



## **Centre for AIDS Reagents**

## **Data Sheet**

<b>REPOSITORY REFERENCE:</b>	ARP2074
NAME:	HIV-1 clone TV1.29
PROVIDED:	10 $\mu$ L purified plasmid DNA, at 1 $\mu$ g/ $\mu$ L.
CLONING SITE:	The HIV-1 env/rev cassette was cloned directly into the cloning site of pcDNA3.1D/V5-His TOPO expression vector, in the correct orientation with the CMV promoter. The size of the insert is 2559 bp.
CLONING VECTOR:	pcDNA3.1D/V5-His TOPO
<b>DESCRIPTION:</b>	A PCR fragment containing full-length env and rev genes was derived from the genomic DNA of infected PBMC. The original virus (HIV-1 Tv1) was obtained from Dr. David Montefiori. The env/rev cassette was cloned into pcDNA3.1D/V5-His TOPO expression vector by a directional cloning approach. A single transformed ampicillin resistant E.coli colong was selected and expanded. Recombinant plasmid carries resistance genes for ampicillin and neomycin.
GENBANK:	U08455
RECOMMENDED STORAGE:	-20°C.
NOTE:	This clone expresses a functional env/rev cassette and can be used to generate psuedotyped infectious virions.
SOURCE:	Dr. John Mascola
<b>REFERENCES:</b>	Nature Medicine, volume 13, no. 9, Sept 2007.









Publications should acknowledge the donor of the reagent and the Programme EVA Centre for AIDS Reagents. Suggested wording can be found on our website in the "Acknowledgements" section at:

www.nibsc.ac.uk/spotlight/centre for aids reagents.aspx

Please also ensure that you send us a copy of any papers resulting from work using reagents acquired through CFAR (this can be electronically or as a paper copy)

