

Supporting information

Safety, immunogenicity and effectiveness of defective viral particles arising in mast cells against influenza in mice

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Table S1. The primer sequences

	Target name	Primers
Influenza A virus	HA	5'- CGCAGTATTCAGAAGAAGCAAGAC -3'
		5'- TCCATAAGGATAGACCAGCTACCA -3'
Mouse	IFN- β	5'- AGATGTCCTCAACTGCTCTC -3'
		5'- AGATTCACTACCAGTCCCAG -3'
	IFN- γ	5'- GACTGTGATTGCGGGGTTGT -3'
		5'- GGCCCGGAGTGTAGACATCT -3'
	ISG56	5'- GACAAGGCAATCACCCCTCTAC -3'
		5'- GTCTTTCAGCCACTTTCTCCA -3'
	TNF	5'- TCACTGGAGCCTCGAATGTC -3'
		5'- GTGAGGAAGGCTGTGCATTG -3'
	IL-6	5'- ACAGAAGGAGTGGCTAAGGA -3'
		5'- CGCACTAGGTTTGCCGAGTA -3'
MCP-1	5'- GCTTCTGGGCCTGCTGTTCA -3'	
	5'- AGCTCTCCAGCCTACTCATT -3'	
β -actin	5'- GAGACCTTCAACACCCCAGC -3'	
	5'- ATGTCACGCACGATTTC -3'	

Table S2. I/T ratios in HD virus isolated from HMC-1 cells and A549 cells following H1N1 infection.

	Time of sample collection	Infectivity titer (I): TCID50/25 μ l	Total titer (T): HA/25 μ l	I/T
HD virus (HMC-1)	30h	6264.84	16	391.5525
HD virus (A549)	18h	6264.84	16	391.5525
LD virus	/	9929102.93	256	38785.55832

Table S3. The safety of mice treated with different doses of H1N1 LD virus and their protective efficacy on mice to fight against IAV. Mice were firstly treated with different doses of H1N1 LD virus and re-challenged with 120TCID50/mouse H1N1 LD virus at 21 days. The survival rates of infected mice were recorded.

TCID50/ mouse	Days post-challenge							Days post-rechallenge (120TCID50/mouse)					
	8	9	10	11	12	13	14	9	10	11	12	13	14
10	5/7	4/7	3/7	3/7	1/7	1/7	1/7	/	/	/	/	/	/
5	6/7	6/7	4/7	3/7	2/7	2/7	2/7	/	/	/	/	/	/
1	7/7	6/7	5/7	5/7	3/7	2/7	2/7	/	/	/	/	/	/
0.5	7/7	7/7	7/7	6/7	5/7	4/7	4/7	/	/	/	/	/	/
0.1	7/7	7/7	7/7	6/7	6/7	6/7	5/7	/	/	/	/	/	/
0.05	7/7	7/7	7/7	7/7	7/7	7/7	7/7	5/7	5/7	3/7	2/7	2/7	2/7
0.01	7/7	7/7	7/7	7/7	7/7	7/7	7/7	6/7	4/7	4/7	3/7	2/7	2/7