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Fab-dimerized glycan-reactive antibodies are a structural category of natural antibodies

Authors
Wilton B. Williams, R. Ryan Meyerhoff, R.J. Edwards, ..., George M. Shaw, Priyamvada Acharya, Barton F. Haynes

Correspondence
wilton.williams@duke.edu (W.B.W.), priyamvada.acharya@duke.edu (P.A.), hayne002@mc.duke.edu (B.F.H.)

In brief
Structural and functional analyses identify a category of glycan-reactive antibodies in macaques and humans that are marked by the dimerization of the antigen-binding fragment. These antibodies are involved in HIV neutralization and also recognize the S2 protein of SARS-CoV-2.

Highlights
- Fab-dimerized glycan-reactive (FDG) natural antibodies (Abs) are prevalent in primates
- Fab-dimerization can occur independent of V_{H} domain-swapping
- Precursor FDG natural Abs acquire neutralization breadth in retroviral infection
- FDG Abs recognize a quaternary epitope in the S2 subunit of SARS-CoV-2 spike
Fab-dimerized glycan-reactive antibodies are a structural category of natural antibodies

Wilton B. Williams,1,2,18,23,‡ R. Ryan Meyerhoff,1,2,18,22 R.J. Edwards,1,2,18 Hui Li,6 Kartik Manne,1 Nathan I. Nicely,1,19 Rory Henderson,1,18 Ye Zhou,15 Katarzyna Janowska,1 Katayoun Mansouri,1 Sophie Gobell,1 Tyler Evangelous,1 Bhavna Hora,1 Madison Berry,7 A. Yousef Abuahmad,1 Jordan Sprenz,1 Margaret Dayton,1 Victoria Stalls,1 Megan Kopp,1 Allen L. Hsu,1 Mario J. Borgia,7 Guillame B.E. Stewart-Jones,6 Matthew S. Lee,19 Naomi Bronkema,10 M. Anthony Moody,1,3,5 Kevin Wiehe,1,2 Todd Bradley,1,20 S. Munir Alam,1,2 Robert J. Parks,1 Andrew Foulger,1 Thomas Ogün,1 Gregory D. Sempowski,1,18 Mattia Bonsignori,1,2 Celia C. LaBranche,4 David C. Montefiori,1,4 Michael Seaman,1,19 Sampa Santra,11 John Perfect,2 Joseph R. Francica,8 Geoffrey M. Lynn,1,14 Baptiste Aussedat,12,21 William E. Walkowicz,12,21 Richard Laga,13 Garnett Kelsoe,1,3 Kevin O. Saunders,1,3,4 Daniela Fera,10

1Duke Human Vaccine Institute, Durham, NC 27710, USA
2Department of Medicine, Duke University, Durham, NC 27710, USA
3Department of Immunology, Duke University, Durham, NC 27710, USA
4Department of Surgery, Duke University, Durham, NC 27710, USA
5Department of Pediatrics, Duke University, Durham, NC 27710, USA
6Perelman School of Medicine, University of Pennsylvania, Philadelphia, PA 19104, USA
7Genome Integrity and Structural Biology Laboratory, NIEHS, NIH, Department of Health and Human Services, Research Triangle Park, NC 27709, USA
8Vaccine Research Center, NIAID, NIH, Bethesda, MD 20892, USA
9Boston Children’s Hospital and Harvard Medical School, Boston, MA 02115, USA
10Department of Chemistry and Biochemistry, Swarthmore College, Swarthmore, PA 19081, USA
11Beth Israel Deaconess Medical Center and Harvard Medical School, Boston, MA 02215, USA
12Memorial Sloan Kettering Cancer Center, New York, NY 10065, USA
13Institute of Macromolecular Chemistry, Czech Academy of Sciences, Prague, Czech Republic
14Avidea Technologies, Inc., Baltimore, MD, USA
15Department of Computer Science, Duke University, Durham, NC 27708, USA
16Department of Biochemistry, Duke University, Durham, NC 27705, USA
17Department of Electrical and Computer Engineering, Duke University, Durham, NC 27708, USA
18These authors contributed equally
19Present address: University of North Carolina-Chapel Hill, Chapel Hill, NC 27599, USA
20Present address: Children’s Mercy Hospital, Kansas City, MO 64108, USA
21Present address: Chemitope Glycopeptide, Brooklyn, NY 11220, USA
22Present address: Division of Interventional Radiology, Department of Radiology, Duke University, Durham, NC 27710, USA
23Lead contact

‡Correspondence: wilton.williams@duke.edu (W.B.W.), priyamvada.acharya@duke.edu (P.A.), hayne002@mc.duke.edu (B.F.H.)

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SUMMARY

Natural antibodies (Abs) can target host glycans on the surface of pathogens. We studied the evolution of glycan-reactive B cells of rhesus macaques and humans using glycosylated HIV-1 envelope (Env) as a model antigen. 2G12 is a broadly neutralizing Ab (bnAb) that targets a conserved glycan patch on Env of genetically diverse HIV-1 strains using a unique heavy-chain (VH) domain-swapped architecture that results in fragment antigen-binding (Fab) dimerization. Here, we describe HIV-1 Env Fab-dimerized glycan (FDG)-reactive bnAbs without VH-swapped domains from simian-human immunodeficiency virus (SHIV)-infected macaques. FDG Abs also recognized cell-surface glycans on diverse pathogens, including yeast and severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) spike. FDG precursors were expanded by glycan-bearing immunogens in macaques and were abundant in HIV-1-naive humans. Moreover, FDG precursors were predominately mutated IgM⁺IgD⁺CD27⁺, thus suggesting that they originated from a pool of antigen-experienced IgM⁺ or marginal zone B cells.
INTRODUCTION

Many pathogens, including human immunodeficiency virus (HIV) envelope (Env) (Doores et al., 2010a; Bonomelli et al., 2011; Cao et al., 2017), yeast (Masuoka, 2004; Hall and Gow, 2013), and severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) spike proteins (Watanabe et al., 2020) express surface glycans. Anti-glycan antibodies (Abs) constitute a subset of natural Abs that are present prior to stimulation by cognate antigen, can exist as different isotypes, and provide the first line of defense against bacterial, fungal, and viral infections (Holodick et al., 2017). In humans, natural Abs can be mutated either after antigen stimulation or by unknown processes as immunoglobulin M (IgM) Abs in the pre-immune B cell repertoire (Holodick et al., 2017; Klein et al., 1998) and have been postulated to be derived from marginal zone and transitional B cells (Casali et al., 1987; Griffin et al., 2011; Quach et al., 2016; Weller et al., 2004; Weill et al., 2009).

HIV-type 1 (HIV-1) Env is heavily glycosylated, thus Abs have limited access to epitopes targeted by broadly neutralizing Abs (bnAbs) that are effective against multiple geographically diverse HIV-1 strains (Doores et al., 2010a; Bonomelli et al., 2011; Cao et al., 2017; Pritchard et al., 2015). HIV-1 Env-reactive bnAbs have been described that used long complementary-determining region (CDR) loops to gain access to bnAb epitopes in the Env protein, while also making stabilizing contacts with surrounding glycans (Walker et al., 2009; Bonsignori et al., 2011, 2017; Pejchal et al., 2011; Garces et al., 2014, 2015; Andrabli et al., 2015; Doria-Rose et al., 2015; Gorman et al., 2016; Fera et al., 2018). Thus, investigators have long sought anti-glycan Abs as a means to turn an effective HIV-1 defense into a vulnerability (Scanlan et al., 2007; Doores, 2015; Crispin et al., 2018; Daniels and Saunders, 2019). However, glycans found on HIV-1 and other pathogens are also present on host molecules (Shackelford and Strominger, 1983; Yu et al., 2010; Blixt et al., 2004), suggesting that the induction of Abs targeting pathogen glycans may be controlled by immune tolerance mechanisms (Scanlan et al., 2007). The Ab 2G12 has been the only example to date of an HIV-1 bnAb that interacts specifically short variable heavy (VH) domain CDR3 loop and a unique immune tolerance mechanisms (Scanlan et al., 2007; QuaÁch et al., 2016). The Ab 2G12 recognizes Man9-enriched glycans but did not neutralize corresponding VH domain swapping for HIV-1 neutralization. FDG Abs, in addition to binding yeast and HIV-1 Env glycans, also recognized a glycan cluster in the S2 subunit of the SARS-CoV-2 spike. Thus, our study emphasized the specificity of FDG Abs for glycosylated proteins and suggested that this natural FDG Ab precursor pool can target diverse human pathogens.

RESULTS

Vaccine-induced HIV-1 Env glycan-reactive Abs

We previously reported a glycopeptide (Man9-V3) mimic of a bnAb epitope containing high-mannose glycoforms on native HIV-1 Env trimer (Alam et al., 2017; Fera et al., 2018). In an effort to induce bnAbs that target high-mannose glycans on HIV-1 Env, four RMs previously immunized with monomeric Man9-V3 (Alam et al., 2017) were subsequently immunized with a multimeric form of Man9-V3 (Figure 1A). After repeated immunizations with monomeric Man9-V3, plasma Abs from two of the four RMs bound Man9-V3; whereas plasma from all four RMs bound Man9-V3 following a single immunization with multimeric Man9-V3 (Figure 1B). From a representative macaque, we isolated a 4-member DH717 Ab clonal lineage; DH717.1 IgG and DH717.2-DH717.4 IgMs were isolated before and after immunization with multimeric Man9-V3, respectively (Figure 1C; Data S3). Recombinant DH717 monoclonal Abs (mAbs) demonstrated glycan-dependent binding to Man9-V3 (Alam et al., 2017), and bound soluble recombinant HIV-1 Env trimers (Saunders et al., 2019). Candida albicans or Cryptococcus neoformans yeast glycans, and high mannose glycans including Man9GlcNAc2 (Figures 2B–2D, S1A, and S1B). DH717 mAbs neutralized env-pseudotyped HIV-1 bearing Envs with Man9-enriched glycans but did not neutralize corresponding wild-type HIV-1 bearing Envs with heterogeneous glycoforms (Figure S1C). Negative stain electron microscopy (NSEM) of DH717.1, DH717.2, and DH717.4 revealed a mixture of canonical Y-shaped Abs, as well as l-shaped Fab-dimerized Abs reminiscent of the l-shaped HIV-1 bnAb 2G12 (Figure 1D). In 2G12, the two Fabs lie side-by-side and in the same plane; thus, both Fabs are seen in a flatwise view via NSEM. In contrast, in DH717, one Fab is seen flatwise, whereas the other Fab is seen edgewise, suggesting that the two Fabs are turned relative to each other 90° about their long axes (Figure 1D). This arrangement was confirmed by 3D negative stain reconstruction of DH717.1 IgG (Figure S1D).

Structural details of Env glycan recognition by DH717 Abs

We obtained a 2.6-Å crystal structure of monomeric DH717.1 Fab complexed with Man9-V3 (Figures 1E and 1G; Data S4).
DH717.1 V\textsubscript{H} interacted with the Man\textsubscript{9}-glycan using three HCDR loops to form a binding pocket into which Man\textsubscript{9}-glycan was inserted (Figures 1G and S1E). No interactions were observed with the peptide itself, reminiscent of the glycan-only binding of 2G12 (Doores et al., 2010b; Murin et al., 2014; Stanfield et al., 2015; Calrese et al., 2003; Chuang et al., 2019). We also determined a 3.5-Å crystal structure of unliganded dimeric DH717.1 Fab (Figures 1F, 1H, and S1G; Data S4). The structure revealed two disulfide-linked Fab dimers in the asymmetric unit, with a disulfide bond linking two Fab via V\textsubscript{H} gene residues cysteine 74 (Figure 1H). DH717.3, which only showed Y-shaped Abs, had a serine at this position (Figure S1F), and disrupting the disulfide in DH717.1 with a cysteine to serine (C74S) mutation resulted in loss of DH717.1 I-shape forms (Figure 1I) and reduction in binding and neutralizing activity (Figures 1J, 1K, and S1H). DH717.1 Fab monomer did not bind recombinant HIV-1 Env trimers, whereas the Fab dimer bound with micromolar affinity, and the DH717.1 IgG mAb bound with nanomolar affinity, thus demonstrating a role for avidity in Env glycan recognition of the DH717.1 IgG mAb bound with nanomolar affinity, thus suggesting a role for host or environmental antigens in their initiation. DH717 B cell lineage was expanded following immunizations with monomeric and multimeric Man\textsubscript{9}-V3 (Francia et al., 2019; Alexander et al., 2000; Alam et al., 2017) as well as a soluble recombinant HIV-1 Env trimmer (Saunders et al., 2019) bearing near-native high mannose glycan patch (Figure 2A). Recombinant mAbs (DH717.5 NGS and DH717.6 NGS) bearing DH717 clonally related mutated IgM V\textsubscript{H} found at week 0 paired with post-vaccination DH717.1 V\textsubscript{H} had minimal or no binding to soluble recombinant Env trimers, Man\textsubscript{9}-V3, Candida albicans yeast antigens, and high mannose glycans (Figures 2B–2E). Additionally, the DH717 mAb bearing the near-germline unmutated common ancestor (UCA) genes weakly bound Candida albicans glycans but did not bind Man\textsubscript{9}-V3, whereas the mature DH717 lineages mAbs demonstrated affinity maturation after Man\textsubscript{9}-V3 immunization as shown by increased binding to Man\textsubscript{9}-V3, soluble recombinant HIV-1 Env trimer, and Candida albicans or Cryptococcus neoformans yeast glycans (Figures 2F–2E).

To define the nature and characteristics of circulating DH717 FDB Abs, we interrogated the transcriptome of blood DH717 B cells. We studied 23 transcriptionally unique B cell clusters in blood DH717 lineage B cells, 1 week after final immunization with the nanoparticle form of Man\textsubscript{9}-V3 (Figure 2F). The DH717 B cells were found in clusters 0 and 4 (Figure 2F) that had 315 enriched genes that achieved statistical significance with a 0.3–1.4 log fold change (Figure 2G; Data S5). B cell receptor signaling pathway was enriched in both clusters 0 and 4, and IgM was the top gene enriched in cluster 4 with statistical significance (log fold change = 0.9, p < 0.001) (Data S5). Interestingly, we found genes for IgM, phosphatases, and FcR-like proteins enriched in clusters 0 and 4, similar to transcripts previously found in human blood IgM memory and marginal zone (MZ) B cells (Weller et al., 2004). Moreover, we found that IgM DH717.3 displayed an IgM*IgD*CD27* phenotype (Figure 2H) in agreement with previously identified human blood marginal zone B cells (Weill et al., 2009; Weller et al., 2004). Additional DH717 IgM Abs displayed IgM* and IgD*CD27* or IgD*CD27* phenotypes (not shown), thus implicating different differentiation pathways for DH717 lineage B cells.

Precursor FDB Abs in HIV-1-naive humans

We interrogated the B cell repertoire of nine HIV-1 naive individuals for candidate FDB Ab precursors that demonstrated Man\textsubscript{9}-V3 or yeast binding, and/or contained I-shaped Abs. Antigen-specific flow sorting revealed that candidate FDB Ab precursors were present at an average frequency of 1 per ~340,000 B cells (Figure 3A), which was considerably greater than the frequencies of bnAb precursors that targeted different HIV-1 Env epitopes (Steichen et al., 2019; Havenar-Daughton et al., 2018). From flow sorting, 37 glycan-reactive Abs used a variety of heavy and light chain genes, although 46% used V\textsubscript{k}2-24 for the light chain (Figure 3B) in contrast to a V\textsubscript{k}2-24 frequency of 0.3%–0.6% in HIV-1 naive individuals (DeKosky et al., 2015), demonstrating immunoglobulin repertoire skewing as previously reported for natural Abs (New et al., 2016). The majority of flow

Figure 1. Characteristics of HIV-1 vaccine-induced Env glycan-reactive neutralizing B cell lineage, DH717
(A) Schedule of Man\textsubscript{9}-V3 immunizations in RMs. (B) Plasma IgG at weeks (WK) post infection from four immunized RMs were tested in ELISA for binding to Man\textsubscript{9}-V3. Data shown are representative from multiple assays and binding titer was reported as log area under the curve (AUC). Vertical dotted lines indicate immunization time points. (C) Representative flow cytometry sort plot of HIV-1 Env-reactive memory B cells detected at WK 26 and 73. Phylogram of V\textsubscript{H} genes isolated from a clonal lineage of DH717 Abs found at WK 25 and 73. (D) NSEM 2D class averages for DH717.1–DH717.4, and HIV-1 bnAbs 2G12 and VRC01, are labeled I or Y according to shape. (E and F) Monomeric (E) and dimeric (F) Fabs are also shown. (G) Crystal structure of DH717.1 Fab monomer (V\textsubscript{H} and V\textsubscript{L} in dark and light teal, respectively) bound to Man\textsubscript{9}-V3. The terminal glycan-moieties of Man 9-V3 (gray sticks) bound a pocket formed by the CDRH1–3 loops (dark red, orange and orange) of the DH717.1 Fab. Inset shows a zoomed-in view of glycans binding. (H) Crystal structure of DH717.1 dimer with dark teal or pink VH, and light teal or pink VL. Inset shows a zoomed-in view of the inter-Fab disulfide (yellow spheres). (I) NSEM of cysteine to serine DH717.1 mutant (C74S). (J) DH717.1 wild-type and C74S mutant IgGs were tested for neutralization of env-pseudotyped HIV-1 bearing EnvVs with heterogeneous [−Kif] or Man\textsubscript{9}-enriched +[Kif] glycans in TZM-bi cells in a single assay. Neutralization titers were reported as IC\textsubscript{50} in μg/mL. These Abs were also tested in a SPR assay for binding soluble stabilized forms of recombinant HIV-1 Env trimers (K); shown are SPR binding curves for each Ab. See also Figure S1 and Data S1, S3, and S4.
Figure 2. The ontogeny and dynamics of HIV-1 vaccine-induced DH717 FDG Abs

(A) Absolute counts of unique V\textsubscript{H} reads for blood-derived DH717 lineage members at longitudinal time points (weeks). *p < 0.0001 (non-parametric Wilcoxon test) for increase in the distribution of estimated mutation frequencies at week 105 compared to week 145. The arrows indicate immunization time points with the initial Man\textsubscript{9}-V3 monomer (black arrow), Man\textsubscript{9}-V3 multimer (red arrow), nanoparticle (green arrow), and the first of five recombinant HIV-1 Env trimers (blue arrow).

(B–D) Recombinant mAbs bearing the week 0 (DH717.5\_NGS and DH717.6\_NGS), near-germline (DH717UCA), and post-vaccination (DH717.1–DH717.4) Ab sequences were tested in ELISA for binding soluble stabilized form of recombinant HIV-1 Env trimer (B), Man\textsubscript{9}-V3 (C), and heat-killed yeast antigens (D). CH65 was used as a negative control mAb, whereas biotinylated trimer-specific mAbs (see Figure S1), and peptide (DH1013)- and Env glycan (2G12)-specific mAbs were used as positive controls. ELISA data shown are averaged from 3–5 separate assays for each mAb tested; error bars represent SEM. Binding was measured at OD\textsubscript{450nm}.

(E) Binding of DH717 mAbs to individual glycans in a luminex binding assay, where glycan reactivity was reported as MFI after background subtraction.

(legend continued on next page)
sorted candidate FDG precursor B cells (76.1%, 405/532) were IgM* IgD*CD27+ B cells (Figure 3C).

Seven FDG Abs were chosen, based on glycan binding activity, for expression as mAbs (Data S3). Three of these, IgG3 DH1005, IgM DH1008, and IgM DH1010, displayed a mix of Y- and I-shaped Abs (Figures 3D–3F), and the I-shaped form of DH1005 bound soluble recombinant HIV-1 Env trimer (Figure 3D). All seven mAbs showed glycan-dependent binding to Man$_{9}$-V3 and bound Candida albicans and Cryptococcus neoformans yeast (Figure 3G). Only IgG3 DH1005 neutralized env-pseudotyped HIV-1 bearing Man$_{9}$-enriched Envs (Figure 3H) but did not neutralize wild-type HIV-1 with Envs bearing heterogeneous glycoforms (not shown), thus exhibiting a neutralization profile similar to DH717 Abs that we classified as FDG precursors. These data demonstrated that FDG Abs were abundant in the B cell repertoire of HIV seronegative humans and may be found in the pool of IgM* IgD*CD27+ marginal zone B cells (Weller et al., 2004; Weill et al., 2009).

DH1010 Fab-dimerized IgM mAb isolated from an HIV-1 naive individual had a free cysteine at position 74 in the V$_{H}$ gene, similar to DH717 mAbs isolated from an HIV-1 vaccinated RM. Of 77,336 immunoglobulin heavy and light chain gene pairs studied in three HIV-1 naive individuals (not shown), none contained a cysteine at position 74. Additionally, none of 201 germline V$_{H}$ genes in the human Clonealyst Ig gene database (Ramesh et al., 2017; see STAR Methods) had C74, and <1% of the Abs in the Abysis database (Swindells et al., 2017) had a V$_{H}$ gene encoding C74. These data demonstrated that human Abs with a cysteine in V$_{H}$ position 74 are rare but they may be selected by Man$_{9}$-V3 and HIV-1 Env trimer as observed in RMs by selection of DH717 Abs.

**Macaque SHIV-induced neutralizing FDG B cell lineages**

Because the unique 2G12 bnAb arose in human HIV-1 infection (Buchacher et al., 1994), we sought to isolate FDG bnAbs from RMs infected with pathogenic SHIVs (Roark et al., 2021). Infection with HIV (Bonsignori et al., 2017) or a SHIV (Roark et al., 2021), with the same Env, elicited plasma bnAbs that targeted a high mannose glycan-containing epitope in humans and RMs, respectively. In one SHIV-infected RM (Roark et al., 2021), we interrogated the glycan-reactive memory B cell repertoires in blood and lymph node at week 52 post-infection, and blood at week 104 post-infection.

**Autologous neutralizing FDG B cell lineage DH898**

A seven-member IgG Ab clonal lineage, termed DH898 (Data S3), isolated from lymph node memory B cells at 52 weeks post SHIV infection, used macaque genes V$_{\mu}$2 paired with V$_{H}$1 encoding a 13-amino acid HCDR3 (Figure 4A). DH898 mAbs mediated Env glycan-dependent neutralization of difficult-to-neutralize env-pseudotyped HIV-1 bearing Env from the autologous SHIV infection and neutralized heterologous env-pseudotyped HIV-1 bearing Env from the autologous SHIV infection and neutralized heterologous env-pseudotyped HIV-1 bearing Env with Man$_{9}$-enriched glycans (Figure 4B) but neutralized none of 119 geographically diverse difficult-to-neutralize env-pseudotyped HIV-1 bearing heterologous Envs with heterogeneous glycoforms (Seaman et al., 2010; not shown). Like 2G12, DH898 mAbs demonstrated glycan-dependent binding to Man$_{9}$-V3 (Figure 4C) and bound yeast glycans (Figure S2A). DH898 mAbs also showed glycan-dependent binding to a soluble recombinant HIV-1 Env trimer (Saunders et al., 2019; Figures S2B and S2C). By NSEM, DH898 mAbs showed a mixture of canonical Y-shaped as well as I-shaped Abs (Calarese et al., 2003; Figures 4D and S2D).

To visualize the DH898 epitope and Fab dimer interface, we determined cryo-EM structures of a soluble recombinant HIV-1 Env trimer bound to recombinant DH898.1 Fab (Figures 4E–4G and S3; Data S4; Videos S1 and S2). We identified three populations—an unliganded trimer, and two other populations, each with a single DH898 Fab dimer bound to two different epitopes. In one population, the Fab dimer bound multiple glycans at Env positions 332, 392, and 396, which included high mannose glycans targeted by 2G12 (Figures 4G and S3; Video S2), and was consistent with DH898 IgG mAbs dependence on glycan at Env position 332 for neutralization of the autologous strain (Figure 4B). In the second population, the Fab dimer bound glycans at Env positions 386, 362, and 276 around the Env CD4-binding site (Figures S3L–S3N; Video S2). In both complexes, the Fab dimer interfaces were well resolved and did not show V$_{H}$ domain-swapping (Figures 4F and S3M). Instead, the DH898 Fabbs dimerized via a side-by-side association of their V$_{H}$ domains, mediated by interfacial aromatic and hydrophobic residues (Figures 4H, S2E, S3O, and S3P). The DH898 Fab dimer interface also involved a glutamate-arginine pair at residues E10 and R12 that were located such that they could form salt bridges between the two V$_{H}$ chains (Figure 4I–4J), analogous to a D72 and R57 aspartate-arginine pair seen in 2G12 (Calarese et al., 2003). Mutations predicted to disrupt the Fab dimer interface yielded a lower proportion of I-shaped Abs and loss of Ab binding activity. Conversely, mutations predicted to strengthen the Fab dimer interface resulted in higher proportion of I-shaped Abs, albeit without any enhancement of binding and neutralizing activity (Figures S2F–S2I), suggesting that the I-shaped conformation is necessary but not sufficient for enhanced function. These data demonstrated that DH898 were FDG Abs and suggested that they could be bnAb precursors.

**Broadly neutralizing FDG B cell lineage DH851**

A second four-member clonal lineage, termed DH851 (Data S3), comprised of one IgM and three IgG Abs, was isolated from week 52 lymph node and blood (IgG DH851.1-DH851.3) and...
Figure 3. Characteristics of HIV-1 Env glycan-reactive neutralizing Abs in the B cell repertoire of naive humans
(A) Repertoire analysis of blood-derived Env glycan-reactive B cells in nine HIV-1 naive individuals. Enriched B cells were studied via different sort strategies listed that used fluorophore-labeled Man$_9$-V3 or soluble stabilized recombinant HIV-1 Env trimers.
(B) Immunogenetics of candidate Env glycan-reactive B cells (N = 37) that were isolated from individuals listed in (A).
(C) Repertoire of Env glycan-reactive B cells with IgM$^+$IgD$^+$CD27$^+$ phenotype among six individuals. Flow cytometry data were analyzed from B cells sorted via strategies listed in (A); strategy “c” was used for LKP04.
(D–F) NSEM 2D class averages of DH1005 Ab with I-shaped Ab conformations denoted by “I” in image. 2D class averages of DH1005-SOSIP complex shows Fab-dimer binding to soluble stabilized recombinant HIV-1 Env trimer. Fc domain is not visible here because it lies outside the circular mask used during class averaging. Shown also are 2D class averages of DH1009 (E) and DH1010 (F) that displayed I-shaped Abs.
(G) Seven Env glycan-reactive mAbs were tested in ELISA for binding Man$_9$-V3 and non-glycosylated aglycone V3 peptide and heat-killed yeast antigens. Data shown are from a representative ELISA. Reference mAbs included peptide-reactive DH1013, Env glycan-reactive DH717.2 and 2G12, trimer-reactive PG T151, and negative control CH65 Abs.
(H) Only DH1005 neutralized multi-clade env-pseudotyped HIV-1 bearing Envs with Man$_9$-enriched glycans [Kif]. Neutralization titers are representative of two assays in TZM-bl cells, and neutralization titers were reported as IC$_{50}$ in µg/mL. Control mAbs were 2G12 and CH65.
See also Data S3 and STAR Methods.
Figure 4. Characteristics of HIV-1 Env glycan-reactive neutralizing B cell lineage, DH898, elicited by SHIV infection in RMs

(A) Phylogram and immunogenetics of DH898 neutralizing Abs isolated from a pathogenic SHIV-infected RM (6163).

(B) DH898 mAbs were tested for neutralization of env-pseudotyped HIV-1 bearing an autologous Env from SHIV infection. MAbs were tested for neutralizing HIV-1 bearing wild-type and glycan-deleted mutant Envs in TZM-bl cells. Data shown are from a representative assay and neutralization titers are IC50 in mg/mL.

(C) DH898 mAbs were tested for binding in ELISA to Man9-V3 (black bars) and non-glycosylated aglycone V3 peptide (gray bars). Binding titer was reported as log AUC. Control Abs tested included peptide-reactive DH1013, Env glycan-reactive 2G12, and negative-control anti-influenza CH65 mAbs. Data shown are representative of four assays.

(D) NSEM of DH898.1 IgG showing a mixture of I-shaped (I) and Y-shaped (Y) Abs.

(E) Segmented cryo-EM map of DH898.1 Fabs bound to a soluble stabilized HIV-1 Env trimer showing a single Fab-dimer bound to the trimer. Colors are as in Figure 1.

(F) Zoomed-in view of the Fab-dimer showing a gap between the two Fabs.

(G) A fitted model showing the Fab-dimer binding to high mannose glycans at Env positions 332, 392, and 396.

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were class-switched IgGs in vivo. See also Figures S2 and S3, Data S3 and S4, and Videos S1 and S2.

DH851 interactions with Env were dominated by and consistent with the observed cross-blocking with 2G12 (Figure 5B). DH851.1, DH851.2, and DH851.3 mAbs showed 26% neutralization breadth against a panel of 119 geographically diverse HIV-1 strains (Figure S4B) and were dependent on glycans at Env position 339, 386, and 392 glycans for neutralization (Figure S4C), which included high mannose glycans targeted by 2G12 (Chuang et al., 2019). Like 2G12, DH851 mAbs bound soluble recombinant HIV-1 Env trimers and Man3–V3 in a glycan-dependent manner, and bound Candida albicans or Cryptococcus neoformans yeast glycans (Figures 5C and S4D–S4G). Both bnAbs 2G12 and IgG DH851.2 cross-blocked each other for binding recombinant HIV-1 Env Abs (Figures S4D and S5).

NSEM of DH851 IgGs (DH851.1–DH851.3) revealed a mixture of Y- and I-shaped Abs (Figures 5F and S4H). In contrast, neither the mAbs bearing the near-germline DH851UCA genes nor the non-bnAb DH851 lineage member, DH851.4, were I-shaped (Figure S4H). The lack of I-shaped configurations for DH851.4 Ab and its inability to neutralize wild-type HIV-1 isolates bearing Env with heterogeneous glycoforms suggested that Fab dimerization was required for heterologous HIV-1 neutralization.

In a 5.6-Å cryo-EM reconstruction of DH851.3 bound to a soluble recombinant HIV Env trimer, we observed Env bound to three side-by-side Fab dimers (Figures 5G, 5H and S5; Data S4). Unlike 2G12, and like the I-shaped DH898 Abs, the DH851.3 Fab dimers did not show Vγ-domain-swapping, with a clear gap between the two Fabs in the cryo-EM reconstruction. The DH851.3 Fab dimers bound Env glycans at positions 241, 289, and 334 located within a previously described Env glycan-containing bnAb epitope near the V3 loop (Kong et al., 2013) and consistent with the observed cross-blocking with 2G12 (Figures 5D and S5). DH851 interactions with Env were dominated by its Vγ domains (Figure S1), which were rotated such that the beta-strands of both chains were nearly parallel (Figures 5J and 5K), as opposed to being angled as seen in DH898 (Figure 4I); nevertheless, the interface featured hydrophobic and aromatic residues similar to 2G12 and DH898 (Figures 5L, S3P, S4I, and S5). DH851.4 and the DH851UCA lacking key hydrophobic residues (Figure S4I), and NSEM did not show Fab-dimerization (Figure S4H). In particular, DH851.4 has a T81R mutation that would place a bulky charged residue in the dimer interface, analogous to the I19R mutation that disrupts Vγ domain-swapping and Fab dimerization in 2G12 (Doores et al., 2010b). In DH851, a dimer strengthening double mutant, A19V+T21L, increased the fraction of I-shaped Abs as expected (Figure S4J).

The structures of DH898 and DH851 Fab dimers suggested a common signature for non-domain-swapped Vγ-Vγ, Fab dimerization mediated by interfacial hydrophobic and aromatic residues. This sequence signature was also observed in DH501 (Data S2A), a glycan-reactive Ab previously isolated from a macaque following HIV-1 Env immunization (Saunders et al., 2017a). NSEM of DH501 IgG showed the presence of I-shaped Abs (Data S2B). Recombinant DH501 Fabs also showed Fab dimer populations (Data S2C), and the previously published DH501 crystal structure (PDB: 5IEE) revealed two Fabs within the unit cell, with their Vγ-domains in a side-by-side orientation reminiscent of the DH851 and DH898 Fab dimers (Data S2D). Mutations designed to strengthen or weaken the crystallographic dimer interface, increased or decreased the proportion of I-shaped DH501 IgG population observed, respectively (Data S2E–S2G), thus confirming that the crystallographic dimer was in fact a biologically relevant Fab dimer. When docked as a rigid body, the DH501 dimer showed good fit to the DH851 cryo-EM map (Data S2H). Analysis of the DH501 crystallographic dimer thus provided high-resolution insights into a non-domain-swapped Fab dimer interface (Data S2J–S2O). These data revealed common sequence and structural features of non-domain-swapped FDG Abs DH898, DH851, and DH501 (Figures S2E and S4I; Data S2) that includes the same Vγ hydrophobic pattern seen in 2G12, involving rare framework mutations.

We isolated another FDG bnAb lineage (DH1003) from a RM infected with a pathogenic SHIV different from the one that elicited DH851 (Data S2P and S3). Sequence analysis of DH1003 lineage Abs showed numerous hydrophobic residues among the key interface residues implicated in Fab dimerization (Data S2Q). Representative DH1003 mAbs showed predominantly I-shaped Abs by NSEM, demonstrated glycan-dependent binding to Man9–V3, bound to Candida albicans, and neutralized autologous and heterologous difficult-to-neutralize HIV-1 isolates bearing Envs with heterogeneous glycoforms (deCamp et al., 2014; Data S2S–S2V). These data demonstrated the reproducibility of FDG bnAb generation in macaque SHIV infections.

**Ontogeny of SHIV-induced FDG Abs**

Class-switched and mutated DH898 and DH851 clonally related Vγ genes were found in peripheral blood cells collected prior to macaque SHIV infection (week 0), suggesting that host or environmental antigens may have initiated Fab Ab lineages. We found 596 DH898 and 235 DH851 clonally related sequences that were predominantly IgM. To determine the binding profile of week 0 clonally related DH851 Vγ sequences, recombinant mAbs bearing week 0 DH851 IgG (DH851.5) or IgM (DH851.6 and DH851.8) clonally related VγDγ1-Vδ Abs were paired with DH851 Vγ genes. These pre-infection DH851 mAbs bound weakly to yeast antigens and soluble recombinant HIV-1 Env trimers, but did not bind Man9–V3; in contrast, post-SHIV-infection DH851
mAbs robustly bound yeast antigens, Man₃₋₇V₃, and soluble recombinant HIV-1 Env trimers (Figures S4D, S4F, and S4G). Pre- and post-infection IgM DH851 lineage Abs showed only Y-shaped populations by NSEM, whereas post-SHIV infection IgG DH851 lineage Abs showed Fab-dimerized, I-shaped populations (Figure S4H). These data suggested that FDG precursors existed as mutated or class-switched non-Fab-dimerized Abs prior to SHIV infection, with lineage stimulation potentially triggered by glycanc-bearing host or environmental antigens.

**Glycan-dependent binding to SARS-CoV-2 spike protein by FDG Abs**

Site-specific glycan analysis demonstrated that oligomannose-type glycans, including high mannose constitutes 28% of the glycans on the recombinant spike (S) protein of SARS-CoV-2, the causative agent of the current COVID-19 pandemic (Watanabe et al., 2020). Because FDG Abs recognized high mannose glycans, we tested them for binding SARS-CoV-2 S protein. FDG mAbs, including bnAbs 2G12 and DH851, bound the SARS-CoV-2 S protein to varying degrees in a glycan-dependent manner (Figures S6A and S6). A 3.1-A˚ cryo-EM reconstruction revealed a 2G12 Fab dimer bound to the S2 subunit of the SARS-CoV-2 S protein, with high-mannose glycans 717 and 801 from one protomer and glycan 709 from the adjacent protomer clustered together to form the 2G12 epitope (Figures 6B, 6C, and S7). The V₄ domain-swapped configuration of the 2G12 Fab dimer and its binding interface with the SARS-CoV-2 S protein were well-resolved, with the bound glycans clearly visible in the cryo-EM map. Deletion of glycan 709 reduced or eliminated 2G12 binding to the panel of FDG Abs tested (Figures 6D, 6E, and S6), thus demonstrating a common epitope for the FDG mAbs on the SARS-CoV-2 S protein. Additionally, we found that 2G12 also bound the SARS-CoV-2 and MERS S ectodomains (Figure S6).

Previous reports indicated that the 2G12 bnAb was not poly-reactive for binding recombinant human autoantigens (Haynes et al., 2005; Liu et al., 2015), but a recent study demonstrated that glycan-dependent HIV-1 Env bnAbs also bound host cells from HIV-1 uninfected individuals in a glycan-dependent manner (Blazkova et al., 2019). Similarly, we found that 12 of 23 (52%) human or macaque FDG Abs and 2G12 bnAb bound HEp2 cells but were either weakly or non-reactive with a panel of human autoantigens (Figure 7). The specificity of FDG mAbs for glycans suggested that HEp2 cell-FDG reactivity was glycan-mediated.

**DISCUSSION**

In this study, we identified a new category of prevalent glycan-reactive Abs that target diverse pathogens including HIV-1, SARS-CoV-2, and yeast. Although the Fab dimerized architectures of these glycan-reactive Abs bear resemblance to the unique glycan-reactive, V₄-domain-swapped HIV-1 bnAb 2G12, they differ from 2G12 in their diverse mechanisms of Fab dimerization that occur without V₄ domain swapping.

FDG Abs isolated prior to SHIV infection or HIV vaccination were mainly mutated IgM, thus raising the hypothesis for MZ or IgM-memory B cells being a source for these Abs. The B cell sources of natural Abs have been well characterized in the spleen and MZ of mice (Baumgarth, 2016; Lalor et al., 1989; Ichikawa et al., 2015), but the phenotype of the B cells that generate natural Abs in peripheral blood of humans are not fully established (Holodick et al., 2017; Griffin et al., 2011). Our data demonstrated that human and RM FDG B cells were predominately IgM"IgD"CD27", in agreement with circulating MZ B cells described in humans (Weller et al., 2004). FDG and 2G12 bnAbs have relatively short HCDR3 segments, target only glycans, and are more common than HIV-1 bnAbs with longer HCDR3s (Haynes and Verkoczy, 2014; Steichen et al., 2019). FDG Abs also used varied immunogenetics, thus indicative of a category of Abs, unlike previous classes of bnAB lineages that use more restricted heavy and light chain genes (Zhou et al., 2013; Bonsignori et al., 2018).

We previously showed that in a clonal bnAB lineage (termed DH270), early lineage members only neutralized HIV-1 bearing...
Man9-enriched Envs, whereas mature bnAbs neutralized autologous and heterologous HIV-1 bearing Envs with heterogeneous glycoforms (Bonsignori et al., 2017), thus demonstrating evolution of Abs within a lineage to achieve bnAb status through recognition of Envs with heterogeneous glycoforms. Similarly, although FDG precursors generally neutralized only HIV-1 bearing Envs with high mannose glycans, FDG bnAbs neutralized HIV-1 bearing Envs with heterogeneous glycoforms.

Our data also demonstrated that FDG and 2G12 mAbs had limited polyreactivity. It was postulated that the recognition of a unique glycan patch on HIV-1 Env was key for 2G12 maturation and neutralization potency (Scanlan et al., 2002). FDG Abs can also recognize the HIV-1 Env glycan patch, thus supporting the hypothesis for affinity maturation of FDG B cell lineages via HIV Env selection. Moreover, we demonstrated that avidity was important for glycan recognition by FDG Abs as vaccine-induced DH717 Fab had minimal Env-binding compared to DH717 IgG. We reported two examples where the Y-shaped forms of the Abs were less optimal for HIV-1 Env glycan recognition. First, 2G12 mutations that disrupted its Fab dimer resulted in loss of glycan binding (Huber et al., 2010). Second, Y-shaped DH851.4 IgM showed decreased binding and limited neutralization capacity compared to the Fab-dimerized DH851.1-DH851.3 IgG Abs within the same lineage. Thus, through Fab dimerization, FDG Abs acquired improved affinity and/or avidity for binding Env glycans, including the high mannose patch targeted by 2G12.

FDG mAbs bound a glycan epitope in the S2 subunit of the recombinant SARS-CoV-2 S protein. Consistent with a quaternary epitope, 2G12 binding was sensitive to SARS-CoV-2 S conformation with cold denaturation of the 2P S ectodomain resulting in reduced 2G12 binding (Edwards et al., 2021). Although these HIV-1 Env-directed FDG mAbs did not neutralize SARS-CoV-2 (not shown), our results suggest that FDG bnAbs may be matured to glycans on SARS-CoV-2 S. It remains unknown whether SARS-CoV-2 S-directed FDG mAbs have non-neutralizing protective effects.

The RM from which the FDG bnAb DH851 clonal lineage was isolated had relatively modest heterologous neutralizing plasma Abs that mapped to glycans in the high mannose patch as well as the peptide motif (GDIR) (Roark et al., 2021). Thus, it is likely that FDG bnAbs that target only glycans were minor components of the B cell repertoire of this macaque.

Other modalities of Fab-dimerization have been recently described, including head-to-head dimerization mediated by the CDR loops and intermolecular Fab dimers (Plath et al., 2016). These Abs have been shown to target non-HIV antigens,
**Figure 7. Polyreactivity profile of FDG Abs**

(A) Indirect immunofluorescence assay testing reactivity of FDG Abs in HEp-2 cells. Each mAb was tested at 50 and 25 μg/mL in duplicate reactions; a representative of the replicate data was reported. Positivity scores were determined relative to RM positive (DH1037) and negative (DH570.30) control mAbs. Staining patterns were identified using the Zeus Scientific pattern guide.

(B) Images of representative mAbs staining of HEp-2 cells. Ab reactivity pattern in HEP-2 cells were described in (A).

(C) Nine autoantigens were tested for reactivity by FDG Abs using a commercially available AtheNA Multi-Lyte ANA kit (Zeus Scientific). Serially diluted mAbs were tested for binding and the data analyzed using an AtheNA software. The dash lines represented the positivity score (121 units), which was consistent across independent experiments. 4E10 and CH65 represented positive and negative control mAbs, respectively. Data shown were for representative FDG mAbs that were reactive with HEp-2 cells (A and B). See also STAR Methods.
including CD20 (Rougé et al., 2020) and malaria parasite (Imkeller et al., 2018), although the interactions they mediate differ substantially from the glycan interactions with the FDG Abs we describe in this study, where intramolecular Fab dimerization within the Ab is a mechanism for enhancing weak glycan interactions via avidity. Mouse B cell lines bearing mature 2G12 exhibited domain-swapped forms of the BCR (Doores et al., 2013), but it remains unknown if the membrane-bound BCR of FDG bnAbs is analogous to BCR cross-linking that results in B cell activation (Yang and Reth, 2010a, 2010b).

It is important to note that FDG bnAbs with the capacity to neutralize autologous and heterologous difficult-to-neutralize HIV-1 bearing heterogeneous glycoforms in this report were only isolated from SHIV-infected RMs. SHIV-induced DH851 and DH1003 FDG bnAbs were class-switched, suggesting B cell maturation events occurred in SHIV infection that did not occur during precursor expansion with vaccination. We have previously shown that HIV-1-infected individuals who make bnAbs have multiple immunoregulatory abnormalities that create a permissive immunological environment for bnAb development (Moody et al., 2016; Bradley et al., 2018; Roskin et al., 2020). It will be of interest to determine if SHIV-infected RMs that make bnAbs have similar immunoregulatory abnormalities and whether FDG bnAbs can be found in chronic HIV-1-infected humans. Our data showed that we expanded FDG B cell lineages via vaccination, thus implying that FDG B cells were not tightly controlled by immune tolerance. Finally, that FDG Abs can bind multiple glycan clusters and use multiple BCRs, combined with FDG precursors being well represented in human B cell repertoire make this new category of glycan-reactive B cells an attractive target to consider for induction by HIV-1 vaccines. Since the submission of this paper, Lee et al. have reported that the FDG Ab 2G12 can neutralize strains of influenza (Lee et al., 2021), further demonstrating the breadth of cross-binding activity of the 2G12 FDG Ab for glycosylated virus pathogens.

Limitations of the study

We evaluated glycan-reactivity of FDG Abs on recombinant HIV-1 Env and SARS-CoV-2 S proteins, which may exclude different glycoforms only found on these surface proteins on the intact viruses in vivo. The frequency of FDG precursors was estimated in peripheral blood cells of nine HIV-1 naive individuals, thus future studies in a larger cohort of human B cells and other immune tissues will be informative about the prevalence of these Abs. We defined blood-derived FDG B cells as MZ-like, thus future studies of spleen-derived MZ B cells will verify this pool of B cells as a source of FDG Abs.

STAR METHODS

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at https://doi.org/10.1016/j.cell.2021.04.042.

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AUTHOR CONTRIBUTIONS


DECLARATION OF INTERESTS

B.A. and W.E.W. are co-founders of Chemitope Technologies, and G.M.L. is a founder of Avidea Technologies that now commercially produce peptides used in our HIV-1 vaccination regimen. The remaining authors declare no competing interests.

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REFERENCES


### Key Resources Table

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**Deposited data**

- DH717.1 Fab monomer in complex with man9 glycan: This paper; PDB: 6VTU
- Crystal structure of the disulfide linked DH717.1 Fab dimer, derived from a macaque HIV-1 vaccine-induced Env glycan-reactive neutralizing antibody B cell lineage: This paper; PDB: 6XRJ
- Cryo-EM structure of DH898.1 Fab-dimer from local refinement of the Fab-dimer bound near the CD4 binding site of HIV-1 Env CH848 SOSIP trimer: This paper; PDB: 7L6M; EMDB: EMD-23145
- Cryo-EM structure of CH8483.D0949.10.17chim.6R.DS.SOSIP.664: This paper; PDB: 7L6O; EMDB: EMD-23124
- Cryo-EM structure of DH883.1 bound to HIV-1 Env CH848 SOSIP trimer: This paper; PDB: 7LUA; EMDB: EMD-23519
- Cryo-EM structure of DH851.3 bound to HIV-1 CH505 Env: This paper; PDB: 7LU9; EMDB: EMD-23518
- Cryo-EM structure of SARS-CoV-2 2P S ectodomain bound domain-swapped antibody 2G12 from focused refinement: This paper; PDB: 7L09; EMDB: EMD-23097
- Cryo-EM structure of SARS-CoV-2 2P S ectodomain bound to two copies of domain-swapped antibody 2G12: This paper; PDB: 7L06; EMDB: EMD-23095
- Cryo-EM structure of SARS-CoV-2 2P S ectodomain bound to one copy of domain-swapped antibody 2G12: This paper; PDB: 7L02; EMDB: EMD-23094
- Cryo-electron microscopy reconstruction of antibody DH898.1 Fab-dimer bound to glycans 332, 392, and 396 of HIV Env CH848 10.17 SOSIP trimer: This paper; EMDB: EMD-23152
- Cryo-electron microscopy local refinement of antibody DH898.1 Fab-dimer bound to glycans 332, 392, and 396 of HIV Env CH848 10.17 SOSIP trimer: This paper; EMDB: EMD-23153
- Recombinant mAbs: This paper; MT470283–MT470354
- Gene expression data: This paper; PRJNA685659

**Experimental models: cell lines**

- HEK293T: ATCC; Cat #CRL3216

**Experimental models: organisms/strains**

- Candida albicans: This paper; SC5314
- Cryptococcus neoformans: This paper; H99

**Software and algorithms**

- Cloanalyst Ig gene database: Ramesh et al., 2017; https://www.bu.edu/computation/alimunology/research/software/
- Cell Ranger Single Cell Software Suite v3.1: 10X Genomics; Zheng et al., 2017a; N/A
- Surat r package v3.2.0: Macosko et al., 2015; Satija et al., 2015; Stuart et al., 2019; N/A
- Unicorn 7.0.2 software: GE Healthcare; N/A

**RESOURCE AVAILABILITY**

**Lead contact**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Wilton B. Williams (wilton.williams@mc.duke.edu).
Materials availability
The data presented in this manuscript, and research materials used in this study are available from Duke University upon request and subsequent execution of an appropriate materials transfer agreement.

Data and code availability
The variable heavy and light chain gene sequences for recombinant mAbs were deposited in GenBank and can be accessed using the following accession numbers; GenBank: MT470283–MT470354. The transcriptome sequencing data were deposited in NCBI Sequence Read Archive (SRA) under Bioproject: PRJNA685659. The atomic models and cryo-EM reconstructions and generated during this study are available at PDB and EMBD (https://www.rcsb.org; http://emsearch.rutgers.edu) under the accession codes PDB: 6VTU, 6XRJ, 7L6M, 7L6O, 7LUJ, 7L09, 7L06, and 7L02 and EMDB: EMD-23145, EMD-23124, EMD-23519, EMD-23518, EMD-23097, EMD-23095, EMD-23094, EMD-23152, EMD-23153.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Rhesus macaques (RMs)
All RMs were maintained in accordance with the Association for Assessment and Accreditation of Laboratory Animals. Research was conducted in compliance with the Animal Welfare Act and other federal statutes and regulations relating to animals and experiments involving animals and adheres to principles stated in the Guide for the Care and Use of Laboratory Animals, NRC Publication, 2011 edition.

Immunization of Indian origin RMs and blood draws were performed at (Bioqual Inc., Rockville, MD). The four animals in this study were as follows: RM5932 (male, 3.5 years old), RM5994 (male, ~4 years old), RM5995 (female, 5.25 years old), and RM5996 (female, 5.25 years old). DH717 lineage mAbs were isolated from RM5996. All vaccine doses were divided into equal volume and injected in the bilateral quadriceps muscles. The RMs were first immunized with a dose-escalation of Man9-V3 monomer (50 μg, 100 μg, and 500 μg) each formulated in 25 μg GLA-SE adjuvant. 50 μg of monomeric Man9-V3 was administered at weeks 0, 6, and 12. 100 μg of monomeric Man9-V3 was administered at weeks 18 and 24. 500 μg of monomeric Man9-V3 was administered at weeks 50 and 56. Dose escalation of monomeric Man9-V3 was then followed by 300 μg of Man9-V3 multimer (~6-mer) scaffolded onto N-(2-Hydroxypropyl) methacrylamide (HPMA) linear copolymer in 25 μg of GLA-SE given at week 71. Subsequently, RMs were immunized with a series of sequential soluble stabilized recombinant HIV-1 Env trimers (termed SOSIPs) (200 μg each) adjuvanted in 1 mg of Hiltonol Poly ICLC adjuvant (Oncovir). HIV-1 Env SOSIP trimers used for immunization were derived from CH848 Env strains previously identified as candidate immunogens for eliciting bnAbs that target glycans on the Env high mannose patch in conjunction with the Env peptide backbone (Bonsignori et al., 2017); 10.17 (week 103), 10.31 (week 109), 80.06 (week 115), 5.41 (week 127) and 25.02 (week 135). RMs were finally immunized with Star-Man9-V3 bearing the PADRE T cell helper epitope (300 μg) in 25 μg GLA-SE at week 143. Blood samples were collected two weeks post-immunization. The RM immunogenicity study in this paper was designed to evaluate the immunogenicity and dose of Man9-V3 monomer and to determine the effect of multimerization of Man9-V3 and later full envelope trimers, as boosts. The rational for the doses used were based on our previous experience with peptide immunogens (Zhang et al., 2016). Thus, we designed a dose escalation immunization study with the Man9-V3 monomer to study its immunogenicity and determine whether increased valency by multimerization improved the immunogenicity of the glycopeptide. Man9-V3 was previously used as a bait to isolate an HIV-1 Env bnAb that targeted N332 glycan in the high mannosone patch in conjunction with the GDIR motif in the Env peptide backbone (Alam et al., 2017), both of which comprise the V3 glycan bnAb epitope. Thus, Man9-V3 was postulated to be a possible prime for bnAb precursors that target Env-glycan containing epitopes, including V3 glycan bnAbs; and the 5-valent soluble recombinant HIV-1 Env trimers were subsequently postulated to be candidate immunogens to boost these bnAb precursor B cells (Bonsignori et al., 2017). Hence, our approach to prime with Man9-V3 and boost with multimerized Man9-V3 and then the Env trimers. However, our studies showed that Man9-V3 and one of the trimers (CH848 10.17) were able to select FDA B cells rather than V3-glycan bnAbs, and are thus candidate immunogens for future vaccine strategies to elicit FDA Abs.

SHIV infection of Indian RMs were performed at (Bioqual Inc., Rockville, MD) and previously described (Roark et al., 2021). HIV-1 transmitted-founder (TF) Envs were derived from viruses that established infection in humans, thus TF Env-bearing SHIVs (Li et al., 2016) provided a novel model for studying HIV-1 infection as described (Roark et al., 2021). Blood and plasma samples were collected for binding and neutralization assays. HIV/SHIV bearing CH848TF Env was referred to as an autologous virus strain. DH851 and DH898 mAbs were isolated from a single macaque (RM6163; female, ~5 years old) that was infected with a SHIV bearing HIV-1 CH848TF Env strain, and DH1003 mAbs were isolated from a different macaque (RM10N011; male, ~8 years old) that was infected with a SHIV bearing HIV-1 BG505.T332N Env strain.

HIV-1 naive humans
Our cohort of nine HIV-1 seronegative adults aged 18–48 years (mix of both males and females) were bled for leukapheresis PBMCs in the summer of 2019 or earlier, thus it is unlikely that they were COVID-19 convalescent subjects. From our observation that FDG mAbs bound glycans on SARS-CoV-2 spike, we hypothesized that FDG Abs can be elicited by SARS-CoV-2 spike high mannosone glycans and this is now currently being actively explored in COVID-19 convalescent subjects.
Candida albicans (SC5314) and Cryptococcus neoformans (H99) preparation

A single colony of yeast was inoculated into 250 mL YPD broth and grown for 2 days at 30°C on a shaking incubator set at 225 rpm. The cells were harvested and washed thrice with PBS. The pellets were resuspended in 11 mL PBS and the cell concentration was determined (C. albicans, 3.40x10^9 CFUs/mL and C. neoformans, 1.62x10^9 CFUs/mL, respectively). To heat-kill the yeast, the cells were incubated for 24h at 60°C in a water bath. Three hundred microliters of the heat-killed yeast were plated onto YPD agar, incubated for 7 days to ensure no viable yeast were present.

**METHOD DETAILS**

**Synthesis of polymer-based glycopeptide (Man₉-V₃) constructs**

The Man₉-V₃ monomer comprising a 30 amino acid V₃-glycopeptide with high-mannose (Man₉GlcNAc₂) glycans at position 301 and 332 based on the clade B JRFL mini-V₃ construct (Pejchal et al., 2011) was synthesized as previously described (Alam et al., 2017). Man₉-V₃ multimer was synthesized by reacting multiple copies of a Man₉-V₃ monomer with a multivalent amino group-reactive HPMA-based co-polymer that was prepared as previously described (Lynn et al., 2015). The concept, design and synthesis of Man₉ multimerized on star-shaped polymers (Star-Man₉-V₃), composed of HPMA-based polymer arms radially extending from a poly(amidoamine) (PAMAM) dendrimer core, with the universal PADRE T helper epitope (Alexander et al., 2000) has been described (Francica et al., 2019). In this study, Man₉-V₃ multimer and Star-Man₉-V₃ (~50nm construct containing multiple Man₉-V₃ units) were used as macromolecular nanosized forms of Man₉-V₃ monomer for immunization of NHPs.

**Expression of recombinant HIV-1 Env proteins**

Production of recombinant HIV-1 Envs, including gp120 monomers, and unstabilized gp140 and stabilized SOSIP trimers was previously described (Francica et al., 2019; Saunders et al., 2017a, 2017b, 2019). Recombinant Envs were used for binding assays and NHP immunizations. The main Env strains for soluble stabilized recombinant HIV-1 Env trimers (SOSIPs) used in this study were previously described (Bonsignori et al., 2017; Saunders et al., 2017a) and the complete nomenclatures are shown below: CH848.3.TFchim.6R.SOSIP664v4.1; CH848.3.D0949.10.17chim.6R.DS.SOSIP664; CH848.3.D0949.10.17chim.6R.DS.SOSIP664_N301A; CH848.3.D0949.10.17chim.6R.DS.SOSIP664_N301AN332A; CH848.3.D0949.10.17chim.6R.DS.SOSIP664V4.1_N133DN138T; CH848.3.D0949.10.17chim.6R.DS.SOSIP664V4.1_N133DN138T; and CH848.3.D0949.10.17chim.6R.DS.SOSIP664v4.1degly4.

We tested wild-type and mutant (N301A, N301AN332A, N133DN138T and degly4) Env trimers listed above to evaluate mAb binding. For HIV-1 Env trimimer immunizations in macaques, the following HIV-1 Env CH848 strains were generated as soluble stabilized recombinant trimers (SOSIPs): CH848 10.17, CH848 10.31, CH848 80.06, CH848 5.41 and CH848 25.02. These trimers constitute a sequential 5-valent series of CH848 Env stabilized trimers that had evolved during human CH848TF infection and were predicted to be involved in the induction of HIV-1 Env bnAbs that target glycans in the high mannose in conjunction with the GDIR motif in the Env sequence. For HIV-1 Env trimer immunizations in macaques, the following HIV-1 Env CH848 strains were generated as soluble stabilized Envs: CH848.3.TFchim.6R.SOSIP664v4.1; CH848.3.D0949.10.17chim.6R.DS.SOSIP664; CH848.3.D0949.10.17chim.6R.DS.SOSIP664_N301A; CH848.3.D0949.10.17chim.6R.DS.SOSIP664_N301AN332A; CH848.3.D0949.10.17chim.6R.DS.SOSIP664V4.1_N133DN138T; CH848.3.D0949.10.17chim.6R.DS.SOSIP664V4.1_N133DN138T; and CH848.3.D0949.10.17chim.6R.DS.SOSIP664v4.1degly4.

**Expression of recombinant Coronavirus spikes**

The SARS-CoV-2, SARS and MERS ectodomain constructs were produced and purified as previously described (Wrapp et al., 2020). Briefly, genes encoding residues 1–1208 of SARS-CoV-2 S (GenBank: MN908947), residues 1–1291 for MERS S (GenBank KJ650098.1) and residues 1–1190 for SARS S (UniProt P59594–1) with proline substitutions at residues 986 and 987 (SARS-CoV-2), 1060–1061 (MERS) and 968–969 (SARS) and a “GSAS” substitution at the furin cleavage site (residues 682–685) for SARS-CoV-2 was synthesized and cloned into the mammalian expression vector p2xH. In all constructs, a C-terminal T4 fibrin trimerization motif, an HRV3C protease cleavage site, a TwinStrepTag and an 8XHisTag were also included. All other mutants were introduced in these backgrounds. Expression plasmids encoding the ectodomains sequences were used to transiently transfect FreeStyle293F cells using Turbo293 (SpeedBioSystems). Protein were purified on the sixth day post transfection from the filtered supernatant using StrepTactin resin (IBA). Affinity purified protein was run over a Superose 6/300 increase column. The purified proteins were flash-frozen in liquid nitrogen and stored at −80°C.

**Isolation of Abs from single B cells**

Antigen-specific memory B cells were isolated from RMs or humans using fluorophore-labeled antigens as hooks in various sort strategies outlined below. Antigens used for sorts were conjugated with Alexa Fluor 647 (AF647), Brilliant Violet 421 (BV421), or VioBright 515 (VB515) fluorophores. With the exception of peptides, immune cells were concurrently incubated with a cocktail of staining Abs.
and fluorophore-labeled antigens. Fluorophore-labeled peptides required an incubation with immune cells prior to addition of the staining antibody cocktail to improve binding efficiency. Fluorophore-labeled peptides were used at a ratio of 2.5 µg per 10 million cells, recombinant gp120 and gp140 proteins at 10 µg per 10 million cells, and SOSIP trimers at 90 pmol per 10 million cells, to stain antigen-specific memory B cells. All incubations took place in approximately 1 mL of buffer per 10 million cells. Flow cytometry sorts were performed using a BD FACSaria II (BD Biosciences, San Jose, CA), and the data were analyzed using FlowJo (Treestar, Ashland, OR). Single cell Env-reactive Abs were CD3 (BD Cat#552852; 5 µl per test)/ CD14 (BioLegend Cat#301832; 2.5 µl per test)/ CD16 (BD Cat#557744; 5 µl per test), and/or surface IgD (Southern Biotech Cat#2030-09; 1 µl per test) negative. Macaque B cells were also CD27+/− (BioLegend Cat#302816; 5 µl per test) and CD20 (BioLegend Cat#302336; 1 µl per test) positive, while human B cells were CD38+/− (Beckman Coulter Cat#IM2371U; 1 µl per test) and CD19 (BD Cat#557791; 1 µl per test) positive.

Single sorted B cells were obtained from RMs and humans using previously described flow cytometry-based sorts (Williams et al., 2017b; Alam et al., 2017; Zhang et al., 2016). Cells were individually sorted into 96 well plates containing lysis buffer and immediately stored at −80, as previously described (Liao et al., 2009). Rhesus macaque and human V(1)D(8)i and V(1)J(1) segments were isolated by single-cell PCR approaches that were previously described (Liao et al., 2009; Alam et al., 2017). Antibody sequences were analyzed using a custom-built bioinformatics pipeline for base-calling, contig assembly, quality trimming, immunogenetic annotation with Clonealyst (https://www.bu.edu/computationalimmunology/research/software/), VDJ sequence quality filtering, functionality assessment, and isotyping. Antigen-specific single B cells were indexed sorted, thus allowing us to perform a backgate analysis to identify individual IgM+IgD+CD27+ cells overlaid against the total sorted B cell populations in FlowJo (Treestar, Ashland, OR). In vaccinated RMs, Env-reactive B cells were sorted using soluble unstimulated recombinant HIV-1 Env trimer (HIV-1 JRFL gp140) bearing Envs with Manα2–6Gal antigen (Kifunensine-treated Env) conjugated to both AF647 and BV421 fluorophores. The antigens for sorts of SHIV-infected B cells used tetramerized Manα2–6Gal and HIV-1 Env SOSIPs. The following flow cytometry sort strategies were used to investigate the repertoire of SHIV-elicited Env-reactive B cells, including glycan-reactive B cells: LN wk52 [Manα2–6Gal V3 peptide (+) and Aglycone V3 peptide (−)]; CH848TF gp120 (+) and CH848TF gp120 N332A (−); and CH848TF SOSIP4.1 (+)]. PBMCs wk52 [CH848TF gp120 (+) and CH848TF gp120 N332A (−)]; and PBMCs wk104 [Manα2–6Gal V3 peptide (+) and Aglycone V3 peptide (−)]; CH848TF SOSIP4.1 (+)]; CH848.d949.10.17 DS.SOSIP (+) and CH848.d949.10.17 DS.SOSIP N332A (−); and CH848.d949.10.17 DS.SOSIP (+) and CH848TF SOSIP4.1 (−)].

In HIV-1 naive humans, Env-reactive B cells were sorted using glycopeptides and HIV-1 Env SOSIPs in the following differential sort strategies: (#1) Manα2–6Gal V3 (−) versus Aglycone V3 peptide (+) in a dual or triple color sort strategy; and (#2) CH848 10.17 DS.SOSIP (+) versus CH848 10.17 DS.SOSIP N332A (−); and (#3) CH848 10.17 DS.SOSIP N133D138T (+) versus CH848 10.17 DS.SOSIP_N133D138T_N332A (−). In three individuals (LPK01-LPK03), we sorted 20M PBMCs using strategy #1 (dual color differential sort) or strategy #2 (total for 2 sorts = 40 million PBMCs). For dual color differential sort strategy #1, we sorted B cells that were reactive with Manα2–6Gal V3 conjugated to one fluorophore, but not the aglycone V3 peptide conjugated to a different fluorophore. In one individual (LPK04), we studied B cells enriched from 100 million leukapheresis PBMCs per experiment using sort strategy #3 (total for 2 sorts = 200 million PBMCs). In five individuals (LPK05-LPK09), we studied B cells enriched from 100 million leukapheresis PBMCs using sort strategy #1 (triple color differential sort); here we used a more stringent approach to sort B cells that were reactive with Manα2–6Gal V3 conjugated to two different fluorophores, but not reactive with aglycone V3 peptide conjugated to a third fluorophore. For subjects LPK04-LPK09, we performed negative B cell enrichment from 100M PBMCs in each experiment using a commercially available B cell enrichment kit according to manufacturer’s protocol (StemCell Technologies; Vancouver, BC, Canada). We calculated the frequency of human FDG B cells as the percent of all sorted B cell-derived recombinant IgGs produced in small-scale transfection that bound Manα2–6Gal V3 but not aglycone V3 peptide and/or yeast antigens in ELISA, relative to the total number of B cells analyzed via flow cytometry sort per individual. The estimated frequency of human FDG B cells was reported as the average frequency of FDG B cells across the nine individuals studied.

Expression of recombinant mAbs
Recombinant mAbs were first generated in human embryonic kidney epithelial cell lines (HEK293T, ATCC, Manassas, VA; Cat#CRL3216) in small scale transfections using linear cassettes encoding antibody heavy and light chain genes that were PCR amplified from single B cells (Liao et al., 2009). For generation of larger quantities commercially-obtained (GeneScript, Piscataway, NJ) plasmids with antibody heavy and light chain genes were used to transfact suspension Expi293i cells using ExpiFectamine 293 transfection reagents (Life Technologies, Gibco; Cat#A14524) as described (Bonsignori et al., 2017). Purified recombinant mAbs were dialyzed against PBS, analyzed, and stored at 4°C. All recombinant mAbs were expressed from plasmids encoding a human or macaque IgG constant region, and were QC’ed in western blot for appropriate heavy and light chain protein expression and/or via size exclusion chromatography (SEC) for a single peak indicative of stable antibody expression.

Expression of DH717 Fabs for structural studies
To purify DH717.1 Fab monomer for crystallization studies, the heavy- and light-chain variable and constant domains were cloned into a modified pVRC-8400 expression vector using NheI and NotI restriction sites and the tissue plasminogen activation signal sequence. The C terminus of the heavy-chain constructs contained a non-cleavable 6x-histidine tag. Fabs were expressed using transient transfection of HEK293T cells using linear polyethenimine (PEI) following manufacturer’s protocols. After 5 days of expression, supernatants were clarified by centrifugation. His-tagged Fabs were loaded onto Ni-NTA superflow resin (Qiagen) equilibrated with Buffer
A (10 mM Tris, pH 7.5, 100 mM NaCl), washed with Buffer A + 350 mM imidazole, and eluted with Buffer A + 350 mM imidazole. FabS were then purified by gel filtration chromatography in Buffer A using a superdex 200 analytical column. The fractions corresponding to monomer were pooled and concentrated. While intermolecular disulfide bonds can occur due to purification artifacts, intramolecular disulfide are likely to be formed in vivo because of the structural proximity of two unpaired cysteines, such as what we see in the DH717.1 Fab dimers.

**HIV-1 Env and glycopeptide binding of mAbs via ELISA**

Envelope binding to recombinant mAbs, and plasma or sera were tested in ELISA as previously described (Williams et al., 2017a; Saunders et al., 2019). Binding was assessed using synthetic peptides, recombinant HIV-1 Env gp120, gp140, and stabilized chimeric SOSIP trimers, and heat-killed yeast antigens. ELISA protocols had modifications that supported binding to different antigens. In general, antigens were directly coated to Nunc-absorb (ThermoFisher) plates overnight at 4°C or captured using streptavidin, AbC-mAb (AVIDITY, Colorado, USA) or PGT151 mAb that were coated to Nunc-absorb plates overnight at 4°C. Unbound proteins were washed away and the plates were blocked with either goat serum-based (SuperBlock) (Liao et al., 2009) or non-serum-based (Alam et al., 2017) blocking media for 1 hour. Serial dilution of serum and mAbs were added to the plate for 60 and 90 minutes, respectively. Binding Abs were detected with species-specific HRP-labeled anti-IgG Fc Abs using 20μl per reaction with 1 hour incubation at room temperature. HRP detection was subsequently quantified with 3,3’5,5’-tetramethylbenzidine (TMB). ELISA binding levels were measured at an optical density of 450 nm (OD450nm) and binding titers were analyzed as area-under-the-curve of the log-transformed concentrations (Log AUC).

Binding to biotin-Man9-V3 and biotin-aglycone-V3 was determined as previously described using streptavidin for capturing each peptide on Nunc-absorb ELISA plates (Alam et al., 2017; Bonsignori et al., 2017). In brief, we coated 30ng of streptavidin in 15μl at 2μg/ml per well of a 384-well Nunc-absorb ELISA plate, sealed and incubated overnight at 4°C. Synthetic peptides (10ng) were added to streptavidin in 10μl at 1μg/ml for one hour at room temperature to facilitate peptide capture. Binding to yeast antigens was tested at 1:400 (Cryptococcus neoformans) and 1:2000 (Candida albicans) dilutions made in sodium bicarbonate buffer; 15μl of each yeast solution was coated per well of a 384-well Nunc-absorb ELISA plate, sealed and incubated overnight at 4°C. Yeast antigen reactivity was performed using ELISA conditions for assessing glycan reactivity on glycopeptides (Alam et al., 2017), HRP-conjugated specie-specific secondary Abs were used to detect antibody binding to synthetic peptides and yeast antigens - Goat anti-human IgG-HRP, 1:15000 dilution (Jackson ImmunoResearch Laboratories, Cat# 109-035-098); and Mouse anti-monkey IgG-HRP, 1:20000 dilution (Southern Biotech, Cat# 4700-05). The yeast dilutions tested in ELISA were found to be the lowest dilutions at which we observed maximal binding by glycan-reactive Abs, such as 2G12 in optimization assays. DH1013 isolated from RM5996 was used as a positive control mAb for peptide binding to Man9/V3 and Aglycone V3 peptides, and 2G12 or DH501 (Saunders et al., 2017a) were used as positive control mAbs for binding Man9/V3, but not Aglycone V3 peptide. 2G12 bnAb was also used as a reference mAb for Env glycan and Candida albicans yeast antigen binding (Doores et al., 2010b).

For ELISAs with HIV-1 Env SOSIPs bearing an Avi-tag, the trimer was captured by the AbC-mAb antibody against C terminus Avi-tag, while non Avi-tagged SOSIP trimers were captured by PGT151 bnAb expressed in 293 cells as described (Saunders et al., 2019; Williams et al., 2017d). In brief, Avi-tag and PGT151 mAbs were coated at 30ng per well of a 384-well plate in 15μl at 2μg/ml. HIV-1 Env SOSIPs (20ng) were added to Avi-tag or PGT mAbs in 10μl at 2μg/ml for one hour at room temperature to facilitate SOSIP capture. PGT151 mAb was used as a positive control for binding Avi-tagged SOSIP, while influenza-specific antibody CH65 (Whittle et al., 2011) was used as a negative control. HRP-conjugated specie-specific secondary Abs were used to detect binding Abs (see above). For non-Avi-tagged SOSIPs that were captured by PGT151 mAb, biotinylated mAbs were used as positive (B-2G12 and B-PGT128) and negative (B-CH65) controls - biotinylated-mAbs were tested at 10μg/ml and 3-fold dilutions. This latter approach was used to test only rhesus mAbs and b-mAbs, since the secondary Abs would not detect human PGT151 mAb used for SOSIP capture; Mouse anti-monkey IgG-HRP (see above) and Streptavidin-HRP, 1:30000 dilution (Thermo Scientific, Ref# 21130). FDG mAbs, DH100-5D1011, bound Man9-V3 but not aglycone V3 peptide in ELISA (Figure 3). Additionally, DH100-5D1011 had varied binding levels to Env SOSIP trimers (not shown). Competitive ELISA to assess cross-blocking of recombinant mAbs were previously described (Alam et al., 2013; Saunders et al., 2017a; Bonsignori et al., 2017). We biotinylated the Abs using the following product: BIOTIN-X-NHS, Cayman Chemicals, Cat# 13316. Here, we studied biotinylated 2G12 and DH851 mAbs in competition ELISAs using CH848TF or CH848 10.17 gp120 Env s.

Competitive inhibition of biotinylated-mAbs was measured as a percent of binding in the presence of a competing non-biotinylated mAb relative to binding in the absence of this competing mAb. A successful assay had positive control blocking ≥ 40%. DH851 blocking on CH848TF gp120 was from a single experiment (Figure 5), but the results were similarly observed for blocking on CH848 10.17 gp120 in three separate experiments (not shown).

**Glycan-dependent binding of FDG mAbs to SARS-CoV-2 spike protein**

Recombinant FDG mAbs were tested for binding SARS-CoV-2 trimer (lot 62KJ) in ELISA in the absence or presence of single monomer D-mannose sugar. SARS-CoV-2 trimeric spike protein was captured by streptavidin to individual wells of a 384-well plate, and serially diluted mAbs were tested for binding. ELISA binding assays were optimized to use SARS-CoV-2 spike protein that was aliquotted and frozen upon production; desired aliquots of frozen spike protein were thawed once at 37°C and stored overnight at room temperature. SARS-CoV-2 spike protein, and Manα-V3 and Aglycone V3 peptides were captured via streptavidin on Nunc-absorb
ELISA plates using PBS-based buffers and assay conditions as previously described (Alam et al., 2017; Bonsignori et al., 2017). HIV-1 CH505TF SOSIP trimer and commercially-obtained constructs of SARS-CoV-2 spike ectodomain (S1+S2 ECD, S2 ECD and RBD) (Sino Biological Inc Cat#40589-V08B1 and 40590-V08B respectively and RBD from Genescript Cat#Z03483) were captured using mouse anti-AVI-tag mAb (Avidity LLC, Aurora, CO). In brief, we coated 30 ng of streptavidin in 15 μl at 2 μg/ml or 30 ng of mouse anti-AVI-tag mAb in 15 μl at 2 μg/ml per well of a 384-well Nunc-absorb ELISA plate, sealed and incubated overnight at 4°C. SARS-CoV-2 spike protein (2 μg/ml) were added to streptavidin in 10 μl per well of a 384-well plate for one hour at room temperature to facilitate protein capture. HIV-1 CH505TF SOSIP, and recombinant SARS-CoV-2 S1+S2 ECD, S2 ECD and RBD spike proteins, at final concentration of 2 μg/ml were added to anti-AVI mAb in 10 μl per well of a 384-well plate for one hour at room temperature to facilitate protein capture. Mouse anti-monkey IgG-HRP (Southern Biotech, Cat# 4700-05) or Goat anti-human IgG-HRP (Jackson ImmunoResearch Laboratories, Cat# 109-035-098) secondary Abs were used to detect mAb bound to the SARS-CoV-2 spike protein. HRP detection was subsequently quantified with 3,3’5,5’-tetramethylbenzidine (TMB) by measuring binding levels at an absorbance of 450 nm; binding titers were reported as Log area under the curve (AUC).

Commercially obtained D-mannose (Sigma, St. Louis, MO) was used to outcompete mAb binding to glycans on SARS-CoV-2; D-mannose solutions were produced in ELISA buffers. Competition ELISAs with D-mannose were performed in two separate assays; each with [0.5M] (data not shown) or [1.0M] D-mannose. Mouse-Human chimeric mAb D001 (SARS-CoV-RBD mAb; Sino Biological Inc Cat#40150-D001) was used as a control mAb; D001 was tested at 2 μg/ml and 2-fold serial dilutions (10x), in contrast to all other mAbs tested at 100 μg/ml and 3-fold serial dilutions (10x). Anti-influenza CH65 mAb was used a negative control mAb. We found that goat serum containing buffer (Superblock) – PBS, 4% (w/v) whey protein (BiPro USA), 15% normal goat serum (Invitrogen), 0.5% Tween 20, and 0.05% sodium azide (Sigma-Aldrich) – inhibited binding of glycan-dependent FDG mAbs to SARS-CoV-2 spike protein, whereas positive control RBD-binding mAb D001 bound the spike protein in the presence of superblock. ELISAs using superblock were previously described (Williams et al., 2017a). Moreover, whereas the 2G12 from Polymun bound to S protein in surface plasmon resonance, it was less potent than the recombinantly expressed 2G12 Abs, and did not bind well in ELISA; commercial 2G12 from Polymun was provided in a maltose (mannose) buffer. D-mannose more competitively inhibited binding to fungal glycans by some FDG Abs. For example, SHIV-elicited DH898 mAbs were not outcompeted for binding as much as vaccine-induced DH717 mAbs. These data suggested that FDG Abs have differences in affinities for free-glycans in solution compared to glycan clusters on HIV-1 Env or the promiscuity in binding different glycoforms. In fact, structural analysis of DH898 mAbs in complex with recombinant mAbs. These data suggested that DH898 mAbs have a higher propensity to bind different glycan arrangements on HIV-1 Env and not only the high mannose patch where the vaccine-induced DH717 Abs, SHIV-induced DH851 bnAbs and 2G12 have been shown to bind.

Neutralization assays
mAb neutralizing activity was assessed in TZM-bl cells as described (Montefiori et al., 2012; LaBranche et al., 2019; Montefiori, 2009). Env-pseudotyped HIV-1 bearing Envs used in SHIVs for infection were referred to as autologous viruses, whereas viruses bearing Envs from global panels of geographically diverse multi-clade strains were referred to as heterologous viruses. Difficult-to-neutralize or tier 2 autologous or heterologous HIV-1 isolates are generally used to assess neutralization potency and breadth of HIV-1 Abs; global panel of HIV-1 strains (deCamp et al., 2014) and 119 heterologous tier 2 isolates (Seaman et al., 2010). Env-pseudotyped HIV-1 bearing Envs with Man9-enriched glycans (termed Kif-treated HIV-1 strains) were grown in cells treated with 25 μM Kifunensine to facilitate high mannose glycan expression on HIV-1 Envs and Kif-treated HIV-1 strains were tested for neutralization in TZM-bl cells as previously described (Saunders et al., 2017a). Env-pseudotyped HIV bearing Envs with Man9-enriched glycans express Env expressed in cell lines, but it remains unknown how additional glycosyltransferases present on the Env generated in vivo might impact the binding specificities and function of our FDG mAbs that preferentially target Man9-glycans. For neutralization assays, a mixture of CH01+CH31 bnAbs is used as a positive control for neutralization of all HIV-1 strains, and murine leukemia virus (MLV) or vesicular stomatitis virus (VSV) were used as negative retrovirus controls. For neutralization assays, a positive for neutralizing antibody activity in a sample is based on the criterion of > 3X the observed background against the MLV negative control pseudovirus. Note that DH898 mAbs were tested for neutralization of all tier 2 HIV-1 strains in the global panel, except CNE8. Additionally, we found neutralization titer variabilities in DH851 mAbs across different batches of mAbs, in agreement with the unstable dimerization of Fab5 for these Abs. Only DH1005 neutralized [Kif]-treated HIV strains (Figure 3), but none of the human FDG mAbs (DH1005-DH1010) neutralized non-[Kif]-treated HIV-1 strains (not shown).

Glycan binding of Abs oligomannose bead immunoassay
Recombinant mAbs were tested for binding glycans in an oligomannose bead immunoassay as described (Saunders et al., 2017a). Custom glycan microspheres were generated with different individual glycans to evaluate mAb binding. Binding of mAb to glycan was determined with A Bio-Plex 200 HTS (Bio-Rad) machine with the Bioplex manager software (Bio-Rad) was used to quantify binding of mAb to glycan, and binding was measured as background-subtracted fluorescence.

Anti-nuclear Ab (ANA) reactivity
mAb reactivity to nine autoantigens was measured using the AtheNA Multi-Lyte ANA kit (Zeus scientific, Cat#A21101). MAbs were 2-fold serially diluted starting at 50 mcg/mL. The secondary antibody provided with the kit was Phycoerythrin-conjugated goat anti-human IgG.
The kit SOP was followed for the remainder of the assay. Samples were analyzed using AtheNA software. Positive (+) specimens received a score > 120, and negative (-) specimens received a score < 100. Samples that score 100-120 were considered indeterminate. FDG mAbs that had ANA reactivity were shown in Figure 7. The following mAbs were negative for autoantigen binding (not shown): DH851.2, DH717.1, DH717.3, DH1005-DH1008, and DH1010-DH1011.

**HEp-2 cell staining**

Indirect immunofluorescence binding of mAbs to human epithelial (HEp-2) cells (ZEUS IFA ANA HEp-2 Test System, Zeus Scientific, Somerville, NJ) was performed per manufacturer’s instructions. Briefly, 20 μL of diluted mAbs (50 μg/mL, and 25 μg/mL) were added to the appropriate wells on the antinuclear antibody (ANA) slide. Slides were incubated for 20 minutes at room temperature in a humid chamber, and then washed with 1X PBS. Goat Anti-Rhesus Ig(H+L) FITC secondary was added at a concentration of 30 μg/mL to each well. The slides were incubated for 20 minutes, then washed twice, dried, fixed with 33% glycerol and coverslipped. Slides were imaged using an Olympus AX70 microscope with a SpotFlex FX1520 camera. Images were acquired on a 40X objective using the FITC fluorescence channel with different acquisition time as indicated on each image. Positivity was determined by comparing Abs of interest to positive and negative non-human primate antibody controls, DH1037 and DH570.30 respectively. Staining patterns were identified using the Zeus Scientific pattern guide found on the website and previously reported (Buchner et al., 2014; Huang et al., 2012; Haynes et al., 2005).

**Next-generation sequencing (NGS) and analysis of Ab genes**

Illumina MiSeq sequencing of Ab heavy chain VDJ sequences was performed on peripheral blood cells as previously described (Zhang et al., 2016; Williams et al., 2017b). Briefly, for each time point, RNA from peripheral cells was divided equally into two separate portions that were used to generate independent cDNA aliquots for sequencing to confirm the absence or presence of Ab sequences of interest. Paired-end sequences were merged using FLASH (Magoc and Salzberg, 2011), quality filtered (Q score > 30 for > 95% of sequence) and deduplicated. V, D, and J gene segment, clonal relatedness testing and reconstruction of clonal lineage trees were performed using the Cloanalyst software package (Kepler, 2013). Immunogenetics information of rhesus and human antibody sequences were assigned by Cloanalyst using Cloanalyst’s default libraries of rhesus and human immunoglobulin genes, respectively (https://www.bu.edu/computationalimmunology/research/software/). B cell clonality was determined based on similar heavy chain rearrangements and CDR3 length as described (Kepler et al., 2014). For Abs that were inferred as being clonally-related by Cloanalyst, and had the same Vκ sequence but different alleles, we manually inspected the Vκ sequence for similar residues in the HCDR3, including the non-templated nucleotides.

The near-germline or unmutated common ancestor (UCA) of Ab lineages were inferred using the Cloanalyst software program with the macaque immunoglobulin gene library. DH851UCA was initially inferred based on the isolation of DH851.1-DH851.3 lineage members. DH851UCA.2 was later inferred using DH851.1-DH851.4. We found that DH851UCA and DH851UCA.2 had one amino acid difference in a framework region of both the heavy and light chain genes; the UCA sequences were deposited in GenBank. Data shown in this study described the characteristics of DH851UCA. For recombinant Ab expression, week 0 mAbs had clonally-related heavy chain genes detected via NGS paired with DH851.1 light chain, while the DH851UCA had the Cloanalyst-inferred heavy and light chain genes.

To exclude the possibility of generating unmutated common ancestor Abs that do not account for the allelic variability observed in the RM model (Ramesh et al., 2017; Corcoran et al., 2016), we generated a NGS dataset as described above that comprised only of VH1 (DH898 lineage), VH2 (DH851 lineage) and VH4 (DH717 lineage) IgM+ repertoires at baseline or week 0 of the RMIs. Here, we sought to find unmutated or least mutated FDG Ab clonal lineage members indicated from macaque baseline PBMCs. Week 0 clonally-related mAbs were generated by pairing the week 0 NGS-derived heavy chain genes with the light chain of the single B cell-derived Ab isolated following infection or vaccination as described (Williams et al., 2015). Clonally-related DH717 Vκ genes identified via Vκ gene deep sequencing post the Star-Manα-V3 boost had higher average mutation frequencies compared to DH717 Vκ genes identified after the initial trimer boost. However, our attempts to express recombinant mAbs bearing the mutated DH717 Vκ genes paired with the DH717.1 Vκ failed, so it remains unknown whether these mutations improved antibody function.

Week 0 DH898 and DH851 FDG lineage precursors that existed prior to SHIV CH848TF infection in macaque RM6163 were evaluated using NGS in two separate experiments. cDNA of isotype-specific VH1 and VH2 genes were amplified from peripheral blood-extracted RNA and split into separate samples for cDNA library preparation and NGS (Run A and B). Sequences were quality filtered and computationally analyzed, and DH898 and DH8511 clonal lineages assessed using Cloanalyst software program. We found 2 IgG and 594 IgM DH898 clonal lineage members among ~3.5 million unique Vκ1 sequences at week 0: IgA (151760), IgG (860757), and IgM (2457675). We found 21 IgA, 66 IgG and 148 IgM DH851 clonal lineage members among ~2.1 million unique Vκ2 sequences at week 0: IgA (516938), IgG (1282669), and IgM (759778).

**High-resolution single cell VDJ and transcriptome sequencing and analysis**

RM VDJ libraries were generated following the manufacturers protocol (Chromium Next GEM Single Cell V(D)J protocol (v1.1, Rev E)) with three modifications. First, 15 μL of cDNA was used as input for the first BCR enrichment (step 4.1a). Second, RM-specific reverse primers were used to amplify BCR transcripts (step 4.1b and 4.3a). Third, we used 9 PCR cycles in the first BCR enrichment (step 4.1) and 11 PCR cycles for the second BCR enrichment (step 4.3). Primers were designed to amplify the BCR sequences from RMs using...
Surface Plasmon Resonance (SPr)

The binding of DH717.1 and DH717.1 C76S mAbs to 5 different CH848 Env’s was assessed by surface plasmon resonance on Bia-Ultima-X (GE Healthcare) in HBS-N 1X running buffer. Biotinylated trimer solutions were immobilized to a level of 300-350 RU via streptavidin. Second round primers (5′ to 3′) were as follows for IgA, IgG, IgE, IgM, IgD, IgK, and IgL: [IgA] GGGGGGGAGTTATGAGCCT, [IgG] TTTGCACCTGTGGTTGCT, [IgE] AGTGAGTGGTACACCACAT, [IgM] CACTGACACTCTCTGGAACT, [IgD] ATGGGAGGTGACACGCT, [IgK] GTGAGTCGCTGTTGACACT, and [IgL] TGACCTGGGAGCT GTGAGCT. The SPR binding curves of 2G12 and DH717.1 mAbs and Fabs against CH848 10.17 DS.SOSIP and CH848 10.17 DS.SO-SIP were measured using a NanoDrop UV-VIS spectrophotometer. Following SEC purification, DH717.1 Fab monomer and dimer stability were assessed using the 1:1 Langmuir model for 2G12 Fab2. The reported kinetic binding curves are representative of 2 datasets.

For affinity measurements, the DH717.1 Fab was purified using size exclusion chromatography (SEC-FPLC) in PBS 1x. Approximately 10mg of DH717.1 Fab was loaded onto a Superdex 200 increase 10/300 column using a 500 µL loop and run at 0.5 mL/min using an Äkta Pure system (GE Healthcare). DH717.1 Fab peaks were collected via fractionation using a 96-well plate and were analyzed using the Unicorn 7.0.2 software (GE Healthcare). Using a linear regression derived from running protein standards (GE Healthcare) of known molecular weight, the molecular weights of the Fab monomer and dimer peaks were estimated. The monomer and dimer peak fractions were then pooled and concentrated using 10k, 0.5mL centrifugal filters (Amicon) and concentrations were measured using a NanoDrop UV-VIS spectrophotometer. Following SEC purification, DH717.1 Fab monomer and dimer stability were assessed using SDS-PAGE gel electrophoresis and re-analysis by SEC-FPLC. SDS-PAGE gel electrophoresis was performed using the BioRad system and 4%-15% TGX stain free gels. 7µg of each DH717.1 Fab fragment was loaded onto the gel under both reducing and non-reducing conditions. 10µg of a Precision Plus Protein standard (BioRad) was also loaded to help verify the size of the previously published macaque Ig libraries (Ramesh et al., 2017) and Cloanalyst default rhesus Ig library (https://www.bu.edu/computationalimmunology/research/software/). Reverse round primers (5′ to 3′) were as follows for IgA, IgG, IgE, IgM, IgD, IgK, and IgL: [IgA] GGGGGGGAGTTATGAGCCT, [IgG] TTTGCACCTGTGGTTGCT, [IgE] AGTGAGTGGTACACCACAT, [IgM] CACTGACACTCTCTGGAACT, [IgD] ATGGGAGGTGACACGCT, [IgK] GTGAGTCGCTGTTGACACT, and [IgL] TGACCTGGGAGCT GTGAGCT. First round primers (5′ to 3′) were as follows for IgA, IgG, IgE, IgM, IgD, IgK, and IgL: [IgA] GGGGGGGAGTTATGAGCCT, [IgG] TTTGCACCTGTGGTTGCT, [IgE] AGTGAGTGGTACACCACAT, [IgM] CACTGACACTCTCTGGAACT, [IgD] ATGGGAGGTGACACGCT, [IgK] GTGAGTCGCTGTTGACACT, and [IgL] TGACCTGGGAGCT GTGAGCT.
of the fragments. The gel was run at 200V for 35-40 minutes in Tris/Glycine/SDS running buffer. The Fab bands were analyzed via the Gel Doc EZ imager (BioRad). For SEC-FPLC re-analysis, 40μg of each DH717.1 Fab monomer and dimer were loaded onto a Superdex 200 increase 10/300 column using a 100 μL loop and run at 0.5 mL/min using the aforementioned Äkta Pure system.

The binding FDG Abs to CoV spikes was assessed by surface plasmon resonance on Biacore T-200 (GE Healthcare) at 25°C with HBS-EP+ (10 mM HEPES, pH 7.4, 150 mM NaCl, 3 mM EDTA, and 0.05% surfactant P-20) as the running buffer. For the data shown in Figure 6E, S proteins were captured via their C-terminal Twin Streptactin tags at ∼1000 RU levels on two alternate flow cells of a streptavidin coated chip, and 200 nM of each FDG antibody was flowed over all four flow cells; the binding curves were double-reference subtracted. The surface was regenerated between injections by flowing over 1M NaCl in 50mM NaOH solution for 10 s with flow rate of 100μl/min.

For determining affinity and kinetics of 2G12 interaction with SARS-CoV-2 spike, single cycle kinetics analysis was performed on the SARS-CoV-2 S protein immobilized on a streptavidin (SA) chip with five injections of 2G12 Fab, prepared by digestion with papain as described previously (Murin et al., 2014), at 3.125 nM, 6.25 nM, 12.5 nM, 25 nM and 50 nM concentrations. The data were fit to a 1:1 Langmuir binding model.

Structural analyses

Crystallography

DH717.1 IgG was expressed and purified as above. IgG samples were digested with papain and further purified to produce Fab fragment. DH717.1 Fab was purified via size exclusion chromatography to separate the Fab monomers and dimers. The DH717.1 Fab monomer was mixed with 3 molar excess of Man9-V3 at a final complex concentration of 15 mg/mL for crystallization trials. The complex crystallized over a 100 μl reservoir of 100mM Na Citrate, pH 5.0, 30% PEG 4000 in 1-2 days at room temperature. The DH717.1 Fab dimer, concentrated to 10 mg/ml in 0.1M HEPES buffer with 0.15M NaCl, was crystallized over a 60 μl reservoir of 0.1M citric acid pH 3.5, 14% PEG 1,000 in a drop composed of 0.5 μl protein plus 0.5 μl reservoir. The crystals were cryoprotected by brief immersion in reservoir solution supplemented with 25% glycerol for the DH717.1 Fab monomer-glycopeptide complex, and 30% ethylene glycol for the DH717.1 Fab dimer, before being flash frozen in liquid nitrogen. Diffraction data of the monomer-glycopeptide complex were collected at NE-CAT beamline 24-ID-E and data of dimer were collected at SER-CAT with an incident beam of 1 Å in wavelength. Data were processed using HKL-2000 (Otwinowski and Minor, 1997). Matthews analysis of the monomer-glycopeptide complex and dimer data suggested 1 Fab and four Fabbs present in the unit cell, respectively (Matthews, 1968). Molecular replacement calculations for the monomer-glycopeptide complex were carried out with PHASER, using published DH270.5 (Protein Data Bank (PDB) ID 5TPP) as the starting model. The Fab model was separated into its variable and constant domains for molecular replacement. A molecular replacement solution for the dimeric DH717.1 Fab with four Fabbs was found using Phaser with search models DH717.1 Fv (above) and the constant region from DH522 UCA (Terwilliger et al., 2008; Williams et al., 2017b). The solution was improved through alternating rounds of manual rebuilding in Coot and reciprocal space refinement in PHENIX, and geometry optimization using Rosetta-Phenix refinement (phenix.rosetta_refine) (Emsley et al., 2010; Adams et al., 2010). The crystal structure of the monomeric DH717.1 Fab complex showed well-defined electron density for the three protein-proximal mannose residues, while the rest of the Man9 glycan and the V3 peptide were disordered.

Negative-stain electron microscopy (NSEM)

MAbs were diluted to 100 μg/ml final concentration with buffer containing 10 mM NaCl, 20 mM HEPES buffer, pH 7.4, 5% glycerol and 7.5 mM glutaraldehyde. After 5-minute incubation, excess glutaraldehyde was quenched by adding sufficient 1 M Tris stock for a final Tris 75 mM for 5 minutes; then samples were stained with 2% uranyl formate. Images were obtained with a Philips 420 electron microscope operated at 120 kV, at 82,000 x magnification and a 4.02 Å pixel size. The RELION program was used for CTF correction, automatic particle picking and 2D class averaging of the single-particle images. For human FDG precursor recombinant mAbs, only DH1005, DH1009 and DH1010 showed a mixture of I- and Y-shaped conformations (Figure 3); the remaining mAbs only showed Y-shaped Abs (not shown).

Cryo-electron microscopy (cryo-EM) of HIV-1 Env complexes

For the DH898.1-bound complex, purified HIV-1 Env was mixed with 6-fold molar excess of recombinantly expressed and purified DH898.1 Fab to obtain a final concentration of 1.5 mg/ml Env, and incubated for 2 hours at room temperature. 2.5 μL of protein was deposited on a Quantifoil-1.2/1.3 holey carbon grid (Electron Microscopy Sciences, PA) that had been glow discharged for 30 s in a PELOCO easyGlow Glow Discharge Cleaning System. After a 30 s incubation in > 95% humidity, excess protein was blotted away for 2.5 s before being plunged into liquid ethane using a Leica EM GP2 plunge freezer (Leica Microsystems). For the DH851 Abs, attempts to obtain a structure of the complex by mixing with Fabs that were either recombinantly produced or obtained by proteolytic digestion of the IgG were unsuccessful, yielding mostly unliganded Env trimer populations and very small fractions of Env bound to Fab dimer (not shown). We next prepared a complex of DH851.3 bound to HIV-1 Env amenable to structural analysis by first making a complex of the Env with DH851.3 IgG, which was purified by size exclusion chromatography. The purified complex was then subjected to proteolytic digestion with HRV3c protease that cleaved at a site engineered between the Fab and Fc regions. Following purification over a protein A column to remove the Fc, the complex was purified by SEC, and concentrated to 1.5 mg/ml. Cryo-EM grids were prepared in a manner similar to the DH898.1 complex.

Cryo-EM imaging was performed on a FEI Titan Krios microscope (Thermo Fisher Scientific) operated at 300 kV, aligned for parallel illumination. Data collection images were acquired with a Gatan K3 detector operated in counting mode with a calibrated physical
pixel size of 1.066 Å with a defocus range between ~1.0 and ~3.5 μm using the Latitude S software (Gatan Inc.). The dose rate used was ~1.0 e-/Å² s to ensure operation in the linear range of the detector. The total exposure time was 4 s, and intermediate frames were recorded every 0.067 s giving an accumulated dose of ~60 e-/Å² and a total of 60 frames per image. A total of 6058 and 3230 images were collected for the DH851.3- and DH898.1-bound complexes, respectively. Cryo-EM image quality was monitored on-the-fly during data collection using automated processing routines.

Data processing was performed within cryoSPARC (Punjani et al., 2017) including particle picking, multiple rounds of 2D classification, ab initio reconstruction, heterogeneous and homogeneous map refinements, and non-uniform map refinements.

For DH898.1 reconstructions, micrographs were imported into cryoSPARC, CTF-corrected, and manually curated to eliminate astigmatism or defocus outliers and micrographs with excessive ice contamination, yielding 2978 final micrographs. Particles were picked using Blob Picker on 500 micrographs with a minimum and maximum particle diameter of 150 and 250 Å for circular blobs, yielding 259,587 particles, which were extracted with a 320-pixel box size and subjected to 2D class averaging with 100 classes. From these, seven classes were chosen for template picking applied to the full final micrograph set, yielding 1,655,400 particles that were extracted with a 256-pixel box size and subjected to multiple rounds of 2D classification and manual selection, multi-class ab initio refinements and multi-class heterogeneous refinements to eliminate junk particles. Early rounds of analysis quickly identified three particle populations consisting of unliganded Env SOSIP trimer, SOSIP trimer with a single Fab dimer bound near the CD4-binding site (CD4bs), and Fab dimers either free or bound to a poorly resolved blob of the approximate size of an Env gp120 subunit. An intermediate stage hetero-refinement with 91,638 particles used eight starting models that included four CD4bs SOSIP-Fab dimer complexes, two unliganded SOSIP trimers, and two Fab-blob complexes. This refinement yielded two junk classes (4,440 particles total), two Fab-blob classes (13,160 particles total), one unliganded SOSIP class (18,058), two CD4bs classes (40,350 total particles), and a new class with a single Fab dimer bound near the V3 glycan supersite of the SOSIP trimer (15,630 particles). We then returned to an earlier, larger stack with 324,010 particles, and subject this to a hetero-refinement using the unliganded SOSIP, the CD4bs complex and the V3 complex as starting models, which yielded unliganded SOSIP classes with 82,025 particles, and CD4bs-complex classes with 72,951 particles, and a V3-complex class with 21,502 particles. These particle stacks for the unliganded SOSIP and the CD4bs-complex were further cleaned to remove junk particles and yielded final reconstructions with 48,127 and 59,012 particles and achieving final global resolutions of 4.9 and 4.7 Å, respectively, as measured by cryoSPARC’s auto-masked, phase-randomized-corrected gold standard FSC calculation at the FSC 0.143 level. The map of the V3-complex class was used to generate ten 2D templates that were used to re-pick the micrographs, yielding 1,827,101 particles extracted with 320-pixel box size. This stack was subjected to 2D classification and manual selection, multi-class ab initio refinements and multi-class heterogeneous refinements to yield a final reconstruction at 6.1-Å resolution with 14,873 particles. Using cryoSPARC’s 3D Variability tool we noted substantial movement of the Fab dimer relative to the SOSIP trimer in the CD4bs complex; therefore we performed particle subtraction and local refinements focused on the fab dimer of the CD4bs-complex and the V3-complex. The local refinements reported the same or slightly worse resolution compared to the homogeneous refinements; however, the density for the Fab dimer was better defined as judged by eye, particularly the appearance of beta-strands within the Fab dimer interface. Local resolution at the FSC 0.5 level was estimated for all final reconstructions and maps were locally filtered and sharpened at 1.5x the reported B-value from the final refinements. Images were created in UCSF Chimera.

For the movies of the DH898.1-bound complex, a homogeneous set of ~122k particles were identified and re-extracted from the original dose-weighted micrographs using a binning factor 2. 3D classification in RELION (Scheres, 2012) was performed imposing C1 symmetry and using different models generated by cryoSPARC as references for alignment. All models were low-pass filtered to 60 Å before refinement. For the binding site located around a CD4 binding site glycan cluster, we identified one class with 42,745 particles that was consistent with the cryoSPARC reference model and subjected them to further 3D auto refinement. After convergence, the refinement was continued with a soft shape mask and resulted in an overall structure at 6.4-Å resolution. To obtain higher resolution, multibody refinement (Nakane et al., 2018) was performed on the refined particles using two individual soft masks (one for the Fab dimer, another for the Env trimer) created with UCSF Chimera (Petersen et al., 2004) and RELION. The overall resolution for the gp120 trimer portion reached 4.7 Å and for the Fab dimer the estimated resolution was 6.7 Å. At this point we switched to un-binned particles and repeated the same processing strategy, but without significant improvements. We then carried out focused refinement on the Env trimer component using the re-extracted particles and imposing C3 symmetry, which lead to an improved map at 4.3-Å resolution. Based on these refinement parameters, CTF-refine and Bayesian-polishing were performed on the particles. Another round of 3D auto refine was performed using the polished particles and the reference model of the entire complex (with a soft shape mask applied), resulting in an overall resolution of 4.8 Å. Multibody refinement was performed once again based on these updated refinements. The final resolution for the gp120 trimer after post-processing was 4.65 Å and 5.8 Å for the Fab dimer. For the V3-binding site, 17,510 particles were classified out using the V3-binding reference model generated by cryoSPARC. The first round of 3D auto refinement yielded an 8.14 Å resolution map, and multibody refinement resulted in a 7.6 Å map for the gp120 portion and 8.4 Å for the Fab dimer. Given the lower resolution of this refinement and the fewer number of particles compared with binding-site1, CTF-refine and Bayesian polishing were not used for this class. In order to visualize the principal motions present in the data identified by the multibody refinement procedure, we reconstructed intermediate volumes using different relative orientations of each body and produced movies using the ‘Volume Series’ tool from ChimeraX (Goddard et al., 2018). Additional editing of the videos was performed with Movavi Video Editor (Movavi) to label the different components and provide visual aids.
For DH851.3 reconstruction, micrographs were imported into cryoSPARC and CTF-corrected. Particles were picked using Blob Picker on 500 micrographs with a minimum and maximum particle diameter of 125 and 250 Å for elliptical blobs, yielding 99,285 particles, which were extracted with a 384-pixel box size, down-sampled to 128 pixel box size, and subjected to 2D class averaging with 100 classes. From these classes, five were chosen for template picking applied to the full micrograph set, yielding 1,278,458 particles that were extracted with a 384-pixel box size and subjected to multiple rounds of 2D classification and manual selection, multi-class ab initio refinements and multi-class heterogeneous refinements to eliminate junk particles. Refinements that imposed C3 symmetry yielded ~4.8-Å overall resolution, but at the expense of losing density in the constant domains, making it difficult to trace the elbow region to determine whether the Fab dimer was or was not domain-swapped. A 3D variability analysis confirmed that there was considerable flexibility within the constant domains, so the C3 refinement was not pursued further. The final non-uniform C1 reconstruction with 52,398 particles yielded a 5.6-Å overall resolution. Particle subtraction and local refinements focused on the Fab dimer did not improve resolution or the definition of the Fab dimer map, as judged by the lack of visible beta-strands.

**Cryo-EM model fitting**

Starting model for the CH848 SOSIP complex was from PDB 6UM6. Starting model for the DH851 complex was a homology model built using PDB 5VN8 as a template by submitting the DH898 sequence to SWISS-MODEL (Waterhouse et al., 2018). Homology models for DH898 and DH851 Fab s were built with SWISS-MODEL using the top reported template. Intermediate models were generated from the starting models using Rosetta (Song et al., 2013) to rebuild 500-1000 new models into the cryo-EM maps and selecting the model with the best combination of Rosetta energy and fit to the map. The cryo-EM half maps of CH848 SOSIP trimer and DH898.1 Fab dimer were used to improve the maps using a density modification program ResolveCryoEM (Terwilliger et al., 2020). The models were then fit manually in Coot (Emsley et al., 2010) following iterative refinement using Isolde (Croll, 2018) and subsequent manual coordinate fitting in Coot. These models were rigid fit into DH898.1 fab dimer bound to CH848 SOSIP trimer complex map, and the interacting glycans were fit manually in Coot followed in refinement in Isolde. DH851 structure was fit manually in Coot followed by refinement in Isolde as well. Structure and map analysis was performed using PyMol and ChimeraX (Goddard et al., 2018).

**Cryo-EM of SARS-CoV-2 complexes**

Purified SARS-CoV-2 spike preparations were diluted to a final concentration of ~1 mg/mL in 2 mM Tris pH 8.0, 200 mM NaCl and 0.02% NaN3, were mixed with 6-fold molar excess of 2G12 Fab and incubated for 2 hours at room temperature. 2.5 μL of protein was deposited on a CF-1.2/1.3 grid that had been glow discharged for 30 s in a PELCO easiGlow Glow Discharge Cleaning System. After a 30 s incubation in > 95% humidity, excess protein was blotted away for 2.5 s before being plunge frozen into liquid ethane using a Leica EM GP2 plunge freezer (Leica Microsystems).

Cryo-EM imaging was performed on a FEI Titan Krios microscope (Thermo Fisher Scientific) operated at 300 kV. Data were acquired with a Gatan K3 detector operated in counting mode using the Leginon system (Suloway et al., 2005). The K3 experimental parameters are 2.5 exposure with 50 frames at 50 ms framerate with a total dose of 66.43 e-/Å2 and a pixel size of ~1.058 Å/px. This system was also energy-filtered with a slit width of 20 eV. A total of 6804 images were collected.

Cryo-EM image quality was monitored on-the-fly during data collection using automated processing routines. Individual frames were aligned and dose-weighted using MotionCor2 (Zheng et al., 2017b) implemented within the Appion pipeline (Lander et al., 2009). Data processing was performed within cryoSPARC (Punjani et al., 2017) including particle picking, multiple rounds of 2D classification, ab initio reconstruction, heterogeneous and homogeneous map refinements, and non-uniform map refinements. Heterogeneous classifications were performed within RELION (Scheres, 2012) using masks produced in Chimera.

Structure of the all ‘down’ state (PDB ID 6VXX) previously published SARS-CoV-2 ectodomain, and a structure of the 2G12 Fab bound to Man1-2 (PDB ID 6N35) were used to fit the cryo-EM maps in ChimeraX. Coordinates were then fit manually in Coot (Emsley et al., 2010) following iterative refinement using Isolde (Croll, 2018) and subsequent manual coordinate fitting in Coot. Structure and map analysis was performed using PyMol and ChimeraX (Goddard et al., 2018).

**Dimerization sequence analysis**

For sequence analysis of DH851 Abs (see Figure S4), 2G12 was used as a positive control, which has four hydrophobic residues participating in contacts within the Fab dimer interface (Calarese et al., 2003); heavy chain residues 19, 77, 79 and 81. Residue 81 in the heavy chain of 2G12 was considered hydrophobic, because the amphipathic glutamine (Q) side chain was participating in a π-N bond with the aromatic phenylalanine (F) residue 77 of the adjacent heavy chain (Calarese et al., 2003). The isolectine I (l) at position 19 is especially important, because a single mutation to arginine (R) at this position is sufficient to completely disrupt Fab-dimerization and domain-swapping in 2G12 (Doores et al., 2010b). The 2G12 interface contains three rare interface residues (asterisks), which were not present in the germline sequence (Huber et al., 2010) and thus represent mutations acquired during antibody affinity maturation. In contrast to the I-shaped 2G12, the Y-shaped antibody VRC01 has only a single hydrophobic residue, no rare residues, and three charged residues in these positions, including an arginine at position 19. The three DH851 Abs that showed I-shaped, Fab-dimerized Abs, DH851.1 to DH851.3, have three or four hydrophobic interface residues, most of them rare, and no charged residues. In contrast, DH851.4 to DH851.6, DH851.8 and the DH851UCA have only a single hydrophobic interface residue and these Abs are all Y-shaped, similar to the negative control VRC01. For DH1003 lineage analysis (see Data S2Q), the specific pattern of hydrophobic residues within the interface correlated with the phylogram branch points, as indicated in the last column, suggesting that evolution of the lineage correlated with Fab-dimerization; this hypothesis will be explored further in subsequent studies.
QUANTIFICATION AND STATISTICAL ANALYSIS

We used a statistical test to look for distribution shifts between two time points (week 105 and 145) and the DH717 lineage cells within the RMs were considered the unit of analysis (Figure 2A). The asterisk indicated significance at $p < 0.0001$, using a non-parametric Wilcoxon test performed in SAS 9.4 (SAS Institute, Cary, NC). In ELISA (Figures 2, S2, and S4), error bars represented the standard error of the mean (GraphPad Prism) for mAb binding levels that was generated from multiple independent assay.
Supplemental figures

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Figure S1. Characterization of HIV-1 vaccine-induced FDG Ab lineage, DH717, related to Figure 1

(A) DH717 mAbs as well as biotinylated (b) control mAbs were tested in ELISA for binding to soluble recombinant HIV-1 Env trimer (CH848 strain) in a single ELISA. Binding titers were reported as Log AUC. Control mAbs targeted different epitopes on HIV-1 Env: both Env glycan and peptide – PGT121, PGT125 and PGT128; linear peptide (V3 loop) reactive – 19B; glycan-only reactive – 2G12; co-receptor binding site – 17B; and CD4 binding site – DH493). (B) Binding titers of DH717 and control mAbs with heat-killed yeast antigens *Candida albicans* or *Cryptococcus neoformans*. MAbs were tested in standard diluent or diluent spiked with 0.5M D-mannose. Binding data were representative of duplicate ELISAs. (C) Kifunensine [Kif] enriches Man9-glycans on recombinant HIV-1 Envs (Saunders et al., 2017a) (see STAR Methods). DH717 mAbs were tested for neutralization against a multi-clade panel of env-pseudotyped HIV-1 bearing Envs with heterogeneous [(-)Kif] or Man9-enriched glycans [(+)Kif]. Neutralization was tested in TZM-bl cells in a single experiment and titers are reported as IC50 in μg/ml. (D) Negative stain 3D reconstruction of DH717 I-shaped Ab. Map shown as transparent surface. Atomic model of two DH717.1 Fabs shown as ribbon diagrams, fit as rigid bodies into the 20Å resolution NSEM map using UCSF Chimera’s automatic fitmap function. As fit, the V\_H cysteine at residue 74 pointed toward one another (arrow) and the terminal sulfurs were 3.5 Å apart. (E) View of DH717.1 monomer crystal structure showed contact with aspartate, V\_H residue 34 (D34), and the man-9 ligand (green); electron density shown as mesh. (F) V\_H sequence alignment of DH717.1-4 with the inferred near-germline unmutated common ancestor (UCA). DH717.1, 0.2, and 0.4 have cysteine at residue 74, whereas DH717.3 and the UCA do not. (G) Close up views of DH717.1 dimer crystal structure, labeled by region. (H) Size exclusion chromatography (SEC) profile of DH717.1 C74S IgG. The SEC runs were performed on a Superose 6 Increase 10/300 column in running buffer composed of 10mM HEPES, pH 7.3, 150mM NaCl, 5% Glycerol, (l) Immobilization of DH717.1 wild-type (black lines) and DH717.1 C74S (gray lines) on anti-Fc surface. At least three technical repeats are shown for each. The Abs were captured on a CMS chip by flowing 200 nM of the Ab over a flow cell immobilized with ~9000 RU of anti-human Fc Ab. Following this immobilization, binding was measured by flowing over 200 nM solution of Env in the running buffer HBS-EP+ that is composed of 0.01 M HEPES pH 7.4, 0.15 M NaCl, 3 mM EDTA, 0.005% v/v Surfactant P20. Env binding was shown in Figure 1K.
Figure S2. Characterization of neutralizing FDG Ab lineage (DH898) elicited by SHIV infection in RMs, related to Figure 4

DH898 mAbs were elicited by pathogenic SHIV (CH848TF) infection in RM6163 (see STAR Methods). (A) DH898 mAbs were tested in ELISA for binding to Man$_3$-V3, and heat-killed yeast antigens *Candida albicans* or *Cryptococcus neoformans*, in the absence (black bars) or presence (white bars) of 1M D-mannose. Control

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mAbs are reactive to Env glycan (DH501 and 2G12), Env peptide (DH1013) or influenza (CH65). Binding titers were reported as Log AUC. Data shown were from a single ELISA that were in agreement with an independent experiment of DH898 mAbs binding to Man9-V3 and yeast antigens in the absence or presence of 0.5M D-mannose. (B) DH898 mAbs were tested in ELISA for binding soluble stabilized recombinant HIV-1 trimers, CH848 10.17 DS.SOSIP wild-type and mutant trimers, in a single experiment. Mutations in CH848 10.17 DS.SOSIP trimers included deletions of potential N-linked glycan sites in the V3 (N301A_N332A) or V1 (N133D_N138T) regions. Mutations also included amino acid insertions that filled glycan holes on the trimer (D230N_H289N). Binding levels were measured at OD450nm. CH848 10.17 SOSIP trimers were captured using PGT151, Biotinylated (B) 2G12, PGT128 (V3 glycan bnAb) and CH65 mAbs were tested as control Abs. (C) Binding levels of DH898 mAbs to CH848TF SOSIPv4.1 in one ELISA, representative of at least two independent experiments. CH848TF SOSIP trimer was captured using anti-AVI mAb. 2G12, PGT151 and CH65 were tested as control Abs. (D) NSEM class averages for DH898 mAbs with lineage member indicated at upper left of each panel. At the lower left corner, each class average was identified as Y-shaped (Y), l-shaped (l), or ambiguous (A). (E) Sequence analysis of key residues within the Fab-dimer interface shows three hydrophobic or aromatic residues in the interface, similar to 2G12. V\textsubscript{H} interface residues were numbered according to standard Kabat numbering (Wu and Kabat, 1970). Amino acids at each position were indicated by their one-letter code and colored according to Taylor (Taylor, 1997), with polar groups orange, hydrophobic and aromatic groups in shades of green to yellow, positively charged groups blue and negatively charged groups red. Residues that are rare for a particular position, i.e., \leq 1% in the abYsis database (Swindells et al., 2017), are indicated with an asterisk. Bottom row indicates VRC01 as a non-Fab-dimerized negative control. (F) Bar graph indicating the fraction of I-shaped Abs for wild-type (blue bars), strengthening mutants (green bars), or disrupting mutants (red bars) in DH898 mAbs. Mutations were engineered in the Ab VH genes. The % I-shaped was estimated by the fraction of particle images that sorted into I-shaped classes as indicated in each of the NSEM 2D class averages shown in G. (G) NSEM 2D class averages from of DH898.4 wild-type (WT), strengthening double mutations T19I and T70F, and disrupting double mutations R64D and F68D. The class averages shown represented the average of \sim 8,000 to 20,000 individual particle images classified and averaged into ten classes, arranged from the most populated class at the top left to the least populated at the bottom right, and marked as I-shaped (I), Y-shaped (Y), or ambiguous (A). (H) DH898.4 wild-type and mutant mAbs were tested in ELISA for binding to HIV-1 CH848TF SOSIP trimer. MAbs were tested in technical replicates within a single ELISA and binding levels measured at OD450nm; error bars represent standard error of the mean. (I) DH898.4 wild-type and mutant mAbs were tested for neutralization against autologous (CH848TF) HIV-1 strain bearing Env with heterogeneous glycoforms as well as heterologous HIV-1 isolates bearing Env Man9\textsubscript{p} enriched glycans [Kif] in TZM-bl cells. These data were generated in a single neutralization assay and titers were reported as IC\textsubscript{50} in \mu g/ml.
Figure S3. DH898.1 complex cryo-EM data processing summary and structural details, related to Figure 4
(A) Representative micrograph; (B) Power spectrum of micrograph, and fitted contrast transfer function; (C) Picked particles in white circles; and (D) Representative 2D class averages. (E) Ab initio volumes that showed three particle populations: free trimer, Fab-dimer bound to glycans near the CD4-binding site (bs), and Fab-dimer bound to glycans near the base of the V3 loop (left to right) seen in top and side views (top to bottom). Dashed line in side view indicated viral membrane location. (F) Top and side view of free soluble stabilized recombinant HIV-1 Env trimer (SOSIP) colored by local resolution. Mesh surface indicates mask used for FSC calculation. Eye indicates viewing direction in side view. (G) HIV-1 Env SOSIP with Fab-dimer bound to glycans near the CD4bs. (H) HIV-1 Env SOSIP with Fab-dimer bound to glycans near the base of the V3 loop. (I) Gold standard Fourier Shell Correlation (FSC) curves for F-H and J-K indicating global resolution ranging from 3.9 – 7.0 Å. (J) Local refinement of Fab-dimer from CD4bs particle set. (K) Local refinement of Fab-dimer only from V3-glycan bound particle set. (L) Segmented cryo-EM map showing Fab-dimer bound near the CD4bs. Gold-standard FSC resolution is indicated below each map. (M) Local refinement of the Fab-dimer bound near to the CD4bs of a HIV-1 Env SOSIP. (N) Close-up view of the epitope near the CD4bs with a HIV-1 Env SOSIP and Fab-dimer models shown as ribbons; glycans shown as sticks; and the cryo-EM map shown as a transparent surface. (O) (left) Local refined, density modified map of the DH898.1 Fab dimer with cryo-EM reconstruction shown as blue mesh and underlying fitted model in cartoon representation, and (right) Zoomed-in view of the Fab dimer interface with select interfacial residues shown as sticks. (P) Views of the Fab dimer interface rotated 90° clockwise (left) or counter-clockwise (right) relate to the view shown in panel O. (Q) Zoomed-in views of Env regions in the apo structure.
Figure S4. Characterization of FDG bnAb lineage (DH851) elicited by SHIV infection in RMs, related to Figure 5

DH851 mAbs were elicited by pathogenic SHIV (CH848TF) infection in RM6163 (see STAR Methods). (A) DH851 mAbs were tested for neutralization against env-pseudotyped HIV-1 bearing EnvVs with Man$_9$-enriched glycans [(+) Kif] and heterogeneous glycoforms [(−) Kif]. Neutralization was tested in TZM-bl cells and titers reported as IC$_{50}$ in μg/ml. (B) Summary of neutralization profile for DH851 and 2G12 bnAbs that were tested against 119 difficult-to-neutralize multi-clade HIV-1 strains in TZM-bl cells (Seaman et al., 2010). (C) DH851 neutralization epitopes mapped on a SHIV bearing wild-type or mutant HIV-1 Ce1176-strain EnVs. Mutant EnVs had deletion of potential N-linked glycosylation sites that constituted the Env glycan-containing bnAb epitope targeted by glycan-only Abs as well as Abs that bound both Env glycans and peptide (GDfR motif) (Kong et al., 2013). While DH851 mAbs also neutralized HIV-1 Ce1176 strain, a mutation in the GDIR motif (D325N) did not impact the neutralization sensitivity of HIV-1 Ce1176 (not shown). (D) DH851 mAbs bearing sequences for clonally-related V$_H$ genes from week 0 (DH851.5, DH851.6 and DH851.8), computationally-inferred near-germline unmutated common ancestor (DH851UCA), and post-infection (DH851.1-DH851.4) Abs were tested in ELISA for binding to soluble recombinant HIV-1 Env trimer (SOSIP) of autologous CH848TF strain in triplicate experiments. Biotinylated (B) PGT151 (Env trimer reactive) and CH65 (influenza reactive) mAbs were tested as controls. Binding levels were measured at OD$_{450}$nm. (E) DH851 mAbs were tested in ELISA for binding to Man$_9$- and non-glycosylated aglycone V3 peptide in triplicate experiments; error bars represent standard error of the mean. 2G1013 (peptide reactive), 2G12 (Env glycan reactive) and CH65 mAbs were tested as control mAbs. (G) DH851 mAbs were tested in ELISA for binding to heat-killed yeast antigens, Candida albicans or Cryptococcus neoformans in triplicate experiments; error bars represent standard error of the mean. DH851 and CH65 were tested as control mAbs. (H) NSEM 2D class averages of DH851 mAbs displaying I- and Y-shaped Abs. (I) Sequence analysis of key residues within the Fab-dimer interface. (J) Bar graph indicating the fraction of I-shaped Abs for wild-type (blue bars) and strengthening mutants (green bars) in DH851 mAbs. Mutations were engineered in the Ab V$_H$ genes. The % I-shaped was estimated by the fraction of particle images that sorted into I-shaped classes as indicated in each of the NSEM 2D class averages from DH851.2 wild-type (WT) and strengthening double mutations A19T and T21L. The class averages shown represented images classified and averaged into five classes, and marked as I-shaped (I) or Y-shaped (Y).
Figure S5. DH851.3 complex cryo-EM data processing summary and structural details, related to Figure 5
(A) Representative micrograph; (B) Power spectrum of micrograph, and fitted contrast transfer function; (C) Picked particles in white circles; and (D) Representative 2D class averages. (E) Ab initio volume of DH851, bound to a soluble stabilized HIV-1 Env trimer (SOSIP) seen in top and side view. Dashed line in side view indicated viral membrane location. (F) Refined 3D map, without filtering or sharpening and colored by local resolution from 5.5 to 12.5 Å, blue to red. (G) Map after local filtering and B-factor sharpening. Fab constant domains are noisy and at this contour level mostly disappear and are only seen as small, disconnected blobs, but greater details of the complex can be seen. (H) Gold standard Fourier Shell Correlation (FSC) curves indicated global resolution ranging from 5.6 to 8.7 Å. (I) 2D class average shows view of Fab in complex with a HIV-1 Env SOSIP looking down the central axis. Red box shows region where the DH851.3 Fab dimer contacts the adjacent SOSIP protomer. (J) Fitted model shown as ribbon diagram with cryo-EM map shown as transparent volume. (K) Close up of contact seen in panel (I) showed that it may involve the framework region 3 of the light chain, and Env residues around 426-431 or 105-112. (L) View of the DH851.3 Fab dimer bound to Env glycans. Cryo-EM reconstruction was shown as a blue mesh, with underlying fitted coordinates in cartoon and stick representation. (M) and (N) Zoomed-in views of the Fab dimer interface shown within the dotted rectangle in the left panel.
Figure S6. Characterization of glycan-dependent binding of FDG Abs to recombinant SARS-CoV-2 spike protein, related to Figure 6

(A) FDG, SARS-CoV-1 RBD (D001), and influenza HA (CH65) mAbs were tested in ELISA for binding to recombinant SARS-CoV-2 Spike (S) protein. mAb binding was assessed in the absence (-) or presence (+) of D-mannose [1M] to determine if free high mannose can outcompete glycans on the S protein for binding. Binding Ab titers reported as Log AUC were shown in Figure 1. (B) FDG mAbs were tested in ELISA for binding to a set of commercially available constructs expressing the SARS-CoV-2 S1 and S2 extracellular domain (left), S2 domain (middle), and the receptor binding domain (right). Black and white bars show binding in the absence and presence of D-mannose [1M], respectively. Binding Ab titers were reported Log AUC. (C) We tested 2G12 mAb for binding to (from top to bottom) soluble stabilized recombinant HIV-1 Env trimer (CH505TF SOSIP), Man9-V3 and non-glycosylated aglycone V3 peptide, and recombinant SARS-CoV-2 spike ectodomain. HIV-1 CH505TF SOSIP was captured using mouse anti-AVI-tag mAb, whereas SARS-CoV-2 ectodomain and peptides (Man9-V3 and Aglycone) were captured using streptavidin. Blue and red symbols indicated binding in the absence and presence of D-mannose [1M], respectively. Binding was measured at OD450nm. All ELISAs (A-C) were done using BSA-based buffers (see STAR Methods). Data shown are from a representative assay. (D) Size-exclusion chromatogram of protein A affinity purified 2G12 IgG. (E) NSEM 2D class averages of (top) 2G12 IgG dimer, (bottom) 2G12 IgG monomer and (right) 2G12 Fab obtained by digesting 2G12 IgG monomer with papain. (F) SPR sensorgrams showing binding of 2G12 IgG dimer (red line) and 2G12 IgG monomer (black line) to the SARS-CoV-2 S protein. (G) We tested FDG mAbs for binding to the unmutated SARS-CoV-2 S and the N709-glycan deleted variant. Binding was assessed by SPR by capturing the unmutated spike and the N709-glycan deleted spike on flow cells 2 and 4 of a streptavidin coated (SA) chip, and flowing over a 200 nM solution of each Ab simultaneously over all four flow cells. Flow cells 1 and 3 were used as reference flow cells for flow cells 2 and 4, respectively. Buffer blanks were run in a similar manner and the sensorgrams were double-referenced by first subtracting the signal from the reference flow cell and then subtracting the reference-corrected buffer blank. CR3022 IgG and ACE-2 tagged with a mouse-Fc region were used as controls. (H) Binding of 2G12 to spike proteins of SARS-CoV-2 (red), SARS-CoV (blue) and MERS-CoV (green). The SARS-CoV-2/2G12 complex is shown with the spike in red and the bound 2G12 as a transparent pink surface, with the glycans contacting 2G12 shown as sticks and the respective Asn residues as spheres. Similar glycosylation was observed for the SARS-CoV and MERS-CoV S proteins (Asn residues shown as spheres). SPR binding data shown for SARS-CoV-2 (red), SARS-CoV (blue) and MERS-CoV (green) S proteins. The data shown are representative of three independent experiments, and for each dataset the graph showed three technical replicates. IgGs were captured on flow cells of a CM5 chip immobilized with human Anti-Fc antibody (8000RU). 200 nM solution of the SARS-CoV-2 spike was flowed over the flow cells. The surface was regenerated between injections by flowing over 3M MgCl2 solution for 10 s with flow rate of 100μl/min.
6804 micrographs; 2,782,625 particles

1,544,666 particles

1,285,982 particles

258,684 particles

1-2G12-bound
107,551 particles

2-2G12-bound
109,774 particles

2-2G12-bound with "partial" occupancy of third binding site
41,917 particles

(legend on next page)
Figure S7. Cryo-EM data processing details for SARS-CoV-2 S protein complex with 2G12, related to Figure 6
(A) Representative micrograph. (B) CTF fit. (C) Representative 2D class averages. (D) Maps for (left) unliganded and (right) 2G12-bound S obtained after 3D classification. (E-G) Refined maps for SARS-CoV-2 S protein bound to (E) 1-2G12, (F) 2-2G12-, and (G) 2-2G12 (with partial occupancy at the third binding site) Fab2 molecules. Red arrow in (G) points to disordered 2G12 Fab2 bound at the third binding site. (H) (Left) Map combining all particles and focusing refinement on the region within the masks that is shown as a gray mesh overlaid on the final refined map shown as a gray surface. (Right) Fourier shell correlation curves. (I) (Left) Cryo-EM reconstruction of 2G12 bound to the SARS-CoV-2 spike colored by local resolution. (Right) Zoomed-in view showing the cryo-EM reconstruction of the bound 2G12 Fab. (J) (Left) Two distinct states were resolved from the cryo-EM data by heterogeneous classification. Density for the two observed states were shown in green and gray. (Right) Cartoon representation of the SARS-CoV-2 S-protein (bright green, bright orange, blue) and the two 2G12 orientations. The axis of rotation hinged around glycan 709 is represented by a gray cylinder. (K) Zoomed-in view of (from left to right) a region in the S2 domain with map shown as blue mesh and fitted model shown as sticks; glycan 709, glycan 801; glycan 717 bound to 2G12. While not in direct contact with the bound antibody, the HR1 helix may play an indirect role in the binding by stabilizing glycan 717 via a stacking interaction with residues N925 and Q926.