

*Supplementary Materials*

# Combinatorial F-G Immunogens as Nipah and Respiratory Syncytial Virus Vaccine Candidates

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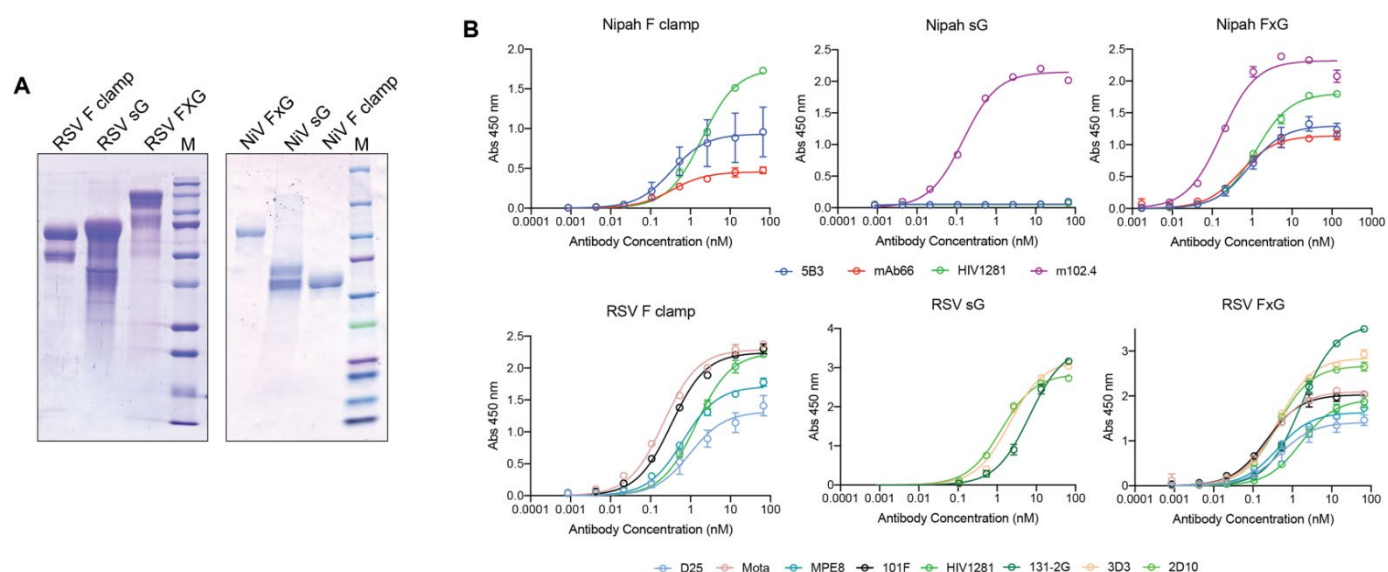
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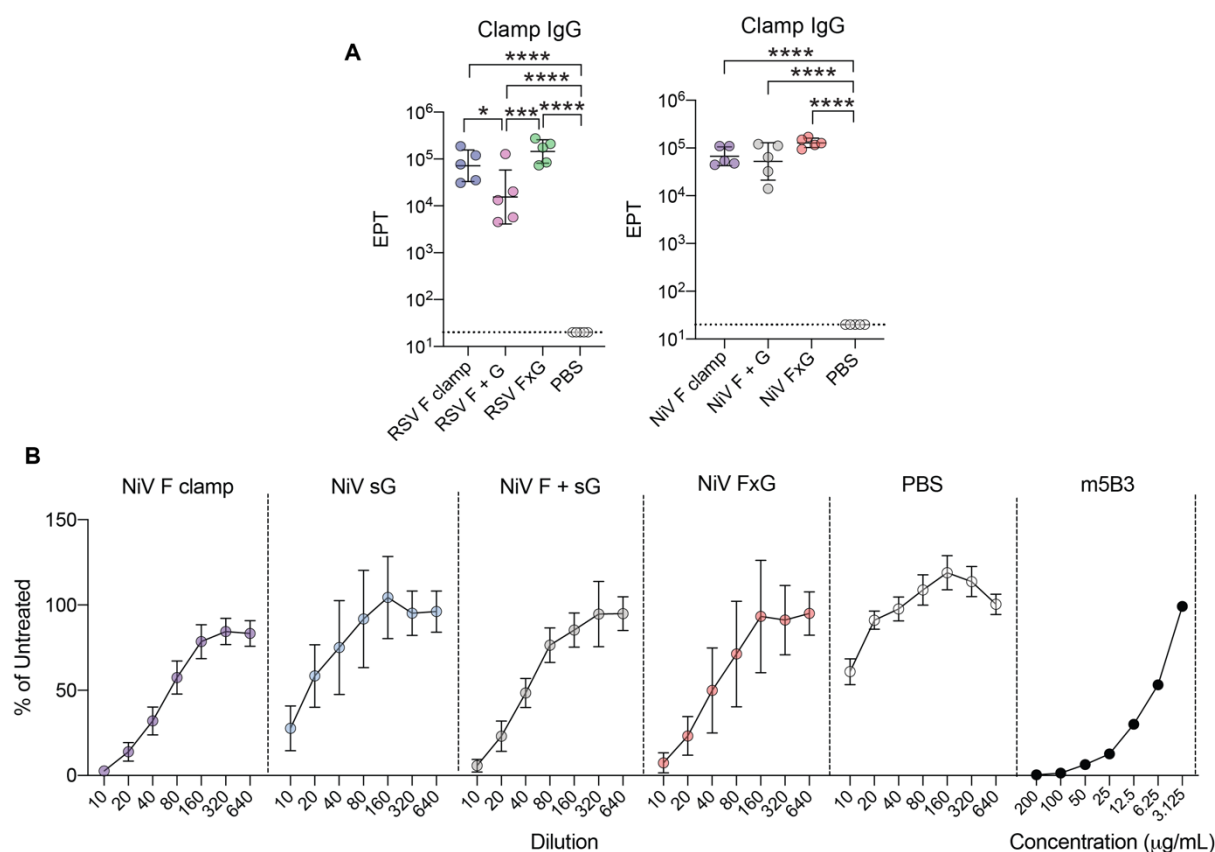
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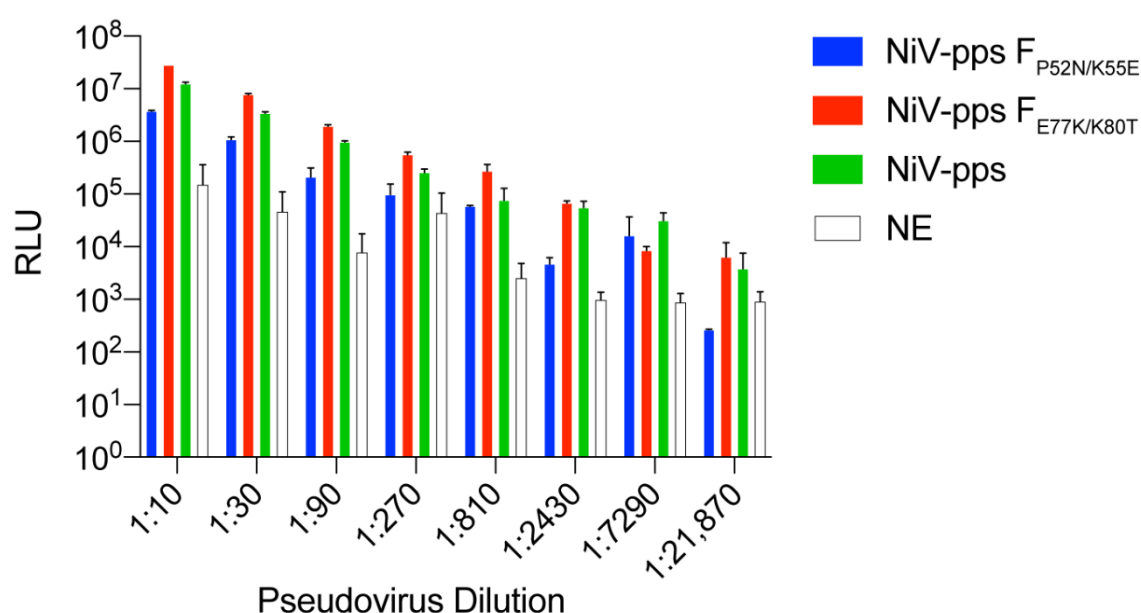
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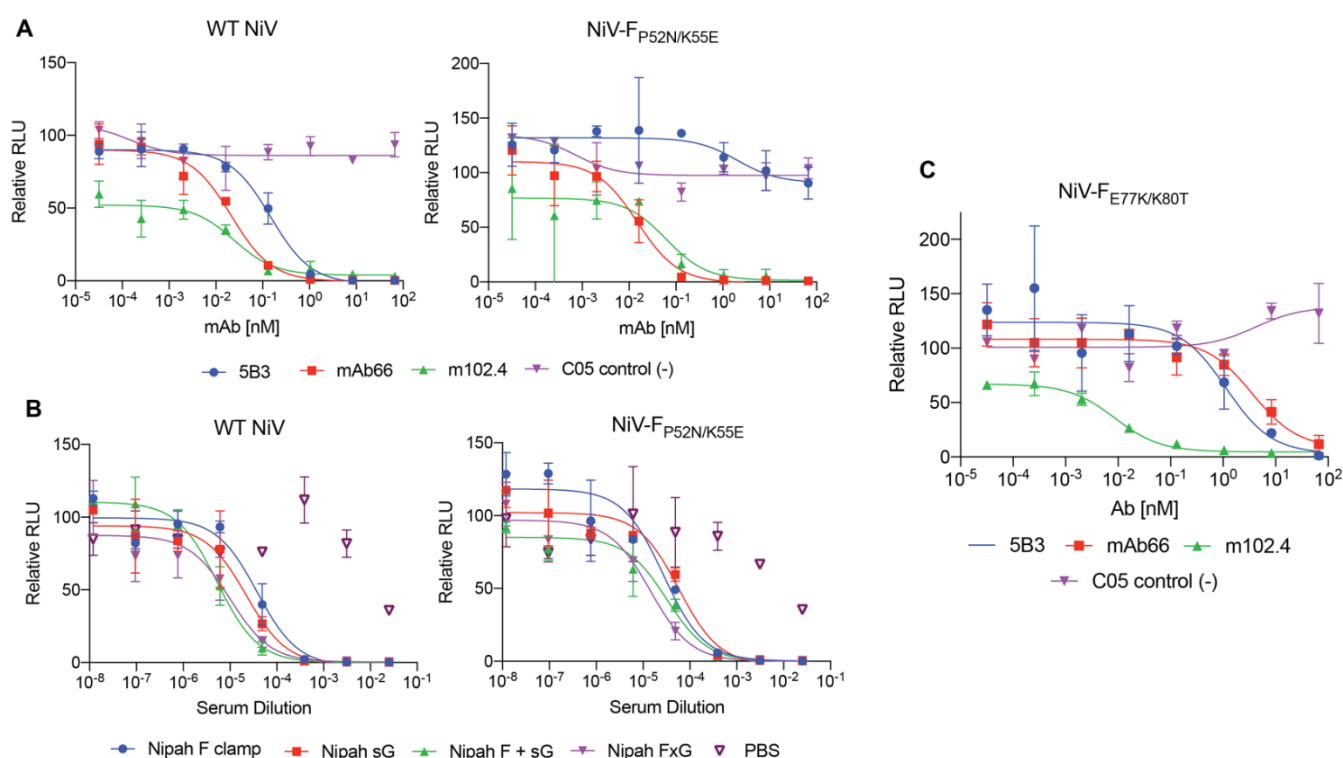
**Figure S1.** (A) SDS-PAGE of RSV and NiV antigens run under reducing conditions. M is Bio-Rad Kaleidoscope molecular marker. (B) Indirect ELISAs of mAbs against NiV and RSV antigens. Data shown of mean of duplicate values with standard deviation. Data fitted with one-site specific binding model to calculate  $K_d$ .



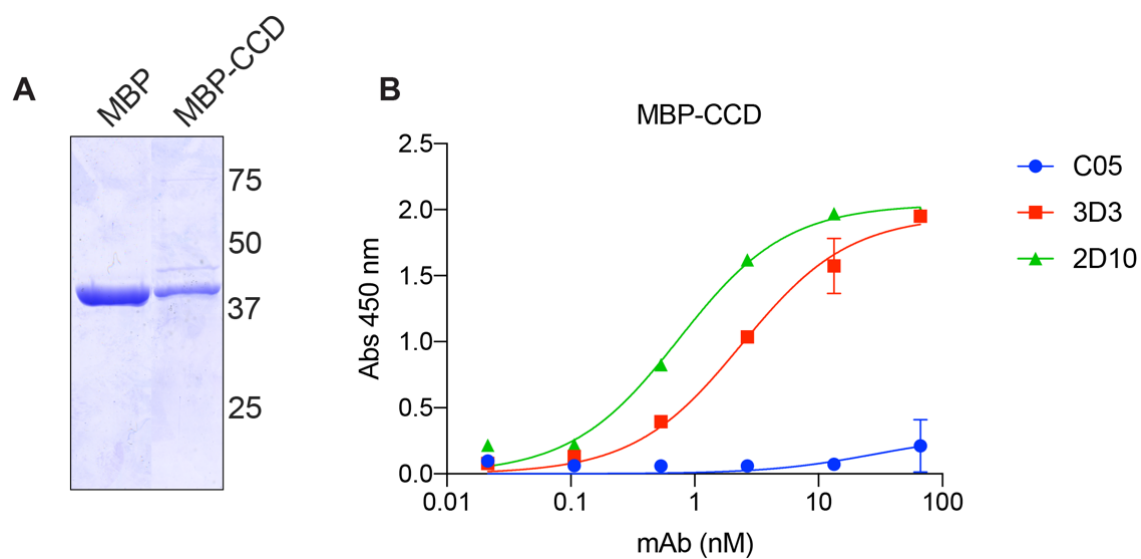
**Figure S2.** (A) Endpoint titer (EPT) of each RSV (left) and NiV (right) groups against the molecular clamp domain, assayed against a non-specific clamp-stabilised protein. Dotted line shows limit of detection and data is expressed as geometric mean with geometric standard deviation (SD). p-values calculated using a one-way Tukey's multiple comparison ANOVA on log transformed values, where \* =  $p < 0.05$ , \*\* =  $p < 0.005$ , \*\*\* =  $p < 0.0005$  and \*\*\*\* =  $p < 0.0001$ . (B) mFITs of NiV groups showing serial dilutions of sera from vaccinated groups as well as positive mAb controls. Each mouse sample was assayed in triplicate. Data is expressed as a percentage of the average luciferase readings seen in untreated controls (no serum) with 50% inhibition dotted line indicated ( $IC_{50}$ ). Values shown are group mean ( $n = 5$ ) with SD.



**Figure S3.** Titration of NiV pseudovirus (NiV-pps) wild-type or mutant F variants on BHK-21 cells. Titer is measured as relative light units (RLU) emitted by firefly luciferase reporter. Non-enveloped (NE) virus is included as background control.



**Figure S4.** NiV pseudovirus neutralisation assays of mAbs (A) and sera (B) against wild-type NiV (WT) or NiV-F<sub>P52N/K55E</sub> mutant knockout. (C) NiV pseudovirus neutralisation assay of mAbs against NiV-F<sub>E77K/K80T</sub> mutant knockout. Data shown is duplicate values of pooled sera or duplicate values of mAbs expressed as percent inhibition relative to virus only readings (relative RLU). C05 is a non-specific antibody control.



**Figure S5.** (A) SDS-PAGE of MBP and MBP-CCD purified protein run under reducing conditions. (B) Indirect ELISA of CCD-specific mAbs or non-specific control (C05) against MBP-CCD protein. Data shown is average of duplicate values with SD and background binding to MBP only subtracted.