Functions of the COPII gene paralogs SEC23A and SEC23B are interchangeable in vivo

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Approximately one-third of the mammalian proteome is transported from the endoplasmic reticulum-to-Golgi via COPII-coated vesicles. SEC23, a core component of coat protein-complex II (COPII), is encoded by two paralogous genes in vertebrates (*Sec23a* and *Sec23b*). In humans, SEC23B deficiency results in congenital dysery-thropoietic anemia type-II (CDAII), while SEC23A deficiency results in a skeletal phenotype (with normal red blood cells). These distinct clinical disorders, together with previous biochemical studies, suggest unique functions for SEC23A and SEC23B. Here we show indistinguishable intracellular protein interactomes for human SEC23A and SEC23B, complementation of yeast Sec23 by both human and murine SEC23A/B, and rescue of the lethality of *sec23b* deficiency in zebrafish by a *sec23a*-expressing transgene. We next demonstrate that a *Sec23a* coding sequence inserted into the murine *Sec23b* locus completely rescues the lethal SEC23B-deficient pancreatic phenotype. We show that SEC23B is the predominantly expressed paralog in human bone marrow, but not in the mouse, with the reciprocal pattern observed in the pancreas. Taken together, these data demonstrate an equivalent function for SEC23A/B, with evolutionary shifts in the transcription program likely accounting for the distinct phenotypes of SEC23A/B deficiency within and across species, a paradigm potentially applicable to other sets of paralogous genes. These findings also suggest that enhanced erythroid expression of the normal *SEC23A* gene could offer an effective therapeutic approach for CDAII patients.

endoplasmic reticulum | vesicular transport proteins | mice | transgenic | SEC23

In vertebrates, ~8,000 secretory proteins (cargos) are cotranslationally inserted into the endoplasmic reticulum (ER) and subsequently transported to the Golgi apparatus via coat protein-complex II (COPII)-coated vesicles before reaching their final destination (endosomes, lysosomes, plasma membrane, or the extracellular space) (1–6). SAR1 and SEC23-SEC24 heterodimers form the inner COPII coat, with the outer coat composed of SEC13-SEC31 heterotetramers. COPII coat assembly begins with GTP-bound SAR1 recruiting cytoplasmic SEC23-SEC24 heterodimers to the ER membrane (7, 8), through direct interaction between SAR1 and SEC23. SAR1-SEC23-SEC24 forms the prebudding complex, which recruits cargo proteins (9–15) either via direct interaction (3, 10, 16–18) or via interaction with intermediary proteins (termed receptors or adaptors) (3, 18–22). Following cargo selection, SEC13-SEC31 heterotetramers are recruited, forming the outer COPII coat. SEC23 functions as a GTPase activating protein stimulating SAR1-GTP hydrolysis, a process required for vesicle fission in vivo (4, 7). The SEC23 GTPase activity is enhanced by SEC31 (23).

The mammalian genome encodes multiple paralogs for SEC23, SEC24, SAR1, and SEC31. These paralogs have been proposed to contribute to COPII coat diversity, facilitating distinct cargo selectivity and tissue-specific secretory repertoires. Alternative splicing has been described for *SEC31* and *SEC24C* (24, 25) and might additionally contribute to COPII diversity and cargo selectivity. Alhough the *SEC23* gene duplication is estimated to have occurred ~615 Mya, the two mammalian paralogs, *SEC23A* and *SEC23B*, encode proteins that are ~85% identical

Significance

In humans, SEC23B deficiency results in congenital dysery-thropoietic anemia type II, a disease of abnormal red blood cell development, while SEC23A deficiency results in cranio-lenticulo-sutural-dysplasia, a disease characterized by bone abnormalities due to defective collagen secretion (but no red blood cell defect). In this study, we show that SEC23A and SEC23B overlap in function, and that the disparate phenotypes of SEC23A/SEC23B deficiency within and across species are likely due to evolutionary shifts in gene-expression programs, rather than distinct functions of the SEC23 paralogs. Our studies provide a rationale for increased SEC23A or SEC23B expression as a therapeutic strategy for congenital dyserythropoietic anemia type II or cranio-lenticulo-sutural-dysplasia, respectively.

at the amino acid sequence level. The sequence identity between the mouse and human orthologs for SEC23A and SEC23B are ~98% and ~97%, respectively.

Mutations in human *SEC23A* result in cranio-lenticulo-sutural dysplasia (CLSD), an autosomal recessive disease characterized by abnormal closure of cranial fontanelles, dysmorphic facial features, skeletal abnormalities, and sutural cataracts (26, 27), features thought to result from defects in collagen secretion. In contrast, loss of SEC23B function results in congenital dyserythropoietic anemia type II (CDAII), an autosomal recessive disease characterized by anemia and increased bi/multinucleated erythroid precursors in the bone marrow (BM) (28).

In mice, germline *Sec23a* deletion results in an open neural tube defect and compromised extraembryonic membranes, likely due to impaired collagen secretion, a phenotype reminiscent of human CLSD (29). In contrast, mice with germline SEC23B deficiency die perinatally, exhibiting massive pancreatic degeneration (30–32) with no anemia or other features of CDAII (30, 33). In both instances, partially compensatory increases in the remaining paralog were shown to be posttranslational, likely due to enhanced stability of SEC23/SEC24 heterodimers (29).

Previous reports suggest unique functions for SEC23A and SEC23B (29, 34–37), potentially due to distinct sets of cargo proteins. Additionally, human SEC23A but not SEC23B was reported to complement yeast SEC23 (35), consistent with distinct function of the two mammalian paralogs.

In surprising contrast to these earlier findings, we now report indistinguishable intracellular interactomes for human SEC23A and SEC23B and also show complementation of yeast SEC23 by both human and murine SEC23A/B paralogs. We further show that the lethality resulting from *sec23b* disruption in zebrafish can be rescued by a *sec23a*-expressing transgene. Finally, we demonstrate that the *Sec23a* coding sequence inserted into the endogenous murine *Sec23b* locus fully rescues the mortality and severe pancreatic phenotype of SEC23B-deficiency in the mouse.

Fig. 1. SEC23A and SEC23B exhibit indistinguishable interactomes in HEK293 cells. (*A*) The constructs shown here, directing expression of SEC23A or SEC23B fused to BirA* and a FLAG tag under the control of a tetracycline-inducible promoter, were stably inserted into Flp-In T-REX HEK293 cells. (*B*) FLAG immunoblotting following tetracycline induction. (*C–F*) Following tetracycline and biotin administration, the full sets of SEC23A and SEC23B interactomes were each determined with BioID, in two independent experiments. Each "interacting protein" is represented by a dot and the spectral counts are indicated on the *x* and *y* axes. Pairwise comparisons of spectral counts for a given protein are shown across paralogs (*C* and *D*) or between biologic replicas for the same paralog (*E* and *F*).

Results

The SEC23A and SEC23B Interactomes Are Indistinguishable. Recruitment of SAR1-SEC23-SEC24 to the cytoplasmic face of the ER membrane forms the inner coat of the COPII vesicle. Interaction of SEC23 with each of the four SEC24 mammalian paralogs (SEC24A–D) provides a potential for functional differentiation between SEC23A and SEC23B. However, we detected no differences in paralog specificity by coimmunoprecipitation (*SI Appendix*, Fig. S1).

To comprehensively characterize the full set of interacting proteins for each SEC23 paralog, we performed two independent "BioID" (proximity-dependent biotinylation) experiments in HEK293 cells equivalently expressing either BirA*-tagged SEC23A, SEC23B, or a GFP control (Fig. 1 *A* and *B*). As expected, known SEC23-interacting proteins were identified, including the COPII structural proteins SEC24, SAR1, SEC13, and SEC31. Comparison of the SEC23A to SEC23B interactomes revealed a remarkably high correlation in two independent experiments (Fig. 1 *C* and *D* and *SI Appendix*, Fig. S2), equivalent to the correlation between biological replicates for SEC23A (Fig. 1*E*) or SEC23B (Fig. 1*F*). Thus, the comprehensive interactomes of SEC23A and SEC23B in HEK293 cells are indistinguishable by BioID.

Mouse and Human SEC23 Paralogs Complement the Yeast Sec23 Protein. Human SEC23A but not human SEC23B was previously reported to complement the yeast Sec23 protein (35), a puzzling finding in light of the above evidence for indistinguishable SEC23A/B interactomes in mammalian cells. We thus retested complementation in a temperature sensitive *sec23-1* mutant yeast strain using both human (h) and mouse (m) SEC23A and -B sequences. Indeed, all four mammalian sequences (hSEC23A, hSEC23B, mSEC23A, and mSEC23B) rescued growth of the temperature-sensitive *sec23-1* mutant yeast at the restrictive temperature (Fig. 2*A*). The discrepancy between these findings and the previous report (35) is likely explained by poorer expression of hSEC23B in yeast compared with hSEC23A (Fig. 2*B* and *SI Appendix*, Fig. S3), potentially due to differences in codon usage and protein stability.

Sec23b Deficiency Does Not Produce an Erythroid Phenotype in Zebrafish. The zebrafish (ZF) genome also contains two *sec23* paralogs (*sec23a* and *sec23b*). SEC23B deficiency in humans results in CDAII, a disease characterized by anemia and increased binucleated erythroid precursors (6), with *sec23b* knockdown in ZF previously reported to similarly result in an increased percentage of circulating binucleated erythrocytes (28). In contrast, SEC23B deficiency in mice results in perinatal mortality and massive pancreatic degeneration but no findings comparable to CDAII (30–33). To address these discordant phenotypes across species, we generated ZF heterozygous for a 53-bp deletion in exon 5 of *sec23b* (*sec23b*^{+/−}) by CRISPR/Cas9 genome editing. No difference in the percentage of circulating binucleated erythrocytes was noted at day 16 between *sec23b*^{−/−} ZF compared with WT and *sec23b*^{+/−} ZF clutchmates (Fig. 3 *A* and *B*). Similarly, day 7 *sec23b*^{−/−} ZF exhibited no difference in hemoglobin content (as assessed by *o*-dianisidine staining) compared with WT controls (Fig. 3*C* and *SI Appendix*, Fig. S4*B*). However, no *sec23b*^{−/−} ZF survived beyond 3 wk of age (Fig. 3*D*), although no gross or histologic abnormalities were detected at these time points (including in the pancreas), with the exception of reduced overall ZF size (Fig. 3*E* and *SI Appendix*, Fig. S4*A*).

Sec23a Expression Rescues the Lethality of Sec23b Deficiency in ZF. To determine if Sec23a could rescue the early mortality of Sec23b-deficient ZF, one-cell–stage embryos generated from *sec23b*^{+/−} intercrosses were injected with a vector (designed to integrate into the ZF genome) expressing Sec23a and GFP or GFP alone under
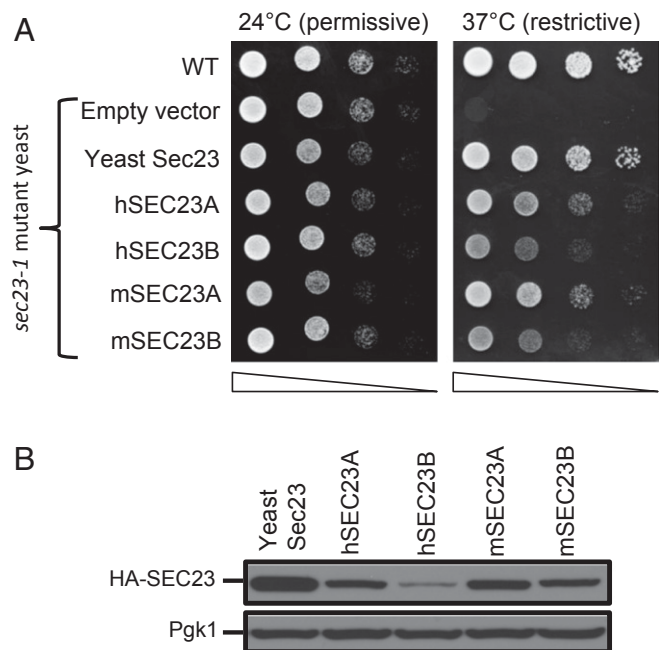

Fig. 2. Human and mouse SEC23A and SEC23B complement the yeast Sec23 protein. (*A*) The coding sequences of the human (h) or mouse (m) SEC23 paralogs (as well as yeast Sec23) were cloned with a 3′ sequence encoding an HA-tag into the pRS416 vector downstream of a *ZEO1* promoter. In contrast to WT yeast, temperature-sensitive *sec23-1* mutant yeast transfected with a control "empty vector" grow at the permissive (24 °C) but not at the restrictive (37 °C) temperature. Growth of temperature-sensitive *sec23-1* mutant yeast expressing h or m SEC23A/SEC23B (or yeast Sec23) was determined at the permissive and restrictive temperatures. Yeast cultures were plated as 10-fold serial dilutions. (*B*) Immunoblotting for HA (with Pgk1 as loading control) detects relative expression for each construct.

the control of a ubiquitin promoter (*SI Appendix*, Fig. S4*C*). Genotyping of transgene-expressing ZF (identified by GFP positivity) (*SI Appendix*, Fig. S4*D*) demonstrated that Sec23a rescues the mortality of Sec23b-deficiency in ZF (Fig. 3 *F* and *G*).

Generation of Mice Expressing Sec23a from the Sec23b Locus. The ZF results suggest at least partial overlap in function between SEC23A and SEC23B. To characterize the capacity of the SEC23A protein to replace SEC23B in mammals in vivo, we substituted *Sec23a* coding sequences for *Sec23b* (at the endogenous *Sec23b* genomic locus). Although efforts to engineer this allele by direct oocyte injection were unsuccessful (*SI Appendix*, Fig. S5*A*), embryonic stem (ES) cells carrying the *Sec23b*[b-a] allele were generated as shown in Fig. 4*A*, and used to obtain mice carrying the *Sec23b*[b-a] allele. *Sec23b*[+/b-a] mice (Fig. 4 *B* and *C*) were observed at the expected Mendelian ratio in N2 progeny (Table 1, *Sec23b*[+/b-a] × *Sec23b*^{+/+}). *Sec23b*[+/b-a] mice were indistinguishable from their WT control littermates, exhibiting normal growth, fertility, and gross appearance, and no abnormalities on standard pathology evaluation.

The Sec23b[b-a] Allele Rescues the Mortality of Sec23b^{−/−} Mice. An intercross between mice heterozygous for the *Sec23b*[b-a] allele generated the expected number of *Sec23b*[b-a/b-a] offspring at weaning (Table 1, *Sec23b*[+/b-a] × *Sec23b*[+/b-a]). In WT mice, SEC23B is the predominantly expressed SEC23 paralog in the pancreas (Fig. 4*D*). Consistent with substitution of the SEC23A coding sequence for SEC23B, no SEC23B immunoreactive material was detected in pancreas tissues of *Sec23b*[b-a/b-a] mice, with high levels of SEC23A protein observed (Fig. 4*D*), in contrast to WT mice, which exhibit the
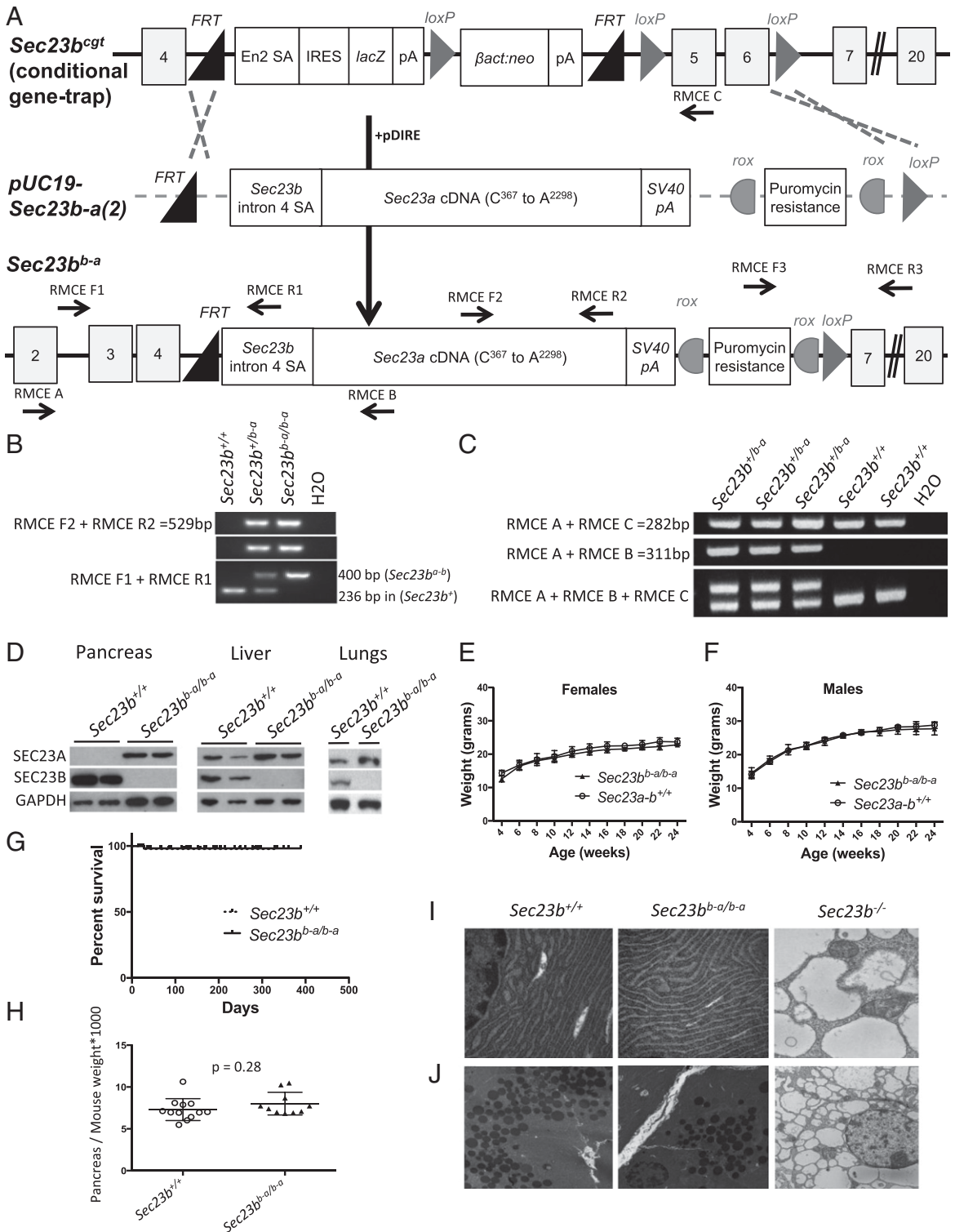
A

Investigator	Genotype	# mono-nucleated RBC	# bi/multi-nucleated RBC	% bi/multi-nucleated RBC	p-value
#1	WT (n = 5)	4119	6	0.145	-
	sec23b+/− (n = 8)	6204	10	0.161	0.79
	sec23b−/− (n = 7)	4536	8	0.176	0.51
#2	WT (n = 4)	2201	4	0.181	-
	sec23b−/− (n = 4)	2135	4	0.187	0.94

B

C

D

Progeny of sec23b+/− intercross	WT	sec23b+/−	sec23b−/−	P-value
8 Weeks	11	19	0	0.0012
4 Weeks	10	18	0	0.0016
3 Weeks	10	35	1	0.0004
18 days	7	10	7	0.34
14 days	2	5	3	0.37
7 days	1	11	4	0.5

E

F

Progeny of sec23b+/− intercross	Injected with	WT	sec23b+/−	sec23b−/−	p-value
4 Weeks	sec23a-p2A-GFP	2	13	3	0.22
4 Weeks	Empty vector	12	33	0	0.0001

G

Fig. 3. Mortality in Sec23b-deficient ZF between 18 and 21 d is rescued by Sec23a. (A and B) sec23b heterozygous (sec23b+/−) ZF with a 53-bp deletion were generated with CRISPR/Cas9. Binucleated erythrocyte counts from peripheral blood of ZF offspring from a sec23b+/− × sec23b+/− intercross were evaluated by two independent investigators blinded to the ZF genotype. (C) sec23b−/− ZF exhibit indistinguishable o-dianisidine staining compared with WT clutchmate controls. Scores 1–3 correspond to staining intensities (3 = highest staining intensity). (D) sec23b−/− ZF (generated from sec23b+/− intercrosses) die between 18 and 21 d. (E) H&E stains of sec23b−/− and WT ZF at day 16. sec23b−/− ZF appear histologically normal. (Magnification: 10×.) (F) One-cell–stage embryos generated from sec23b+/− ZF intercrosses were injected with a vector (designed to integrate in the ZF genome) that expresses Sec23a and GFP or GFP alone as control. Genotyping at 4 wk of age demonstrates rescue of sec23b−/− mortality by Sec23a. (G) PCR genotyping assay demonstrating 2 sec23b−/− ZF, expressing Sec23a from a ubiquitin promoter, alive at 4 wk of age.

opposite pattern. SEC23B was also undetectable in SEC23b^{b-a/b-a} liver and lung tissues (Fig. 4D).

The Sec23b^{b-a} Allele Rescues the Pancreatic Phenotype of Sec23b^{−/−} Mice. Sec23b^{b-a/b-a} mice exhibited normal growth (Fig. 4 E and F) and fertility, normal gross appearance, no abnormalities on histopathologic evaluation (Fig. 5 and SI Appendix, Fig. S6), and normal overall survival (up to 1.5 y of follow-up) (Fig. 4G). In contrast to the pancreatic degeneration and distended ER observed in Sec23b^{−/−} mice (32, 33), pancreas tissue from Sec23b^{b-a/b-a} mice exhibited normal weight (Fig. 4H) and histology (Fig. 5A),

indistinguishable from pancreas of WT littermate controls. The Sec23b^{b-a/b-a} pancreatic ER also appeared normal by transmission electron microscopy (Fig. 4I), with intact acinar cell zymogen granules (Fig. 4J), in contrast to the appearance in Sec23b^{−/−} mice (Fig. 4 I and J). Other secretory tissues that were perturbed in SEC23B-deficient mice (including salivary glands, nasal glands, gastric epithelium, and intestinal cells) (32) also appeared entirely normal histologically in Sec23b^{b-a/b-a} mice (Fig. 5 B–E).

Taken together, these findings demonstrate that the SEC23A protein fully rescues the SEC23B-deficient phenotype in mice,

Fig. 4. Mice expressing *Sec23a* from the *Sec23b* locus (*Sec23b^{b-a/b-a}*) exhibit normal growth and survival. (*A*) Design of the replacement vector pUC19-Sec23b-a(2), which was coelectroporated with pDIRE into *Sec23b^{+/cgt}* embryonic stem cells. Arrows indicate locations of the genotyping primers. (*B*) PCR demonstrates the expected band sizes for correct *Sec23b^{b-a}* 5′ and 3′ insertions. (*C*) RT-PCR identifies the expected splicing of *Sec23b* exon4 into the *Sec23a* cDNA. (*D*) SEC23B is the predominantly expressed SEC23 paralog in the WT mouse pancreas. In contrast, mice with insertion of the *Sec23a* coding sequence into the *Sec23b* locus have no detectable SEC23B in their pancreas tissues by Western blot, with a high level of SEC23A detected. Similarly, SEC23A is the exclusively expressed SEC23 paralog in other tissues from *Sec23b^{b-a/b-a}* mice, including liver and lung. (*E* and *F*) *Sec23b^{b-a/b-a}* female and male mice exhibit normal growth compared with WT littermate controls. (*G*) *Sec23b^{b-a/b-a}* mice exhibit normal survival compared with WT littermate controls. (*H*) Pancreas tissues from *Sec23b^{b-a/b-a}* mice exhibit normal weights compared with WT littermate controls. (*I* and *J*) Normal appearance of ER (*I*) and zymogen granules (*J*) in *Sec23b^{b-a/b-a}* (and WT *Sec23b^{+/+}*) compared with *Sec23b^{-/-}* pancreas tissues by electron microscopy. [Magnification: 25,000× (*I*); 7,900× (*J*).]

Table 1. Mouse crosses with the $Sec23b^{b-a}$ allele

Mouse crosses and genotypes	Expected, %	Observed at weaning, %	Observed at E14.5, %	Observed at E13.5, %
$Sec23b^{+/b-a} \times Sec23b^{+/+}$*				
$Sec23b^{+/+}$	50	47.8 (155)		
$Sec23b^{+/b-a}$	50	52.2 (169)		
$Sec23b^{+/b-a} \times Sec23b^{+/b-a}$†				
$Sec23b^{+/+}$	25	23.4 (67)		
$Sec23b^{+/b-a}$	50	50.5 (145)		
$Sec23b^{b-a/b-a}$	25	26.1 (75)		
$Sec23a^{+/-} \times Sec23b^{+/b-a}$‡				
$Sec23a^{+/+}\ Sec23b^{+/+}$	25	29.1 (23)		
$Sec23a^{+/+}\ Sec23b^{+/b-a}$	25	21.5 (17)		
$Sec23a^{+/-}\ Sec23b^{+/+}$	25	29.1 (23)		
$Sec23a^{+/-}\ Sec23b^{+/b-a}$	25	20.3 (16)		
$Sec23a^{+/-}\ Sec23b^{+/b-a} \times Sec23a^{+/-}$§				
$Sec23a^{+/+}\ Sec23b^{+/+}$	16.7	16.1 (14)	11.1 (2)	12.2 (5)
$Sec23a^{+/+}\ Sec23b^{+/b-a}$	16.7	19.5 (17)	11.1 (2)	9.8 (4)
$Sec23a^{+/-}\ Sec23b^{+/+}$	16.7	28.7 (25)	22.2 (4)	53.6 (22)
$Sec23a^{+/-}\ Sec23b^{+/b-a}$	16.7	35.6 (31)	55.6 (10)	24.4 (10)
$Sec23a^{-/-}\ Sec23b^{+/+}$	16.7	0 (0)	0 (0)	0 (0)
$Sec23a^{-/-}\ Sec23b^{+/b-a}$	16.7	0 (0)	0 (0)	0 (0)

No. of mice is expressed in parentheses.
*Observed at weaning, $n = 324$; $P = 0.73$ (P value calculated for $Sec23b^{+/b-a}$ vs. WT mice).
†Observed at weaning, $n = 287$; $P = 0.89$ (P value calculated for $Sec23b^{b-a/b-a}$ vs. all other genotypes).
‡Observed at weaning, $n = 79$; $P = 0.48$ (P value calculated for $Sec23a^{+/-}\ Sec23b^{+/b-a}$ vs. all other genotypes).
§Observed at weaning, $n = 87$ ($P < 0.0001$); observed at E14.5, $n = 18$ ($P = 0.037$); observed at E1.5, $n = 41$ ($P = 0.0007$). P values calculated for $Sec23a^{-/-}\ Sec23b^{+/b-a}$ mice vs. all other genotypes.

when expressed under the endogenous regulatory elements of the *Sec23b* gene. Although mice homozygous for the latter genetically engineered *Sec23b* locus (expressing only SEC23A and no intact SEC23B protein) appear entirely normal and indistinguishable from WT controls, a subtle phenotypic difference or key contribution from the residual SEC23B N-terminal tail retained in the SEC23B-A protein (Fig. 4*A* and *SI Appendix*, Fig. S5*B*) cannot be excluded. Nevertheless, these data demonstrate complete (or near complete) overlap in function between the two SEC23 paralogs.

The *Sec23b^{b-a}* Allele Fails to Complement Disruption of *Sec23a* in *Sec23a^{−/−}* Mice. Mice with germline SEC23A deficiency die at midembryogenesis (29). To test the capacity of the *Sec23b^{b-a}* allele to rescue the lethality of mice homozygous for an inactivated endogenous *Sec23a* allele (*Sec23a^{−/−}* mice), *Sec23b^{+/b-a}* mice were crossed with *Sec23a^{+/−}* mice. Although the expected number of *Sec23b^{+/b-a} Sec23a^{+/−}* pups from a *Sec23b^{+/b-a} × Sec23a^{+/−}* cross were present at weaning (Table 1, *Sec23a^{+/−} × Sec23b^{+/b-a}*), no *Sec23b^{+/b-a} Sec23a^{−/−}* progeny were observed at weaning or at embryonic day (E) 13.5 (Table 1, *Sec23a^{+/−} Sec23b^{+/b-a} × Sec23a^{+/−}*). Thus, SEC23A expression under control of *Sec23b* transcriptional regulatory sequences (*Sec23b^{b-a}*) fails to complement loss of expression from the *Sec23a* locus.

The *SEC23B/SEC23A* Expression Programs Have Shifted During Evolution. Although the above data suggest functional equivalence for SEC23A and SEC23B in yeast, ZF, and mice, the disparate phenotypes of SEC23B deficiency in humans and mice remain unexplained. To address this question, the relative expression of *SEC23B*:*SEC23A* in WT tissues from both humans and mice was determined by qRT-PCR. This ratio was higher in human BM (9.7) compared with human pancreas (1.8), although higher in mouse pancreas (12.7) compared with mouse BM (2.6), consistent with the pathology of SEC23B deficiency in these two species. The differential expression of SEC23A/B in human and murine erythroid cells is consistent with previously published results (38, 39). In addition, a previous analysis of chromatin occupancy in erythroid cells demonstrated a state of transcriptional repression at the human

SEC23A locus relative to the mouse (40). SEC23B deficiency also results in a salivary gland defect in mice (Fig. 5*B*), although this has not been observed in humans. Consistent with this pattern, SEC23B is the predominantly expressed paralog in the murine salivary gland (32), while in humans the *SEC23B*:*SEC23A* mRNA expression ratio is ~0.27 (www.gtexportal.org/).

Taken together, these results suggest that the disparate human/mouse phenotypes are due to an evolutionary shift in the tissue-specific expression programs between mouse and human for these two functionally equivalent proteins.

Discussion

The genes encoding a number of components of the COPII machinery have been duplicated during vertebrate evolution, with evidence for unique protein function for the corresponding paralogous proteins. In the case of SEC23, the distinct and highly specific phenotypes of SEC23A and SEC23B deficiency in humans and mice, together with evidence for paralog-specific contributions to collagen (29) and EGFR (36) transport, as well as a potential role for SEC23B in oncogenesis (41), suggest unique functions for each of these paralogous proteins.

However, our data demonstrate indistinguishable interactomes for SEC23A and SEC23B in HEK293 cells and complementation of yeast Sec23 function by both human and murine SEC23A and SEC23B paralogs. We also show that transgenic germline *sec23a* expression rescues the mortality of *sec23b* disruption in ZF and that SEC23A rescues the mortality of SEC23B-deficient mice when expressed under the control of the endogenous *Sec23b* transcription machinery. These data strongly suggest complete overlap in function between the SEC23A and SEC23B proteins, despite their origin via an ancient gene duplication preceding the origin of vertebrates. The high degree of sequence conservation between SEC23A/B (~85% amino acid identity) is also consistent with our findings, suggesting strong selection at the level of protein function.

Gene duplications are frequent evolutionary events, with the duplicated copies most commonly accumulating loss-of-function mutations with subsequent disappearance from the genome (42–44).
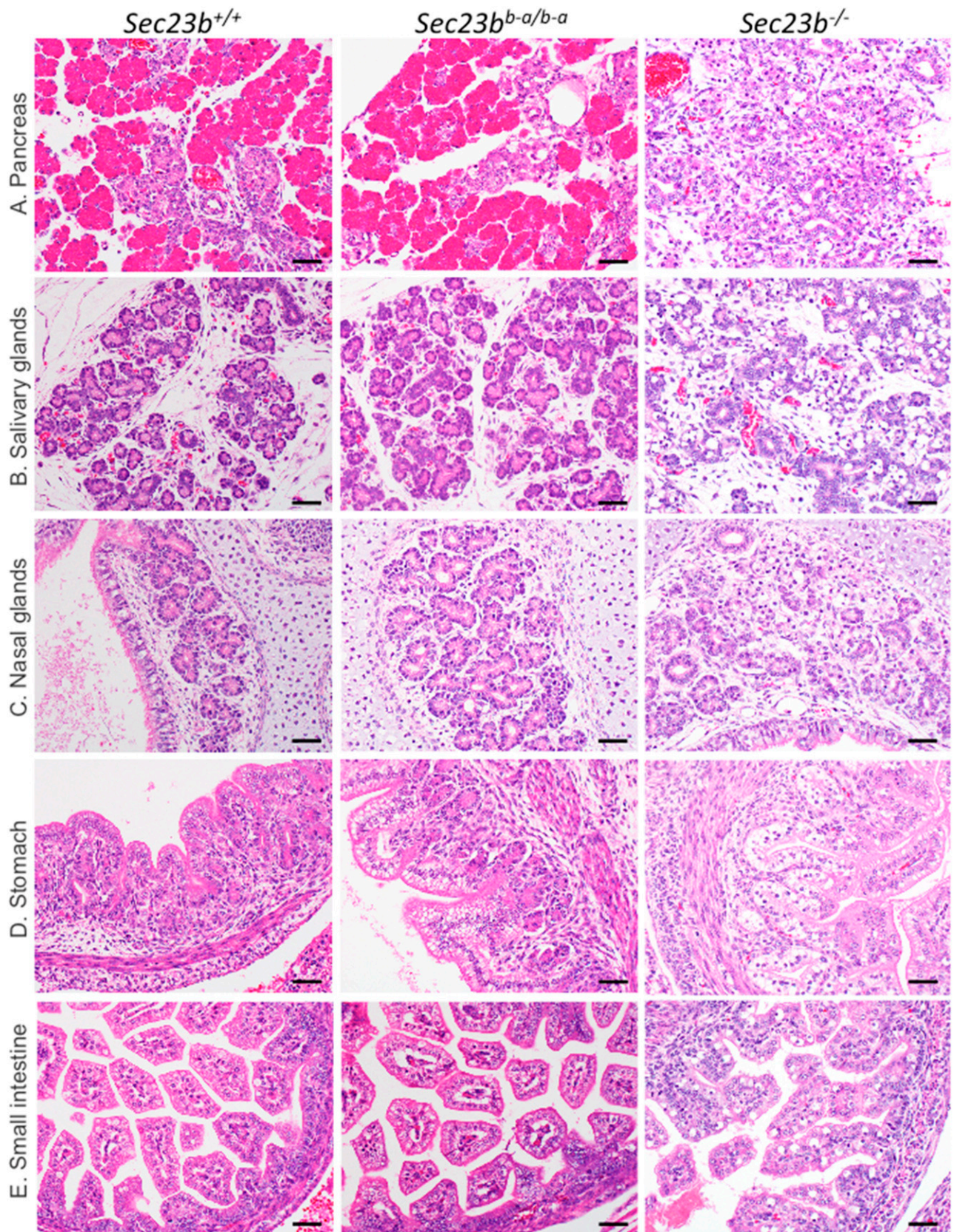
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Fig. 5. Mice expressing *Sec23a* from the *Sec23b* locus exhibit normal pancreas histology. (*A–E*) In contrast to *Sec23b*$^{-/-}$ mice, *Sec23b*$^{b-a/b-a}$ mice exhibit normal (*A*) pancreas, (*B*) salivary glands, (*C*) nasal glands, (*D*) stomach, and (*E*) small intestine histology, similar to WT (*Sec23b*$^{+/+}$) littermate controls. (Scale bars, 40 μm.)

Less frequently, paralogous gene copies become fixed either through neofunctionalization (acquisition of a new function by one of the paralogous gene copies), or subfunctionalization (the division of the functions of the ancestral gene between the paralogous copies). The latter process can occur either at the level of protein function or gene expression. A recent analysis suggests that subfunctionalization of expression evolves slowly and is rare among recently duplicated genes pairs, although is a more common mechanism among duplicated transcription factors (45). Consistent with this model, functional equivalence for a number of transcription factor paralogs has been demonstrated in mice by gene replacement (46–49), although overlap is often incomplete (46, 49).

Our evidence for functional equivalence between the SEC23A and SEC23B proteins suggests that the evolutionary subfunctionalization of these two genes occurred at the level of their gene-expression programs rather than protein function, with select cell types manifesting greater dependence on expression of one or the other paralog. Further shifts in the *SEC23A* and *SEC23B* expression programs during mammalian evolution could account for the disparate phenotypes of SEC23A and SEC23B deficiency in mice and humans. Consistent with this hypothesis, human SEC23B is the predominantly expressed paralog in the BM, consistent with the phenotype of CDAII, while SEC23A is the more strongly expressed paralog in calvarial osteoblasts (50), consistent with the CLSD phenotype. In contrast, SEC23B is the predominant paralog in the mouse pancreas, consistent with the pancreatic phenotype observed in murine SEC23B deficiency (30–32).

Similar subfunctionalization at the level of gene expression might also explain highly tissue-restricted phenotypes for other genetic diseases due to loss-of-function in paralogous genes encoding other COPII subunits or other core components of the cellular machinery. This paradigm might also explain other examples of highly discordant phenotypes observed between animal models and humans for various other diseases (51).

Finally, the apparent complete rescue of the SEC23B-deficient phenotype in mice by activating SEC23A expression in SEC23B-dependent tissues suggests that therapeutic strategies to increase the expression of SEC23A in human hematopoietic precursor cells could ameliorate the CDAII phenotype. Such approaches could include delivery of modified CRISPR/Cas9 or transcription factor constructs designed to specifically activate the SEC23A gene (52–55). This paradigm could also be applicable to a number of other human disorders due to mutations in functionally overlapping paralogous genes.

Methods

Evolutionary Sequence Analysis of *Sec23* Paralogs. The gene tree for SEC23A/SEC23B was examined on Ensembl (https://useast.ensembl.org/Homo_sapiens/Gene/Compara_Tree?collapse=12002157%2C12002179%2C12001773%2C120-01917%2C12001992%2C12002187%2C12001885%2C12001925%2C12002033-%2C12002044%2C12002005%2C12001805%2C12001978%2C12002051%2C1-2002189%2C12002084%2C12002151%2C12001996%2C12002037%2C12002020-91%2C12001765%2C12002146%2C12002175%2C12001947%2C12001985%-2C12001831;db=core;g=ENSG00000100934;r=14:39031919-39109646). This analysis estimates the *Sec23* duplication as having occurred ~615 Mya.

Coimmunoprecipitation. FLAG-tagged SEC23A and SEC23B, and RFP-tagged SEC24A, SEC24B, SEC24C, and SEC24D were cloned into the PED plasmid. Expression vectors were transfected into COS cells using FUGENE HD (Promega) per the manufacturer's instructions. FLAG coimmunoprecipitation with anti-FLAG M2 beads (Sigma) was performed as per the manufacturer's instructions.

Generation of Stable Cell Lines. The coding sequences of human *SEC23A* and *SEC23B* were cloned into the pENTR vector (K2400-20; Invitrogen) per the manufacturer's instructions and then into the pcDNA5-pDEST-BirA-FLAG-C-term vector by Gateway cloning. Bait proteins (SEC23A or SEC23B) were stably expressed in Flp-In T-REx HEK293 cells as previously described (56). Stable pools of integrated cells were selected with 200 μg/mL hygromycin. For BioID experiments, cells were treated with tetracycline (1 μg/mL) to induce gene expression and with biotin (50 μM) for 24 h and were subsequently harvested for analysis, as previously described (57).

Affinity Purification and BioID Mass Spectrometry. Cell lysates were prepared and biotinylated proteins were captured on streptavidin beads and subjected to trypsin digestion, as previously described (57). Tryptic peptides were suspended in 5% formic acid and stored at −80 °C until analysis by MS. Tryptic peptides were analyzed on a TripleTOF 5600 (Sciex) instrument, as previously described (56). Two biological replicates were analyzed for each sample and for controls, and data analysis was performed as follows. Raw MS files were converted to mzXML and searched using the Mascot and Comet search engines. A protein database consisting of the human and adenovirus complements of RefSeq v57 supplemented by common contaminants and epitope-tagged sequences was appended with an equal number of decoy sequences. For peptide identification, a 30-ppm precursor ion and 0.15-Da fragment ion tolerance were specified, and Asn and Gln deamidation and Met oxidation were allowed as variable modifications for searches with Comet and Mascot, essentially as described. Search results were further processed using PeptideProphet to generate a list of peptides, which were then assembled to protein lists using ProteinProphet. The resulting protein lists were then processed using Prohits (default parameters) to generate a spectral count matrix. The spectral count matrix was formatted according to CRAPome/REPRINT specifications (manually) and uploaded onto reprint-apms.org. Protein interactions were scored using two enrichment scoring approaches, empirical fold-change score (FC) and SAINT (SP). A list of high-confidence interactions was generated by applying the following filters: FC ≥ 10; SP ≥ 0.96; abundance ≥ 2. Finally, high-scoring interactions were visualized as a network (using the network generation utility of REPRINT). Raw abundances of filtered interactions in each replicate were plotted against others to evaluate correlation of prey abundances.

The MS dataset has been assigned the MassIVE ID MSV000081703 and is available for FTP download at ftp://MSV000081703@massive.ucsd.edu. The dataset was assigned the ProteomeXchange Consortium (proteomecentral.proteomexchange.org/cgi/GetDataset) identifier PXD008180.

Sec23 Complementation in Yeast. The coding sequences of the mouse and human *SEC23* paralogs, (as well as control yeast *sec23*) were cloned into the yeast expression vector pRS416, with a 3′ sequence encoding an HA tag. The sequence integrity of all constructs was confirmed by Sanger sequencing. Vectors with the following promoters were used: the *ZEO1* constitutive promoter and the *CUP1* copper-inducible promoter, resulting in a range of expressions of the SEC23 proteins. The latter vectors were transduced into a temperature-sensitive *sec23-1* yeast mutant in the BY4741 background (58), which expresses the functional endogenous yeast Sec23 protein at the permissive temperature (24 °C) but not at the restrictive temperature (37 °C).

Generation of Sec23b-Deficient ZF Using CRISPR/Cas9. The *sec23b* locus was identified in the ZF genomic sequence assembly (59). A guide RNA (gRNA) targeting the ZF *sec23b* exon 5 sequence (ggtggacacatgtctggagg) was cloned into the pDR274 vector, as previously described (60). gRNA was then in vitro transcribed using the T7 Quick High Yield RNA Synthesis kit (New England Biolabs) and purified using the RNA Clean and Concentrator kit (Zymo Research). gRNA was coinjected with Cas9 RNA, generated as previously described (60) into single-cell ZF embryos. ZF were genotyped for indels with primers ZF *sec23b* CRISPR F and ZF *sec23b* CRISPR R (*SI Appendix*, Table S1) spanning the expected CRISPR cut site. ZF heterozygous for a 53-bp deletion in exon 5 of *sec23b* (*sec23b⁺/⁻*) were generated. *sec23b⁺/⁻* ZF were intercrossed to generate Sec23b-deficient ZF (*sec23b⁻/⁻*). Blood collected from WT, *sec23b⁺/⁻*, and *sec23b⁻/⁻* ZF was stained with the HEMA 3 kit (Fisher) and evaluated under light microscopy by two independent investigators blinded to the ZF genotype.

Generation of Transgenic ZF Expressing *sec23a* Under the Control of a Ubiquitin Promoter. The ZF *sec23a* cDNA was cloned downstream of the ubiquitin promoter into the JS218 vector (61) (SEC23A-GFP). The correct *sec23a* sequence was confirmed by Sanger sequencing. A p2A sequence separates the *sec23a* cDNA and the GFP coding sequence, resulting in ubiquitous *sec23a* and *GFP* expression from the same mRNA molecule (*SI Appendix*, Fig. S4C).

SEC23A-GFP or JS218 (in which GFP alone is expressed ubiquitously) were coinjected with transposase mRNA, as previously described (61), into one-cell–stage embryos generated from *sec23b⁺/⁻* ZF intercrosses to generate transgenic ZF ubiquitously expressing SEC23A and GFP or GFP alone. The latter ZF were genotyped with primers ZF *sec23b* CRISPR F and ZF *sec23b* CRISPR R (listed in *SI Appendix*, Table S1).

Cloning the *Sec23b-a* dRMCE Cassette into pUC19. The *Sec23b-a* cassette was generated by assembling the following sequences from the 5′ to 3′ direction: an FRT sequence, the endogenous *Sec23b* intron 4 splice acceptor sequence, *Sec23a* from C[367] to A[2298] of the cDNA coding sequence (encoding murine SEC23A starting at Arg123), and the SV40 polyA sequence present in the *Sec23b*[cgt] allele (30, 62). The above *Sec23b-a* cassette was inserted into pUC19 at the HindIII and SalI restriction sites, producing pUC19-Sec23b-a(1) (*SI Appendix*, Fig. S5A) and a puromycin resistance cassette flanked by *rox* sites was inserted 3′ of the SV40 polyA sequence using PmeI and AscI restriction sites added to the vector by PCR, resulting in pUC19-Sec23b-a(2) (Fig. 4A), with sequence integrity confirmed by Sanger sequencing.

Plasmid Purification and Microinjections. The pDIRE plasmid driving expression of both FLPo and iCRE was obtained from Addgene (#26745). The pCAGGS-FLPo and pCAGGS-iCre plasmids, which contain the CAG promoter/enhancer driving the expression of FLPo and iCre recombinase, respectively, were generated as previously described (63). Plasmids were purified using the Qiagen EndoFree Plasmid Maxi Kit (#12362; Qiagen) or the Machery-Nagel NucleoBond Xtra Maxi EF kit (#740424.10; Machery-Nagel), per the manufacturers' instructions.

Zygotes generated from the in vitro fertilization of C57BL/6J oocytes with sperm from *Sec23b*^{+/−} male mice (33), were coinjected with pUC19-Sec23b-a(1) and pDIRE plasmids and then implanted in pseudopregnant foster mothers. Pups were genotyped at 2 wk of age. All microinjections were performed at the University of Michigan Transgenic Animal Model Core.

Transient Electroporation of ES Cells. The ES cell clone EPD0237_3_G06 heterozygous for the *Sec23b*[cgt] allele (33) was obtained from the European Conditional Mouse Mutagenesis Program (62) and coelectroporated with pUC19-Sec23b-a(2) and either pDIRE or a combination of pCAGGS-FLPo and pCAGGS-iCre, as previously described (63). One week postelectroporation, individual ES cells were plated in 96-well plates. Cells were expanded and plated in triplicates for DNA analysis, frozen stocks, and puromycin selection. Following puromycin selection, 288 single cell ES clones were evaluated demonstrating correct targeted insertion of the dRMCE replacement vector in 13 of 288 independent clones.

Generation of *Sec23b*[+/b-a] Mice. Three independent EPD0237_3_G06 ES cell clones correctly targeted with pUC19-Sec23b-a(2) were cultured, expanded, and then microinjected into albino C57BL/6 blastocysts to generate *Sec23b*[+/b-a] mice (Fig. 4A). The chimeric progeny were crossed to B6(Cg)-Tyr[c-2J]/J mice (Jackson Laboratory stock #000058) and germline transmission was confirmed by genotyping the progeny for the *Sec23b*[+/b-a] allele. The *Sec23b*[+/b-a] allele was maintained on a C57BL/6J background.

Mouse Genotyping. Genotyping of the *Sec23b*[b-a] allele was done on genomic DNA extracted from tail biopsies using primers RMCE F2 and RMCE R2, which detect the dRMCE replacement construct. These primers are located in *Sec23a* exons 10 and 14, respectively, and therefore should not yield a detectable product from the WT *Sec23a* allele. To identify potential random insertions of the dRMCE allele, PCR genotyping was also performed with primers RMCE F1 and RMCE R1 to confirm correct insertion of the Sec23b-a(1) and Sec23b-a(2) alleles at the 5′ end, and with primers RMCE F3 and RMCE R3 to confirm correct placement of the Sec23b-a(2) allele at the 3′ end. All mice with a visible PCR product with primers RMCE F2 and RMCE R2 also had PCR products with primers RMCE F1-RMCE R1 and primers RMCE F3-RMCE R3, effectively excluding random insertion.

Genotyping for the *Sec23b*[b-a] allele was also performed by RT-PCR on RNA isolated from mouse tail biopsies, using a three-primer competitive PCR assay, with a forward primer (RMCE A) located in exon 2 of *Sec23b* and two reverse primers, one located in the *Sec23a* cDNA (RMCE B) and one located in exon 5 of *Sec23b* (RMCE C). This competitive PCR assay should generate a 282-bp band from the WT allele and a 311-bp band from the *Sec23b*[b-a] allele (Fig. 4C), which were resolved on a 2% (wt/vol) agarose gel. The location of the primers is indicated in Fig. 4A and *SI Appendix*, Fig. S5A and the primer sequences are listed in *SI Appendix*, Table S1.

RT-PCR. RNA was isolated from tissues using the RNeasy Mini kit (Qiagen), and reverse transcription was performed using the SuperScript first strand synthesis system for RT-PCR (Invitrogen) with random primers.

Western Blot and Antibodies. Total cell lysates were prepared and Western blots were performed as previously described (33). Rabbit anti-SEC23A and anti-SEC23B antibodies were generated as previously described (33), and mouse anti-GAPDH and anti–RAL-A antibodies were obtained from Millipore and BD Biosciences, respectively. Anti-FLAG, anti-RFP, and anti-HA antibodies were obtained from Cell Signaling Technology, Abcam, and Sigma, respectively.

Complete Blood Counts. Peripheral blood was collected from the retro-orbital venous sinuses of anesthetized mice as previously described (30), and complete blood counts were performed using the Advia120 whole-blood analyzer (Siemens), as previously described (33).

Animal Care. Mice and zebrafish were housed at the University of Michigan and all animal care and procedures were in accordance with the Principles of Laboratory and Animal Care, established by the National Society of Medical Research. Mice had free access to water and food and were kept in cages in a 12-h light/dark cycle. Zebrafish were maintained in a recirculating system in a 14-h light/10-h dark cycle. All animal protocols used in this study were approved by the University of Michigan Committee on the Use and Care of Animals.

Histology, Cytology, *o*-dianisidine Staining, and Electron Microscopy. For histological analysis, mouse tissues or ZF embryos were harvested and immediately fixed in 4% paraformaldehyde or aqueous buffered zinc formalin (Z-fix, Anatech). Tissues were then routinely processed, paraffin-embedded, sectioned, and stained with H&E. BM smears were prepared using a brush technique, air dried, and stained with a modified Romanowsky staining (Diff-Quik) technique. ZF embryos were stained with *o*-dianisidine, as previously described (64). For electron microscopy, mouse pancreas tissues were prepared as previously described (30, 31) and sections were examined on a Philips CM100 electron microscope. Images were recorded as previously described (30, 31).

Statistical Analysis. The statistical significance of deviation from the expected Mendelian ratio of mouse crosses was assessed using the χ^2 test. The statistical significance of the differences between parameters among different genotype groups was assessed using the Student's *t* test. The difference in overall survival between mice of distinct genotypes was assessed using the Kaplan–Meier method.

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