UBE2O remodels the proteome during terminal erythroid differentiation


INTRODUCTION: The reticulocyte–red blood cell transition is a canonical example of terminal differentiation. The mature red blood cell has one of the simplest cellular proteomes known, with hemoglobin remarkably concentrated to ~98% of soluble protein. During reticulocyte maturation, the proteome is remodeled through the programmed elimination of most generic constituents of the cell, in parallel with abundant synthesis of cell type–specific proteins such as hemoglobin. The mechanisms that drive rapid turnover of soluble and normally stable proteins in terminally differentiating cells remain largely unclear.

RATIONALE: The ubiquitin-proteasome system (UPS) was discovered in reticulocytes, where it is highly active. However, its function in this developmental context has not been established. UBE2O is an E2 (ubiquitin-conjugating) enzyme that is co-induced with globin and expressed at elevated levels late in the erythroid lineage. We identified an anemic mouse line with a null mutation in Ube2o, and used multiplexed quantitative proteomics to identify candidate substrates of UBE2O in an unbiased and global manner. We found that the protein compositions of mutant and wild-type reticulocytes differed markedly, suggesting that UBE2O-dependent ubiquitination might target its substrates for degradation to effect remodeling of the proteome.

To test whether UBE2O was sufficient for proteome remodeling, we engineered a non-erythroid cell line to inducibly express UBE2O above its basal level. Upon induction, we observed the decline of hundreds of proteins from these cells, in many cases the same proteins as those eliminated from reticulocytes. Overexpression of an active-site mutant of UBE2O did not show these effects. Therefore, a major component of the specificity underlying differentiation-linked proteome remodeling appears to be carried by UBE2O itself. These results also indicate that UBE2O may function as a hybrid enzyme with both E2 and E3 (ubiquitin-ligating) activities. In support of this model, candidate substrates identified by proteomics were ubiquitinated by purified UBE2O without the assistance of additional specificity factors.

RESULTS: The most prominent phenotypes of the Ube2o mutant are an anemia characterized by small cells with low hemoglobin content (microcytic hypochromic anemia), and a defect in the elimination of ribosomes, the latter being a key aspect of reticulocyte maturation. When we added recombinant UBE2O protein to reticulocyte lysates from the null mutant, ubiquitin was conjugated primarily to ribosomal proteins. Moreover, immunoblot analysis and quantitative proteomics revealed elevated levels of multiple ribosomal proteins in mutant reticulocytes. Sucrose gradient analysis indicated the persistence not only of ribosomal proteins but of ribosomes themselves during ex vivo differentiation of mutant reticulocytes. Accordingly, ribosomes were eliminated upon induction of UBE2O in non-erythroid cells. The elimination of organelles from reticulocytes, as exemplified by that of mitochondria, was not affected in the Ube2o mutant, indicating the specificity of its effects on programmed protein turnover.

Free ribosomal proteins were ubiquitinated by purified UBE2O, which suggests that these proteins are true substrates of the enzyme. However, UBE2O substrates are diverse in nature and not limited to ribosomal proteins. Individual domains of UBE2O bound substrates with distinct specificities. Thus, the broad specificity of UBE2O reflects the presence of multiple substrate recognition domains within the enzyme.

Proteasome inhibitors blocked the degradation of UBE2O-dependent substrates in reticulocytes, although UBE2O does not form polyubiquitin chains. Rather, UBE2O adds single ubiquitin groups to substrates at multiple sites. Proteasome inhibitor treatment ex vivo led to depletion of the pools of many amino acids; this result implies that the flux of ubiquitinated substrates through the reticulocyte proteasome is sufficient to supply amino acids needed for late-stage translation of mRNA. In late erythropoiesis, several ubiquitin-conjugating enzymes and ligases are induced together with Ube2o while most components of the UPS disappear. We propose that the UPS is not simply amplified during erythroid maturation, but is instead broadly reconfigured to promote remodeling of the proteome.

CONCLUSION: A highly specialized UPS is expressed in the reticulocyte and is used to remodel the proteome of these cells on a global scale. UBE2O, a hybrid E2-E3 enzyme, functions as a major specificity factor in this process. In reticulocytes, and perhaps in other differenti- ated cells such as in the lens, the induction of ubiquitinating factors may drive the transition from a complex to a simple proteome.
UBE2O remodels the proteome during terminal erythroid differentiation

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During terminal differentiation, the global protein complement is remodeled, as epitomized by erythrocytes, whose cytosol is ~98% globin. The erythroid proteome undergoes a rapid transition at the reticulocyte stage; however, the mechanisms driving programmed elimination of preexisting cytosolic proteins are unclear. We found that a mutation in the murine Ube2o gene, which encodes a ubiquitin-conjugating enzyme induced during erythropoiesis, results in anemia. Proteomic analysis suggested that UBE2O is a broad-spectrum ubiquitinating enzyme that remodels the erythroid proteome. In particular, ribosome elimination, a hallmark of reticulocyte differentiation, was defective in Ube2o−/− mice. UBE2O recognized ribosomal proteins and other substrates directly, targeting them to proteasomes for degradation. Thus, in reticulocytes, the induction of ubiquitinating factors may drive the transition from a complex to a simple proteome.

The autosomal recessive hem9 mutation was identified in a systematic ethynitrosourea screen for mouse mutants (9). Homozygous hem9 mutants have a form of anemia characterized by small cells (microcytosis), reduced concentrations of hemoglobin (hypochromia), and elevated RBC counts (erythrocytosis) (Table 1). Reciprocal transplantation studies demonstrated that these phenotypes are intrinsic to the hematopoietic system (fig. S1). Histological characterization of peripheral blood smears (fig. S2, A to B) confirmed the phenotype of hypochromic anemia.

We mapped hem9 to a ~309-kb interval on chromosome 11. A single gene in this interval, Ube2o, was expressed in erythroid precursors far above its basal level in other cell types (4, 10, 11). Sequencing revealed a nonsense mutation predicted to truncate the C-terminal 168 amino acids. UBE2O protein was undetectable by immunoblot analysis of reticulocyte-rich blood from Ube2o−/− animals (fig. S2C). To validate the mutation, we bred hem9 mutants to two Ube2o gene-trap alleles. Compound heterozygotes lacked UBE2O protein and exhibited an RBC phenotype similar to that of hem9 homozygotes (fig. S2D). Thus, hem9 is a null allele of Ube2o (hereafter Ube2o−/−).

UBE2O−/− reticulocytes showed reductions in many ubiquitin conjugate species (Fig. 1A). Given that hundreds of ubiquitinating factors are expressed in mammals, it is unusual to find such a dominant role of a single component. Unassembled and misfolded globins are preferentially ubiquitinated, as seen in β-thalassemia, where unpaired α-globin precipitates cause cellular damage, ineffective erythropoiesis, hemolysis, and anemia (12). Purified UBE2O ubiquitinated purified α-globin, although some ubiquitination of endogenous α-globin persisted in the Ube2o−/− mutant (fig. S3, A and B).

We tested whether UBE2O-dependent degradation of excess α-globin might contribute to the Ube2o−/− phenotype by crossing a deletion of the β-globin loci (HbB′′′ allele), which causes a β-thalassemia intermedia phenotype (13), with the Ube2o−/− line. However, UBE2O deficiency actually increased hemoglobin levels in the HbB′′′−/− mutant (Table 1). Mitigation of the anemia in the double mutant was accompanied by increased numbers of RBCs (Table 1), which were smaller in mean volume than controls and had diminished hemoglobin content.

UBE2O deficiency markedly reduced precipitated hemoglobin aggregates that are associated with β-globin deficiency, and accordingly diminished the accumulation of insoluble α-globin, without affecting the steady-state α/β-globin ratio for soluble protein (fig. S4, A to D). Furthermore, double mutants showed reduced splenomegaly, a major consequence of β-thalassemia caused by increased RBC destruction and heightened splenic erythropoiesis. RBC life span was also increased in the double mutants, which is consistent with a reduction in harmful unpaired α-globin chains (fig. S4E).

In summary, although α-globin is a UBE2O substrate, defective ubiquitination and degradation of α-globin could not fully explain the Ube2o−/− phenotype; instead, the loss of UBE2O ameliorated phenotypes attributable to α-globin excess. The reduced globin levels of the mutant appear to reflect attenuation of translation and α- and β-globin, as indicated by ribosome profiling (fig. S5, A and B). In the mutant, eukaryotic initiation factor 2α (eIF2α) phosphorylation was induced, which is suppressive of translation and may therefore account for the reduction of globin expression (fig. S5C).

Ribosomal proteins are abundantly ubiquitinated by UBE2O

To identify additional substrates of UBE2O, we added recombinant enzyme or a catalytically inactive mutant (UBE2O-C1037A; hereafter UBE2O-CA), at physiological levels, to reticulocyte lysates from null animals. Prior to the reaction, endogenous E2 activity was quenched chemically. Recombinant UBE2O formed ubiquitin conjugates robustly, whereas UBE2O-CA was inactive (Fig. 1B). Newly formed conjugates were identified by liquid chromatography–tandem mass spectrometry (LC-MS/MS) analysis (table S1). Although numerous targets were identified, including α- and β-globin (fig. S6A), the major class of targets was ribosomal proteins (RPs), comprising 87% of spectral counts of all ubiquitinated peptides that were specific to wild-type UBE2O, other than those from UBE2O itself (Fig. 1C). The two most frequently identified ubiquitinated RPs were RPL29 and RPL35, which had been suggested to interact with UBE2O by high-throughput affinity purification mass spectrometry (14). Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analysis (15) of all 105 UBE2O targets (table S1) showed a strong selection for RPs (adjusted P value = 3.65 × 10−4).
ubiquitinates RPs in reconstituted case were biotin-ubiquitin and autoubiquitinated E1. In the absence of did not have conjugating activity in lysates, as the only bands evident in this were supplemented with biotin-tagged ubiquitin and ubiquitin-activating streptavidin (proteins.

antibody to ubiquitin. Each sample is from a different mouse whose polyacrylamide gel electrophoresis (PAGE) and immunoblotted with an

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ing conditions, RPs remained the major class of proteins.

Fig. 1. Ube2o−/− reticulocytes are deficient in eliminating ribosomal proteins. (A) Proteins from reticulocyte lysates were resolved by SDS–polyacrylamide gel electrophoresis (PAGE) and immunoblotted with an antibody to ubiquitin. Each sample is from a different mouse whose reticulocytes were induced by serial bleeding. Ub-HBA represents the ubiquitinated form of α-globin. (B) Ubiquitination was reconstituted in Ube2o−/− reticulocyte lysates by means of recombinant UBE2O. Reactions were supplemented with biotin-tagged ubiquitin and ubiquitin-activating enzyme (UBE1) and were incubated for 45 min at 37°C. Samples were resolved by SDS-PAGE. Proteins were electrophoblated and visualized using streptavidin–horseradish peroxidase. W, UBE2O-WT (wild type); C, UBE2O-CA (the catalytically inactive CI037A mutant, used as a control). UBE2O-CA did not have conjugating activity in lysates, as the only bands evident in this case were biotin-ubiquitin and autoubiquitinated E1. In the absence of lysate, UBE2O was autoubiquitinated, as previously reported (27). (C) UBE2O ubiquitinates RPs in reconstituted Ube2o−/− lysates. Ubiquitin conjugates (see Fig. 2A) purified by NeutrAvidin-biotin pull-down were digested on beads with trypsin. Peptides containing diglycine-modified lysines (ubiquitination sites) were identified and mapped by LC-MS/MS. All RP ubiquitination events were unique to the UBE2O-WT sample from those of the UBE2O-CA sample from those of the UBE2O-WT reaction for a given diglycine peptide over six replicate experiments. (D) Levels of RPs from Ube2o−/− and wild-type reticulocytes assessed by immunoblotting (α denotes an antibody). GAPDH and α-spectrin are loading controls; 100 μg of protein was loaded per lane. (E) Volcano plot of quantitative proteomics analysis, representing the relation of the log2 of the fold change [log2(Ube2o+/+/Ube2o−/−)] and the log2 of the P value adjusted using Benjamini-Hochberg correction. STRING analysis (16) also showed a definitive cluster of RPs (fig. S6, B and C). When conjugate purification was repeated under denaturing conditions, RPs remained the major class of ubiquitination targets (fig. S6D and table S1).

Thus, the ubiquitination pattern does not reflect preexisting UBE2O-independent ubiquitin modifications that cofractionated with UBE2O-dependent modifications simply because they were present on the same ribosome.

A hallmark of the transition from the reticulocyte to the erythrocyte stage of differentiation is the elimination of ribosomes (17). The associated mechanism is poorly understood (18, 19). We probed reticulocyte lysates from wild-type and

Table 1. Ube2o+/− mice have hypochromic, microcytic anemia. Erithroepoietic parameters in Ube2o+/− (Ube2o+/−) and genetic interaction between Ube2o−/− and the β-globin–deficient Hbbth3 allele. Spleen:BW, ratio of spleen weight to body weight; RBC, red blood cells; HGB, hemoglobin; MCV, mean cell volume; MCH, mean corpuscular hemoglobin; RETIC ABS, absolute reticulocytes; Chr, mean reticulocyte hemoglobin content. All studies were performed in 8-week-old female littermates, with n = 6 animals in each group. Significance was calculated by one-way ANOVA with Tukey multiple-comparisons test. *P < 0.05 versus wild type; †P < 0.05 versus Hbbth3+/− Ube2o+/−; ‡P < 0.05 versus Hbbth3+/− Ube2o−/−. Values are means ± SD. (See also fig. S3F.)
Ube2o−/− mice with antibodies to seven RPs (Fig. 1D). In all cases, we observed elevated levels in the mutants, consistent with a broad defect in RP elimination in Ube2o−/− mice.

To search globally for proteins that may be targets of UBE2O, we compared the proteomes of wild-type and Ube2o−/− reticulocytes by tandem mass tagging (TMT) mass spectrometry (20) (fig. S7). Of 1235 proteins quantified (Fig. 1E and table S2), 183 were elevated in mutant reticulocytes. The results suggested that UBE2O, when induced to high levels, might execute a broad program of ubiquitination and degradation. KEGG pathway enrichment analysis yielded a highly significant enrichment of ribosomes (Fig. 1E). Ferritins (FTL1 and FTH1) were also elevated in Ube2o−/− reticulocytes (Fig. 1E). The increase in ferritin levels appears to reflect translational control, because ribosome profiling studies showed elevated levels of translating mRNA for FTL and FTH1 in the mutant (fig. S8A). Ribosome pausing was prominent at iron-responsive elements with the FTH1 and FTH1 mRNAs in wild-type but not mutant reticulocytes (fig. S8B); this result suggested that Ube2o−/− reticulocytes are responding to increased cytosolic iron or heme levels and that reticulocyte iron deficiency is unlikely to be the cause of the anemia. Accordingly, the heme-regulated eIF2α kinase (HRI), a sensor of iron deficiency, was not induced in Ube2o−/− mutants, nor did a null mutation in the HRI-encoding gene modify the anemic phenotype (fig. S9).

**Ube2o−/− mice are deficient in ribosome elimination**

We next tested whether UBE2O governs the level of RPs (or, more narrowly, that of free RPs, which are typically unstable (21)). Sucrose gradient analysis showed that Ube2o−/− reticulocytes have elevated levels of ribosomes, particularly 80S monosomes (Fig. 2A and fig. S10). When cultured ex vivo, reticulocytes mature into erythrocytes over 48 to 72 hours (19, 22), with both ribosomes and mitochondria being eliminated. Cultured reticulocytes were analyzed by flow cytometry using MitoTracker Deep Red, which stains mitochondria, and thiazole orange, which stains RNA; in reticulocytes, the most abundant RNA is ribosomal (rRNA) (Fig. 2B). After 72 hours of ex vivo differentiation, the median fluorescence intensity of thiazole orange staining in Ube2o−/− reticulocytes was higher than in the wild type by a factor of 2.5, whereas MitoTracker staining was higher in the mutant only by a factor of 1.6. The lack of significant UBE2O involvement in mitochondrial elimination is consistent with previous indications that maturation-associated mitophagy, an autophagic process, is ubiquitin-independent in reticulocytes (19, 23).

Polysome profiles from wild-type reticulocytes cultured ex vivo showed that ribosome elimination was nearly complete by 31 hours; by contrast, the 80S monosome peak remained prominent in wild-type and Fth1−/− reticulocytes (Fig. 2A), consistent with delayed turnover of multiple RPs during maturation (fig. S11). Thus, ribosome levels are elevated in Ube2o−/− reticulocytes because of a defect in ribosome elimination during terminal differentiation. Despite the persistence of this peak, few polysomes were evident in the gradient profile, perhaps because of low levels of mRNA or translational initiation factors at this stage of differentiation.

**UBE2O is sufficient to drive ribosome elimination in non-erythroid cells**

Having found that UBE2O is critical for ribosome elimination in reticulocytes, we tested whether it is sufficient for elimination in a non-erythroid cell, as might be expected if UBE2O recognized RPs directly. We integrated the full-length UBE2O open reading frame into Flp-In T-REx 293 cells under a doxycycline-inducible promoter. The UBE2O-C4 mutant, integrated at the same locus, was used as a control (Fig. 3A and fig. S12, A and B). UBE2O was induced for 48 to 72 hours to recapitulate the period of terminal erythroid maturation (fig. S12A). 293-E2O cells remained viable, with no indication of induced cell death (fig. S12C). The effect of UBE2O expression on the proteome was assessed by mass spectrometry, which quantified 7807 proteins (fig. S13, A and B, and table S3). Upon UBE2O induction, we identified 858 proteins present at substantially lower levels (≥50%) than in UBE2O-CA by day 3 (Fig. 3B), with a clear preferential effect on basic proteins (Fig. 3D). Thus, ectopic UBE2O induction led to extensive proteome remodeling.

Among proteins whose levels were reduced more than 50% upon UBE2O induction, KEGG pathway enrichment analysis identified a highly significant down-regulation of RPs (Fig. 3, B and C). This was confirmed by immunoblotting for RPL29, RPL35, and RPL23A (Fig. 3E). RPs were not degraded at comparable rates, inconsistent with what would be expected for an autophagic mechanism. Ribosome degradation might proceed in steps, with RPs showing the fastest degradation more likely to be direct substrates of UBE2O. We also observed significant enrichment in ribosome biogenesis and rRNA processing pathways, which are nonfunctional in reticulocytes because they are enucleate (Fig. 3B and fig. S13, C to E). These effects of UBE2O are likely direct, because multiple ribosomal and nuclear proteins were ubiquitinated by recombinant UBE2O when added to extracts of human embryonic kidney (HEK) 293 cells and analyzed by mass spectrometry (fig. S14, A to C, and table S1). After UBE2O induction, polysome profiles showed a strong reduction of the 80S monosome peak (Fig. 3F). Although reticulocytes cannot produce ribosomes de novo, a reduction of ribosome levels in 293-E2O cells could reflect either degradation of preexisting ribosomes or interference with ribosome synthesis. To discriminate between these possibilities, we suppressed the transcription of new rRNA with a selective inhibitor of RNA polymerase I, CX-5461 (24, 25). Under these “chase” conditions, UBE2O drove the destabilization of RPs after 24 to 48 hours of inhibitor treatment, with no effect observed in the catalytic null mutant (fig. S15). Therefore, preexisting RPs can be eliminated by UBE2O.

In summary, the 293-E2O cell data recapitulate a critical aspect of normal erythroid differentiation and confirm the inducible breakdown of ribosomes—complexes that are thought to be highly stable under basal conditions. Furthermore, UBE2O is sufficient to drive ribosome elimination in non-erythroid cells in the absence of reticulocyte-specific factors, which suggests that the control of erythroid ribosome turnover may be dependent principally on UBE2O itself.

**UBE2O executes a broad program of ubiquitination**

On the basis of proteomic data on reticulocytes and 293-E2O cells, we selected seven potential nonribosomal targets of UBE2O for confirmation by immunoblotting. We observed strongly elevated levels of all of these proteins in mutant reticulocytes, including a chaperone for α-globin,
AHSP (α-hemoglobin stabilizing protein) (Fig. 4A). AHSP was strongly stabilized in a time-course study using ex vivo cultured Ube2o−/− reticulocytes (Fig. S16A). However, elevated AHSP levels do not explain the erythroid phenotype of these mutants, because an Ahsp−/− mutation did not alleviate the Ube2o−/− phenotype (Fig. S16, B and C). In summary, immunoblot analysis validated the proteomics data and thus suggested that an extensive set of RPs and non-RPs is eliminated from reticulocytes under the control of UBE2O.

Our finding that UBE2O is sufficient to drive the degradation of its target proteins suggests that it may mediate direct recognition of these proteins. However, E3 rather than E2 enzymes typically mediate substrate recognition (26). The molecular mass of UBE2O is 143 kDa (Fig. 4B), whereas typical E2 enzymes are of 20 to 25 kDa. It therefore seemed likely that UBE2O may function as an E3 enzyme fused to an E2 (27–29). To test this model, we reconstituted the ubiquitination of candidate UBE2O substrates with purified components.

When partially purified ribosomes were tested by immunoblot analysis, we observed that ubiquitination of RPs was efficient relative to that of histone H2B, a model substrate of UBE2O (Fig. 4C). The ubiquitin acceptor proteins in this reaction were confirmed as RPs by LC-MS/MS (Fig. S18). In summary, several in vivo ubiquitination sites mapping (fig. S17) were confirmed as RPs by LC-MS/MS (Fig. 4C).

Ribosomal proteins were assayed, as substrates. Indeed, when tested individually, recombinant RPL35, RPL36A, and RPL37 all proved to be UBE2O substrates (Fig. 4D). Other proteins predicted from proteomics data to be UBE2O substrates were all efficiently ubiquitinated, whereas calmodulin, a negative control, was not (Fig. 4D). UBE2O added multiple ubiquitin groups to these substrates, although almost exclusively in the form of multi-mono ubiquitin modification (Fig. S18). In summary, several in vivo targets of UBE2O were also modified by this enzyme in a purified system, which suggests that UBE2O, in addition to being an E2 enzyme, is also an E3—a specificity-determining ubiquitination factor.

In addition to its E2 (or UBC) domain, UBE2O has four distinct evolutionarily conserved regions: CR1, CR2, CR3, and a predicted coiled-coil (CC) element (Fig. 4B). When truncated forms of UBE2O were assayed, the importance of the tested domains for conjugation rates varied depending on the substrate tested, consistent with CR1 and perhaps other conserved domains being substrate specificity elements (Fig. 4C). The hypothesis that CR1 and CR2 are substrate recognition domains was assessed more directly by testing them for interaction with UBE2O substrates in pull-down binding assays. Both CR1 and CR2 bound UBE2O substrates, although their binding specificity differed (Fig. 4E). CR1 and CR2 contain acidic patches, which may be involved in their binding to basic proteins. A confirmed acidic substrate, AHSP, did not show significant binding to CR1 or CR2 (Fig. 4E), nor was its ubiquitination dependent on these domains (Fig. 4C). These results indicate that UBE2O is capable of multiple modes of substrate recognition.

In summary, UBE2O has the expected features of an E2-E3 hybrid enzyme in which ubiquitin is charged by the UBC domain of UBE2O, to be subsequently donated to proteins that are recognized in most cases by the CR1 and CR2 domains.

Ribosomal proteins are degraded by the proteasome

Previous studies have shown that regulated ribosome degradation occurs via autophagy in yeast (30–33). In reticulocytes, however, inhibitors of autophagy do not affect ribosome breakdown (19). Selective autophagy in reticulocytes is mediated by NIX, but this factor drives the elimination of mitochondria rather than ribosomes (19, 23, 34).

To test whether RPs are eliminated by the proteasome, we incubated reticulocytes with the proteasome inhibitors PS-341 and epoxomicin. After ex vivo differentiation, treated wild-type reticulocytes retained thiazole orange staining, resembling Ube2o−/− reticulocytes that had not been exposed to proteasome inhibitor (Fig. 5A and fig. S19A). These data suggest that RPs are degraded by the proteasome. In fact, treated wild-type reticulocytes retained their thiazole orange staining more fully than did untreated Ube2o−/− reticulocytes (Fig. 5A), suggesting the existence of alternative pathways of ribosome elimination that are UBE2O-independent but proteasome-dependent. In contrast to the proteasome, p97—
which dissociates components of several multi-subunit complexes in a ubiquitin-dependent manner (35)—had no detectable effect on ribosome elimination (fig. S20).

Ubiquitin depletion is evident in reticulocytes treated with the highest concentrations of proteasome inhibitors (fig. S19B). To address this potential caveat, we reconstituted RP degradation in a cell-free reticulocyte lysate system, which permitted supplementation of free ubiquitin. The degradation of RPs was reconstituted by adding back UBE2O to null reticulocyte lysate. RPs were not destabilized by UBE2O-CA, or under conditions of proteasome inhibitor treatment, while maintaining free ubiquitin levels (Fig. 5B and fig. S19C). However, the stabilizing effect of the inhibitors, although apparently complete for RPL29 and RPS23, was partial in the case of RPL35 and RPL36A, as judged from the intensity of the major protein band (Fig. 5B and fig. S19C). For both proteins, a high-molecular weight species appeared upon proteasome inhibition, which likely represented ubiquitinated forms of RPL35 and RPL36A. To confirm that proteasome inhibitors completely blocked RP degradation, we treated the reaction with USP21 to deubiquitinate high-molecular weight forms of the protein. This treatment concentrated RPL35 into a single band that was constant in intensity over the time course of incubation (fig. S19D).

The importance of the proteasome in remodeling the proteome in reticulocytes was further tested by metabolomic analysis. Amino acids such as Lys and Arg, which are abundant in RPs, were indeed depleted by proteasome inhibitors (Fig. 5C). We therefore considered whether UBE2O might cooperate with other ubiquitinating factors that may be induced in late erythroid differentiation. Using RNA-sequencing data for erythroid cells taken from progressive stages of differentiation (6–8), we identified eight E2 and E3 enzymes that are induced in late erythroid cells in parallel with UBE2O (Fig. 5D). We hypothesize that these enzymes work in concert to globally remodel the erythroid proteome. Expression of the vast majority of ubiquitinating enzymes is extinguished during this period (fig. S21, A and B), so that the UPS is not simply amplified during erythroid differentiation, but is instead deeply reconfigured. The unique transformation of the UPS in these cells may reflect a strategy to eliminate thousands of cellular proteins from reticulocytes, thus enabling extreme concentration of globin.

Discussion

In many cases of terminal cellular differentiation, the proteome is remodeled on a global scale to generate a highly specific cellular phenotype. The extreme case is erythroid differentiation, where a single species, hemoglobin, is concentrated to ~98% of soluble protein (7). Accomplishing this rapid transformation of the cellular phenotype requires not only highly active protein synthesis, but also degradation of a vast set of preexisting, otherwise relatively stable, soluble proteins. This degradation must also be selective, so as to spare properly assembled globin and other components of the mature RBC. Despite this active degradation, most components of the ubiquitin pathway are down-regulated transcriptionally during erythroid differentiation. However, a small subset of ubiquitin-conjugating enzymes and ubiquitin ligases is induced, among them UBE2O. We propose that UBE2O plays a major role in proteomic remodeling, with tens and perhaps hundreds of proteins being eliminated through its activity.

As an E2-E3 hybrid enzyme, UBE2O functions autonomously and can thus remain active well into the reticulocyte stage, promoting maturation at a time when most E2 enzymes have been lost. The ability of UBE2O to promote reticulocyte maturation also depends on its broad substrate specificity; multiple substrate recognition domains enable the recognition of diverse substrates and may, through avidity, allow coordinate low-affinity substrate interactions to be productive of ubiquitination. In an accompanying paper, Yanagitani et al.
degradation, UBE2O-dependent ubiquitination
multiubiquitin chains are required to initiate
ings show that ubiquitination of RPs, in either
Nguyen et al previously been observed (44, 45),

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UBE2O prefers basic substrates, including RPs,

Significance was calculated by two-sample

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ubiquitin ligase. Because quality-control ligases are

ubiquitinated RPs, in either their free or assembled state, can be used to

indeed, ribosome ubiquitination has previously been associated with the suppression of their turnover via autophagy (33).

Contrary to the longstanding paradigm that multiquibuitin chains are required to initiate degradation, UBE2O-dependent ubiquitination

events efficiently targeted substrates to the pro-

traversable to α-globin excess, as seen for a model of β-thalassemia. UBE2O is thus a potential ther-

apeutic target not only for mixed-lineage leukemia and other cancers, as recently proposed (44, 45), but also for β-thalassemia, one of the most prev-

ilegant inherited diseases worldwide.

REFERENCES AND NOTES


(36) report that UBE2O is a quality-control ubiquitin ligase. Because quality-control ligases are characteristically broad in their specificity, a ligase of this type may be pre-adapted to serve in global proteome remodeling when induced to high levels.

UBE2O prefers basic substrates, including RPs, and this specificity may underlie a defining feature of the reticulocyte-erythrocyte transition: the elimination of ribosomes (37). Although the ubiquitination of ribosomes and of free RPs has previously been observed (21, 37–39), our findings show that ubiquitination of RPs, in either their free or assembled state, can be used to promote the elimination of ribosomes from cells. Indeed, ribosome ubiquitination has previously been associated with the suppression of their turnover via autophagy (33).

Contrary to the longstanding paradigm that multiquibuitin chains are required to initiate degradation, UBE2O-dependent ubiquitination

Fig. 5. The ubiquitin-proteasome system in late erythroid differentiation. (A) Wild-type and Ube2o−/− reticulocytes were differentiated ex vivo at 37°C for 48 hours with proteasome inhibitors (50 nM epoxomicin and 50 nM PS-341) or DMSO vehicle control, then analyzed by fluorescence-activated cell sorter. Left: Wild-type reticulocytes treated with proteasome inhibitors versus DMSO control. Right: Both wild-type and Ube2o−/− reticulocytes treated with proteasome inhibitors and DMSO. (B) Reconstitution of RP degradation in Ube2o−/− reticulocyte lysates. Ubiquitin levels were supplemented to prevent depletion of free ubiquitin upon proteasome inhibition. Proteasomes were inhibited by adding PS-341 (50 nM) and epoxomicin (50 nM) together. GAPDH is a loading control; 100 µg of protein was loaded per lane. (C) Wild-type reticulocytes were treated with proteasome inhibitors (50 nM epoxomicin and 50 nM PS-341) and differentiated ex vivo for 48 hours. Quantitative metabolomic profiling of treated and untreated wild-type reticulocytes showed a depletion of multiple free amino acids due to proteasome inhibition, with Lys and Arg most strongly affected. Significance was calculated by two-sample t test. *P < 0.05, **P < 0.01, ***P < 0.001. (D) An ensemble of ubiquitin ligases and ubiquitin-conjugating enzymes is induced in late erythroid differentiation. Stages are arranged in a temporal progression. Stage 1 includes proerythroblasts and early basophilic erythroblasts; stage 2, early and late basophilic erythroblasts; stage 3, polychromatophilic and orthochromatophilic erythroblasts; stage 4, late orthochromatophilic erythroblasts and reticulocytes. The induction of globin mRNA is shown for comparison. Raw RNA-sequencing data are taken from (6).


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**SUPPLEMENTARY MATERIALS**

www.sciencemag.org/content/357/6350/eaan0218/suppl/DC1

Materials and Methods
Supplementary Text
Figs. S1 to S21
Tables S1 to S3

References (77–78)
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UBE2O remodels the proteome during terminal erythroid differentiation


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Removing orphan proteins from the system

The degradation of excess subunits of protein complexes is a major quality-control problem for the cell. How such "orphans" are recognized and tagged for degradation is poorly understood. Two papers identify a protein quality-control pathway that acts on some of the most abundant protein complexes in the human body: hemoglobin and ribosomes (see the Perspective by Hampton and Dargemont). Yanagitani et al. show that the central player in this process is an unusual enzyme (UBE2O) that recognizes substrates and tags them for destruction. Other quality-control pathways tend to use separate factors for target selection (often a chaperone), ubiquitin donation (an E2), and ubiquitin conjugation (an E3). Encoding all three activities in a single factor whose function can be reconstituted in a purified system provides a tractable route to detailed mechanistic and structural dissection. Nguyen et al. show the importance of the UBE2O pathway in the differentiation of red blood cells. _Science_, this issue p. 472, p. eaan0218; see also p. 450