Forcible destruction of severely misfolded mammalian glycoproteins by the non-glycoprotein ERAD pathway

Satoshi Ninagawa,1,3,4 Tetsuya Okada,1 Yashihi Sumimoto,1 Satoshi Horimoto,1 Takehiro Sugimoto,1 Tokiro Ishikawa,1 Shinichi Takeda,2 Takashi Yamamoto,5 Tatashi Suzuki,6 Yukiko Kamiya,3,4 Koichi Kato,3,4,7 and Kazutoshi Mori1

1Department of Biophysics, Graduate School of Science, Kyoto University, Sakyo-ku, Kyoto 606-8502, Japan
2Department of Radiation Genetics, Graduate School of Medicine, Kyoto University, Sakyo-ku, Kyoto 606-8501, Japan
3Institute for Molecular Science and Okazaki Institute for Integrative Bioscience, National Institutes of Natural Sciences, Myodaiji, Okazaki 444-8787, Japan
4Department of Mathematical and Life Sciences, Graduate School of Science, Hiroshima University, Higashi-Hiroshima 739-8526, Japan
5Glycometabolome Team, Systems Glycobiology Research Group, RIKEN Global Research Cluster, Wako, Saitama 351-0198, Japan
6Department of Pharmaceutical Sciences, Nagoya City University, Mizuho-ku, Nagoya, 467-8603, Japan
7Graduate School of Engineering and EcoTopia Science Institute, Nagoya University, Chikusa-ku, Nagoya 464-8603, Japan

Glycoproteins and non-glycoproteins possessing unfolded/misfolded parts in their luminal regions are cleared from the endoplasmic reticulum (ER) by ER-associated degradation (ERAD)-L with distinct mechanisms. Two-step mannose trimming from Man9GlcNAc2 is crucial in the ERAD-L of glycoproteins. We recently showed that this process is initiated by EDEM2 and completed by EDEM3/EDEM1. Here, we constructed chicken and human cells simultaneously deficient in EDEM1/2/3 and analyzed the fates of four ERAD-L substrates containing three potential N-glycosylation sites. We found that native but unstable or somewhat unfolded glycoproteins, such as ATF6α, ATF6α(C), CD3-δ−ΔTM, and EMCl, were stabilized in EDEM1/2/3 triple knockout cells. In marked contrast, degradation of severely misfolded glycoproteins, such as null Hong Kong (NHK) and deletion or insertion mutants of ATF6α(C), CD3-δ−ΔTM, and EMCl, was delayed only at early chase periods, but they were eventually degraded as in wild-type cells. Thus, higher eukaryotes are able to extract severely misfolded glycoproteins from glycoprotein ERAD and target them to the non-glycoprotein ERAD pathway to maintain the homeostasis of the ER.

Introduction

Soluble or transmembrane proteins in the ER possessing unfolded or misfolded parts in their luminal regions are retrotranslocated back to the cytoplasm and degraded by the proteasome, a series of events termed ER-associated degradation (ERAD)-L (Xie and Ng, 2010; Smith et al., 2011; Brodsky, 2012). ERAD-L deals with both glycoproteins (gpERAD) and non-glycoproteins (non-gpERAD). Misfolded parts of non-glycoproteins are most likely recognized by molecular chaperones represented by BiP. In contrast, glycoprotein degradation relies on the structure of N-glycans present on the surface of ERAD-L substrates. GlcMan,GlcNAc2 attached to the asparagine residue is trimmed by glucosidase I and II to Man9GlcNAc2, which had been thought to have no mannosidase activity (Mast et al., 2005), whereas subsequent trimming from Man8GlcNAc2 to Man7GlcNAc2 was performed mostly by EDEM3 and EDEM1, EDEM2, and EDEM3. These experiments were performed in chicken DT40 cells using conventional homologous recombination and in human HCT116 cells using the transcription activator-like effector nuclelease method. Our results were surprising (Ninagawa et al., 2014). Initial trimming from Man9GlcNAc2 to Man8GlcNAc2 was conducted by EDEM2, which had been thought to have no mannosidase activity (Mast et al., 2005), whereas subsequent trimming from Man8GlcNAc2 to Man7GlcNAc2 was performed mostly by EDEM3 and to a slight extent by EDEM1. Erman1 appeared to play only a minor role in gpERAD.

Here, by constructing and analyzing DT40 and HCT116 cells simultaneously deficient in EDEM1/2/3, we show that different ERAD pathways are used depending on the severity of...
misfolding in the ERAD-L substrates and that non-gpERAD has a much more positive role in protein quality control in the ER than its previously proposed role as a backup for gpERAD under ER stress (Ushio et al., 2013), as indicated by the finding that non-gpERAD is able to clear severely misfolded glycoproteins from the ER even in the absence of pharmacological ER stress.

Results and discussion

We constructed EDEM1/2/3 triple KO (TKO) in both DT40 (gEDEM-TKO; Fig. S1) and HCT116 (hEDEM-TKO; Fig. S2 A) cells, in which functional EDEM1, EDEM2, or EDEM3 mRNA was not expressed (Fig. 1, A and C). gEDEM-TKO and hEDEM-TKO cells grew slightly more slowly than respective wild-type (WT) cells (Fig. 1, A and C). However, ER stress was not elicited in these cells because the levels of three ER stress marker proteins, BiP, ATF4, and XBP1S, were not elevated (Fig. S2 B). The N-glycan profiles of total cellular glycoproteins revealed that M9 level was markedly elevated in gEDEM-TKO and hEDEM-TKO cells, similarly to WT DT40 and HCT116 cells treated with kifunensine, an inhibitor of α1,2-mannosidases in the ER (Fig. 1, B and D), as expected from our previous results (Ninagawa et al., 2014). Accordingly, degradation of endogenous ATF6α, an ERAD-Lm (m for membrane protein) substrate containing three potential N-glycosylation sites (Fig. 3 A) was blocked almost completely in gEDEM-TKO and hEDEM-TKO cells (Fig. 1, E and F). The extent of this blockade was comparable with that observed in WT cells treated with MG132, a proteasome inhibitor, or kifunensine (Ninagawa et al., 2014). We also found that hATF6α(C)-TAP, an artificial ERAD-Ls (s for soluble protein) substrate containing the luminal region of human ATF6α flanked by the N-terminal signal sequence and C-terminal tandem affinity purification tag (Fig. 3 A), was stabilized in gEDEM-TKO cells (Fig. 1 G). During SDS-PAGE, hATF6α(C)-TAP in gEDEM-TKO cells migrated slightly more slowly than in WT cells because of the defect in mannose trimming, and this difference in migration was lost after endoglycosidase H (EndoH) treatment (Fig. 1 H). Instability of the native protein ATF6α in WT cells may result from intrinsically disordered parts found in its luminal region (Rosenbaum et al., 2011).

We next examined the effect of EDEM-TKO on degradation of the null Hong Kong (NHK) variant of α1-protease inhibitor (α1PI), the most frequently used ERAD-Ls substrate (Sifers et al., 1988). Similarly to the case of hATF6α(C)-TAP, NHK in gEDEM-TKO cells migrated slightly more slowly than in WT cells during SDS-PAGE, and this difference in migration was lost after EndoH treatment (Fig. 2 A). In the case of NHK-QQQ, the nonglycosylated version of NHK and its migration was lost after EndoH treatment (Fig. 2 A). In the case of NHK-Δ219–270 mutants were detected as a doublet, although slower bands were predominant (Fig. S3 A). When cells were lysed directly in SDS sample buffer, hATF6α(C)-myct1-302 and the Δ280–298 mutant were detected as a single band, whereas the Δ111–119, 182–194 and Δ219–270 mutants were detected as a doublet, although slower bands were predominant (Fig. S3 A). When cells were lysed with digitonin and then mixed with 2× SDS sample buffer, faster bands of hATF6α(C)-myct1-302 and the Δ280–298 mutant appeared, and levels of faster bands of the Δ111–119, 182–194 and Δ219–270 mutants increased (Fig. S3 A). We interpret this appearance of faster bands upon lysis with digitonin to represent partial deglycosylation of hATF6α(C) derivatives by peptide:N-glycanase (PNGase), an enzyme responsible for deglycosylation of glycoproteins dislocated from the ER to the cytosol (Misaghi et al., 2004), for three reasons. First, both bands were sensitive to EndoH treatment (Fig. S3 B); second, the appearance of the faster band of the Δ111–119, 182–194 mutant was significantly inhibited by Z-VAD-fmk (Fig. S3 C), which inhibits PNGase; and third, the faster band of the Δ111–119, 182–194 mutant did not appear in mouse embryonic fibroblast (MEF) cells deficient in PNGase but did appear in MEF cells deficient in the cytosolic endo-β-N-acetylglucosaminidase (ENGase; Fig. S3 D; Huang et al., 2015). The time course of the appearance of the
Figure 1. **Effect of EDEM-TKO on N-glycan profiles and ATF6α degradation.** (A) RT-PCR to amplify cDNA corresponding to gEDEM1/2/3 mRNA in DT40 cells of various genotypes and doubling times of WT and gEDEM-TKO cells. (B) Isomer composition of N-glycans prepared from total cellular glycoproteins of WT, WT treated with kifunensine (Kif, 10 µg/ml, 6 h), and gEDEM-TKO cells. This experiment was completed once. (C) RT-PCR to amplify cDNA corresponding to hEDEM1/2/3 mRNA in WT and hEDEM-TKO cells and their doubling times. (D) Isomer composition of N-glycans prepared from total cellular glycoproteins of WT, WT treated with kifunensine (10 µg/ml, 12 h), and hEDEM-TKO cells. This experiment was completed once. (E) Cycloheximide chase to determine the degradation rate of endogenous gATF6 (only one ATF6 gene in chicken genome) in WT and gEDEM-TKO cells using anti-gATF6 (n = 3). (F) Pulse-chase to determine the degradation rate of endogenous hATF6α in WT and hEDEM-TKO cells using anti-hATF6α (n = 3). (G) Cycloheximide chase to determine the degradation rate of transfected hATF6α(C)-TAP in WT and gEDEM-TKO cells using anti–c-myc (n = 3). (H) Immunoblotting of cell lysates prepared from WT and gEDEM-TKO cells expressing transfected hATF6α(C)-TAP with or without EndoH treatment using anti–c-myc. hATF6α(C)-TAP* denotes the nonglycosylated hATF6α(C)-TAP.
of faster bands in PNGase sensitivity assay in cell lysates indicated that the Δ111–119, 182–194 and Δ219–270 mutants were more misfolded than hATF6α(C)-myct1-302 and the Δ280–298 mutant (Fig. S3 E).

It was also previously reported that a misfolded protein is more susceptible to trypsin digestion than a folded protein (Izawa et al., 2012; Xu et al., 2013). The results of trypsin sensitivity assay suggested that the Δ111–119, 182–194 and Δ219–270 mutants were more misfolded than hATF6α(C)-myct1-302 and the Δ280–298 mutant (Fig. S3 E).
mutants were more misfolded than hATF6α(C)-myc1-302 and Δ280–298 mutant (Fig. S3 F).

To substantiate our hypothesis further, we determined whether severely misfolded glycoproteins are degraded by non-gpERAD in human cells also. For this, we focused on mouse CD3-δ-ΔTM, an ERAD-Ls substrate containing three potential N-glycosylation sites (Fig. 4 A; Bernasconi et al., 2010), which is localized in the ER (Fig. 4 B), because a part of hATF6α(C)-myc1-302 was secreted into medium in HCT116 cells (not depicted), contrary to its full retention in the ER of DT40 cells. The results of the PNGase sensitivity assay in cell lysates suggested that the two insertion mutants were more misfolded than WT (Fig. 4 C). We found that degradation of exogenously expressed mCD3-δ-ΔTM was delayed in hEDEM-TKO cells (Fig. 4 D) to a similar extent to that in WT cells treated with MG132 (Fig. 4 E), kifunensine (not depicted), or by 7 aa (Pro-Ala-Pro-Ala-Pro-Ala-Pro) between 33 and 34 aa (mCD3-δ-ΔTM<33-7aa-34>) or 70 and 71 aa (mCD3-δ-ΔTM<70-7aa-71>), present in the second or sixth β-sheet, respectively, in the case of human CD3-δ (Fig. S3 G) resulted in similar degradation of the mutant proteins in the two types of cells, except for a delay at the early chase period in hEDEM-TKO cells (Fig. 4 F), despite their localization in the ER (Fig. 4 B).

We further found that the type I transmembrane human protein EMC1 containing three potential N-glycosylation sites (Fig. 5 A), a subunit of the ER membrane protein complex consisting of six subunits (EMC1–EMC6; Jonikas et al., 2009), behaved like ATF6α. Endogenous hEMC1 turned over slowly in WT HCT116 cells, and its turnover was delayed in hEDEM-TKO cells (Fig. 5 B). Exogenously expressed Flag-tagged hEMC1 was localized in the ER (Fig. 5 C) and behaved very similarly to endogenous hEMC1 (Fig. 5 E). Importantly, however, its Δ35–220 mutant, which lacks most of the pyrroloquinoline quinone (PQQ) domain...
containing the β-propeller repeat (Kopec and Lupas, 2013), was localized in the ER (Fig. 5 C), was more sensitive to trypsin and therefore more misfolded than hEMC1-Flag (Fig. 5 D), and was degraded similarly in the two types of cells, except for a delay at the early chase period in hEDEM-TKO cells (Fig. 5 E).

Our results identified a previously unnoticed system operating in the ER. Chicken and human cells can degrade severely misfolded glycoproteins in the absence of EDEM-mediated mannose trimming. This novel quality control cannot be easily discerned by knocking down EDEM1/2/3 singularly, and was accordingly identified for the first time here after our construction of EDEM-TKO cells in combination with our comparison of the fates of ATF6α and NHK. Based on the results reported for recombinant human ERmanI (Aikawa et al., 2012), we consider that ERmanI cannot trim Man9GlcNAc2 present in native but unstable or somewhat unfolded glycoproteins, leading

Figure 4.  Effect of EDEM-TKO on mCD3-δ-ΔTM degradation.  [A] Schematic structures of WT and two insertion mutants of mCD3-δ-ΔTM.  [B] Immunofluorescence of HCT116 cells expressing transfected WT or insertion mutant of mCD3-δ-ΔTM using anti-HA.  [C] PNGase sensitivity assay of WT and two insertion mutants of mCD3-δ-ΔTM expressed in HCT116 cells by transfection (immunoblotting with anti-HA).  The asterisk denotes partially deglycosylated mCD3-δ-ΔTM.  [D] Pulse-chase to determine the degradation rate of transfected WT mCD3-δ-ΔTM in WT and hEDEM-TKO cells using anti-HA (n = 3).  [E] Pulse-chase to determine the degradation rate of transfected WT mCD3-δ-ΔTM in WT cells and those treated with 40 μM MG132 using anti-HA.  This experiment was completed once.  [F] Pulse-chase to determine the degradation rate of transfected insertion mutants of mCD3-δ-ΔTM in WT and hEDEM-TKO cells using anti-HA (n = 3).  [D–F] Means ± SD are shown.  [D and F] *, P < 0.05; **, P < 0.01.
Figure 5. Effect of EDEM-TKO on hEMC1 degradation. (A) Schematic structures of WT and deletion mutant of hEMC1. (B) Pulse-chase to determine the degradation rate of endogenous hEMC1 in WT and hEDEM-TKO cells using anti-hEMC1. This experiment was completed once. (C) Immunofluorescence of HCT116 cells expressing transfected hEMC1-Flag or hEMC1-ΔPQQ-Flag using anti-Flag. (D) Trypsin sensitivity assay of hEMC1-Flag and hEMC1-ΔPQQ-Flag expressed in HCT116 cells by transfection (immunoblotting with anti-Flag). The data shown are from a single representative experiment out of two repeats. (E) Pulse-chase to determine the degradation rate of transfected hEMC1-Flag and hEMC1-ΔPQQ-Flag, as well as endogenous hEMC1 in WT and hEDEM-TKO cells using anti-hEMC1 (n = 3). Means ± SD are shown. *, P < 0.05; **, P < 0.01. (F) Model. Newly synthesized polypeptides of non-glycoproteins are subjected to productive folding. Folded non-glycoproteins are secreted, whereas unfolded or misfolded proteins are targeted to non-gpERAD for proteasomal degradation. Newly synthesized polypeptides of glycoproteins are subjected to N-glycosylation and productive folding. Folded glycoproteins are secreted, whereas native but unstable glycoproteins, somewhat unfolded glycoproteins, and severely misfolded glycoproteins are subjected to EDEM-mediated mannose trimming, followed by proteasomal degradation via gpERAD. In addition, severely misfolded glycoproteins are subjected to non-gpERAD via protein X.
to marked stabilization of ATF6α in EDEM2 single KO cells (Ninagawa et al., 2014), but can trim Manα1GlcNAc2 present in severely misfolded glycoproteins, such as NHK, to produce Manα1GlcNAc2 in EDEM2 single KO cells, which is trimmed to Manα1GlcNAc2 by EDEM3/1 for proteasomal degradation (Fig. 2 F). Indeed, degradation of NHK was markedly blocked by kifunensine, which inhibits both ERman1 and EDEM1/23, at the early chase period (1.5 h) in HCT116 cells (Fig. 2, G and H). This also explains the apparent blockade of NHK degradation by kifunensine observed at the early chase periods (1–2 h) in previous studies (Hosokawa et al., 2001; Hirao et al., 2006). Complete blockade at 3-h chase (Liu et al., 1999) was not reproduced in either recent studies (Pan et al., 2011; Fujimori et al., 2013) or here (Fig. 2, G and H).

This protein structure–based decision is made only in higher eukaryotes because yeast does not possess an efficient non-gpERAD (Jakob et al., 2001; Kostova and Wolf, 2005). In marked contrast, degradation of NHK-QQQ is much more efficient than that of NHK in both mammalian (Ninagawa et al., 2008) and DT40 cells (compare Fig. 2 B with Fig. 2 D). This explains why the delay in NHK degradation at the early chase periods in EDEM−TKO cells was canceled in the later chase periods (Fig. 2, D and G).

The next obvious question is what molecules extract severely misfolded glycoproteins from gpERAD and target them to non-gpERAD (protein X in Fig. 5 F). Identification of the switch will further enhance our understanding of the protein quality control operating in the ER.

Materials and methods

Construction of plasmids

Recombinant DNA techniques were performed according to standard procedures (Sambrook et al., 1989). The integrity of all constructed plasmids was confirmed by extensive sequencing analyses. Site-directed mutagenesis was performed using Dpn I. p3*flag-CMV-14 Expression Vector (Sigma-Aldrich) was used to express a protein tagged with Flag at the C terminus. Platinum transcription activator-like effector nuclease plasmids to knock out HEDEM family genes were constructed previously (Ninagawa et al., 2014).

Reagents

Puromycin (0.5 µg/ml), histidinol (1 mg/ml), mycophenolic acid (15 µg/ml), blasticidin S (25 µg/ml), G418 (2 mg/ml for DT40 and 0.6 mg/ml for HCT116), and Zeocin (1 mg/ml) were used for the selection and maintenance of drug-resistant clones. Kifunensine was purchased from Cayman Chemical Company, EndoH from EMD Millipore, cycloheximide from Sigma-Aldrich, MG132 from Peptide Institute, Z-VAD-fmk from Promega, trypsin from Nacalai Tesque, and digitonin from Wako Pure Chemical Industries.

Cell culture, transfection, and N-glycan profiling

DT40 cells were cultured and transfected as described previously (Ninagawa et al., 2011). HCT116 cells (ATCC CCL-247) and MEF cells were cultured in Dulbecco’s modified Eagle’s medium (glucose 4.5 g/liter) supplemented with 10% fetal bovine serum, 2 mM glutamine, and antibiotics (100 U/ml penicillin and 100 µg/ml streptomycin) at 37°C in a humidified 5% CO2/95% air atmosphere and transfected using Lipofectamine 2000 (Invitrogen) or X-tremeGENE 9 (Roche) according to the manufacturers’ instructions. Doubling times were determined by counting cell numbers every 24 h for a total of 120 h for DT40 cells, and every 48 h for a total of 192 h for HCT116 cells. Each value represents the mean of triplicate determinations. Pyridylamination and structural identification of N-glycans of total cellular glycoproteins were performed as described previously (Horimoto et al., 2013; Ninagawa et al., 2014).

Immunological techniques

Immunoblotting analysis was performed according to the standard procedure (Sambrook et al., 1989) as described previously (Ninagawa et al., 2011). Chemiluminescence obtained using Western Blotting Luminol Reagent (Santa Cruz Biotechnology, Inc.) was detected using an LAS-3000mini Lumino Image analyzer (Fujifilm). Rabbit anti–chicken ATF6 (Horimoto et al., 2013) and anti–human ATF6α (Haze et al., 1999) antibodies were raised previously. Rabbit polyclonal anti-EMC1 antibody, which reacts with both chicken and human EMC1, was raised against human EMC1 (529–943 aa) fused to glutathione S-transferase. Anti-c-myc antibody (9E10) was obtained from Wako Pure Chemical Industries, anti-Flag antibody from Sigma-Aldrich, anti-HA antibody from Recenttec, anti-KDEL antibody from Medical and Biological Laboratories, anti-ATF4 and anti-XBP1 antibodies from Santa Cruz Biotechnology, Inc., and anti-α1PI antibody from Dako.

Pulse-chase experiments using 9.8 MBq/dish EASY TAG EXPRESS Protein labeling mix [35S] (PerkinElmer) and subsequent immunoprecipitation using anti–human ATF6α, anti-α1PI, anti–human EMC1, anti-Flag, or anti-HA antibody and protein A-coupled Sepharose beads (GE Healthcare) were performed according to procedures described previously (Ninagawa et al., 2011).

For immunofluorescence analysis, HCT116 cells transfected using X-tremeGENE 9 were fixed by incubation on ice for 6.5 min in methanol/acetic acid (1:1). Anti-Sec61β antibody was obtained from Proteintech. Secondary antibodies labeled with FITC and CF594 were obtained from ICN Biomedicals and Biotium, respectively. Microscope images were obtained at room temperature with 40 magnification using a DM IRE2 and confocal software (both from Leica).

PNGase sensitivity assay

HCT116 cells at 24 h after transfection or DT40 cells at 16 h after transfection were washed with PBS, collected, and suspended in 200 µl buffer A (50 mM Tris/HCl, pH 8.0, containing 1% NP-40 [for mammalian cells] or 1% digitonin [for chicken cells]), 150 mM NaCl, protease inhibitor cocktail [Nacalai Tesque], and 20 µg MG132). Lysates were left on ice for 20 min and clarified by centrifugation at 14,000 rpm for 10 min at 4°C. After incubation for various periods on ice, cleared lysates were mixed with 2x SDS sample buffer containing 100 mM dithiothreitol and 2 µM Z-VAD-fmk, boiled for 5 min, and subjected to SDS-PAGE followed by immunoblotting.

Trypsin sensitivity assay

DT40 cells or HCT116 cells transfected were washed with PBS three times, collected, and suspended in 250 µl (for DT40 cells) or 400 µl (for HCT116 cells) buffer B (50 mM Tris/HCl, pH 8.0, containing 1% NP-40, 150 mM NaCl, 2 µM Z-VAD-fmk, and 20 µM MG132). Lysates were left on ice for 20 min and clarified by centrifugation at 14,000 rpm for 10 min at 4°C. Cleared lysates were incubated with various concentrations of trypsin for 15 min at 4°C and then mixed with 2x SDS sample buffer containing 100 mM dithiothreitol and 10x protease inhibitor cocktail (Nacalai Tesque), boiled for 5 min, and subjected to SDS-PAGE followed by immunoblotting. The presence of equal amounts of proteins in cleared lysates before trypsin digestion was confirmed by immunoblotting with anti-β-actin antibody.
Southern blot hybridization
Southern blot hybridization was performed according to standard procedures (Sambrook et al., 1989) as described previously (Ninagawa et al., 2011). Specific probes were prepared as described previously (Ninagawa et al., 2014) and labeled with digoxigenin. Subsequent reaction with anti-digoxigenin antibody (Roche) and treatment with the chemiluminescent detection reagent CDP-star (GE Healthcare) were performed according to the manufacturers’ specifications. Chemiluminescence was visualized using an LAS-3000mini Lumino Image analyzer (Fujifilm).

Genomic PCR
Homologous recombination in HCT116 cells was confirmed by genomic PCR using a pair of primers described previously (Ninagawa et al., 2014).

RT-PCR
Total RNA prepared from WT DT40 cells or various KOs (~5 x 10⁶ cells) and from WT HCT116 cells or various KOs (~3 x 10⁶ cells) by the acid guanidinium/phenol/chloroform method using ISOGEN (Nippon Gene) was converted to cDNA using Moloney murine leukemia virus reverse transcription (Invitrogen) and random primers. The full-length open reading frame of gEDEM1, gEDEM2, gEDEM3, hEDEM1, hEDEM2, or hEDEM3 was amplified using PrimeSTAR HS DNA polymerase (Takara Bio Inc.) and a pair of primers described previously (Ninagawa et al., 2014). Southern blot hybridization was performed according to standard procedures (Sambrook et al., 1989) as described previously (Ninagawa et al., 2014). We further checked by Southern blotting (Fig. S1 B), and RT-PCR analysis revealed the expected expression of each mRNA (Fig. S1 C). We further constructed TKO cells deficient in gEDEM1, gEDEM2, and gEDEM3 (Fig. S1 A). Homologous recombination was checked by Southern blotting (Fig. S1 B), and RT-PCR analysis revealed the expected expression of each mRNA (Fig. S1 C).

Construction of gEDEM-TKO DT40 cells
We previously constructed gEDEM1, gEDEM2, or gEDEM3 single KO DT40 cells (Ninagawa et al., 2014). We then constructed double KO cells deficient in gEDEM1 and gEDEM2, gEDEM2 and gEDEM3, and gEDEM1 and gEDEM3 (Fig. S1 A). Homologous recombination was checked by Southern blotting (Fig. S1 B), and RT-PCR analysis revealed the expected expression of each mRNA (Fig. S1 C). We further constructed TKO cells deficient in gEDEM1, gEDEM2, and gEDEM3 (Fig. S1 A, right). Homologous recombination was checked by Southern blotting (Fig. S1 B), and RT-PCR analysis revealed the expected expression of each mRNA (Fig. S1 C).

Construction of hEDEM-TKO HCT116 cells
We previously constructed hEDEM1, hEDEM2, or hEDEM3 single KO HCT116 cells (Ninagawa et al., 2014). We disrupted the hEDEM2 gene in hEDEM3 single KO cells and then disrupted the hEDEM1 gene in the resulting hEDEM2/3 double KO cells to create hEDEM-TKO HCT116 cells (Fig. S2 A).

Online supplemental material
Fig. S1 shows the generation of DT40 cells deficient in two or three of gEDEM1/2/3. Fig. S2 shows the generation and characterization of hEDEM-TKO cells and the effect of MG132 on the stability of hATF6(C) derivatives in DT40 cells. Fig. S3 shows analysis of folding status of hATF6(C) derivatives in DT40 cells. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201504109/DC1. Additional data are available in the JCB DataViewer at http://dx.doi.org/10.1083/jcb.201504109.dv.

Acknowledgments
We thank Ms. Kaoru Miyawaga for her technical and secretarial assistance and Ms. Yukiko Isono (IMS) for her help in the analyses of N-glycans. We are grateful to people in the Yamate-Area Facility, Center for Radioisotope Facilities, and Okazaki Research Facilities, National Institutes of Natural Science.

This work was financially supported in part by grants from the Ministry of Education, Culture, Sports, Science and Technology of Japan (26291040 to K. Mori, 26840065 to S. Ninagawa, 23221005 and 15K06996 to T. Okada, 15K18529 to T. Ishikawa, 26110725 to T. Suzuki, 26102518 to Y. Kamiya, and 24249002, 25102008, and 15H02491 to K. Kato) and by the Joint Studies Program (2014-2015) of the Institute for Molecular Science.

The authors declare no competing financial interests.

Submitted: 24 April 2015
Accepted: 7 October 2015

References


