

# AccuSpan™ HCV RNA Linearity Panel

2410-0166 / Batch #10612186

## OVERVIEW

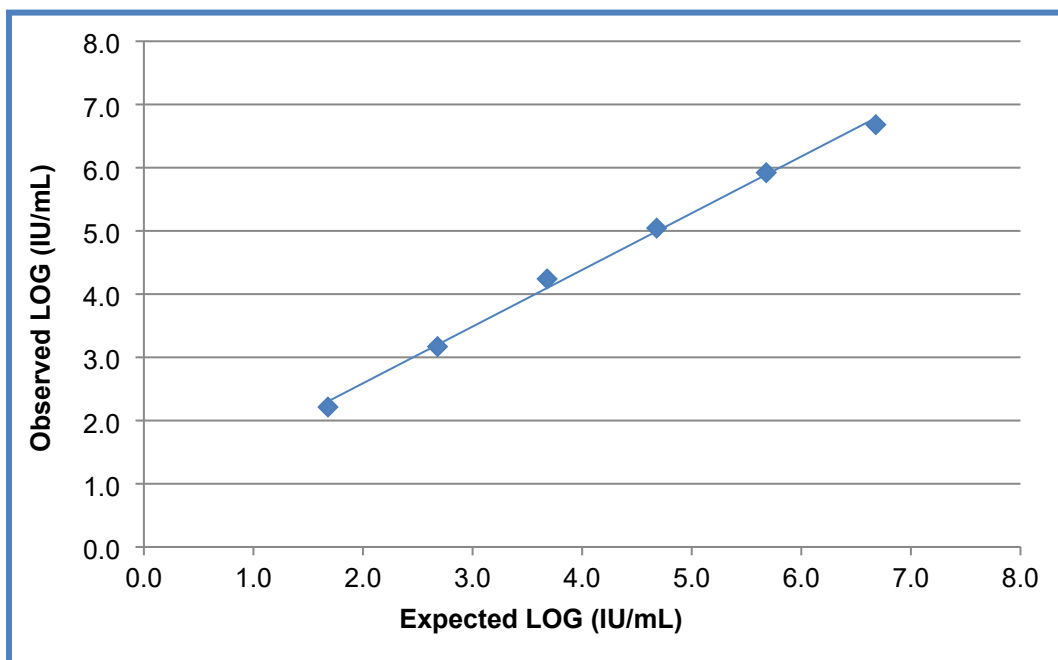
The AccuSpan™ HCV RNA Linearity Panel (2410-0166 / Batch #10612186) is an eight-member panel made from serial dilutions of high titer HCV positive plasma with established reactivity for HCV (Hepatitis C) RNA. This panel consists of seven members representing serial log dilutions of HCV positive plasma in HCV RNA negative diluent and one negative member prepared from the diluent. The diluent was prepared from normal human plasma that was filtered through a 0.2 micron filter. Sodium azide (0.09%) was added as a preservative.

Results are reported for each panel member on specific test methods. Linearity is shown graphically by plotting observed results against expected results. The WHO International Standard was tested in the same run as the AccuSpan HCV RNA Linearity Panel members. Both expected and observed results for the standards are reported; expected results for the WHO standards are the WHO assigned values.

For Research Use Only. Not for use in diagnostic procedures. Data are offered for informational purposes. LGC / SeraCare Life Sciences does not claim that others can duplicate test results exactly.

CAUTION: These materials have not been treated and should be considered biohazardous. Follow Universal Precautions.

AccuSpan™ HCV RNA Linearity Panel



HCV RNA results were obtained using the Roche COBAS® AmpliPrep/COBAS® TaqMan® HCV Test, v2.0 test method. Results are the mean of three replicates. A line of best fit is shown.

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## HCV RNA

Panel Member	Roche COBAS® AmpliPrep / COBAS® Taqman® HCV Test, v2.0 (log IU/mL) <sup>1,2</sup>	Abbott m2000 Realtime HCV (log IU/mL) <sup>1,2</sup>
01	<b>6.68</b> <sup>3</sup>	<b>6.69</b> <sup>3</sup>
02	<b>5.92</b>	<b>6.04</b>
03	<b>5.04</b>	<b>4.88</b>
04	<b>4.24</b>	<b>3.95</b>
05	<b>3.17</b>	<b>2.91</b>
06	<b>2.21</b>	<b>2.03</b>
07	<b>1.69</b> , TND, <1.18*	<b>1.26</b> , <1.08, <1.08*
08	TND	TND
Test Date	13-Apr-2022	04-Apr-2022
Test Site	RL	RL
Test Kit Range	15 to 100,000,000 IU/mL 1.18 to 8.00 log IU/mL	12 to 100,000,000 IU/mL 1.08 to 8.00 log IU/mL
Kit Part Code	NA	NA
Kit Lot No.	H22428	526576
Kit Exp. Date	31-Mar-2023	30-Apr-2023

<sup>1</sup>Results are reported as log International Units per mL (log IU/mL); positive/reactive results are noted in bold red.

<sup>2</sup>Results are reported as the mean result of triplicate testing.

<sup>3</sup>Panel member #1 was tested at a 1:10 dilution and results were corrected for the dilution factor.

\*Interpretation variability based on triplicate testing, individual replicates reported.

TND = Target Not Detected; RL = Reference Lab; NA = Not Available

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## 5th WHO International HCV RNA Standard (14/150)

Sample ID	Expected Values (log IU/mL)	Observed Values on Roche COBAS® AmpliPrep/COBAS® TaqMan® HCV Test, v2.0	
		(log IU/mL) <sup>1</sup>	% Difference <sup>2</sup>
01	4.60	<b>4.37</b>	-5.1
02	4.00	<b>3.77</b>	-5.8
03	3.70	<b>3.70</b>	0.0
04	3.00	<b>3.00</b>	-0.2
Test Date		13-Apr-2022	
Test Site		RL	
Test Kit Range		15 to 100,000,000 IU/mL 1.18 to 8.00 log IU/mL	
Kit Part Code		NA	
Kit Lot No.		H22428	
Kit Exp. Date		31-Mar-2023	

<sup>1</sup>WHO standards were tested in the same test run as the AccuSpan™ HCV RNA Linearity Panel members. Samples were run in singlet. Positive/reactive results are noted in bold red.

<sup>2</sup>Percentage difference is how much the observed concentration differs from the expected concentration. Values calculated for reference only. Laboratories may use the data to apply a correction factor to the test results.  
RL = Reference Lab; NA = Not Available

The package insert for this panel can be found at [www.seracare.com](http://www.seracare.com).

A printed copy of the package insert or data sheet may be requested by email at [CDx-Info@LGCGroup.com](mailto:CDx-Info@LGCGroup.com) or by phone at 508.244.6400.