

Pioneering Viral Vector Tools & Translational Medical Research

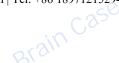
Test Report for BC-RV-CVS EnvA461

1. Test Report Information

				
		Product Informa	tion	
Product Name:	CVS-EnvA-△G-EGFP		Product Number:	BC-RV-CVS EnvA461
Batch Number:	BC-RV240402	Blan.	Titer:	RV:2.00E+08 IFU/mL
	c &	Helper Virus Inf	ormation	8/2
Virus 1	rAAV-EF1 α -DIO-mCherry-F2A-TVA		Product Number	BC-0061
Batch Number:>	BC230825-0061-9 (diluted)		Titer;	AAV:7.9E+11 vg/mL
Virus 2	rAAV-EF1 α -DIO-N2cG		Product Number :	BC-0442
Batch Number:	BC240202-0442-9 (diluted)		Titer:	AAV: 2. 02E+12 vg/mL
Virus 3	rAAV-hSyn-SV40 NLS-Cre		Product Number :	BC-0159
Batch Number:	BC231110-0159-9 (diluted)		Titer:	AAV:3.03E+11 vg/mL
		Test Methodol	ogy	;n C
Test Site:	Primary Motor Cortex (M1), AP=+1.54, ML=+1.60, DV=-1.60		Animal Species:	C57 BL/6 mouse
Test Methods:	Day0	TVA:G:Cre were mixed at a ratio of 1:2:1 and injected at 100 nl per animal.		
	Day14 BC-RV-CVS EnvA462 was i per animal.		462 was inj	ected into M1 at 150 m
	Day21 Mice sacrificed, tissue samples collected.			
Injection	2024. 05. 17		Perfusion Date:	2024. 06. 07

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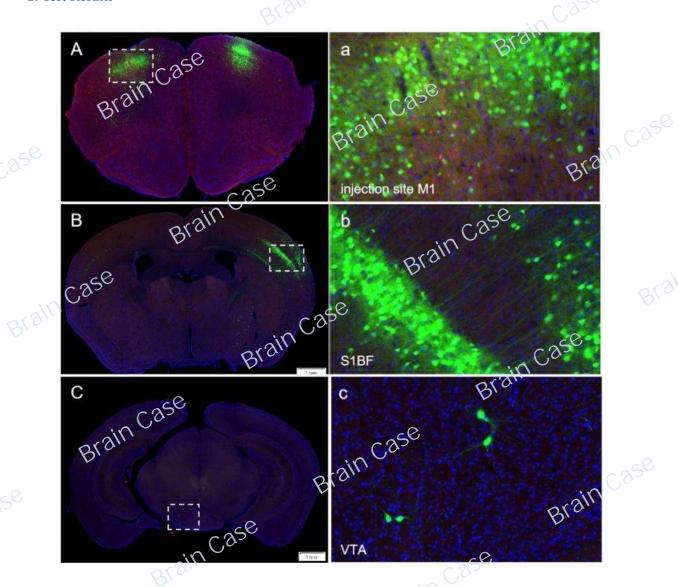
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2. Test Results



The RV helper viruses (TVA:G:Cre = 1:2:1) were injected into the primary motor cortex (M1). After a 14-day expression period, RV was injected at the same site. The arrival of the same site. standard housing conditions for an additional 7 days before being perfused for tissue collection. Brain tissues were subsequently fixed, sectioned, and imaged.

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Strong green fluorescence signals were observed in M1, S1BF, VTA, and other brain regions, confirming the effectiveness of the RV-based monosynaptic retrograde tracing strategy.

4. Signatures

Injection Performed by:

孝志林

Imaging/Scanning Performed by:

Reviewed by:

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