Stabilization of TM Trimer Interactions during Activation of Moloney Murine Leukemia Virus Env

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The transmembrane subunit (TM) of the trimeric retrovirus Env complex is thought to direct virus-cell membrane fusion by refolding into a cell membrane-interacting, extended form that subsequently folds back on itself into a very stable trimer of hairpin-like TM polypeptides. However, so far there is only limited evidence for the formation of a stable TM trimer during Env activation. Here we have studied the oligomer composition and stability of an intermediate and the fully activated form of Moloney murine leukemia virus (Mo-MLV) Env. Activation of Mo-MLV Env is controlled by isomerization of its intersubunit disulfide. This results in surface subunit (SU) dissociation and TM refolding. If activation is done in the presence of an alkylator, this will modify the isomerization-active thiol in the SU of Env and arrest Env at an intermediate stage, the isomerization-arrested state (IAS) of its activation pathway. We generated IAS and fully activated Envs in vitro and in vivo and studied their states of oligomerization by two-dimensional blue native polyacrylamide gel electrophoresis (PAGE) and nonreducing sodium dodecyl sulfate (SDS)-PAGE. The IAS Env was composed of trimers of SU-TM complexes, whereas the activated Env consisted of SU monomers and TM trimers. When the oligomers were subjected to mild SDS treatment the TM trimer was found to be 3.5 times more resistant than the IAS oligomer. Thus, this demonstrates that a structural conversion of TM takes place during activation, which results in the formation of a stable TM trimer.

Viral membrane fusion proteins such as the influenza virus hemagglutinin (HA) and retrovirus envelope protein (Env) are made as trimers of a type I transmembrane protein in the endoplasmic reticulum of the infected cell (13, 38, 39). They become potentiated for activation by limited proteolytic processing of their ectodomain by furin when they pass the Golgi complex on their way to the site of virus budding in the cell (5, 11, 17, 24, 28, 32, 37). The cleavage generates a surface subunit (SU) and a transmembrane subunit (TM) of each fusion protein precursor. The SUs carry the receptor binding activity and the TMs the fusion activity. Studies on HA have shown that the latter activity is expressed by refolding of the TMs (30). Analysis of the native HA crystal structure demonstrated that the TM (HA2) subunits had a long central α-helix with which they formed a triple-stranded coiled-coil fusion protein (38). The N-terminal fusion peptide of HA2 was buried in a cavity on the surface of the HA stem and linked by a shorter α-helix and a loop to the N terminus of the long helix. Analysis of the crystal structure of a proteolytic product of activated HA showed that it consisted of a trimer of HA2 ectodomains devoid of the fusion peptide. The short helix and the loop, which connected the fusion peptide to the central helix of native HA2, extended the latter at its N-terminal end (1). Furthermore, in the middle of the central helix of native HA2 a loop had been formed that allowed chain reversal and packing of the C-terminal region of the HA2 polypeptide into the grooves of the coiled coil in an antiparallel orientation. From these data a model was created for how the HA2 subunits mediate membrane fusion. According to this, the HA2 subunits first extend the central helix. This will expel the fusion peptide on the top of the molecule, where it can interact with the cell membrane. At a second stage, the HA2 subunits make the back folding reaction. This will orient the C-terminal membrane anchors of the HA2 subunits into the same direction as their N-terminal fusion peptides and thereby facilitate approximation and fusion of the viral and the cell membranes.

The HA2 activation is suppressed in native HA by the associated SU HA1. Triggering follows when the HA1 subunit is displaced by the effects that the low pH has on the HA structure in endocytosed virus (10, 15). HA2 has been proposed to be arrested in its kinetic folding pathway in the native HA (2). Acid triggering overcomes the kinetic block and allows HA2 to fold into a stable conformation, e.g., the trimer of hairpin-like HA2 polypeptides. This has been supported by the increased thermostability of the activated TM trimer compared to native HA, the ability to trigger HA under neutral conditions by unspecified protein perturbation treatments such as heat and urea incubations, and the fact that recombinant HA2 ectodomain, produced in the absence of the HA1 subunit, folds into the low-pH form (2, 4, 26).

The Env of retrovirus is thought to control the membrane fusion by using its fusion protein SU and TM subunits in a similar way as influenza virus. This is corroborated by several findings. First, although there is no atomic structure of the native Env, analyses of crystals of ectodomain fragments of the retrovirus TM protein have demonstrated a similar thermostable complex of three hairpin-like polypeptide structures, as is found in the low-pH form of HA2 (3, 6, 16). Second, TM peptides corresponding to the interacting regions in the TM hairpin can effectively inhibit Env-mediated membrane fusion, suggesting a prehairpin-to-hairpin conversion of TM during
Env activation (9, 14, 18, 22, 35). Third, Env, which normally is activated by receptor binding or by the receptor and subsequent acid treatment in the endosome, can also be triggered by nonspecific protein perturbation (31, 33). This suggests that TM is metastable in native Env but folds into a stable trimeric complex as a result of activation. However, so far there is only limited evidence for the actual formation of a stable TM trimer during the activation of the complete Env complex. Indeed, there is only one report showing that the TM of the avian leukosis virus (ALV) Env converts into a trimer that resists sodium dodecyl sulfate (SDS) at up to 60°C upon activation (21). Here we have studied the stability of the Env trimer of Moloney murine leukemia virus (Mo-MLV).

Mo-MLV Env activation is controlled by isomerization of the intersubunit disulfide (25, 34). This is linked to a C-X-C motif in SU, where the other Cys carries a free thiol. Upon receptor binding, the latter becomes activated to attack the intersubunit disulfide and rearrange it into a disulfide isomer within the motif. This results in dissociation of the SU, refolding of TM, and membrane fusion. However, activation in the presence of an alkylator will, after initial receptor-induced changes in Env, modify the thiol before it has time to attack the intersubunit disulfide. The Env will now be blocked in isomerization and arrested at an intermediate stage of its activation process (isomerization-arrested state [IAS]) (34, 36). However, the fusion activity of Env can be rescued by reducing the intersubunit disulfide with dithiothreitol (DTT). The native state of Env is stabilized by Ca²⁺ ions, and receptor-induced isomerization involves removal of the suppressing Ca²⁺ ions from Env (34). Consequently, isomerization, i.e., Env activation, can also be induced in vitro by Ca²⁺ depletion and unspecific protein destabilization such as incubation in Ca²⁺-free medium, lysis with a buffer containing a nonionic detergent and EDTA, SDS, or treatment with urea or heat.

In the present work we activated Mo-MLV Env using solubilization of virus particles with mild detergent and EDTA in the presence or absence of alkylator to generate Env in the IAS or fully activated Env and determined their oligomeric state and stability toward dissociation by a low concentration of SDS. The native Env trimer could not be analyzed in this assay because SDS is a potent inducer of the disulfide isomerization reaction. We found that Env in the IAS was composed of SU-TM trimers and the fully activated form of TM trimers. Significantly, the TM trimers were 3.5 times more resistant toward dissociation than the SU-TM trimers of the IAS form of Env. This suggests that the TM of Mo-MLV refolds during Env activation into a form that supports stable trimerization.

MATERIALS AND METHODS

Cells and virus. XC (ATCC CCL-165, LGC Promochen, Boras, Sweden) and MO-V-3 (G. Schmidt, GSF-National Research Center for Environment and Health, Neuherberg, Germany) cells were maintained in Dulbecco's modified Eagle's medium ( Gibco BRL) (1 g/liter glucose for XC cells and 4.5 g/liter glucose for MO-V-3 cells) supplemented with 10% fetal calf serum, 20 mM HEPES, and 1-glutamine (growth medium). [35S]Cys-labeled wild-type Mo-MLV was prepared in MO-V-3 cells by overnight labeling with 90 μCi/ml [35S]Cys (GE Healthcare Biosciences, Buckinghamshire, United Kingdom) in the presence of 25 μM unlabeled cysteine as described previously (25). Virus in culture supernatant was purified by sedimentation in a step gradient of 1 ml 50% (wt/wt) and 4.5 ml 20% (wt/wt) sucrose in 50 mM HEPES, 100 mM NaCl (pH 7.4) (HN buffer) containing 1.8 mM CaCl₂ for 1 h at a relative centrifugal force at maximum radius of 2.2 × 10⁶ × g (26 k rpm) and 4°C in a Beckman SW 41 rotor.

In vitro activation of Mo-MLV Env, generation of IAS, DTT treatment, and cross-linking. Gradient-purified [35S]Cys-labeled Mo-MLV was activated in HN buffer containing one of the following combinations: 0.15% Triton X-100 and 9 mM EDTA, 0.15% Triton X-100 and 1.8 mM CaCl₂, 9 mM EDTA only, 1.8 mM CaCl₂ only, 36 mM octyl-β-D-glucopyranoside (octylglucoside [OG]) and 9 mM EDTA, or 36 mM OG only. Incubation was for 0, 15, 30, 60, or 90 min at 37°C. At the indicated time 20 mM N-ethylmaleimide (NEM) was added and the incubation continued up to 180 min at 37°C. Env blocked at the IAS was generated by including 20 mM NEM prior to activation incubation at 37°C. In order to reduce the intrasubunit disulfide bond of the TM subunit, TM trimers were generated from [35S]Cys-labeled Mo-MLV by incubation in HN buffer containing 0.15% Triton X-100 and 9 mM EDTA for 40 min at 37°C, and then 33 mM DTT was added, the incubation was continued for 20 min at 37°C, and finally 80 mM NEM was added. Cross-linking of activated Env and Env in the IAS was performed on [35S]Cys-labeled Mo-MLV that had been incubated for 20 min at 37°C in HN containing 36 mM OG and 9 mM EDTA without and with 20 mM NEM, respectively. Dithiobis(succinimidyl)propionate (DSP) (Pierce Co., Rockford, IL) in the range from 0 to 800 μM was added and the samples incubated for 30 min at room temperature. Excessive reagent was blocked by addition of 45 mM glycine and 5 min at room temperature. The oligomeric state of the proteins was analyzed, without further purification, by blue native polyacrylamide gel electrophoresis (BN-PAGE) in the absence or presence of an excess of SDS and by nonreducing SDS-PAGE.

Receptor-induced isomerization. Virus was bound to receptor-carrying XC cells grown on 24-well dishes (BD Falcon, Franklin Lakes, NJ) by spin inoculation. Briefly, the cells were washed once in phosphate-buffered saline (PBS), gradient-purified [35S]Cys-labeled Mo-MLV diluted in growth medium supplemented with 8 μg/ml Polybrene (250 μg/ml) and 1.8 mM CaCl₂ for 1 h at a relative centrifugal force at maximum radius of 1.3 × 10⁶ × g (2,900 rpm) in a Beckman JS 5.9 rotor. The cells were washed once in PBS, 1 ml growth medium was added, and the cells were incubated for 30 min at 37°C to allow receptor-induced activation of the Env complex. The cells were washed twice in ice-cold PBS and extracted with a buffer containing 0.15% Triton X-100, 0.75 M 6-amino-hexanoic acid, 9 mM EDTA, and 10 mM NEM for 10 min at 60°C in HN containing 36 mM OG and 9 mM EDTA. The cells were then transferred to an Eppendorf tube, insoluble matter was removed by low-speed centrifugation, and the supernatant was incubated for 30 min at 4°C.

Limited SDS treatment of Env protein complexes. Aliquots of virus-XC cell extracts, in vitro-activated Env (with or without DTT treatment), or Env in the IAS were incubated in the presence or absence of SDS (1.5 mol SDS/mol Triton X-100) for 10 min at room temperature. The samples were then split in two and prepared directly for BN-PAGE and nonreducing SDS-PAGE.

Electrophoresis and quantitations. BN-PAGE was performed essentially as described previously (20, 27). Solubilized samples were mixed with an equal volume of 2× BN sample buffer containing 100 mM morpholinepropanesulfonic acid (MOPS), 100 mM Tris-HCl, 40% glycerol, and 0.1% Serva blue G (pH 7.7) and incubated at room temperature for 10 min. The indicated samples from the cross-linking experiments were supplemented with 12 g/ml SDS and incubated at room temperature for 3 min at 70°C. All samples were separated on 4.5 to 16% acrylamide gel (of which 2.6% comprised bisacrylamide), gradient gels containing 50 mM bis-Tris (pH 7.0) and 0.5 M 6-amino-hexanoic acid at 4°C min at 200 V for 80 with 50 mM MOPS—50 mM Tris-HCl (pH 7.7) containing 0.002% Serva blue G as the cathode buffer and the same buffer without Serva blue G as the anode buffer. When cell extracts or SDS-containing samples were analyzed, 0.01% Serva blue G was used in the cathode buffer. 3-Methylated molecular mass marker proteins (CFA 626) and unlabeled standard proteins (HMW-native) from GE Healthcare Biosciences (Buckinghamshire, United Kingdom) were used. For two-dimenional (2D) BN-PAGE/nonreducing SDS-PAGE a sample was separated on BN-PAGE (0.75-mm by 10-cm by 8-cm gel), and the middle part of the lane was cut out and incubated first in 1 ml HN buffer containing 0.1 M NEM for 30 min on ice and then in 1 ml sample buffer containing 120 g/liter SDS, 0.19 M Tris-HCl (pH 8.0), 93 g/liter sucrose, 14 mM EDTA, 0.6 g/liter bromophenol blue, 0.4 g/liter methanethione, and 20 mM NEM for 30 min at room temperature. The gel strip was then fitted on an SDS-polyacrylamide gel (1 mm by 10 cm by 10.5 cm) (13% acrylamide, of which 2.6% comprised bisacrylamide) and electrophoresed under nonreducing conditions. All gels were dried and exposed to phosphor-imager screens (BAS-MS2025; Fujifilm, Science Imaging Scandinavia, Nacka, Sweden) and the labeled proteins visualized (and quantitated) using a Molecular Imager (BAS1800; Bio-Rad Laboratories, Hercules, CA). The extent of Env isomerization was estimated as described previously (29). The percentage of oligomeric TM was calculated as TM₁₀₀ in the presence (TM oligo) or absence (TM oligo-SDS) of SDS (see Fig. 4 for identities of spots). The percentage
of oligomeric SU-TM was calculated in the corresponding way. The difference in SDS sensitivity between the two complexes (S) was estimated as $S = \frac{(SU-TM_{\text{oligo}} - SDS) - (SU-TM_{\text{oligo}} + SDS)}{(TM_{\text{oligo}} - SDS) - (TM_{\text{oligo}} + SDS)}$.

RESULTS

Lysis-induced SU-TM disulfide isomerization. We chose to activate viral Env in our in vitro experiments by membrane lysis. We first studied the separate effects of Triton X-100 and EDTA in HN buffer on the intersubunit disulfide isomerization reaction. For this purpose, $[^{35}\text{S}]\text{Cys}$-labeled Mo-MLV was prepared in MOV-3 cells and purified by ultracentrifugation in a sucrose step gradient. The virus was then incubated in HN buffer with either Triton X-100 and EDTA, Triton X-100 and Ca$^{2+}$, or EDTA only for 0 to 180 min at 37°C. As a control we used incubation in HN buffer containing Ca$^{2+}$ (Fig. 1). At the indicated times, further isomerization was blocked by adding NEM and all samples incubated for a total of 180 min. The viral proteins were analyzed directly, i.e., without immunoprecipitation, by nonreducing SDS-PAGE. This revealed the disulfide isomerization as SU and TM subunits released from the covalently linked SU-TM complexes (Fig. 1A). The analysis showed that the EDTA treatment was much more effective at causing isomerization than Triton X-100 (Fig. 1B). Indeed, incubation with the nonionic detergent induced only slightly more isomerization than incubation in the control buffer. However, the greatest effect was obtained by combining Triton X-100 and EDTA in the HN buffer. A similar pattern was found for the Triton X-100-related detergent Nonidet P-40 (data not shown). We also tested OG, an unrelated mild detergent, in the HN buffer. This was shown to be much more efficient than Triton X-100 and caused efficient isomerization even in the absence of EDTA (Fig. 1C). We concluded that mild detergents differed significantly in their effects on Env activation. In subsequent in vitro experiments we activated Env using HN buffer containing EDTA and either Triton X-100 or OG.

Subunit structures of IAS and activated Envs. The oligomeric structures of IAS and activated Envs were compared by BN-PAGE. In contrast to SDS-PAGE, BN-PAGE preserves many noncovalent interactions between subunits in oligomeric complexes (27). Thus, $[^{35}\text{S}]\text{Cys}$-labeled Mo-MLV was lysed in OG-EDTA HN buffer in the presence or absence of NEM for 20 min at 37°C to generate IAS and activated Envs. The samples were then analyzed by BN-PAGE and nonreducing SDS-PAGE. The latter analyses showed the isomerization-arrested SU-TM complexes of IAS Env (Fig. 2A, lanes 1 and 2) and the isomerization-released SU and TM subunits of the activated Env (Fig. 2B, lanes 1 and 2). In the BN-PAGE analysis, most of the IAS Env migrated as a large complex and the rest as apparent SU-TM complex monomers (Fig. 2C, lane 1). The corresponding analysis of activated Env revealed the SU as monomers, whereas the TM could not be identified (Fig. 2D, lane 1). To further characterize the oligomeric complexes, we cross-linked the lysed samples with increasing concentrations of DSP and subsequently subjected them to SDS treatment. SDS treatment of un-cross-linked IAS Env lysate dissociated all large SU-TM oligomers seen in the BN-PAGE into apparent SU-TM complex monomers (Fig. 2C, lane 2). These migrated slightly faster than the monomers of the corresponding

FIG. 1. In vitro activation of SU-TM disulfide isomerization. $[^{35}\text{S}]\text{Cys}$-labeled Mo-MLV was incubated in HN buffer containing either Triton X-100 and EDTA, Triton X-100 and Ca$^{2+}$, EDTA, OG and EDTA, or OG and Ca$^{2+}$ for 0 to 180 min at 37°C. A control incubation was done in HN buffer containing Ca$^{2+}$. NEM was added, all samples were incubated for a total of 180 min, and viral proteins were analyzed by nonreducing SDS-PAGE. Shown are a phosphorimage of a gel with samples incubated in HN buffer with Triton X-100 and EDTA (A) and quantifications of the isomerization efficiencies under all conditions tested (B and C). In the gel analyses SU-TM complexes and free SU and TM subunits are indicated together with other viral proteins to the right and molecular weight standards to the left. The isomerization efficiencies ($\pm$ standard deviations; $n = 6$) are given as percentages of complete isomerization.
Upon cross-linking with increasing DSP concentrations, the SU-TM complex was found as increasing amounts of apparent dimers and trimers (Fig. 2C, lanes 3 to 6). This was reflected by the formation of very slowly migrating material in the nonreducing SDS-PAGE (Fig. 2A, lanes 3 to 6). The BN-PAGE analysis of the cross-linked and SDS-treated activated Env lysate showed only SU monomers with slightly increased mobility (Fig. 2D, lanes 3 to 6). However, the nonreducing SDS-PAGE analysis resolved the TM subunit as dimers and trimers with increasing cross-linking (Fig. 2B, lanes 3 to 6). We concluded that IAS Env was a trimer of the SU-TM complex and that the activated Env dissociated into SU monomers and a trimer of TM. The nonreducing SDS-PAGE analyses of the cross-linked samples also showed that the viral matrix (MA) protein was present as apparent dimers in virus lysed with or without NEM (Fig. 2A and B, lanes 5 and 6). This was confirmed by 2D nonreducing SDS-PAGE/reducing SDS-PAGE (data not shown).

Nonreducing SDS-PAGE analysis in the second dimension reveals the TM trimer in BN-PAGE of activated Env. Our aim was to compare the stabilities of the trimeric interaction in SU-TM trimers of IAS Env and in TM trimers of activated Env by a limited SDS treatment. An obvious approach would be to treat the respective trimers with a small amount of SDS, cross-link, and analyze the pattern of cross-linked oligomers compared to that in untreated controls by nonreducing SDS-PAGE. Unfortunately, mild SDS treatments proved to affect cross-linking per se more than the oligomeric status of the proteins. Therefore, we investigated whether BN-PAGE could be used for the separation and detection of not only the IAS Env trimer but also the TM trimer in its native state if the analysis was complemented with nonreducing SDS-PAGE in the second dimension. In this gel system proteins that keep the same oligomeric state through both dimensions will, in the nonreducing SDS-PAGE, form a diagonal front of faster- or slower-migrating spots depending on their size. Because BN-PAGE, in contrast to nonreducing SDS-PAGE, uses a gradient gel, the front will have a curved appearance. In contrast, proteins that are oligomeric in BN-PAGE and dissociate into subunits in nonreducing SDS-PAGE will migrate faster and form spots below the curved front. Thus, samples of IAS and activated Envs, prepared as described above, were subjected to separate BN-PAGE and nonreducing SDS-PAGE and to 2D BN-PAGE/nonreducing SDS-PAGE. The nonreducing SDS-PAGE showed the complete formation of isomerization-arrested SU-TM complexes in IAS Env and free SU and TM subunits in activated Env (Fig. 3A, lanes 1 and 2). The BN-PAGE demonstrated that most of the SU-TM complexes were associated into trimers, whereas only free SU and apparent capsid (CA) were clearly detectable in activated Env (Fig. 3B, lanes 1 and 2). However, occasionally, as in this experiment, we were able to discriminate a diffuse band in front of SU which was not present in the IAS Env sample and thus fit TM trimers. Indeed, we found that almost all apparent TM in the second-dimension gel of the 2D BN-PAGE/nonreducing SDS-PAGE migrated as being derived from such an oligomer (Fig. 3D). Monomeric TM subunits were expected to migrate in the lower part of the curved front formed by the spots of the apparently monomeric SU, CA, MA, and nucleocapsid proteins of the virus. The identities of the apparent TM and SU spots were confirmed by analyses of the IAS Env, which contained only covalently linked SU-TM complexes (Fig. 3C). These were mostly migrating below the 2D gel front, as expected for oligomers in the BN-PAGE dimension (Fig. 3C [lower contrast]). The composition of the TM oligomer was determined...
by cross-linking activated Env and subjecting it together with an un-cross-linked control sample to 2D BN-PAGE/nonreducing SDS-PAGE (Fig. 3E and F). We found that the predominant TM spot in the second-dimension gel was virtually absent and instead a strong spot appeared above, which migrated slightly slower than the 46-kDa marker protein (not shown), between SU and CA. This fit a trimer of the 15-kDa TM. Below the apparent TM trimer spot there was a weak spot, migrating slightly slower than CA, which fit TM dimers. The cross-linked TM trimer migrated somewhat slower than would be expected in the first dimension. This could be due either to the shape of the complex or to lower-than-normal Coomassie blue stain binding. The 2D gel analysis of the cross-linked sample also revealed the apparent MA dimers described above, now comigrating with CA monomers. We concluded that virtually all TM of the activated Env formed TM trimers and that the 2D gel analysis could be used as an assay to reliably detect trimeric TM in its non-cross-linked form.

The TM trimer of activated Env is more stable than the SU-TM trimer of IAS Env. To compare the stabilities of the SU-TM trimer of IAS and the TM trimer of activated Env, we prepared a virus lysate with Env partially in the IAS and partially in the activated form and subjected them to a limited SDS treatment. However, this was not possible with samples lysed with OG because of the high critical micellar concentration of this detergent. Therefore we had to switch to HN buffer.

FIG. 3. Detection of TM trimers of activated Env by 2D BN-PAGE/nonreducing SDS-PAGE. Samples of [\textsuperscript{35}S]Cys-labeled Mo-MLV were lysed in OG-EDTA HN buffer in the presence or absence of NEM for 20 min at 37°C to generate IAS Env and activated Env. A portion of the latter sample was subsequently cross-linked with DSP. Samples were subjected to 1D and 2D PAGE. (A and B) Phosphorimages of nonreducing SDS-PAGE and BN-PAGE of IAS Env (lanes 1) and activated Env (lanes 2). (C and D) 2D BN-PAGE/nonreducing SDS-PAGE of IAS Env (C) and activated Env (D). (E and F) 2D BN-PAGE/nonreducing SDS-PAGE of un-cross-linked (E) and cross-linked (F) activated Env. (C') Cutout of the marked region of the second-dimension gel with SU-TM complexes at lower contrast. The oligomeric states of the complexes in the first dimension (BN-PAGE) are indicated. The marker lane (1D) in the 2D BN-PAGE/nonreducing SDS-PAGE contains the respective sample separated in nonreducing SDS-PAGE only. Viral proteins and their oligomeric state (C) in the first dimension (BN-PAGE) are indicated.
with Triton X-100, which has a much lower critical micellar concentration. Thus, a labeled virus sample was partially activated in Triton X-100–EDTA HN buffer before NEM was added, and the remaining unactivated Env was chased into the IAS form. One half of the sample was subjected to mild treatment with SDS and the other half left untreated. The samples were analyzed by 2D BN-PAGE/nonreducing SDS-PAGE. Shown are phosphorimages of the gel analyses of untreated (A) and SDS-treated (B) samples. Viral proteins and their oligomeric state in the first dimension (BN-PAGE) are indicated to the right. (A’, A”, B’ and B”) Cutouts with the oligomeric state of the proteins in the first dimension indicated. (C) Relative amounts of the trimeric forms of the SU-TM complexes and the TM subunits in the two samples as percentages of the total SU-TM and TM, respectively (mean ± standard deviation; n = 6).

The stability of the TM trimer of activated Env is dependent on its intrasubunit disulfide. The TM subunit of MLV contains a single disulfide between C86 and C93 (7). This covalent linkage is necessary for the high thermostability of the trimer of hairpins found in the crystallized TM ectodomain fragment, which is presumed to reflect the structure of TM in fully activated Env (6, 7). Therefore we tested whether DTT treatment of the TM trimers of in vitro-activated Env would destabilize the oligomer. We prepared a virus lysate that was fully activated in vitro and then incubated with DTT before it was subjected to mild SDS treatment. The sample was analyzed by 2D BN-PAGE/nonreducing SDS-PAGE, which showed that more than half of the TM had shifted from the oligomer location into monomers (Fig. 5). This indicated that the stability of the in vitro-activated TM trimer was dependent on the intrasubunit disulfide and suggested structural similarities with the crystallized form.

Receptor activation of Env results in the formation of a stable TM trimer. Incubation of Mo-MLV with receptor-pos-
Receptor triggering of Env results in the formation of stable TM trimers. [35S]Cys-labeled Mo-MLV was bound by centrifugation to cultures of XC cells at 4°C and then incubated for 30 min at 37°C. NEM was added and the virus-cell samples lysed with Triton X-100–EDTA HN buffer. One sample was then mildly treated with SDS and the other one left untreated. Both samples were analyzed by 2D BN-PAGE/nonreducing SDS-PAGE. Shown are phosphorimages of the gel analyses of the untreated sample (A) and the SDS-treated sample (B). Marker lanes (1D), cutouts, and labeling as in Fig. 4. Note that A’ and B’ are shown in higher contrast to better visualize the TM subunit. (C) Relative amounts of the oligomeric forms of the SU-TM complexes and the TM subunits in the two samples as percentages of the SU-TM and TM, respectively (mean ± standard deviation; n = 4).

DISCUSSION

Our present analyses show that activation of the Mo-MLV Env results in the formation of a TM trimer, which was much more resistant toward SDS dissociation than the trimer of the IAS activation-intermediate form of Env. This provides direct biochemical evidence that a structural conversion of TM takes place during activation, which results in the formation of a stable TM trimer. Corresponding evidence has earlier been obtained only for the TM of the ALV Env (21). Together with previous structural analyses of TM ectodomain fragments and of TM peptide dependent inhibition studies, our analyses of oligomer stability provide strong support for an HA-like activation model for retrovirus Env also.

Earlier we showed that Mo-MLV Env in the IAS, in contrast to native Env and Env arrested at a late stage by lipid membrane modification, was able to complex with externally added TM peptides and thus inhibit reactivation of the IAS upon reduction of the intersubunit disulfide with DTT (35). Additionally, the native Env, in contrast to the IAS Env, has been found to hide the isomerization-active CXXC-thiol and intersubunit disulfide for external modification (34). These data together with the present study showing that the oligomeric interactions of IAS Env are significantly weaker than those of the activated TM trimer suggest that IAS Env is an open structure with its TM subunits possibly in an extended prehairpin conformation. This might correspond to the postulated HA activation intermediate where the fusion peptide connecting the α-helix and loop have extended the central helix of HA2 and expelled the fusion peptide, but the jackknife-like turn of HA2 has not yet taken place (30). Although we have not yet localized the fusion peptide in the Mo-MLV IAS Env, such an intermediate might represent a crucial step, especially in Mo-MLV Env activation. The intermediate would allow the TM to establish an interaction with the target membrane before the contact via SU is lost by isomerization of the intersubunit...
disulfide and before the TM undergoes its backfolding reaction into a stable trimer. In the case of ALV, which is activated by receptor binding and low pH in forming prehairpin and hairpin conformations of a retroviral envelope glycoprotein. J. Virol. 70:8201–8209.


and refolding induced by the synergistic effects of receptor binding and low pH is coupled to infection. J. Virol. 78:1403–1410.


