single donor in the United States.
4. No preservatives have been added.
5. All material has been tested and found negative for Anti-HIV1/2, HIV p24 Ag, Anti HCV by FDA cleared tests.

ZeptoMetrix LLC • 25 Kenwood Circle, Franklin, MA 02038 • Tel (508) 553-5800 • Fax (508) 520-1525
 This Product was manufactured in a facility which has a Quality Management System that is ISO 13485 certified.
 For Customer Support, please visit www.zeptometrix.com or email zepto.customerservice@antylia.com

PIHBV11058 Rev04
 Effective Date: 08/13/2021