

# Structural and functional studies on the extracellular domain of BST2/tetherin in reduced and oxidized conformations

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HIV-1 and other enveloped viruses can be restricted by a host cellular protein called BST2/tetherin that prevents release of budded viruses from the cell surface. Mature BST2 contains a small cytosolic region, a predicted transmembrane helix, and an extracellular domain with a C-terminal GPI anchor. To advance understanding of BST2 function, we have determined a 2.6 Å crystal structure of the extracellular domain of the bacterially expressed recombinant human protein, residues 47–152, under reducing conditions. The structure forms a single long helix that associates as a parallel dimeric coiled coil over its C-terminal two-thirds, while the N-terminal third forms an antiparallel four-helix bundle with another dimer, creating a global tetramer. We also report the 3.45 Å resolution structure of BST2(51-151) prepared by expression as a secreted protein in HEK293T cells. This oxidized construct forms a dimer in the crystal that is superimposable with the reduced protein over the C-terminal two-thirds of the molecule, and its N terminus suggests pronounced flexibility. Hydrodynamic data demonstrated that BST2 formed a stable tetramer under reducing conditions and a dimer when oxidized to form disulfide bonds. A mutation that selectively disrupted the tetramer (L70D) increased protein expression modestly but only reduced antiviral activity by approximately threefold. Our data raise the possibility that BST2 may function as a tetramer at some stage, such as during trafficking, and strongly support a model in which the primary functional state of BST2 is a parallel disulfide-bound coiled coil that displays flexibility toward its N terminus.

coiled coil | crystal structures | HIV | innate immunity | restriction factor

Viral infection can induce a type I interferon response, which in turn stimulates the expression of many genes that encode innate immunity factors (1). One of the upregulated proteins, bone marrow stromal cell antigen 2 (BST2/tetherin/CD317/HM1.24), inhibits the release of HIV-1 and other enveloped viruses from the surface of cells in which it is expressed (2–10). HIV-1 virions retained at the cell surface in the presence of BST2 are fully formed and mature but remain attached to the cell surface by a protease-sensitive linkage (11, 12) that contains BST2 (13, 14). The retained virions can subsequently be internalized and have been visualized in early and late endosomes, and they may ultimately be degraded following fusion with lysosomes (12).

HIV-1 escapes BST2-mediated restriction through the activity of its accessory protein viral protein U (Vpu) (15–17). Most, but not all (18), studies have found that Vpu reduces cell surface levels of BST2, and the two proteins appear to interact through their transmembrane (TM) domains (19–25). Various mechanisms for BST2 surface downregulation have been proposed, including lysosomal degradation, proteasomal degradation, and/or sequestration/retargeting of BST2 to the trans-Golgi network (reviewed in ref. 8). The general importance of BST2 is indicated by the findings that other viruses use different proteins to overcome BST2 restriction, including the Env/glycoproteins of Ebola virus and some strains of SIV and HIV-2, (4, 26, 27), the negative

factor (Nef) protein of other SIV strains (24, 28, 29), and the K5 protein of Kaposi's Herpes Sarcoma virus (30, 31).

The mature BST2 protein has an unusual architecture, comprising a small cytosolic domain and a TM helix at the N terminus (residues 22–44), a glycosylphosphatidylinositol (GPI) anchor at the C-terminal Ser160 residue, and an approximately 105-residue-long disulfide-rich coiled coil structure predicted for the extracellular domain (32–34). This configuration appears central to function, with the leading model being that virions are held at the cell surface by a bridge in which the extracellular domain of one or, more likely, multiple molecules of BST2 spans the gap between plasma membrane and the membrane of the otherwise detached viral particle (13, 14, 34). It has been argued that the GPI modification targets BST2 to cholesterol-rich regions of the plasma membrane (33), which may concentrate the protein at sites of viral budding. However, removal of the GPI anchor reportedly does not release virions from their tethered state (14). Finally, functions for BST2, independent of viral restriction, have also been proposed in trafficking and signaling (33), and in organization of the subapical actin cytoskeleton (35).

It has been proposed that BST2 forms a parallel dimeric coiled coil that is stabilized by C53–C53, C63–C63, and C91–C91 disulfide bonds. This model is supported by nonreducing SDS/PAGE analyses of mutant proteins isolated from the surface of HEK293T cells (32, 34) and by a recent crystal structure of residues 89–147 of the BST2 extracellular domain (36). Both the N-terminal TM helix and the C-terminal GPI can be incorporated into HIV-1 virions, and it has therefore been suggested that BST2 either forms an antiparallel dimer (14) or that it forms a parallel dimer whose tethering orientation is not critical for viral restriction (34). We also note that the importance of BST2 dimerization has been disputed (37). Remarkably, HIV-1 restriction is preserved (at approximately 10% levels) in cells that express an engineered protein that was designed to maintain the overall BST2 architecture but does not conserve the specific amino acid residue identities (34). Finally, the efficiency of BST2-mediated HIV-1 inhibition appears to be concentration-dependent, because virions that contain BST2 can be released under

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Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, [www.pdb.org](http://www.pdb.org) [ID codes 3nwh (reduced BST2) and 2xg7 (oxidized BST2)].

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conditions of low BST2 expression (34). These considerations emphasize the importance of determining the oligomeric state(s) of BST2. In an effort to understand the mechanistic basis for BST2 activity better, we have determined crystal structures and solution oligomerization states of oxidized and reduced BST2 proteins and tested the restriction activity of a structure-based mutant that cannot switch oligomerization states when reduced.

## Results

**Reduced BST2 Crystal Structure.** We determined a structure of the human BST2 extracellular domain (residues 47–152) produced in *Escherichia coli* and crystallized in the presence of reductant to 2.6 Å resolution and  $R/R_{\text{free}}$  values of 26.0%/27.8% (Fig. 1 and Table S1). All but a few of the BST2 residues at the termini of the four molecules in the asymmetric unit were visible, although the central regions (residues 92–117) have relatively weak density and high B values.

Subunits are arranged as parallel dimers that splay apart slightly to form an antiparallel 4-helix bundle over the N-terminal 40 residues and generate a 245 Å-long tetramer in the asymmetric unit (Fig. 1A). The remaining two-thirds forms a dimeric coiled coil, with residues 118–150 displaying the standard “knobs-into-holes” packing of *a* and *d* heptad repeat residues seen in classical coiled coils (38–40), and residues 92–117 packing more loosely to display some but not all of the *a* and *d* residue knobs-into-holes interactions (Fig. S1). Equivalent packing for the C-terminal two-thirds was reported by Hinz et al. (36).

Although each of the subunits forms one continuous helix that is devoid of sharp turns, the helices do not all superpose closely over their entire length. The two parallel dimers in the asymmetric unit are very similar to each other and superpose with a root mean square deviation of 0.79 Å over all C $\alpha$  atoms (Fig. 1B), but the two subunits within one parallel dimer show an approximately 30° deflection in the vicinity of residue 90, near the tetramer-dimer transition (Fig. 1C). This asymmetry results from small differences spread over several residues and may indicate that the BST2 sequence encodes an asymmetric dimeric structure, although we prefer the explanation that the molecule has inherent flexibility.

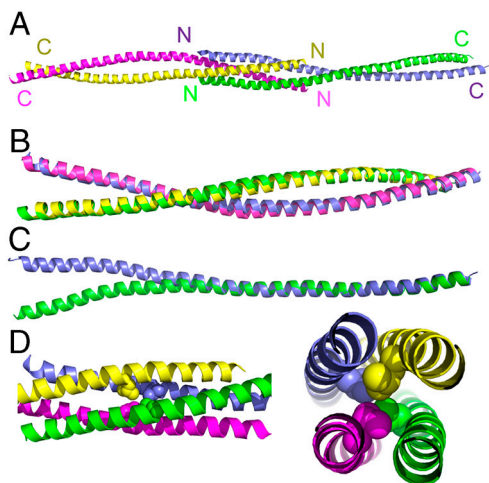
C53 and C63 are contained within the N-terminal four-helix bundle while C91 is immediately C-terminal to the bundle. All three cysteines are reduced in this structure, whereas biochemical data indicate that BST2 forms disulfide bonds in the oxidizing

environment of the cell exterior (34). Residues C53 and C63 are too far from their partners to form disulfide bonds without disrupting packing of the four-helix bundle region. In contrast, neighboring C91 residues could form a disulfide bond without substantial movement of their main chains from the crystal structure. We have built C91 residues with their sulfur atoms separated by 4.7 Å in this structure because the density, which is relatively poor in this region, does not indicate formation of a disulfide.

**Oligomeric States in Solution.** Equilibrium analytical ultracentrifugation (AUC) was used to determine the association state of recombinant BST2 proteins in solution. This method provides shape-independent estimates of mass and is therefore ideally suited for studies of highly extended assemblies such as BST2. These studies employed slightly longer BST2(47–154) constructs in order to include two tyrosine residues and thereby increase optical absorbance. The AUC data showed that the recombinant protein is a tetramer in solution under the reducing conditions used in crystallization, over an initial concentration range of 0.5–2.0 mg/ml (41.7–166.7 μM). Equilibrium protein distributions could be globally fit to a single species tetramer model (Fig. S2) and also indicated a tetrameric association when the molecular weight was allowed to float (Fig. 2A).

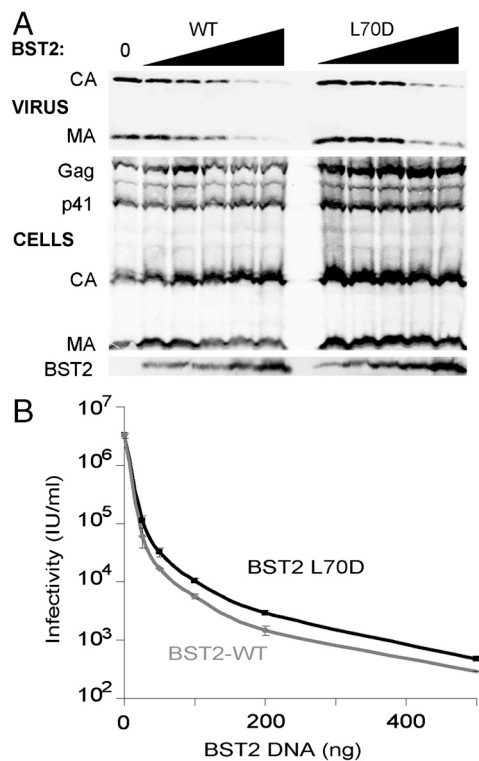
To determine if oligomerization was altered by oxidation, purified reduced BST2 was dialyzed for one week against identical buffer that lacked reductant. Mass spectrometry showed that this protein was a fully disulfide-linked dimer (Fig. S3). The protein eluted from a sizing column primarily as a well-defined peak but also included a broad peak at shorter retention times that presumably corresponded to oxidized aggregates. AUC showed that protein from the main peak is dimeric over the concentration range tested (0.5–2.0 mg/ml), either by fitting to a dimer (Fig. S2) or by allowing the molecular weight to float (Fig. 2B). These data indicate that BST2 forms a stable tetramer under reducing conditions and converts into a stable dimer upon oxidation.

**Oxidized Crystal Structure.** To determine the BST2 extracellular domain structure under oxidizing conditions, we expressed secreted BST2(51–151) in HEK293T cells and completed purification and crystallization in the absence of reductant. Crystals of Endo-H treated BST2(51–151) diffracted anisotropically to a maximum resolution of 3.45 Å resolution. This structure, with one dimer in the asymmetric unit, was determined by molecular replacement using residues 89–151 of the reduced structure as a search model. The refined model ( $R/R_{\text{free}} = 26.9\%/29.8\%$ ) comprises residues 72–151 for one of the two molecules in the asymmetric unit and residues 77–151 for the other molecule. The oxidized and reduced structures overlap closely from residue 90 to the C terminus but diverge toward the N terminus where the reduced protein tetramerizes (Fig. 3). For example, the C $\beta$ –C $\beta$  distance of juxtaposed A88 residues, which occupy an *a* coiled coil position, is 4.2 Å in the oxidized structure and 7.4 Å in the reduced structure. Clear electron density, confirmed by kicked omit maps, is observed for a disulfide bridge linking C91 residues validating the oxidized state of the molecule (Fig. 3). Two sugar moieties extending from N92 and positioned perpendicular to the C91–C91 disulfide could be located in the electron density (Fig. S4). Finally, additional N-terminal residues could be modeled in a helical conformation (Fig. S5), although due to the modest resolution, gaps in the density, and appearance of disorder, we have not included those residues in the deposited model. Thus, the dimeric coiled coil extends into the region that is tetrameric in the reduced structure, and residual density suggests that the coiled coil continues most of the way to the N terminus of this construct, albeit with pronounced flexibility.



**Fig. 1.** Structure of reduced BST2(47–152). (A) The four molecules in the asymmetric unit. Yellow/magenta and blue/green subunits form parallel dimers. (B) Overlap of the two parallel dimers. (C) Overlap of the blue and green subunits on residues 90–152. (D) Orthogonal views showing the L70 side chain buried at the center of the tetramer.





**Fig. 4.** Inhibition of HIV-1 release and infectivity by BST2 proteins. (A) Western blot of samples from 293 T cells that expressed HIV-1 $_{\Delta Vpu}$  and either lacked BST2 (lane 1, unrestricted control) or expressed increasing levels of wild-type BST2 (lanes 2–6) or BST2(L70D) (lanes 7–11). Panel 1, levels of virion-associated viral Gag proteins released into the supernatant (VIRUS, anti-CA and anti-MA). Panel 2, intracellular Gag protein levels (CELLS, anti-CA and anti-MA). Panel 3, intracellular BST2 levels (CELLS, anti-BST2). The first lane of Panel 1 contains 25-fold less sample than the other lanes. Quantification showed that, on average, cells transfected with BST2(L70D) expressed  $1.4 \pm 0.3$ -fold more protein than those transfected with the wild-type BST2 ( $n = 5$ ,  $\pm$ s.d.). Nevertheless, cells expressing BST2(L70D) released  $2.5 \pm 1.3$ -fold more virion-associated CA and MA protein ( $n = 10$ ,  $\pm$ s.d.). (B) Graph showing HIV-1 titers in the presence of increasing quantities of expression vectors. Wild-type BST2 (light curve), BST2(L70D) (dark curve). Each data point corresponds to a sample analyzed in the Western blots shown in panel A. On average, viral titers were  $1.9 \pm 0.1$ -fold higher in cells expressing BST2(L70D) mutant (vs. wild-type BST2,  $n = 10$ ,  $\pm$ s.d.).

While our manuscript was in preparation, a structure of the bacterially expressed C-terminal two-thirds of BST2 (residues 80–147) was reported by Hinz and colleagues (36). This structure showed formation of the C91–C91 disulfide bond, was ordered over residues 89–147, and superimposes closely with our BST2 structures (RMSD = 0.5–0.8 Å over essentially all C $\alpha$  atoms). This reinforces the impression that the C-terminal two-thirds of the BST2 extracellular domain forms a dimeric coiled coil over both the well packed C-terminal third and over the less regularly associated central third.

Hinz et al. (36) also reported small angle X-ray scattering (SAXS) analysis of dimeric BST2(80–147) and BST2(47–159) proteins. Their data indicate an elongated structure that has a bend at about one third from the N terminus. An attractive possibility is that the N-terminal third of the molecule is flexible, with one specific possibility being that the coiled coil conformation continues toward the N terminus but can hinge about residues approximately 80–90. This apparent flexibility is consistent with (i) our finding that residues in this region adopt different conformations in our reduced and oxidized structures (Fig. 3) and in different subunits in the reduced structure (Fig. 1C), (ii) the appearance of electron density maps for our oxidized BST2(51–151) structure (Fig. 3 and Fig. S5), (iii) the protein’s protease sensitiv-

ity at residue 80 (36), and (iv) the retention of tethering activity when an HA tag is introduced following residue 82 (36).

In addition to a hinging motion, it is possible that the N-terminal third of the BST2 extracellular domain has internal flexibility. This would be consistent with the absence of favored buried coiled coil residues in this region of the sequence and our finding that L70D, which corresponds to a buried *d* position in a simple coiled coil model derived from our crystal structures, expresses and purifies like the wild-type sequence and retains activity in a viral restriction assay (Fig. 4). The distribution of cystine residues is compatible with formation of a standard coiled coil throughout the length of the molecule because these residues would occupy *a* (C53), *d* (C63), and *d* (C91) positions in the model extended from the crystal structure. Disulfides can typically be accommodated in these positions, although they are expected to cause some structural perturbation when introduced into an *a* site (41). Just one disulfide is required for stability because constructs lacking any two cystines retain viral restriction activity, albeit with reduced potency (34). The conservation of all three cystines in a region of presumably loose-packed coiled coil is probably not simply due to a requirement for stability because coiled coils can be very stable in the absence of any disulfides. Rather, the cysteine conservation likely reflects a balance between requirements of stability and flexibility for optimal BST2 function.

The model in which BST2 forms a coiled coil from residue 47 to 152 predicts an overall length of 140 Å. This compares with the finding from SAXS measurements that the BST2(47–159) construct has an overall length of 170 Å (36). This 30 Å length difference might be explained by extended conformation(s) for residues 153–159, which are not present in any of the crystal structures plus a few additional residues at the ends of the molecule, which tend to be poorly defined in electron density. Partial unwinding of helices of the N-terminal disulfide-bound portion of the coiled coils is also a possibility.

A number of observations indicate that the N-terminal region of the BST2 extracellular domain is functionally important, including the findings that activity is lost when all disulfide bonds are disrupted (34) or when seven residues within 62–73 are mutated (36). Although we find that the L70D protein retains activity, thereby indicating that formation of a stable reduced tetramer is not essential for viral restriction, the restriction activity is reduced by approximately twofold compared to wild-type, even though expression of the mutant increases slightly. This reduction in potency indicates that L70 contributes to restriction, likely by stabilizing a functional conformation. Indeed, the activity reduction in the L70D mutant approaches the 10-fold loss of activity seen in an “artificial” BST2 molecule that replaces essentially all of the amino acid residues (34). L70D may diminish restriction activity by impacting the conformation or stability of the parallel coiled coil conformation. Alternatively, L70D may diminish activity by inhibiting formation of the reduced BST2 tetramer that, though not absolutely essential for HIV-1 restriction, might contribute to the efficiency of restriction or to some other BST2 function. It is striking that BST2 forms a stable tetramer in solution, indicating that the four-helix bundle is an energetically stable conformation, although we note that isolated coiled coil peptides can adopt nonphysiological association states (42). In principle, tetramerization could occur at several different stages of the BST2 “life cycle.” One possibility is that BST2 could become reduced upon localization to a cellular compartment that induces disulfide bond reduction (43, 44), in which case tetramerization could serve as a signal that alters protein trafficking (e.g., promoting BST2 recycling or retention).

Finally, the structure raises the general question, why is BST2 dimeric? Because the leading model is that virions are tethered by the TM helix and GPI anchors binding plasma and viral membranes, one can question why a single isolated subunit is not

sufficient for activity. There are a number of possible explanations, which are not mutually exclusive. Consistent with a report that the GPI anchor is dispensable for restriction (14), one possibility is that virions can be tethered by the two TM helices of a dimer embedding separately in plasma and virion membranes. A second possibility is that dimerization provides a more structured extracellular domain that is better able to fold and resist proteolysis. Third, formation of an extended coiled coil, rather than a more flexible isolated single subunit, may optimally separate the TM helices and GPI anchors, thereby increasing the chance that virions will bud between them and become tethered to the cell. Finally, dimerization may strengthen the tether through avidity effects that result from having two attachments to each membrane. Avidity effects may also explain why higher concentrations of BST2 molecules produce more efficient virion retention (34) because higher order BST2 dimer-dimer interactions could be stabilized in the constrained environment of the cell surface. Although important questions remain regarding the structure of the tether that connects virus and cell, our data indicate that a BST2 parallel coiled coil is likely to be a fundamental component.

## Methods

**Structure Determination of Reduced BST2.** Human BST2(47–152) was expressed in *E. coli*, purified under reducing conditions, and concentrated to 9–16 mg/ml for crystallization (see *SI Text* for details). Crystals grew in sitting drops comprising 2  $\mu$ l of protein solution and 2  $\mu$ l of reservoir solution (18% PEG-MME 2K, 0.1 M BisTris pH 7.2, 4  $\mu$ M TCEP). Crystals were transferred to reservoir solution made up with 30% MPD and plunged into liquid nitrogen. Data were collected from a selenomethionine-substituted crystal at SSRL beamline 7.2 and processed with *HKL2000* (45). Eight of the expected twelve Se sites were located using the auto solve option in *Phenix* (46) and phased to 3.0 Å resolution. The model was built using *COOT* (47) and refined to 2.6 Å with *REFMAC5* (48, 49) using translation/libration/screw (TLS) parameters (50) and programs in the CCP4 suite (51). The crystallized construct includes six additional residues (GIDPFT) at the N terminus that are not visible in electron density maps. The following residues lack defined density; molecule A (K47); B (K152); C (K47, K152); D (K47, A48, N49, K151, K152).

**Structure Determination of Oxidized BST2.** Human BST2(51–151) was expressed in HEK293T cells, purified, and concentrated to approximately 80 mg/ml (Bradford) for crystallization (see *SI Text* for details). MRC crystallization plate sitting drops were set up using a Mosquito crystallization robot by mixing 100 nl of protein solution with an equivalent amount of reservoir. Stacks of platelike crystals grew using a reservoir of 8% PEG8000, 100 mM Tris-HCl pH 8.5 at 18 °C. Crystals were vitrified in crystallization reservoir supplemented with 25% glycerol for data collection at the Diamond Light Source I04 beamline. Data were integrated and scaled using the programs *MOSFLM* (52) and *SCALA* (53) of the CCP4 suite. The structure of BST2(51–151) was determined by molecular replacement using *PHASER* (54) and residues 89–151 of reduced BST2(47–152) as the search model. 20 cycles of rigid-body refinement were followed by 10 cycles of positional refinement using *REFMAC5* (48, 49) with the occupancy of C91 residues set to 0.01.  $2mF_o - DF_c$  and  $mF_o - DF_c$  maps indicated the existence of the C91–C91 disulfide and helical density was visible N-terminal to the template. Model building used *COOT* (47). Final refinement cycles used secondary structure restraints between hydrogen bonded helical N-O atoms. *Phenix.refine* (55) was used to generate secondary structure restraints and to calculate maximum likelihood

averaged kick omit maps (56). The refined model lacks a substantial portion of the N terminus, which could not be reliably built, although residual density is present, particularly in proximity to neighboring molecules in the lattice (Fig. S5).

**Analytical Equilibrium Ultracentrifugation.** Data were collected at 4 °C in an Optima XL-A centrifuge (Beckman). Reduced BST2(47–154) samples were in a buffer containing 20 mM Tris pH 7.4, 300 mM NaCl, and 4 mM TCEP. Alternatively, BST2(47–154) was oxidized by extensive (1 week) dialysis against the same buffer lacking reductant. Formation of a disulfide-linked dimer upon oxidation was verified by mass spectrometry (Fig. S3). Oxidized and reduced protein samples were centrifuged at 12,000 rpm and 16,000 rpm with initial protein concentrations of 2.1, 1.06, and 0.53 mg/ml (166.7  $\mu$ M, 83.3  $\mu$ M, 41.7  $\mu$ M). Data were globally fit to ideal single species models with fixed or floating molecular masses using the non linear least squares algorithms in the *HETEROANALYSIS* software (57). Protein partial specific volumes and solvent densities were calculated with the program *SEDNTERP* (version 1.09) (58).

**BST2 Restriction of Virus Release and Infectivity.** 293T and HeLa-TZM reporter cells (obtained from Drs. J. C. Kappes and X. Wu through the AIDS Research and Reference Reagent Program, Division of AIDS, NIAID, NIH) were maintained using standard procedures. Wild-type and mutant BST2 proteins were expressed from a pCAG vector under the control of a constitutively active CMV promoter. The HIV-1( $\Delta$ Vpu) version of the HIV-1 molecular clone NL4-3 has been described previously (12). Virus was produced by calcium phosphate transfection of semiconfluent six well-plates of 293T cells with 1  $\mu$ g of viral DNA and with increasing quantities of vectors expressing wild-type BST2 WISP10-435 or BST2(L70D) WISP10-436 (0, 25, 50, 100, 200, and 500 ng) or an empty pCAG control vector.

For Western blotting experiments, virus-containing media was harvested 40 h posttransfection. Virions (1 ml) were pelleted through a 20% sucrose cushion at 15,000  $\times$  g, and the pellet was resuspended in 30  $\mu$ l 1X SDS/PAGE loading buffer (12  $\mu$ l loaded/lane). Cells from the same samples were harvested, pelleted, lysed in 60  $\mu$ l SDS-loading buffer, and boiled for 10 min (6  $\mu$ l loaded/lane). Levels of virion-associated CA and MA proteins in the culture media (Virus) and cellular Gag-derived and BST2 proteins (Cells) were analyzed by Western blotting using a mouse anti-BST2 (kindly provided by Chugai Pharmaceutical, 1:500 dilutions) or our own rabbit anti-HIV CA and MA antisera (both mixed at 1:1,000). The secondary antibodies were antimouse IgG (goat) polyclonal conjugated to Alexa680 (1:10,000, Invitrogen) and antirabbit IgG (donkey) polyclonal conjugated to IRdye800 (1:10,000, Rockland). Western blots were visualized and quantified using an Odyssey scanner (Li-Cor Biosciences).

Infectious titers were determined from viral supernatants by the numbers of  $\beta$ -galactosidase-positive foci formed on HeLa-TZM indicator cells. Titters provided in Fig. 4B show the mean  $\pm$  the range from two independent repetitions of the experiment.

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