

Table S1. Primers used in this study

Primer names	Primer sequences (5' - 3')
D117L-T-RNA-F	<u>TAATACGACTCACTATAGGGACAATCGCCAAATATCCCGGAA</u>
D117L-T-RNA-R	GTGGTGAGGCTCAGGTTGTTGTA
D117L-crRNA1-F	<u>GAAATTAATACGACTCACTATAGGGGATTAGACTACCCCAAAAAACGAAGGGGACTAAAACGAGC</u> AAAATCAAAATGCCCAGGA
D117L-crRNA1-R	TCCTGGGCATTTTGATTTTGCTCGTTTTAGTCCCCTTCGTTTTTGGGGTAGTCTAAATCCCCTATAG TGAGTCGTATTAATTC
D117L-crRNA2-F	<u>GAAATTAATACGACTCACTATAGGGGATTAGACTACCCCAAAAAACGAAGGGGACTAAAACGAG</u> CAAAATCAAAATGCCCAGG
D117L-crRNA2-R	CCTGGGCATTTTGATTTTGCTCAGTTTTAGTCCCCTTCGTTTTTGGGGTAGTCTAAATCCCCTATAG TGAGTCGTATTAATTC
D117L-crRNA3-F	<u>GAAATTAATACGACTCACTATAGGGGATTAGACTACCCCAAAAAACGAAGGGGACTAAAACCTTATA</u> GTAAACGATGGCAACGAT
D117L-crRNA3-R	ATCGTTGCCATCGTTTACTATAAGTTTTAGTCCCCTTCGTTTTTGGGGTAGTCTAAATCCCCTATAG TGAGTCGTATTAATTC
RPA-D117L-F1	<u>GAAATTAATACGACTCACTATAGGGACAATCGCCAAATATCCCGGAACAACCTGCGATTC</u>
RPA-D117L-R1	AGAATACCGGAAATGGTGGTGAGGCTCAGGTTG
RPA-D117L-F2	<u>TAATACGACTCACTATAGGGTTTTAGCCATAACAATCGCCAAATATCCCGGAAC</u>
RPA-D117L-R2	CGGAAATGGTGGTGAGGCTCAGGTTGTTGTACA
PRRSV-UF	TAATACGACTCACTATAGGGGCCCTGCCAYCACG
PRRSV-UR	TCGCCCTAATTGAATAGGTGA

PEDV-N-F1	TAATACGACTCACTATAGGGGGGCCGTCGACATGGCTTCTGTCAGTTTTCAGGATCG
PEDV-N-F1	GGGCCGATATTCATTTCTGTATCGAAGATCTCGTTGATAATTTCAAC
PPV4-F1	TAATACGACTCACTATAGGGCTTTGCTTTGTCCAACGCAGA
PPV4-R1	TAGATGTCCTGGCACAGATACTTGAC
PRV-gC-F1	TAATACGACTCACTATAGGGCGAGACCGAGGGCGTCTACAC
PRV-gC-R1	GCCCATCATCAGCGCCTGC
CSFV-UF	CTGGGTGGTCTAAGTCCTGAGTA
CSFV-UR	GATCAACTCCATGTGCCATGTA
PCV3-F1	TAATACGACTCACTATAGGGCTGTTATTTTGGATGATTTTATG
PCV3-R1	CACAGCCGTTACTTCACCC
SIV-HA-F1	AAACCAATGATAGGGCCAA
SIV-HA-R1	TTGCACTACACAGTTACCAC
p72-UF	TGCTCATGGTATCAATCTTATCG
p72-UR	CCACTGGGTTGGTATTCCTC
p72-TaqMan	5' FAM-TTCCATCAAAGTTCTGCAGCTCTT-TAMRA 3'

Note: T7 promoter sequences are underlined, and LwCas13a binding sequence is bold and italic.

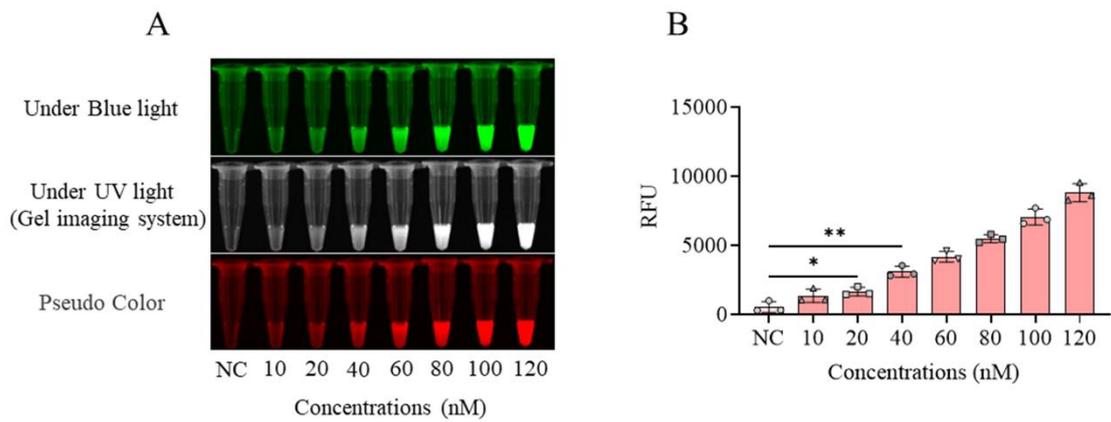


Fig S1

Figure S1. Optimization of RNA probe 5'-6-FAM-ssRNA-BHQ1-3' concentration. A, The blue and ultraviolet light signals mediated by crRNA1 at different concentrations of RNA probe. B, The fluorescence signal intensity mediated by crRNA1 at different probe concentrations by using microplate reader at different concentrations of RNA probe. * $P < 0.05$, ** $p < 0.01$ versus a single no target RNA control (NC) (n=3).

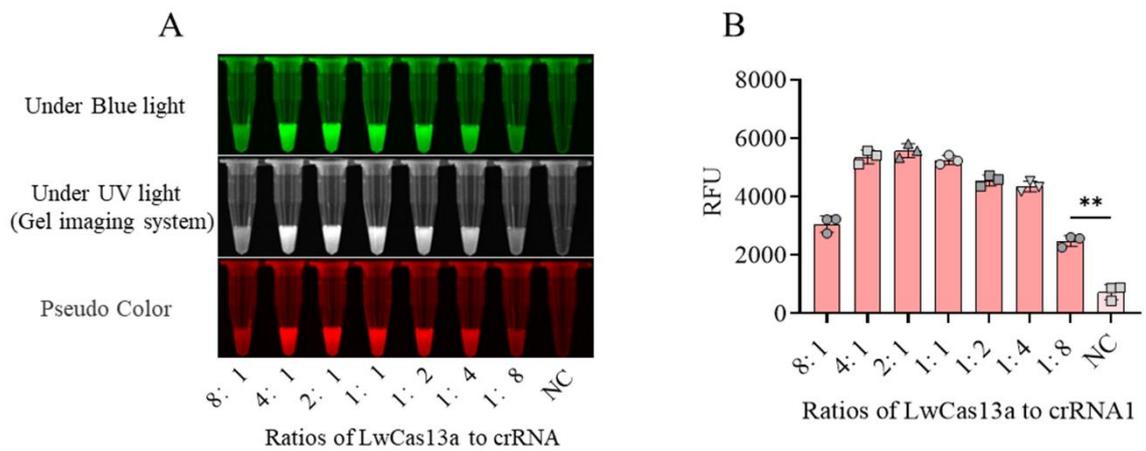


Fig S2

Figure S2. Optimization of the ratio between LwCas13a protein and crRNA1. A, The blue and ultraviolet light signals mediated by crRNA1 at the indicated ratios of LwCas13a : crRNA1. B, The fluorescence signal intensity mediated by crRNA1 at the indicated ratios of LwCas13a : crRNA1. ** $P < 0.01$ versus no target RNA control (NC) (n=3).