The Transmembrane Domain of Influenza Hemagglutinin Exhibits a Stringent Length Requirement to Support the Hemifusion to Fusion Transition

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Abstract. Glycosylphosphatidylinositol-anchored influenza hemagglutinin (GPI-HA) mediates hemifusion, whereas chimeras with foreign transmembrane (TM) domains mediate full fusion. A possible explanation for these observations is that the TM domain must be a critical length in order for HA to promote full fusion. To test this hypothesis, we analyzed biochemical properties and fusion phenotypes of HA with alterations in its 27–amino acid TM domain. Our mutants included sequential 2–amino acid (Δ2–Δ14) and an 11–amino acid deletion from the COOH-terminal end, deletions of 6 or 8 amino acids from the NH2-terminal and middle regions, and a deletion of 12 amino acids from the NH2-terminal end of the TM domain. We also made several point mutations in the TM domain. All of the mutants except Δ14 were expressed at the cell surface and displayed biochemical properties virtually identical to wild-type HA. All the mutants that were expressed at the cell surface promoted full fusion, with the notable exception of deletions of >10 amino acids. A mutant in which 11 amino acids were deleted was severely impaired in promoting full fusion. Mutants in which 12 amino acids were deleted (from either end) mediated only hemifusion. Hence, a TM domain of 17 amino acids is needed to efficiently promote full fusion. Addition of either the hydrophilic HA cytoplasmic tail sequence or a single arginine to Δ12 HA, the hemifusion mutant that terminates with 15 (hydrophobic) amino acids of the HA TM domain, restored full fusion activity. Our data support a model in which the TM domain must span the bilayer to promote full fusion.

Key words: hemagglutinin • hemifusion • transmembrane domain • glycosylphosphatidylinositol anchor • SNARE proteins

Introduction
Influenza virus fusion is mediated by the hemagglutinin (HA)1 trimer (for reviews see Stegmann, 1994; Gaudin et al., 1995; Hughson, 1995; Hernandez et al., 1996). Each HA monomer is composed of two subunits: HA1, which contains the receptor binding and major antigenic sites, and HA2, which is primarily responsible for fusion. HA2 contains an NH2-terminal fusion peptide, a region of high α-helical propensity, a 27–amino acid transmembrane (TM) domain, and a 10–amino acid cytoplasmic tail. Studies have demonstrated the importance of the fusion peptide as well as structural changes within the trimeric coiled coil for fusion (Carr and Kim, 1993; Bullough et al., 1994; Steinhauser et al., 1995; Qiao et al., 1998, 1999).

In previous work, we demonstrated that replacing the TM and cytoplasmic tail domains of HA with a glycosylphosphatidylinositol (GPI) anchor generated an HA trimer that could promote only hemifusion (Kemble et al., 1994). This suggested that the TM domain plays an important role during the hemifusion to fusion transition (Kemble et al., 1994; Melikyan et al., 1995b, 1997a; Blumenthal et al., 1996; Nussler et al., 1997; Chernomordik et al., 1998). Other studies have demonstrated that replacing the HA TM domain (and/or cytoplasmic tail) with those from foreign proteins, both viral and nonviral, has no effect on fusion (Roth et al., 1986; Dong et al., 1992; Schrot-Diez et al., 1998; Melikyan et al., 1999). Although the aforementioned conclusions have been based on studies using different assays, the collective findings suggest that there may not be any specific sequence requirements for the HA TM domain to support full fusion (Roth et al., 1986; Dong et al., 1992; Schrot-Diez et al., 1998; Melikyan et al., 1999). However, the HA TM domain may re-
quire a minimal length to promote the hemifusion to fusion transition. The major goal of this study was to test the latter hypothesis.

Materials and Methods

Mutagenesis

HA mutants were generated in HA cDNA (X-cl strain) present in the pTM1 vector using the Quik-Change Site-Directed Mutagenesis kit (Stratagene) according to the manufacturer’s instructions. Oligonucleotide primers with stop codons in the TM domain were used to generate cytoplasmic tail-α HA (Tail-HA) and then, sequentially, two-amino acid deletions from the COOH-terminal end of the TM domain (Δ2–Δ14; see Fig. 1). Oligonucleotide primers were also used to create the following additional HA mutants (in the Tail-HA construct): a deletion of 6 amino acids from the NH2-terminal end of the TM domain (Δ185–190; NΔ6); a deletion of 6 amino acids from the NH2-terminal end and 2 amino acids from the COOH-terminal end of TM (Δ185–190; Δ210–211; NΔ6Δ2); a deletion of 12 amino acids from the NH2-terminal end of TM (Δ185–196; NΔ12); deletions of 6 or 8 amino acids from the central region of TM (Δ195–200; MΔ6Δ6; and 195–202; MΔ8Δ8); a deletion of 11 amino acids from the COOH-terminal end of TM (Δ1011–111; Δ11); 6 single point mutants (S194L, N194A, G204A, G204L, W185A, and W188A); and two double point mutants (W185A/W188A and S194L/G204L) in the HA TM domain. The point mutants were made in the context of the full-length HA construct (i.e., containing the cytoplasmic tail). We also engineered two HA mutants that contained the TM domain of Δ2 HA (amino acids 185–189), followed by either the entire cytoplasmic tail sequence of HA (Δ2 Tail-HA) or a single arginine (Δ12Arg HA). In this paper we use the term GPHI-HA to refer to the construct BHA-PI (KS) described in Kemble et al. (1994). When GPHI-HA is expressed, a nine-amino acid sequence from the decay-accelerating factor GPI anchor addition signal, containing a lysine to serine substitution, remains with the HA ectodomain (Kemble et al., 1994). GPI-HA was subcloned into the pTM1 vector. Mutant HA cDNAs were sequenced to confirm that the desired mutations had, but that second site mutations had not, been introduced.

Expression of Wild-Type HA and Mutant HAs

CV-1 cells (CCL 70; American Type Culture Collection) were maintained in Iscove’s modified Dulbecco’s medium (IMDM; GIBCO BRL) containing 10% supplemented calf serum (SCS; Hyclone Laboratories, Inc.), 50,000 U penicillin, 50,000 μg streptomycin (GIBCO BRL), and an additional 146 mg glutamine (GIBCO BRL) per 0.5 liter. Wild-type (WT) and mutant HAs were expressed using the vaccinia virus T7 RNA polymerase transient transfection system (Fuerst et al., 1986). Confluent monolayers of CV-1 cells were infected with modified vaccinia ankara (MVA; a gift of Barber and Moss, National Institutes of Health, Bethesda, MD) at a multiplicity of infection of 10 PFU per cell and incubated at 37°C for 1 h with intermittent rocking. After removing the virus inoculum, the cells were washed once with Dulbecco’s PBS (Cellgro; Fisher Scientific) and then transfected with cDNA using 12.0 μl TransIT (Panvera) per 6-cm dish according to the manufacturer’s instructions. Unless otherwise stated, we used 5.0 μg cDNA per 6-cm dish except for Δ10, Δ12, and NΔ12, for which we used 7.5 μg cDNA per 6-cm dish. After a 5-h incubation (at 37°C in a 5% CO2 incubator), the DNA/TransIT mixture was replaced with IMDM, and the cells were incubated at 31°C for 15–20 h.

Metabolic Labeling

CV-1 cells expressing WT and mutant HAs were metabolically labeled with [35S]Translabel (ICN Biomedicals) essentially as described previously (Kemble et al., 1993). After transfection, the cells were incubated for 2 h in Cys/Met–DMEM (GIBCO BRL) containing 2% SCS at 37°C. The medium was then replaced with 1.25 ml Cys/Met–DMEM containing 50–75 μCi [35S]Translabel and 2% SCS, and the cells were incubated at 31°C for 14–18 h.

Cell Surface Biotinylation, Immunoprecipitations, and Western Blot Analyses

Biotinylation of cell surface proteins was performed as described previously (Qiao et al., 1999). Trypsin cleavage of HA0 was performed as described previously (Qiao et al., 1998) except that 10 μg/ml trypsin in PBS was used and incubation was for 10 min at room temperature (RT). Cells were then treated with 50 μg/ml soybean trypsin inhibitor (STI; Sigma-Aldrich) for 10 min at RT. CV-1 cells expressing WT and mutant HAs were washed with PBS, lysed in a cell lysis buffer containing 1% NP-40 and protease inhibitors as described previously (Delos et al., 2000), and then immunoprecipitated as described (Kemble et al., 1993). Immune complexes were suspended in SDS gel loading buffer containing 0.14 M β-mercaptoethanol, boiled for 5 min, and separated by 10% SDS-PAGE. These gels were transferred to nitrocellulose, blocked using superblotto (0.5% Tween 20, 3% [wt/vol] BSA, 18% [wt/vol] glucose, 1% [wt/vol] milk, and 10% glycerol in PBS) and probed with antibodies to HA (C-HA1) or streptavidin-HRP (Pierce Chemical Co.) as described previously (Qiao et al., 1999).

Sucrose Gradient Analysis

CV-1 cells expressing WT and mutant HAs were treated with trypsin, then STI, and lysed as described above. Cell lysates were layered on continuous 3–30% sucrose (wt/vol) gradients. After centrifugation, 12 395 μl fractions were collected and prepared as described previously (Qiao et al., 1998).

C-HA1 Conformational Change Assay

Transfected CV-1 cells were metabolically labeled overnight as described above. After treatment with trypsin and STI (see above), the cells were incubated at 37°C for 10 min in fusion buffer (100 mM NaCl, 10 mM Hepes, 10 mM MES, 10 mM succinate, and 2 mg/ml glucose) adjusted to the indicated pH. After renaturation with pH 7.0 fusion buffer, the cells were lysed and immunoprecipitated with the C-HA1 antibody as described previously (Kemble et al., 1993). Samples were analyzed by SDS-PAGE and PhosphorImager analysis (Molecular Dynamics).

RBC Labeling, Binding, and Lipid and Content Mixing Assays

Freshly collected human RBCs were either colabeled with octadecylrhodamine B chloride (R18) and carboxyfluorescein (CF; Molecular Probes, Inc.) or labeled with CF only (Melikyan et al., 1999). WT and mutant HA-expressing cells were treated with trypsin and STI as described above, washed once with PBS+ (PBS containing 0.1 g/liter CaCl2, and 0.1 g/liter MgCl2), and incubated with a solution containing 0.05% labeled RBCs for 15 min at RT. Unbound RBCs were removed by three washes with PBS+ and fusion was triggered by incubating the HA-expressing cells with pH 5.0 fusion buffer at 37°C for either 2 or 5 min (times indicated in the figure legends). The pH 5.0 solution was replaced with pH 7.0 fusion buffer, and the cells were examined with a fluorescence microscope. Where indicated, cells were treated for 1 min at RT with either 0.1 or 0.5 mM chloromazine (CPZ) in fusion buffer, pH 7.0, and then returned to PBS+. Images were collected using an Axioplan 2 microscope (Carl Zeiss, Inc.) equipped with a C4742-95 CCD camera (Hamamatsu), and Openlab (Improvement) software, and were prepared using Adobe Photoshop.

Preparation of Microsomal Membranes

Transfected CV-1 cells were biotinylated and treated with trypsin and STI as described above. The cells were then released from their dishes by incubation for 10 min at RT in 1.0 ml PEEG (PBS containing 0.5 mM EDTA, 0.5 mM EGTA, and 10 mM glucose), transferred to a 1.5-ml Eppendorf tube, pelleted at 325 g for 2 min at 4°C, resuspended in 0.8 ml DBH buffer (10 mM Tris-HCl, pH 7.5, 1 mM MgCl2), and incubated for 5 min on ice (to induce cell swelling). The cells were then passed 10 times through a 25-gauge needle. Surose was added to bring the solution to a final 1.18 M (wt/wt) concentration by the addition of 2.0 M sucrose (in DBH), and the suspension was overlaid with 3.0 ml of 0.25 M sucrose (in DBH). The nuclei were pelleted by centrifugation in an SW55 rotor at 192,000 g for 90 min at 4°C. The interface containing the microsomal membrane fraction was collected, transferred to a tube containing 4.0 ml of 0.25 M sucrose (in DBH), and centrifuged as before. The pellet containing the microsomal membranes was collected.

Carbonate Extraction

The pellet containing the microsomal membranes was resuspended in 0.3 ml 50 mM TEA, pH 7.5 (triethanolamine, pH adjusted with acetic acid). The pH was adjusted to 11.0 by the addition of 0.1 volume 1 M Na2CO3 (pH 11.0), and the suspension was incubated on ice for 20 min. Mem-
branes were layered on top of a 0.68-mI sucrose cushion (0.2 M sucrose, 20 mM Hepes-NaOH (pH 11.0), 150 mM potassium acetate, 2.5 mM magnesium acetate) and centrifuged at 135,000 g in a TLS-55 rotor for 20 min at 4°C. The supernatant was collected and reneutralized by the addition of 30 mM HCl and designated the “supernatant fraction.” The pellet was washed with 0.5 ml lysis buffer containing protease inhibitors, incubated on ice for 20 min, centrifuged to clear debris at 16,000 × g for 10 min at 4°C, and transferred to a fresh 1.5-ml Eppendorf tube and designated “pellet fraction.” FACS® Analysis

Transfected CV-1 cells (6-cm dishes) were released from the dish with PEEG (as described above) and centrifuged to a 1.5-ml Eppendorf tube. The cells were then washed twice with cold PBS® containing 0.02% azide (PBSA) and centrifuged at 325 g for 2 min at 4°C. The cells were resuspended in 0.2 ml cold PBSA containing 2% SCS and incubated for 30 min on ice with 1.7 ml of 1.0 mg/ml Site A mAb. The cells were then washed twice with PBSA as described above, resuspended in PBSA containing 2% SCS, and incubated with 1.0 ml FITC-conjugated goat anti-mouse IgG for 30 min on ice. The cells were then washed twice with cold PBSA, resuspended in PBS containing 2% paraformaldehyde, and analyzed by FACS® at the University of Virginia Core Facility, using a FACSScan™ flow cytometer (Becton Dickinson). Endo F Treatment

Transfected CV-1 cells (6-cm dishes) were metabolically labeled and treated with trypsin and STI as described above. The cells were released from the dish by a brief treatment with PEEG (as described above) and transferred to a 1.5-ml Eppendorf tube. Lysates were prepared and HA immunoprecipitated with the Site A mAb as described above. After the immunoprecipitation, 50 μl N-Glycosidase F buffer (1% octylglucoside, 0.2% SDS, 40 mM Tris, pH 8.0, 5 mM EDTA, and 1% β-mercaptoethanol) was added to the protein A–agarose (PAA) beads. The beads were then washed twice with PBSA as described above, resuspended in PBSA containing 2% SCS, and incubated with 1.0 μl FITC-conjugated goat anti-mouse IgG for 30 min on ice. The cells were then washed twice with cold PBSA, resuspended in PBS containing 2% paraformaldehyde, and analyzed by FACS® at the University of Virginia Core Facility, using a FACSScan™ flow cytometer (Becton Dickinson).

Figure 1. HA TM domain truncation mutants. Line diagram of the HA gene. The region encompassing the TM domain (gray box) is expanded below. Deletion mutations are shown as sequential removal of two amino acids from the COOH-terminal end of the TM domain (mutants Δ2–14) starting with the tail construct. Nα12 HA lacks the NH2-terminal 12 amino acids of the TM domain as well as the cytoplasmic tail. GPI-HA has been aligned with the correct ectodomain sequences of the TM truncation mutants.

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higher molecular weight band seen in some of the gradients corresponds to intracellular HA0.

To address whether the mutant HAs change conformation at the same pH as WT HA, HA-expressing cells were briefly incubated at the indicated pH, lysates were prepared, and HA was immunoprecipitated using C-HA1, a conformation-specific antibody (White and Wilson, 1987). As seen in Fig. 3 B, Δ2–Δ12 HA changed conformation with a pH dependence similar to that of WT HA.

Fusion Activity of Δ2–Δ12

We evaluated the fusion activity of Δ2, Δ4, Δ6, Δ8, Δ10, and Δ12 HA using a dye transfer assay. RBCs colabeled with a lipid dye (R18) and a soluble content dye (CF) were bound to HA-expressing cells, and fusion was induced as described in Materials and Methods. After 5 min at 37°C and pH 5, the cells were returned to neutral pH medium and examined with a fluorescence microscope. As seen in Fig. 4, both dyes transferred efficiently to cells expressing Δ2, Δ4, Δ6, Δ8, and Δ10 HA. A different phenotype was seen for cells expressing Δ12 HA: whereas we observed efficient lipid dye transfer (97%), content dye transfer was severely restricted (<10 vs. 97% for WT HA). As expected, we did not observe transfer of R18 or CF by WT or mutant HA-expressing cells at neutral pH or at low pH if the cells had not been pretreated with trypsin to process HA0 (data not shown).

Given the striking observation that Δ10 HA mediated robust lipid and content mixing whereas Δ12 HA mediated robust lipid mixing with minimal content mixing, we...
also tested a mutant lacking the 11 COOH-terminal residues of the TM domain (Δ11 HA). Δ11 HA exhibited biochemical properties similar to WT HA (Table I). Whereas Δ11 HA mediated efficient lipid mixing (Fig. 4), it was significantly impaired in its ability to mediate content dye transfer (Fig. 4).

The density of HA at the cell surface can influence the fusion phenotype (Ellens et al., 1990; Melikyan et al., 1995a; Blumenthal et al., 1996; Danieli et al., 1996). Therefore, we took care to analyze the fusion phenotype of mutant HAs expressed at comparable levels to a known amount of WT HA such that the level of WT HA expression at the cell surface was equivalent to that of 3.5 μg WT HA cDNA.

Figure 4. Quantitation of lipid mixing and content mixing. CV-1 cells expressing WT or mutant HAs were prepared for fusion as indicated in Materials and Methods, except that fusion was triggered at pH 5.0 for 2 min at 37°C and renaturalized. The amount of cDNA used per transfection is indicated. Lipid and content mixing events were averaged from 4–12 random fields (mean ± SEM). Percent lipid dye transfer (hatched bars) was determined by dividing the number of cells receiving lipid dye by the number of cells with bound RBCs in each field. Percent content dye transfer (black bars) was determined by dividing the number of cells receiving content dye by the number of cells with bound RBCs in each field. Relative surface expression of HA in fluorescence units (FU), was obtained by FACS® analyses, and is presented as the mean fluorescence intensity per cell normalized to that of 3.5 μg WT HA cDNA.

Table I. Summary of Results: Effects of Truncation Mutations in the HA TM

| HA          | Trimer cleavage | Trypsin binding | RBC binding | Lipid mixing | Content mixing | ConfΔ |
|-------------|----------------|-----------------|-------------|--------------|----------------|-------|
| WT (3.5 μg) | +              | +               | ++          | ++           | ++             | 5.0   |
| WT (0.5 μg) | +              | +               | +           | ++           | ++             | 5.0   |
| TailΔ       | +              | +               | +           | ++           | ++             | 5.0   |
| Δ2          | +              | +               | +           | ++           | ++             | 5.0   |
| Δ4          | +              | +               | +           | ++           | ++             | 5.0   |
| Δ6          | +              | +               | +           | ++           | ++             | 5.0   |
| Δ8          | +              | +               | +           | ++           | ++             | 5.0   |
| Δ10         | +              | +               | +           | ++           | ++             | 5.0   |
| Δ11         | ND             | +               | +           | +            | --             | ND    |
| Δ12         | +              | +               | +           | --           | --             | 5.0   |
| NΔ12        | ND             | +               | +/−         | --           | --             | ND    |
| Δ12Tail     | ND             | +               | +           | ++           | ++             | 5.0   |
| GPI-HA      | +              | +               | +           | +            | +              | 5.0Δ  |

The transfection efficiency of the HA TM domain mutants ranged from 61 to 78% (SEM was <5%). Expression at the cell surface is indicated in Fig. 4. ND, not done. 3.5 μg WT refers to the higher amounts of cDNA used in transfections and with which comparison with TailΔ HA and Δ2–Δ10 should be made. 0.5 μg WT refers to the lower amount of cDNA used in transfections and with which comparison with Δ11, Δ12, NΔ12, and Δ12Tail should be made.

*Kemble et al., 1993.

NΔ12 HA, was then examined for biochemical properties. Like all of the other truncation mutants, NΔ12 HA was expressed at the cell surface, was processed by trypsin into HA1 and HA2, and exhibited a faster migrating HA2 subunit than tail-HA (Table I, and data not shown). By all of these criteria, NΔ12 HA resembled Δ12 HA. However, it was not as well expressed at the cell surface as Δ12 HA, as determined by FACS® analysis and RBC binding (~80% compared with Δ12 HA; Fig. 4 and Table I). In terms of fusion with RBCs, NΔ12 HA mediated significant lipid mixing, albeit less than seen with Δ12 HA (63 vs. 97%). With respect to content mixing, NΔ12 HA mediated <5% dye transfer similar to the behavior of Δ12 HA and GPI-HA (Figs. 4 and 5). Neither Δ12 HA nor NΔ12 HA promoted significant content mixing (>10%) even after 60 min of incubation at 37°C at either pH 4.8 or 5.0 (data not shown).

Comparison of the Lipid and Content Mixing Ability of D10 HA and WT HA

Given the dramatic decrease in content mixing ability between Δ10 HA and Δ12 HA (and Δ11 HA), we examined the fusion activity of Δ10 HA in more detail. For this purpose, we compared the lag times, initial rate, and final extent of lipid mixing with WT HA and Δ10 HA at different pH values. At all pH values tested, the lag time before the onset of dye transfer was equivalent for Δ10 HA and WT HA (Table II). There was no difference in either the initial rate or the final extent of lipid mixing for Δ10 HA and WT HA at pH 5.0 and 5.25. At pH 5.5, the latter parameter was somewhat lower for Δ10 HA. Hence, the lipid mixing properties of Δ10 HA were very similar to those of WT HA at all pH values tested. In addition, when incubated at the suboptimal pH of 5.25 for 2 min at 37°C, Δ10 HA mediated content mixing to the same extent as WT HA (data not shown).

Effect of CPZ on Content Mixing

Previous work has shown that treatment with 0.1 mM CPZ, a membrane-permeable amphipathic reagent that
partitions preferentially into the inner leaflet of the plasma membrane, efficiently induces full fusion in cases of “stunted fusion” caused by performing fusion experiments under suboptimal conditions (Melikyan et al., 1997a). Stunted fusion is operationally defined as lipid mixing without substantial content mixing due to the formation of small or transient fusion pores. It is thought to occur after hemifusion. In contrast, higher concentrations of CPZ (0.4–0.5 mM) are needed to induce GPI-HA to promote content mixing, and the extent of content mixing seen with GPI-HA in the presence of 0.4–0.5 mM CPZ never reaches that seen with WT HA (Melikyan et al., 1997a). Therefore, we assessed the effects of 0.1 and 0.5 mM CPZ on the ability of Δ12 HA, NΔ12 HA, and GPI-HA to promote content transfer. After binding double-labeled RBCs (R18 and CF) to HA-expressing cells, fusion was triggered by lowering the pH to 5.0 for 5 min at 37°C. The medium was reneutralized, and CPZ was added to the cells at neutral pH. After a 1-min incubation at RT, the CPZ solution was replaced with PBS+. The percentage of R18-stained HA-expressing cells that became labeled with CF was then determined. In the absence of CPZ, ∼1, 7, and 3% of cells expressing NΔ12 HA, Δ12 HA, and GPI-HA, respectively, received aqueous dye (Fig. 6). The addition of 0.1 mM CPZ increased content dye transfer to ∼5, 8, and 4%, respectively. Addition of 0.5 mM CPZ induced a greater extent of CF transfer: ∼20, 27, and 22%, respectively. Representative images of aqueous dye transfer before and after the addition of either 0.1 or 0.5 mM CPZ are shown in Fig. 7. Hence, cells expressing NΔ12 HA and Δ12 HA respond similarly to CPZ as do cells expressing GPI-HA in terms of their ability to promote aqueous dye transfer. The presence of R18 in the RBC membrane augments the transfer of aqueous contents to GPI-HA–expressing cells (Markosyan et al., 2000). Therefore, we assessed content dye transfer from RBCs that were not labeled with R18. As expected, content dye transfer under these conditions was less than with double-labeled RBCs (data not shown). Most importantly, Δ12 HA and NΔ12 HA still responded similarly to CPZ as did GPI-HA: a brief treatment with 0.5 but not 0.1 mM CPZ increased content dye transfer (data not shown).

Membrane Association of Δ12 HA

Given the striking phenotype of HA lacking 12 amino acids in the TM domain (lipid, but not content, mixing), we explored how Δ12 HA is anchored in the membrane. Like WT HA, Δ12 HA (as well as GPI-HA) was resistant to carbonate extraction (Fig. 8A). Given that some GPI-anchored proteins associate with cholesterol and sphingomyelin-rich detergent-insoluble membrane fractions (DIGs; Simons and Ikonen, 1997; Friedrichson and Kurzchalia, 1998; Varma and Mayor, 1998), we examined the solubility

Table II. Lipid Mixing of WT versus Δ10 HA

| pH | HA   | Lag time | Initial rate of fusion | Final extent |
|----|------|----------|------------------------|--------------|
| 5.0| WT   | 45       | 1.0                    | 100 ± 4      |
|    | Δ10  | 50       | 1.13                   | 104 ± 1.3    |
| 5.25| WT   | 65       | 1.01                   | 83 ± 6       |
|     | Δ10  | 70       | 0.96                   | 83 ± 6       |
| 5.5| WT   | 80       | 0.74                   | 71 ± 4       |
|    | Δ10  | 80       | 0.67                   | 58 ± 6       |

Lag time indicates the amount of time before initial spread of R18. Initial rate of fusion is the change in intensity over time (∆D/∆t) for the initial rise in fluorescence relative to WT HA at pH 5.0. Final extent is the percent intensity values (± SEM) relative to WT HA at pH 5.0 at the highest extent of fusion. The data were calculated from two independent experiments. See Materials and Methods for details.
of Δ12 HA and NΔ12 HA in Triton X-100 at 4°C before and after treating cells with methyl β-cyclodextrin to remove cholesterol (Scheiffele et al., 1997). Proteins that associate with lipid raft microdomains are often relatively insoluble in 1% Triton X-100 in the cold (Simons and Ikonen, 1997). Cholesterol depletion can increase the solubility of these proteins in Triton X-100, presumably by disrupting the raft microdomains (Scheiffele et al., 1997). Both Δ12 HA and NΔ12 HA (data not shown) were readily solubilized by Triton X-100 at 4°C, suggesting that they do not associate with DIGs. In contrast, both WT HA and GPI-HA were partially insoluble in Triton X-100 at 4°C, and depletion of cholesterol appeared to increase their solubility (Fig. 8 B).

Figure 6. CPZ induces transfer of aqueous dye (CF) from RBCs to hemifused NΔ12, Δ12, and GPI-HA–expressing cells. CV-1 cells expressing WT and mutant HAs were prepared as described in the legend to Fig. 4, except that fusion was triggered for 5 min at pH 5.0 and 37°C before renaturalization. After triggering fusion, these cells were exposed to either 0.1 or 0.5 mM CPZ for 1 min at room temperature. The CPZ solution was replaced with PBS+ and the cells were observed as above. In control experiments, virtually all WT HA–expressing cells (0.5 μg WT HA, inset graph) were stained with CF in the absence of CPZ. Percent content mixing was determined by dividing the number of cells receiving CF by the number of cells with bound RBCs. Error bars show the SEM for four to five independent experiments (mean ± SEM). Only 0.5 mM CPZ promoted significant content dye transfer between labeled RBCs and cells expressing NΔ12 HA, Δ12 HA, and GPI-HA.

Effect of MβCD on Fusion by Δ12 HA and NΔ12 HA

Because of the general interest in glycoprotein localization to plasma membrane microdomains (Scheiffele et al., 1997; Hooper, 1999), we asked whether treatment of cells expressing WT HA, GPI-HA, Δ12 HA, or NΔ12 HA with 20 mM MβCD influenced their fusion activity. Treatment of WT HA, GPI-HA, Δ12 HA, and NΔ12 HA–expressing cells with MβCD did not affect the fusion phenotype. WT HA still mediated efficient lipid and content dye transfer, whereas the mutant HAs still demonstrated significant lipid mixing but little or no content mixing (Fig. 8). Similar results were obtained using WT HA of the Japan strain (H2N2; Melikyan et al., 1999).

Figure 7. Effect of CPZ on content mixing. CV-1 cells expressing GPI-HA, NΔ12, and Δ12 HA cDNA were processed for fusion with CF-labeled RBCs in the absence or presence of the indicated amount of CPZ, as described in the legend to Fig. 6. Only 0.5 mM CPZ is able to promote significant content dye transfer (see Fig. 6).
Why D12 HA May Cause Lipid, but Not Content, Mixing

We have considered two general models for why D12 HA mediates only lipid mixing whereas D10 HA mediates content mixing as well. In the first model (Fig. 9 A, 1), we consider that D12 HA has recruited specific (e.g., shorter fatty acyl chain) lipids around it such that it spans a thinned bilayer. The lipids in such a thinned bilayer may not be competent to promote the hemifusion to fusion transition. In the second model, we consider that the TM domain of D12 HA is simply too short to span a bilayer; it may be anchored either perpendicularly (Fig. 9 A, 2a), obliquely (Fig. 9 A, 2b), or parallel (Fig. 9 A, 2c) to the membrane normal. To test between these models, we analyzed a mutant HA in which we added back the hydrophilic cytoplasmic tail (10 amino acids) to the end of D12 HA. We reasoned that if D12 HA spans a thinned bilayer (model 1), addition of the cytoplasmic tail should not affect its fusion phenotype. If, however, D12 HA does not span a bilayer (model 2), then addition of the cytoplasmic tail may force D12 HA to span a bilayer and it may therefore be able to support full fusion. As seen in Fig. 10 A, the mutant D12Tail HA clearly promotes full fusion.

Additional HA TM Domain Mutants

To ascertain whether we could detect any specific TM domain sequences needed for HA to promote full fusion, we constructed additional point and deletion mutations.
within the HA TM domain (Fig. 11). We mutated the tryptophans (to alanines) within the highly conserved WILW sequence at the beginning of the HA TM domain. We mutated a serine at position 194, since this residue is the analogue of a glycine implicated as being important for the fusion activity of Japan HA (Melikyan et al., 1999). We also mutated a glycine at position 204 (singly and in combination with Ser 194), since glycines near the middle of the TM domain have been reported to be important for fusion mediated by the vesicular stomatitis virus envelope glycoprotein (VSV G; Cleverley and Lenard, 1998). We also deleted six or eight amino acids at different locations within the TM domain. All mutants were examined for expression at the cell surface, for their ability to be cleaved by trypsin into HA1 and HA2, for RBC binding, and for fusion (both lipid and content mixing). Most of the mutants were also examined for trimer formation and the pH dependence of the conformation change (Table III). As seen in Table III, by all of the criteria examined, these additional mutants in the HA TM domain behaved virtually

Figure 9. (A) Model of possible interactions between the Δ12 (and NΔ12) TM domains and membranes: (1) The truncated TM domains of Δ12 and NΔ12 may span a thinned bilayer; (2a) the TM domain may project into, but not span, the bilayer in a perpendicular orientation relative to the membrane surface; (2b) the TM domain may project obliquely into the bilayer; (2c) the TM domain may be anchored at the surface parallel to the lipid bilayer. HA is presented as a monomer for clarification. The TM domain is depicted as an α-helix, but it may adopt other structures. (B) Models of HA and SNARE-mediated hemifusion to fusion transition. Multipliers of HA trimers promote hemifusion (mixing of the outer, but not inner leaflets of the lipid bilayer). Subsequent interactions between the fusion peptides (white) and the TM domains (dark gray), either alone or in concert, with the hemifusion diaphragm may promote full fusion. In the case of the SNAREs, the TM domains of the t- (gray) and v-SNARE (black) may perform analogous functions.

Figure 10. Addition of the cytoplasmic tail or a single arginine to Δ12 HA restores fusion. CV-1 cells transfected with 0.5 µg WT, 7.5 µg Δ12, 5.0 µg Δ12Tail, and 5.0 µg Δ12Arg were prepared for fusion as described in Materials and Methods. Fusion was triggered at pH 5.0 for 2 min at 37°C. Images presented in A and B are from separate experiments. Δ12Tail and Δ12Arg were expressed at the cell surface at levels comparable to that using 0.5 µg WT HA cDNA. The COOH-terminal sequences of Δ10, Δ11, Δ12, and Δ12Arg HA mutants are: Δ10 WILWSFASCFLLCVV Δ11 WILWSFASCFLLC Δ12 WILWSFASCFLLC Δ12Arg WILWSFASCFLLCR.
Additional TM domain deletion and point mutants. Line diagram of the HA gene. In detail is the region surrounding the HA TM domain (gray box). Point mutations were made in the context of full-length HA and are indicated in large font. Deletion mutations were made in the context of tail HA and are indicated as spaces. Δ10, Δ12, and NΔ12 HA are included as a reference.

the same as WT HA. Most importantly, all of the 12 additional HA TM domain mutants exhibited efficient lipid and efficient content mixing.

Discussion

Cells expressing the ectodomain of HA linked to the membrane via a GPI anchor (GPI-HA) promote lipid, but not content, mixing (Kemble et al., 1994). The behavior of GPI-HA indicates that the HA TM domain, which is predicted to be 27 amino acids in length, plays an important role in the hemifusion to fusion transition. In this study, we tested whether there is a length or sequence requirement for the TM domain to facilitate this important step in the fusion cascade. To do this, we first engineered stop codons into the TM domain, generating mutant HAs with sequential 2–amino acid deletions from the COOH-terminal end (up to 14 amino acids). We also engineered mutant HAs lacking 6 or 12 amino acids from the NH2-terminal end, mutants lacking 6 or 8 amino acids from the central region, and a mutant lacking 11 amino acids from the COOH-terminal end. These HA mutants lacked the cytoplasmic tail, which has been shown to be dispensable for the fusion pore initiation and for fusion pore enlargement (Markosyan et al., 2000). In the present study, we found that two mutant HAs, Δ12 HA and NΔ12 HA, are severely restricted in their ability to mediate mixing of a 376–mol wt content dye, CF. Since the level of CF mixing seen with Δ12 HA and NΔ12 HA is similar to that seen with GPI-HA (Fig. 6 B), it is likely that Δ12 HA and NΔ12 HA are unable to efficiently promote the hemifusion to fusion.

Δ12 HA and NΔ12 HA Mediate Hemifusion

Hemifusion is functionally defined as the merger of the outer, but not the inner, leaflets of the fusing bilayers, such that aqueous continuity is not established. Many investigators have proposed that biological fusion events proceed through a hemifusion intermediate (Palade, 1975; Pinto da Silva and Nogueira, 1977; Kalderon and Gilula, 1979; Lucy and Ahkong, 1986; Chernomordik et al., 1987; Nanavati et al., 1992).

Studies with GPI-HA indicate that progression to a fusion pore, as monitored by the transfer of small content dyes, does not occur when GPI-HA–expressing cells are induced to fuse with RBCs (Kemble et al., 1994; Melikyan et al., 1995b; Nussler et al., 1997). Combinations of dye transfer and electrophysiological assays also indicated that GPI-HA does not induce fusion pores in planar membranes (Melikyan et al., 1995b; Razinkov et al., 1999). However, more recent electrophysiological studies indicate that under certain conditions (e.g., pH 4.8 with membrane-labeled RBCs), GPI-HA can induce fusion pores during fusion with RBCs, but that these pores occur less frequently than with WT HA, never enlarge, and are strongly influenced by the presence of lipid dyes in the target membrane (Markosyan et al., 2000). These observations indicate that the TM domain is required for efficient fusion pore initiation and for fusion pore enlargement (Markosyan et al., 2000). In the present study, we found that two mutant HAs, Δ12 HA and NΔ12 HA, are severely restricted in their ability to mediate mixing of a 376–mol wt content dye, CF. Since the level of CF mixing seen with Δ12 HA and NΔ12 HA is similar to that seen with GPI-HA (Fig. 6 B), it is likely that Δ12 HA and NΔ12 HA are unable to efficiently promote the hemifusion to fusion.

Table III. Summary of Results: Effects of Point Mutations and Additional Deletions in the HA TM Domain

| HA mutants | Percentage of cells FU | Trypsin cleavage | RBC binding | Lipid mixing | Content mixing | ConfΔ |
|------------|-----------------------|------------------|-------------|--------------|---------------|-------|
| W185A      | 76                    | 299              | ND          | ++           | ++            | +     |
| W188A      | 50                    | 270              | ND          | ++           | ++            | +     |
| W185A/     | 62                    | 137              | +           | ++           | ++            | +     |
| W188A      |                        |                  |             |              |               |       |
| S194A      | 68                    | 204              | ND          | ++           | ++            | +     |
| S194L      | 66                    | 213              | +           | ++           | ++            | +     |
| S194L/     | 49                    | 130              | ND          | ++           | ++            | +     |
| G204A      | 22                    | 224              | ND          | ++           | ++            | +     |
| G204L      | 79                    | 337              | +           | ++           | ++            | +     |
| NΔ6        | 38                    | 152              | +           | ++           | ++            | +     |
| NΔ6Δ2      | 38                    | 87               | +           | ++           | ++            | +     |
| MidΔ6      | 30                    | 145              | +           | ++           | ++            | +     |
| MidΔ8      | 46                    | 179              | +           | ++           | ++            | +     |

Percentage of cells indicates the expression efficiency (SEM ± 5.9). Fluorescence units (FU) indicates the mean fluorescence as detected by FACS® analysis (SEM ± 24). ND, not done.
transition and are unable to support pore enlargement. Additional evidence that Δ12 HA and NΔ12 HA are blocked at the stage of hemifusion, and not at stunted fusion, is that 0.5 mM CPZ is required to induce appreciable content dye transfer (Fig. 6), as has been seen with GPI-HA (Melikyan et al., 1997a). We propose that Δ12 HA and NΔ12 HA are protein mimetics of GPI-HA.

Length Requirement of the HA TM Domain

We have uncovered a surprisingly stringent length requirement for the HA TM domain to be able to (efficiently) promote the hemifusion to fusion transition. HAs harboring a 17–amino acid (predicted) TM domain promote full fusion, whereas an HA with a 16–amino acid (predicted) TM domain is severely impaired in promoting full fusion and HAs with 15–amino acid (predicted) TM domains appear to arrest at hemifusion.

The finding that there is a stringent length requirement of 17 amino acids for the HA TM domain to efficiently promote the hemifusion to fusion transition suggests that HAs with TM domains ≳17 amino acids are anchored differently in the bilayer than fusion-impaired HAs that have shorter TM domains (≈16 amino acids). Using a synthetic peptide representing the transmembrane segment of X:31 HA, Tatulian and Tamm (1999) have recently shown that the WT HA TM domain (27–amino acid predicted) spans DMPC/DMPG bilayers as an α-helix that aligns roughly perpendicular to the bilayer normal. As discussed in Results (with reference to Fig. 9 A), we have considered two general models for how the fusion-defective TM domain mutants (with TM domains ≲16 amino acids) are anchored in the bilayer. The first (Fig. 9 A, 1) envisions that the short (≈16 amino acids) TM domains are aligned like the WT HA TM domain (as a perpendicular α-helix), but to span the bilayer, they have had to recruit specific lipids (e.g., with short fatty acyl tails) around them. Such lipids may not be able to adopt the necessary curvature to allow fusion to progress beyond hemifusion (Melikyan et al., 1997a; Chernomordik et al., 1998). The second general model proposes that the fusion-incompetent TM domains (≈16 amino acids) cannot span the bilayer (Fig. 9 A, 2a, 2b, and 2c). Our finding that addition of the hydrophilic 10–amino acid cytoplasmic tail sequence to a TM domain (predicted) TM domain is severely impaired in promoting full fusion. Hence, although we cannot exclude the possibility that there may be a sequence motif that is important for the X:31 HA TM domain to promote the hemifusion to fusion transition, we have not found such a

Sequence Requirements of the HA TM Domain

We also asked whether we could identify any specific residues within the HA TM domain that are required to promote the hemifusion to fusion transition. During analysis of four additional truncation mutants (of six or eight amino acids) and eight point mutants at different locations in the TM domain (Table III), we were unable to uncover any specific sequence requirement for fusion. In particular, two highly conserved tryptophan residues within the WILW motif at the NH2-terminal end of the TM domain, which appear to be important for targeting HA to the apical surface of epithelial cells (Scheiffele et al., 1997; Lin et al., 1998), do not appear to be required for fusion. We also examined the requirement for fusion of particular residues within the interior of the HA TM domain based on two reports concerning the role of glycine residues within the TM domain of viral fusion proteins. Cleverley and Lenard (1998) suggested the importance of glycine residues within the TM domain of VSV G. Substitution of both glycine residues within the TM domain of VSV G was reported to result in a hemifusion phenotype (Cleverley and Lenard, 1998). Furthermore, a study using Japan HA, which contains two glycine residues within the TM domain, demonstrated that mutation of the more NH2-terminal glycine to a leucine (G520L HA) caused a restricted hemifusion phenotype (no pore formation and no transfer of lipid or content dye; Melikyan et al., 1999). We have made the equivalent mutations within the TM domain of X:31 HA (Table III). Our results suggest that neither a serine residue at position 194 (the analogue of G520 in Japan HA) nor a glycine residue at position 204, either individually or jointly, is necessary for fusion (Table III).

If one models the WT HA TM domain as an α-helix, there is a short face of four polar residues (two cysteines and two serines). However, several of our mutant HAs disrupt this motif (i.e., leave only two polar residues), but do not impair fusion. Hence, although we cannot exclude the possibility that there may be a sequence motif that is important for the X:31 HA TM domain to promote the hemifusion to fusion transition, we have not found such a
motif. The differences we observe in the sequence requirements for fusion with X:31 HA (H3N2 subtype) and that reported for Japan HA (H2N2 subtype) may be due to differences in the subtype of HA or the techniques used.

**Possible Parallel Roles for the Fusion Peptide and the TM Domain in the Hemifusion to Fusion Transition**

Four indirect lines of evidence suggest that although they have different net hydrophobicities, the HA fusion peptide and the TM domain may play parallel roles in the hemifusion to fusion transition. First, we have recently demonstrated that replacement of the glycine at the first position of the HA fusion peptide with a serine (Ser HA) arrests HA fusion at the hemifusion stage (Qiao et al., 1999). Second, recent work using synthetic peptides corresponding to the fusion peptide and the TM domain indicates that these two peptides have similar effects on synthetic bilayers (Han et al., 1999; Tatulian and Tamm, 1999). Both domains appear to order their respective lipid environments, thus decreasing the amount of water bound at the water–bilayer interface as well as increasing the surface hydrophobicity. A third line of evidence that the fusion peptide and the TM domain may be playing parallel roles in breaking the hemifusion diaphragm is the observation that in the lowest energy state of the protein, the fusion peptide and the TM domain are predicted to be very closely opposed (Chen et al., 1999). A fourth line of indirect evidence is that studies on the topology of synthetic versions of the HA fusion peptide as well as studies with HAs containing mutations in the fusion peptide are consistent with the notion that the first 18 residues of the fusion peptide embed in the target bilayer and are important for fusion (Gray et al., 1996; Macosko et al., 1997; Danielli, T., and J.M. White, unpublished data). Hence, the two membrane-interactive domains of HA may have similar length requirements (~17 amino acids) to efficiently promote full fusion.

**Sequence Requirements of the TM Domains of Other Viral Fusion Proteins**

We have not observed any specific sequence requirements within the X:31 HA TM domain for full fusion. However, other investigators have suggested that there are specific sequence requirements for fusion within the TM domain of other viral fusion proteins. As discussed above, specific TM domain sequence requirements have been suggested for the VSV G glycoprotein (Cleverley and Lenard, 1998) and the HA from the Japan strain of influenza (Melikyan et al., 1999). In addition, mutation of a specific proline residue within the Moloney murine leukemia virus envelope glycoprotein, to either alanine, glycine, or valine, diminished syncytia formation (Taylor and Sanders, 1999). Finally, mutation of a conserved arginine residue to leucine within the TM domain of the HIV gp160 eliminated HIV-1 envelope–mediated syncytia formation (Owens et al., 1994). These findings suggest that for some viral fusion proteins, specific TM domain sequences may be important for fusion. Even for cases where there are specific sequence requirements, we propose that the length of the TM domain will be a critical determinant of fusion for all viral fusion proteins.

**Possible Relevance to SNARE-mediated Fusion**

Recent structural and biochemical data indicate that the SNARE ( soluble N-ethylmaleimide–sensitive factor [NSF] attachment protein receptor) complexes, key players in intracellular fusion events, share similarities with many viral fusion proteins (Hanson et al., 1997; Hohl et al., 1998; Poirier et al., 1998; Skehel and Wiley, 1998; Sutton et al., 1998). Formation of a “SNAREpin” structure consisting of a four-helix coiled coil domain is thought to force the fusogenic membranes together (Katz et al., 1998; Weber et al., 1998). The v- and t-SNARE proteins are each anchored into their respective membranes by a TM domain (21–24 residues in length; Fig 9 B, bottom). It may therefore be the case that the TM domain of one SNARE performs a function similar to that of the viral fusion peptide while the TM domain of the other SNARE acts in an manner equivalent to the TM domain of the viral fusion protein (Fig. 9 B). Therefore the length of SNARE TM domain anchors may be a critical determinant of fusion. Recent evidence suggests that this may indeed be the case. An analysis of the behavior of *Caenorhabditis elegans* mutants has suggested that the TM domain of the t-SNARE, unc-64 must span its bilayer to function properly (Saijee et al., 1998). And, very recently McNew et al., using lipid anchors of varying length, provided evidence that the TM domain of the v-SNARE must be anchored in both bilayer leaflets to promote fusion (in this case, lipid mixing; McNew et al., 2000). Hence, it seems likely that the length of the TM domains of both viral fusion proteins and SNAREs will be an important determinant of fusion, and in some cases, of the hemifusion to fusion transition (Fig. 9 B).

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