AGR2 Function Requires a Unique Endoplasmic Reticulum Localization Motif

Aparna Gupta, Aiwen Dong, and Anson W. Lowe
Departments of Medicine and the Stanford Digestive Disease Center
Stanford University, Stanford, California 94305
Running head: Functional AGR2 requires a specific ER localization motif
Address correspondence to: Anson W. Lowe, Alway Building, Rm. M211, 300 Pasteur Drive, Stanford, CA 94305-5187. FAX: 650-723-5488; E-mail: lowe@stanford.edu

Background: Whether AGR2 promotes adenocarcinoma growth as a secreted or ER localized protein is not known.

Results: A unique C-terminal motif, KTEL, is required for AGR2 ER localization and function.

Conclusions: Not all ER localization motifs are interchangeable.

Significance: Specific ER localization signals may be required for protein function. AGR2's tumor promoting effects are mediated from the ER and not as a secreted protein.

SUMMARY

Soluble proteins are enriched in the ER by retrograde transport from the Golgi that is mediated by the KDEL receptors. In addition to the classic C-terminal KDEL motif, a variety of sequence variants are also capable of receptor binding that result in ER localization. Although different ER localization signals that exhibit varying affinities for the KDEL receptors exist, whether there are functional implications was unknown. The present study determines whether AGR2 requires a specific ER localization signal to be functionally active. AGR2 is expressed in most human adenocarcinomas and serves a role in promoting growth and the transformed phenotype. Using two different cell lines in which AGR2 induces expression of either the EGFR ligand amphiregulin or the transcription factor CDX2, only the highly conserved wild-type C-terminal KTEL motif results in the appropriate outcome. Deletion of the KTEL motif results in AGR2 secretion and loss of AGR2 function. AGR2 function is also lost when ER residence is achieved with a C-terminal KDEL or KSEL instead of a KTEL motif. Thus variations in ER localization sequences may serve a specific functional role, and in the case of AGR2 is served specifically by KTEL.

INTRODUCTION

Soluble proteins are targeted to the endoplasmic reticulum (ER) by two major mechanisms. The first is signal peptide directed translocation of newly translated proteins into the ER. Second, the proteins remain enriched in the ER by KDEL receptor mediated retrieval of proteins that have entered the intermediate compartment or the Golgi apparatus (1-3). Whether a protein is retrieved back to the ER is determined by four carboxy-terminal amino acids that mediate binding to one of three KDEL receptors in higher vertebrates. The classic endoplasmic reticulum localization motif in higher vertebrates is KDEL (4). Subsequent work led to a Prosite motif ([KRHQS]-[DENQ]-E-L>) that encompasses a set of carboxy-terminal peptide sequences that result in ER residence. The implications of why different carboxyl-terminal sequences capable of ER localization exist are not clear, but recent studies have demonstrated that affinity for the three known KDEL receptors is affected by which localization motif is employed (5,6). Whether different ER localization motifs also carry functional significance is not known.

Anterior Gradient Homolog 2 (AGR2) encodes a 17kDa protein that is highly conserved in vertebrates. AGR2 was first described in Xenopus laevis where its expression is responsible for the development of a glandular organ called the cement gland (7,8). A significant role in tissue regeneration was established for AGR2 in salamanders where it functions in nerve dependent limb regeneration (9,10). AGR2 is also expressed by secretory cells in the normal murine intestine (11). In humans, enhanced AGR2 expression was first described in breast cancer, which was
followed by similar observations in most human adenocarcinomas, including those derived from the esophagus, pancreas, lung, ovary, and prostate (12-19). Both in vitro and in vivo studies have demonstrated that AGR2 promotes tumor growth and metastasis (11,14,20).

Recent studies have provided insights into AGR2’s mechanism of action. AGR2 expression in esophageal and lung adenocarcinoma cells induces expression of the EGF receptor ligand amphiregulin (AREG), which is responsible for the transformed phenotype observed in cultured cells in vitro (21). In addition, AGR2 stimulation of AREG expression, required activation of the Hippo signaling pathway co-activator, YAP1. Thus AGR2 expression promotes tumor growth and the transformed phenotype by affecting the Hippo and EGF signaling pathways.

The induction of AREG expression also provides a means to identify structural requirements for AGR2 activity, including protein domains that are essential for its biologic action. The AGR2 amino terminus contains a sequence motif characteristic of signal peptides, which results in protein targeting to the cell’s secretory pathway. Indeed, several studies have proposed that AGR2 secretion from the cell is necessary for its action (7,10,14,16). In addition, yeast two-hybrid screens identified AGR2 binding proteins that naturally occur on the cell surface (9,22).

Whether AGR2 binding to the identified receptors results in a biological response, however, has yet to be established.

Immunocytochemistry of AGR2 expressing cells, however, reveals an intracellular pattern that is most consistent with an ER distribution (11,21). AGR2’s carboxy-terminus contains a tetra-peptide sequence, KTEL, that is conserved in all vertebrates from Xenopus to humans (Treefam accession TF321449 (23)). Although the sequence does not agree with the Prosite consensus sequence for ER residence (4,24), a recent study by Raykhel et al. demonstrated that the KTEL motif does result in binding to the three known KDEL receptors, which results in ER localization (5). The study also demonstrated that the KTEL motif results in lower affinities for the three known KDEL receptors when compared to proteins terminating with a KDEL sequence.

This study addresses two questions concerning AGR2 biology and the functional significance of endoplasmic reticulum localization signals. The first is whether ER residence of AGR2 is necessary for its function. The second is whether endoplasmic retention by the KTEL sequence is absolutely required for AGR2 function, as suggested by its high conservation in all species where AGR2 is expressed, or whether other ER localization signals may serve a similar role.

**EXPERIMENTAL PROCEDURES**

**Cell lines** - IEC-6, a rat small intestinal jejunal cell line (ATCC, Manassas, VA) was cultured in Dulbecco's modified Eagle's media with 4mM L-glutamine, 1.5 g/L sodium bicarbonate, 4.5g/L glucose, and supplemented with 0.1U/ml bovine insulin and 10% fetal bovine serum (Hyclone, Thermo Fisher Scientific) (25). The IEC6 stable cell line expressing AGR2-KTEL, AGR2-KDEL, AGR2-KSEL and AGR2-STOP were transfected with pcDNA3.1 expression vectors (Invitrogen, Carlsbad, CA) and cultured in the presence of 2mg/ml of G418 (Mediatech, Inc., Manassas, VA). Human OE33 esophageal adenocarcinoma cells were obtained from Sigma-Aldrich (St. Louis, MO) and cultured in RPMI 1640 with 10% FBS. Transient transfection for different AGR2 mutants was used for OE33 cells. The GFP-KTEL and RFP-KTEL expression vectors were transiently transfected into both IEC-6 and OE33 cells.

**Antibodies** - Mouse monoclonal anti-CDX2Biogenex (San Ramon, CA). A mouse monoclonal anti-GRP78 (HNGC symbol HSPA5) was obtained from Enzo Life Sciences (Farmingdale, NY). The antibody was generated against the peptide SEKDEL derived from HSPA5 (also known as GRP78), which reacts against all KDEL proteins, and is thus referred to as an anti-KDEL antibody. Rabbit anti-human AGR2 antisera was produced as previously described (21).

**Expression vectors** - All recombinant DNA constructs utilized the pcDNA3.1 expression vector (Invitrogen, Carlsbad, CA) and stable expression was achieved using G418 selection. The AGR2 constructs utilized the human AGR2 gene (GeneBank Accession NM_006408). In vitro mutagenesis was performed using the QuikChange® Site-Directed Mutagenesis Kit (Stratagene, Santa Clara, CA). Dr. Erik Snapp (Albert Einstein College of Medicine, Bronx, NY)
graciously provided the GFP- and RFP-KDEL expression vectors (26). The KDEL terminus was then mutated to KTEL for the purposes of this study.

Quantitative real-time PCR - RNA levels were quantified using real-time PCR. qPCR primers included: rat AGR2: 5’-TCAGTCTCGGAATCTGTCTTTT-3’, 5’-TCTTTAACGTTGA-3’; rat CDX2: 5’-GCGGCAACCTTTTGTAATGGAT-3’, 5’-ACTCAGTTTCTCTCTGATGTTGA-3’; rat β-actin: 5’-TGAAACAGCGATTGTCAACAACTG-3’, 5’-ATACAGGGACAACACACAGCTGGAT-3’; human AGR2: 5’-ATGAGTGCCCACACAGTCAA-3’, 5’-GGACATACTGGCCATCAGGA-3’; human AREG: 5’-GTGGTGCTGTCGCTCTTGATA-3’, 5’-ACTCACAGGGGAAATCTCACT-3’; human β-actin: 5’-GAGCGCGGCTACAGCTT-3’, 5’-TCCTTAATGTCACGCACGATTT-3’.

Total RNA was isolated from cells using TRIzol reagent (Invitrogen). First-strand cDNA was synthesized from total RNA using Superscript II reverse transcriptase (Invitrogen) with random hexamer primers. Real-time PCR reactions were performed using IQSYBR Green Supermix and the iCycler iQ detection system (Bio-Rad, Hercules, CA).

Protein Immunoblotting - Protein concentrations of cell lysates was spectrophotometrically determined (NanoDrop 2000, Thermo Fischer Scientific, Wilmington, DE). Protein samples were resolved using 4-12% Bis-Tris gel (Invitrogen) and transferred to PVDF membranes (Millipore Corp., Bedford, MA). Membranes were blocked for 1 hour using 5% non-fat milk in buffer containing 20mM Tris, 137mM NaCl, 0.15 Tween 20, pH 7.6; followed by incubation with primary antibodies (anti-AGR2, 1:2000; anti-CDX2, 1:50; anti-β-actin, 1:5000; anti-KDEL, 1:1000). Detection was achieved with the appropriate secondary antibodies followed by chemiluminescence (GE Healthcare, Piscataway, NJ). For detection of AGR2 in the culture media, equal number of cells was plated in regular culture media. After 1 day, the media was changed to serum free media. Twenty-four hours later the media was collected and concentrated using a spin column that retains 90-95% of 17 kDa protein (MICROCON YM-30, Millipore), followed by immunoblotting for secreted AGR2. The concentrated media was stained with colloidal Coomassie Brilliant Blue R-250 (Invitrogen) to normalize for the amount of protein loaded for each cell line. All immunoblots were quantified using a flatbed scanner and ImageJ (http://rsbweb.nih.gov/ij/index.html).

Immunocytochemistry - 25,000 cells were plated in 8 well chamber slides (Applied Scientific South San Francisco, CA). The cells were fixed in 4% paraformaldehyde for 10 minutes, followed by permeabilization using 0.1% saponin, 10% FBS in PBS for 20 min. PBS with 10% FBS was then used for blocking for 20 min. Primary antibodies were diluted in PBS with 10% FBS and incubated for 2 hours (anti-AGR2 1:300; anti-cdx2 1:5, anti-KDEL 1:50) at room temperature. Anti-mouse or anti-rabbit IgG conjugated to either Alexa594 or Alexa488 (Invitrogen) was used as a secondary antibody. Imaging was achieved with a Nikon TS-1 microscope equipped for confocal laser scanning fluorescence microscopy (Nikon C1 system) or with standard fluorescence with a Nikon Eclipse E600 microscope.

Miscellaneous Methods - Secreted AREG in solution was measured using the Amphiregulin Duo-set ELISA kit (DY262, R&D Systems, Minneapolis, MN.). Quantification of immunofluorescent images was achieved using ImageJ (http://rsbweb.nih.gov/ij/index.html). The image threshold before counting cells utilized the algorithm by Li (http://pacific.mpi-cbg.de/wiki/index.php/Auto_Threshold).

RESULTS

Expression of secreted and ER retained AGR2 isoforms

Our previous studies have utilized RNA interference to define AGR2’s function in tumor cells. To explore the functional significance of the carboxy-terminal sequence, a cell line with no AGR2 expression was used in this study. The presence of wild-type AGR2 containing a KTEL carboxyl-terminal domain may confound experiments when mutant proteins are expressed in the same cells. For the present study, function was prospectively monitored by the AGR2-induced appearance of new gene transcription, which provided a cleaner background to assess functional determinants. IEC-6 cells are derived from normal rat intestinal crypts and represent...
non-transformed cells that were first propagated as sustained primary cultures (25). Previous studies have shown that IEC-6 cells can be induced to differentiate through the expression of various developmental genes (27-30). No AGR2 expression is detected in wild-type IEC-6 cells when evaluated with quantitative real-time PCR, protein immunoblotting, or immunocytochemistry (Figure 1).

Stable wild-type AGR2 expression was achieved in IEC-6 cells to produce IEC-6:AGR2-KTEL. In addition to wild-type AGR2, IEC-6 cells were also transfected with DNA constructs in which the carboxy-terminal sequence was mutated from KTEL to KDEL (T173D, IEC-6:AGR2-KDEL) or a STOP was inserted before the KTEL (K172STOP, IEC-6:AGR2-STOP). The cells exhibited no gross phenotypic changes. Consistent with other proteins retained in the ER, immunofluorescence for AGR2 in IEC-6:AGR2-KTEL and IEC-6:AGR2-KDEL cells revealed punctate cytoplasmic staining consistent with an ER distribution (Figure 1A-B). ER localization was confirmed when the cells were also probed with an anti-KDEL antibody that binds HSPA5 (also known as GRP78), an ER resident protein, which revealed a similar staining pattern (Figure 1E-F). In contrast, IEC-6:AGR2-STOP cells (Figure 1C) showed no cytoplasmic staining for AGR2 and were similar to IEC-6 cells transfected with the vector control (Figure 1D). Protein immunoblotting of cell lysates also revealed the presence of AGR2 protein for IEC-6:AGR2-KTEL and IEC-6:AGR2-KDEL cells but not in the IEC-6:AGR2-STOP cells. In contrast, AGR2 was detected in the media only for the IEC-6:AGR2-STOP cells, revealing that it is secreted and not retained (Figure 1H). Thus IEC-6:AGR2-KTEL and IEC-6:AGR2-KDEL cells exhibited ER localization in IEC-6 cells.

Only KTEL retained AGR2 induces CDX2 in IEC-6 cells
A screen for markers of intestinal development revealed that expression of the transcription factor CDX2 was markedly induced upon expression of wild-type AGR2 in IEC-6 cells. Real-time quantitative PCR did not detect CDX2 expression in wild-type IEC-6 cells, but IEC-6:AGR2-KTEL cells showed a dramatic induction of CDX2 RNA (Figure 2A). CDX2 protein production was confirmed with protein immunoblotting (Figure 2B). Immunocytochemistry revealed the expected nuclear localization for CDX2, although the distribution among cells was heterogenous (Figure 2C). Thus CDX2 expression in IEC-6 cells is a downstream result of AGR2 expression and can be used as a functional readout.

In contrast to the IEC-6:AGR2-KTEL cells, the IEC-6:AGR2-KDEL and IEC-6:AGR2-STOP cells did not show significant CDX2 RNA or protein expression (Figure 2), despite the presence of AGR2 RNA (Figure 2A) and protein (Figure 1H, J). Thus secreted AGR2 (IEC-6:AGR2-STOP) was not able to induce CDX2 expression. In addition, only ER localization by KTEL resulted in CDX2 expression. When a single amino acid substitution was performed changing KTEL to KDEL, CDX2 expression was not induced despite the presence of AGR2 in the ER.

KTEL and KDEL-carboxy terminal proteins share the same receptor
Whether wild-type AGR2 is being retained in the ER by the same mechanism as other KDEL-retained ER resident proteins was explored using competition experiments. Previous studies in yeast revealed that overexpression of a carboxy-terminated HDEL protein results in secretion of ER resident proteins through competition with the ERD2 receptor (1,31). The HDEL receptor (ERD2) represents the sole determinant for ER localization in yeast, as there is only one HDEL receptor. IEC-6:AGR2-KTEL cells that express CDX2 were transiently transfected with a previously described GFP-KDEL construct. The GFP-KDEL construct (also known as ER-GFP) is an inert protein with no interacting partners and does not undergo glycosylation or form disulfide bridges (26). Transient transfection of GFP-KDEL in IEC6:AGR2-KTEL cells resulted in a 7-fold reduction in CDX2 expression as determined by the fraction of cells that express nuclear CDX2 expression on immunofluorescence (Figure 3B, E). Expression of a KTEL carboxy-terminated GFP, GFP-KTEL, also induced a 3.8 fold decrease in CDX2 nuclear expression (Figure 3C, E). Consistent with the decrease CDX2 protein detected by immunofluorescence, a concomitant decrease in CDX2 RNA was also observed using real-time quantitative PCR of cells transfected with either GFP-KDEL or GFP-KTEL (Figure

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To ensure that the observed results were not due to the GFP protein itself, RFP tagged proteins were similarly expressed in the IEC6:AGR2-KTEL cells; resulting in a similar outcome as measured by immunocytochemistry or CDX2 RNA levels (Figure 3G-H).

As observed for other ER resident proteins, overexpression of GFP-KDEL is expected to displace AGR2 if they share the same KDEL receptor. With the expression of either GFP-KDEL or GFP-KTEL, greater amounts of AGR2 are secreted into the culture media (Figure 3Q). Protein immunoblotting of the culture media also showed increased secretion of soluble HSPA5 ER proteins as detected with anti-KDEL antibodies (Figure 3Q). The results indicate that both KTEL and KDEL terminated proteins bind to the same receptors. The loss of function observed with KDEL terminated AGR2 is not due to binding a different receptor.

Expression of KTEL terminated AGR2 is required for secretion of amphiregulin in human esophageal adenocarcinoma cells

AGR2 is normally expressed in human esophageal adenocarcinoma OE33 cells where it induces the expression of the EGF receptor ligand, amphiregulin (AREG) (21). When AGR2 is overexpressed in OE33 cells, AREG expression is also induced. Thus OE33 cells provided an opportunity to study the role of AGR2’s carboxy-terminal KTEL sequence in a tumor cell line that normally expresses the protein.

The same AGR2 DNA constructs previously used with the IEC-6 cells were also transiently transfected into OE33 cells to determine whether ER localization by the carboxy-terminal KTEL sequence was similarly required for efficient AREG induction. Expression of wild-type KTEL carboxy-terminated AGR2 enhanced AREG RNA and protein secretion by 12.5 and 1.9-fold, respectively (Figure 4C-D). Wild-type AGR2 over-expression resulted in higher levels of AREG secreted into the culture media (Figure 4D). Expression of KDEL carboxy-terminated AGR2, however, induced AREG RNA expression and protein secretion by only 4 and 1.3-fold, respectively, while the AGR2 (STOP) construct resulted in a 4.7 and 1.1-fold difference from the control. Thus the functional significance of the carboxy-terminal ER localization motif is conserved in a tumor cell line in which AGR2 expression is normally observed.

Whether AGR2’s induction of AREG expression also requires binding to the KDEL receptors was evaluated in OE33 cells through over-expression of the GFP-KDEL and GFP-KTEL constructs. Constructs terminating in either KDEL or KTEL motifs resulted in a decrease in AREG protein secretion of 2.4 and 3.2-fold, respectively (Figure 4E).

**ER localization signals with similar physical characteristics as KTEL cannot support AGR2 function**

The substitution of KDEL for KTEL is predicted to result in a significant change in physical characteristics that may affect AGR2 function. Thus an alternative construct was evaluated in which a carboxy-terminal sequence of KSEL replaced the KTEL ER localization motif.

Previous work demonstrated that KSEL exhibits affinity for the KDEL receptors and results in ER localization (5). In addition, the substitution of serine for threonine is predicted to be the least disruptive. AGR2-KSEL expression in IEC-6 cells resulted in an ER distribution similar to that observed with anti-KDEL antibodies (Fig. 5A-B). KSEL terminated AGR2, however, did not induce CDX2 RNA or protein (Fig. 5C-F). When AGR2-KSEL is expressed in OE33 cells, there is 4.7-fold less induction of AREG RNA compared to transfection with AGR2-KTEL (Fig. 5G-H).

Both IEC-6 and OE33 cells express all three known KDEL receptors.

To determine which KDEL receptors were being affected by the competition experiments, real-time PCR was performed on both cell lines to determine the relative levels of the 3 known KDEL receptors, KDELR1, KDELR2, and KDELR3. Both IEC-6 and OE33 cells expressed all 3 receptors, although in different proportions (Figure 6). Similar to what has been previously reported for HeLa cells, RNA for KDELR2 was most abundant compared to the other two receptors.

KTEL Carboxy-terminated proteins sort to the same ER compartment as KDEL terminated proteins
Additional experiments were performed to evaluate sorting by the KTEL motif independent of the AGR2 protein. GFP and RFP with either a carboxy-terminal KTEL or KDEL were expressed together by co-transfection in CHO cells. Subsequent confocal immunocytochemistry revealed that both motifs resulted in colocalization of the GFP and RFP signals in a pattern consistent with the ER (Figure 7 A-C).

Additional experiments were also performed using IEC-6 cells to determine whether AGR2 that is terminated with either KTEL or KDEL will colocalize with other ER resident proteins as determined using the anti-KDEL antibody. In both scenarios, IEC-6 cells that express either AGR2 with a KTEL or KDEL carboxy-terminus co-localized with ER resident proteins as determined with the anti-KDEL antibody (Fig. 7D-I). Thus ER localization by KTEL is required for AGR2 function, and a preferential distribution within the ER was not discernible.

**DISCUSSION**

Previous work established that AGR2 expression in esophageal and lung adenocarcinoma cell lines induces expression of the EGF receptor ligand, AREG. For the present study, AGR2 was discovered to induce the *de novo* expression of CDX2 in IEC-6 cells. AREG and CDX2 expression thus provided an opportunity to evaluate the protein domains necessary for AGR2 function. Published studies to date have reported an extracellular role for AGR2 as a secreted protein as well as an intracellular role as a protein chaperone or stress response factor (7,10,14,16,32,33). Among the factors favoring an intracellular role based in the ER is the presence of a carboxy-terminal motif, KTEL, that is conserved from *Xenopus* to humans.

Using immunofluorescence studies of myc-tagged AGR2, Raykhel et al. previously demonstrated that AGR2 is retained in the ER. The same study utilized biomolecular fluorescence complementation to show that AGR2 binds to all three isoforms of the KDEL receptor, although with lower affinity than controls possessing a KDEL carboxy-terminal sequence (5). Among the factors favoring an intracellular role based in the ER is the presence of a carboxy-terminal motif, KTEL, that is conserved from *Xenopus* to humans.

The present study demonstrates that AGR2 residence within the ER is required for its function as reflected by AREG expression in OE33 adenocarcinoma cells or CDX2 expression by IEC-6 intestinal cells. Converting AGR2 to a secretory protein by deleting the KTEL terminus nullified the functional readout in both cell lines. The results are consistent with experiments in which application of recombinant AGR2 to the culture media was not able to induce either AREG or CDX2 expression (personal observations).

ER residence alone, however, is not sufficient for AGR2 function as conversion of the KTEL sequence to the classic KDEL motif failed to induce AREG or CDX2 expression. The results demonstrate that differences in ER localization signals may serve an essential functional role.

The results of the competition experiments are consistent with AGR2 ER localization that is mediated by the same receptors as those responsible for retaining KDEL terminated proteins. Previous studies revealed that the single yeast counterpart of the KDEL receptor, ERD2, can be saturated by high expression of a HDEL-terminated ligand, resulting in secretion of ER resident proteins such as GRP78 (HSPA5) (1). In addition, saturation of the yeast ERD2 receptor with HDEL-terminated ligand inhibited yeast growth (31), a finding also observed in the present study. Similar to the yeast studies, overexpression of an independent protein that terminates in either KDEL or KTEL compromised AGR2 function and resulted in its secretion. Consistent with the previous work of Raykhel et al. where higher affinity for the KDEL receptors was exhibited by KDEL compared to KTEL terminated fluorescent proteins, a consistent effect was observed in the inhibition of AGR2 function or promotion of AGR2 secretion (Figures 3E, F, G, and Q).

The distribution of KTEL-terminated proteins, including AGR2, was similar to that of other ER resident proteins as determined using the anti-KDEL antibodies. When considered in the context of the competition experiments, the data would support involvement of the previously described KDEL receptors. Because the generation of isoform specific anti-KDEL receptor antibodies have been unsuccessful (5), whether AGR2 preferentially associates with a particular isoform
cannot be determined. Given that the KDEL receptors were identified by sequence homology to the yeast ERD2 gene or to each other, it is possible another receptor exists that is capable of binding both KDEL and KTEL terminated proteins and is required for AGR2 function. If this is true, however, the distribution of such a receptor must result in a distribution for AGR2 that is similar to the other ER proteins retrieved by the KDEL receptors.

Several possibilities may account for the diversity of ER localization signals. One hypothesis is that KDEL receptor binding influences AGR2 function by affecting its subcellular location or promoting a protein complex that includes the receptors. The major difference between the various ER localization sequences described to date is the different affinities displayed for the three KDEL receptors. Recent studies have suggested that different affinities for the KDEL receptor may result in changes in subcellular distribution that are dynamic and dependent on situational conditions such as the cell density or growth media (6).

Protein complexes that include chaperones bound to the KDEL receptors have also been described, which may mediate a function other than the retrograde transport of proteins back to the ER (34). Of particular interest are studies showing that ligand-bound KDEL receptors also bind p38 mitogen-activated protein kinases (MAPKs) and c-Jun amino-terminal kinases (JNKs), which suggests potential involvement in physiological and pathological processes (35). High throughput binding assays have also revealed additional associations between the KDEL receptors and proteins not known to serve a role in retrograde transport, but the functional implications are currently unclear (36-38).

We hypothesize that the ER localization motif serves a biofunctional role, which accounts for the diversity in motifs observed. The carboxy-terminal domain is required for both ER localization and enzymatic function. KDEL receptor binding of soluble lumenal ER proteins such as the thioredoxins is pH dependent, where the higher acidity of the Golgi apparatus results in receptor association. Once the receptor transports its bound ligand back to the ER where the pH is less acidic, the ligand disassociates and becomes a soluble protein (39). Similar to other ER thioredoxins, soluble AGR2 is likely to bind to substrate(s) that are currently being actively pursued. We hypothesize that the ER localization signal is critical for AGR2's conformation and substrate binding in the ER as a soluble protein. Thus carboxy-terminal alteration to KDEL or KSEL retains ER localization, but disrupts AGR2's function as an enzyme in the ER. The highly conserved amino acid sequence of the AGR2 ER localization signal in all species supports an important functional role.

In summary, the present work establishes that AGR2 localization in the ER by KTEL is required for its ability to induce AREG or CDX2 expression. ER residence alone, however, is not sufficient, as a specific localization signal is required for AGR2 function. Thus the diversity in ER localization signals serves an essential functional role, and provides a rationale for AGR2's highly conserved carboxy-terminal motif. As AGR2 expression is enhanced in almost all human adenocarcinomas, the current findings provide insights into potential mechanisms, the subcellular location required for AGR2 function, and potential therapeutic strategies for cancer.

REFERENCES


**FOOTNOTES**

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**FIGURE LEGENDS**

**Figure 1.** - AGR2 expression in rat intestinal IEC-6 cells. (A-D) Immunofluorescence images of IEC-6 cells transfected with AGR2 constructs terminating with a carboxy-terminal KTEL (A), KDEL (B), or a STOP inserted before the wild-type KTEL (C) sequence. (D) Cells transfected with the expression vector alone. For panels A-D: (red) anti-AGR2 antibodies and (blue) nuclear DAPI staining. (E-F) Representative IEC-6 cell transfected with wild-type AGR2 and labeled with anti-KDEL (E, green) and anti-AGR2 (F, red) antibodies. The white bar in the images equals 10 µm. (G) Real-time quantitative PCR determination of AGR2 RNA from transfected IEC-6 cells. Error bars represent 1 ± S.D. (H) Protein immunoblots of transfected IEC-6 cell lysates (anti-AGR2 and anti-β-actin) or concentrated media (anti-AGR2). (I) Coomassie Blue stained SDS-PAGE of culture media depicted in panel H to show protein loading. (J) Graph of AGR2 protein levels as determined by densitometry of the immunoblot in (H). The values are normalized by the corresponding β-actin band density.

**Figure 2.** - AGR2 expression in IEC-6 cells induces CDX2 expression and requires a carboxy-terminal KTEL sequence. (A) Real-time quantitative PCR determination of AGR2 RNA (grey columns, left ordinate) and CDX2 RNA (black columns, right ordinate) levels in transfected IEC-6 cells. Error bars represent 1 ± S.D. (n=2) (B) Protein immunoblots of IEC-6 cell lysates for CDX2 and β-actin. (C-F) Immunofluorescence images of IEC-6 cells transfected with the same AGR2 constructs as depicted in Fig. 1A, but probed with an anti-CDX2 antibody (red) or DAPI nuclear stain (blue). (C) IEC-6:AGR2-KTEL; (D) IEC-6:AGR2-KDEL; (E) IEC-6:AGR2-STOP; (F) IEC-6:VECTOR. The white bar represents 50 µm.

**Figure 3.** - CDX2 induction is mediated by binding to the KDEL receptors. The effects of expression with GFP and RFP proteins terminated at the carboxy-terminus with KDEL or KTEL on CDX2 expression in IEC-6:AGR2-KTEL cells. (A-D) show anti-CDX2 (red) and DAPI (blue) staining in IEC-6 cells transfected with (A) AGR2 alone, (B) AGR2 + GFP-KDEL, (C) AGR2 + GFP-KTEL, (D) expression vector alone and no AGR2. The white bar represents 50 µm. (E) Proportion of CDX2 positive cells for the constructs shown in panels A-D as quantified with IMAGE J software. The total number of cells counted as determined by DAPI staining is listed in parentheses below the abscissa. (F) Real-time
quantitative PCR of CDX2 RNA normalized to β-actin for the same cells shown in (E). (G) Proportion of CDX2 positive cells for similar constructs as shown in (E) except that RFP was substituted for GFP. (H) Real-time quantitative PCR of CDX2 RNA normalized to β-actin for the same cells shown in (G). (I-L, 400X magnification) Controls for GFP transfection efficiency. Phase contrast (I,K) and GFP fluorescence (J,L) of IEC6-AGR2 cells transfected with GFP-KDEL (I,J) or GFP-KTEL (K,L). (M-P) Controls for RFP transfection efficiency. Phase contrast (M,O) and RFP fluorescence (N,P) of IEC6-AGR2 cells transfected with RFP-KDEL (M,N) or RFP-KTEL (O,P). (Q) Culture media of the cells transfected with the GFP constructs were evaluated with protein immunoblotting with anti-AGR2 and anti-KDEL (HSPA5) antibodies. Below each lane is a column graph depicting the density of the bands. (R) Coomassie Blue staining of equal amounts of culture media as a loading control.

**Figure 4.** AGR2 induction of AREG in OE33 esophageal adenocarcinoma cells requires a carboxy-terminal KTEL sequence. (A) Real-time quantitative PCR of total AGR2 RNA in OE33 cells transiently transfected with AGR2 constructs terminating with a carboxy-terminal KTEL, KDEL, or a STOP sequence (before the wild-type KTEL). Vector designates transfection with the expression vector alone. (B) Immunoblots of cell lysates for total AGR2 and β-actin of transfected cells. The AGR2 antisera recognizes both the endogenous and transiently expressed constructs of AGR2. Under each lane is a column graph depicting the total AGR2/β-actin ratio determined by densitometry. (C) Real-time quantitative PCR of total AREG RNA in the same cells as (A). (D) Determination of AREG protein levels in the culture media by ELISA. (E) Determination of AREG concentration in the media of wild-type OE33 cells that have been transiently transfected with GFP-KDEL and GFP-KTEL constructs. (F-I) Controls for GFP transfection efficiency. Phase contrast (F, H) and GFP fluorescence (G, I) of OE33 cells transfected with GFP-KDEL (F, G) or GFP-KTEL (H, I). The error bars for all column graphs represent 1±S.D.

**Figure 5.** Mutation of KTEL to KSEL results in compromised AGR2 induction of CDX2 and AREG. (A-B) Immunofluorescence labeling of AGR2-KSEL expressing IEC-6 cells with anti-AGR2 (A) and anti-KDEL (B) antibodies. The white bar represents 10µm. (C) Quantified RNA levels determined by real-time qPCR for AGR2 (left ordinate) and CDX2 (right ordinate) for AGR2-KTEL and AGR2-KSEL transfected IEC-6 cells. (D) Proportion of CDX2 positive cells in IEC-6 cells transfected with the AGR2-KTEL or AGR2-KSEL constructs. (E-F) Representative images from IEC-6 cells transfected with the AGR2-KTEL (E) or AGR2-KSEL (F) constructs and stained for CDX2 (red) and DAPI (blue). The white bar represents 100µm. (G) Quantified RNA levels determined by real-time qPCR for AGR2 (left ordinate) and AREG (right ordinate) derived from AGR2-KTEL and AGR2-KSEL transfected OE33 cells. (H) OE33 data from (G) that depicts AREG expression normalized to total AGR2 expression. Error bars on all graphs represent 1±S.D.

**Figure 6.** IEC-6 and OE33 cells express all three KDEL receptor isoforms. Real-time quantitative PCR for the three KDEL receptors (KDELR1, KDELR2, and KDELR3) in human OE33 and rat IEC-6 cells. The ordinate represents RNA levels normalized to β-actin. The error bars represent 1±S.D. of three determinations.

**Figure 7.** Carboxy-terminal KDEL and KTEL proteins are both sorted to the same compartment. (A-C) Confocal imaging of CHO cells transfected with RFP-KDEL (A) and GFP-KTEL (B) in CHO cells. (C) Merged RFP/GFP image. (D-I) IEC-6 cells transfected with AGR2-KTEL (D-F) or AGR2-KDEL (G-I) and probed with anti-AGR2 (D, G) or anti-KDEL (E, H) antibodies. (F, I) Merged images of the preceding two panels.
Figure 1

A. B.

C. D.

E. F.

G. H.

I. J.
Figure 2

A. 

<table>
<thead>
<tr>
<th></th>
<th>KTEL</th>
<th>KDEL</th>
<th>STOP</th>
<th>VECTOR</th>
</tr>
</thead>
<tbody>
<tr>
<td>AGR2 RNA</td>
<td>70</td>
<td>50</td>
<td>40</td>
<td>30</td>
</tr>
<tr>
<td>CDX2 RNA</td>
<td>80</td>
<td>60</td>
<td>50</td>
<td>40</td>
</tr>
</tbody>
</table>

p = 0.0012

p = 0.015

B. 

KTEL  KDEL  STOP  VECTOR

- CDX2
- β-actin

C. 

D. 

E. 

F.
Figure 3

A. ARG2 only + GFP-KDEL + GFP-KTEL - AGR2

B. HSPA5

C. 188

D. 98

E. 62

F. 49

G. 38

H. 28

I. 14

J. 7

K. ARG2 only + RFP-KDEL + RFP-KTEL

L. NO AGR2

M. CDX2 mRNA (2^(-ΔCT) x 1000)

N. CDX2 positive cells (%)

O. ARG2 only + GFP-KDEL + GFP-KTEL

P. NO AGR2

Q. CDX2 mRNA (2^(-ΔCT) x 1000)

R. CDX2 positive cells (%)

S. p < 0.0001

T. p = 0.0002

U. p = 0.0067

V. p = 0.0067

W. p = 0.026

X. p = 0.0053

Y. p = 0.0003

Z. p < 0.0001

AA. p < 0.0001

BB. p < 0.0001

CC. p < 0.0001

DD. p < 0.0001

EE. p < 0.0001

FF. p < 0.0001

GG. p < 0.0001

HH. p < 0.0001

II. p < 0.0001

JJ. p < 0.0001

KK. p < 0.0001

LL. p < 0.0001

MM. p < 0.0001

NN. p < 0.0001

OO. p < 0.0001

PP. p < 0.0001

QQ. p < 0.0001

RR. p < 0.0001

SS. p < 0.0001

TT. p < 0.0001

UU. p < 0.0001

VV. p < 0.0001

WW. p < 0.0001

XX. p < 0.0001

YY. p < 0.0001

ZZ. p < 0.0001

AAA. p < 0.0001

BBB. p < 0.0001

CCC. p < 0.0001

DDD. p < 0.0001

EEE. p < 0.0001

FFF. p < 0.0001

GGG. p < 0.0001

HHH. p < 0.0001

III. p < 0.0001

JJJ. p < 0.0001

KKK. p < 0.0001

LLL. p < 0.0001

MMM. p < 0.0001

NNN. p < 0.0001

OOO. p < 0.0001

PPP. p < 0.0001

QQQ. p < 0.0001

RRR. p < 0.0001

SSS. p < 0.0001

TTT. p < 0.0001

UUU. p < 0.0001

VVV. p < 0.0001

WWW. p < 0.0001

XXX. p < 0.0001

YYY. p < 0.0001

ZZZ. p < 0.0001
Figure 4

A. 

B. 

C. 

D. 

E. 

F. 

G. 

H. 

I.
Figure 5

A. 

B. 

C. 

D. 

E. 

F. 

G. 

H. 

Figure 5
Figure 6

![Normalized RNA (2^{-\Delta CT} \times 1000)](image)

- **KDEL1**, **KDEL2**, **KDEL3**
- **OE33**
- **IEC-6**

Legend:
- Light gray bars: OE33
- Black bars: IEC-6
Figure 7