

1 **Supplementary material to**

2 **SARS-CoV-2 Neutralization in Commercial Lots of Plasma-derived Immunoglobulin**

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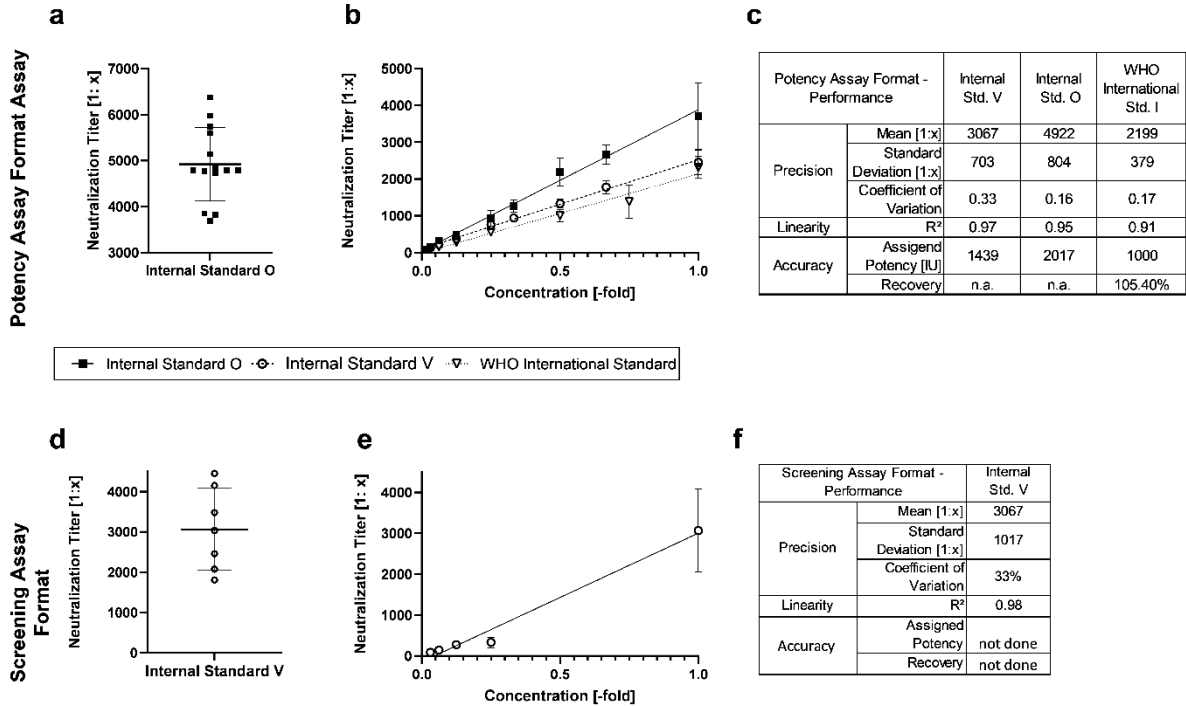
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2 Supp.Fig. 1 Retrospective SARS-CoV-2 microneutralization assay characteristics for potency assay format (a-c)

3 and screening assay format (d-f). a and d intermediate precision of internal standards O and V measured as fold

4 dilution as calibrator in each experimental run. b and e linearity data of the two assay formats. Depicted is the

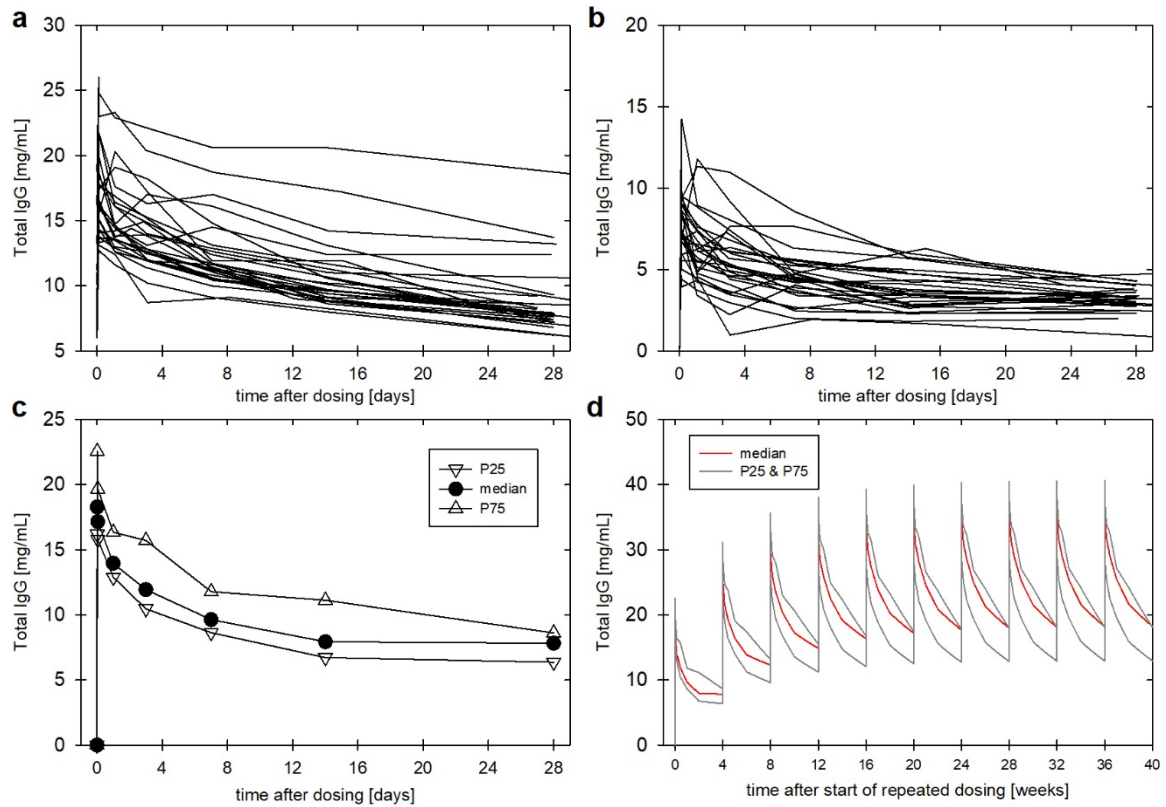
5 mean result ± SD per dilution level for each standard tested. c and f tabular results of retrospective analyses on

6 potency and screening assay formats for precision, linearity and accuracy, where applicable

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2 Supp. Fig. 3: PK data processing for superposition analyses. **a** reference dataset of the observed individual time  
 3 courses of untransformed IgG-levels on 4-weekly repeated dosing of IVIG (median dose: 0.406 g/kg-BW) [40].  
 4 **b** individual time courses of the baseline-adjusted IgG-levels on 4-weekly dosing of IgG. **c** reference single-dose  
 5 profiles, i.e. time courses of the median, 25th (P25), and 75th (p75) percentiles of the data from panel b dose-  
 6 normalized by g/kg-BW. **d** approximated time courses of the dose-normalized IgG-levels (mg/ml per g/kg-BW  
 7 dosed) throughout ten 4-weekly repeated doses of IVIG derived from the median, P25 and P75 reference profiles  
 8 (see panel c) using a superposition cascade with time-staggered dosing