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Organic Solute Transporter α-β Protects Ileal Enterocytes From Bile Acid–Induced Injury

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SUMMARY

Enterocyte bile acid stasis in early postnatal life results in bile acid–induced cytotoxicity, oxidative stress, and a restoration response. Blocking enterocyte apical membrane bile acid uptake protects against ileal epithelial injury.

BACKGROUND & AIMS: Ileal bile acid absorption is mediated by uptake via the apical sodium-dependent bile acid transporter (ASBT), and export via the basolateral heteromeric organic solute transporter α-β (OSTα-OSTβ). In this study, we investigated the cytotoxic effects of enterocyte bile acid stasis in Ostα−/− mice, including the temporal relationship between intestinal injury and initiation of the enterohepatic circulation of bile acids.

METHODS: Ileal tissue morphometry, histology, markers of cell proliferation, gene, and protein expression were analyzed in male and female wild-type and Ostα−/− mice at postnatal days 5, 10, 15, 20, and 30. Ostα−/−/Asbt−/− mice were generated and analyzed. Bile acid activation of intestinal Nrf2-activated target genes in neonatal Ostα−/− mice. Expression of reduced nicotinamide adenine dinucleotide phosphate oxidase-1 and Nrf2-anti-oxidant responsive genes were increased significantly in neonatal Ostα−/− mice at these postnatal time points. Bile acids also activated Nrf2 in Drosophila enterocytes and enterocyte-specific knockdown of Nrf2 increased sensitivity of flies to bile acid–induced toxicity. Inactivation of the Asbt prevented the changes in ileal morphology and induction of anti-oxidant response genes in Ostα−/− mice.

CONCLUSIONS: Early in postnatal development, loss of Ostα leads to bile acid accumulation, oxidative stress, and a restitution response in ileum. In addition to its essential role in maintaining bile acid homeostasis, Ostα-Ostβ functions to protect the ileal epithelium against bile acid–induced injury. NCBI Gene Expression Omnibus: GSE99579. (Cell Mol Gastroenterol Hepatol 2018;5:499–522; https://doi.org/10.1016/j.jcmgh.2018.01.006)

Keywords: Ileum; Reactive Oxygen Species; Nuclear Factor Erythroid-Derived 2-Like 2; Neonate; Drosophila.

In the hepatocyte, bile acids are synthesized from cholesterol, conjugated to taurine or glycine, and then secreted into bile for storage in the gallbladder or passage into the small intestine. During a meal, bile acids are present
in high concentration in the small intestine where they act as detergents to facilitate absorption of fats, fat-soluble vitamins, and cholesterol. After reaching the distal small intestine, bile acids are almost quantitatively reabsorbed by ileal enterocytes, and carried back to the liver in the portal circulation for uptake and resecretion into bile. This enterohepatic circulation functions to safely store and then promptly deliver bile acids to the intestinal lumen, while limiting the systemic distribution and potential cytotoxicity of these amphipathic molecules. The major membrane carriers that maintain the enterohepatic circulation of bile acids are known and include specific transporters expressed on the sinusoidal and canalicular membranes of the hepatocyte, and on the apical brush border and basolateral membranes of the ileal enterocyte. In ileum, the apical sodium-dependent bile acid transporter (ASBT) (gene symbol: \( SLC10A2 \)) and the heteromeric organic solute transporter \( \alpha-\beta \) (OST\( \alpha \)-OST\( \beta \)) (gene symbols: \( SLC51A-SLC51B \)) mediate these steps, respectively. ASBT mutations in human beings or inactivation of the Asbt in mice yields a primary bile acid malabsorption phenotype that is characterized by impaired intestinal bile absorption, induction of hepatic bile acid synthesis, and increased fecal loss of bile acids in the absence of ileal histologic or ultrastructural changes. Inactivation of Ost\( \alpha \) in mice also impairs intestinal bile acid absorption. However, unlike Asbt\(-/-\) mice or patients with ASBT mutations, Ost\( \alpha \)/- mice show a complex phenotype that includes a paradoxic reduction in hepatic bile acid synthesis and ileal hypertrophy. The changes in bile acid metabolism are associated with altered gut-liver bile acid signaling through the farnesoid X receptor (FXR)-fibroblast growth factor (FGF) pathway, and inactivation of FXR in Ost\( \alpha \)-/- mice reverses the reductions in hepatic Cyp7a1 expression and bile acid synthesis. By contrast, the altered ileal morphology in Ost\( \alpha \)-/- mice is not affected by inactivation of FXR. The ileal changes observed in Ost\( \alpha \)-/- mice, which include villous blunting, are typically associated with epithelial damage and subsequent healing. Although adult Ost\( \alpha \)-/- mice do not show overt symptoms of intestinal injury, such as increased inflammatory gene expression in ileum, bleeding, or diarrhea, newborn Ost\( \alpha \)-/- mice show a small postnatal growth deficiency, and this may coincide with the onset of injury or initiation of an adaptive response.

The cause of the altered ileal morphology in Ost\( \alpha \)-/- mice is unclear, but may involve bile acids. Conjugated bile acids are fully ionized at physiological pH and are largely membrane-impermeable in the absence of transporters. As such, conjugated bile acid accumulation and intracellular toxicity occurs when membrane transporters facilitate their uptake and their cellular export is inhibited. Bile acid–induced damage is best described for forms of liver disease and has been studied for hepatocytes and the biliary epithelium. For example, in progressive familial intrahepatic cholestasis type 2, inherited mutations in the bile salt export pump (gene symbol: \( ABCB11 \)) blocks canalicular bile acid export, leading to the accumulation of bile acids and subsequent hepatocellular damage. The molecular mechanisms implicated in bile acid–induced hepatocyte injury include induction of endoplasmic reticulum stress and mitochondrial damage. However, more recent studies have suggested that intracellular conjugated bile acids indirectly stimulate hepatocyte chemokine expression and induce liver injury via a hepatocyte-initiated inflammatory response. Beyond the liver and biliary tract, there is a substantial flux of bile acids across the ileal epithelium. Surprisingly little is known regarding how the ileal enterocyte is protected against bile acid cytotoxicity. In Ost\( \alpha \)-/- mice, we hypothesize that continued ileal enterocyte Asbt-mediated bile acid uptake in the absence of a mechanism for efficient export will increase intracellular bile acid levels. This in turn can promote bile acid–induced injury and drive the apparent epithelial damage observed. The ontogeny of bile acid synthesis and transport has been carefully described in rats and mice, with an abrupt induction of ileal Asbt expression coinciding with concentrative bile acid uptake between postnatal days 17 and 21. If bile acids are important for the morphologic changes observed in Ost\( \alpha \)-/- mice, then Asbt-mediated bile acid uptake into the ileal enterocytes should precede or coincide with the intestinal injury. To test this hypothesis, we investigated the temporal relationship in Ost\( \alpha \)-/- mice between the intestinal adaptive response and initiation of active ileal bile acid absorption, and whether inactivation of the Asbt is protective in Ost\( \alpha \)-/- mice. The potential mechanisms underlying the postulated bile acid–induced injury also were explored with a reductionist approach using the Drosophila model organism.

**Materials and Methods**

**Animals, Treatments, and Tissue Collection**

The Institutional Animal Care and Use Committees at the Wake Forest School of Medicine and Emory University approved these experiments. The Ost\( \alpha \)-/- and Asbt\(-/-\) mice were generated as previously described. The mice were backcrossed onto a C57BL/6J background for 8 generations and compared with wild-type (WT) littermates on the same background. The Ost\( \alpha \)/Asbt\(-/-\) mice were generated by cross-breeding the corresponding null mice and compared with lines generated from WT, Ost\( \alpha \)-/-, and Asbt\(-/-\) littermates as controls. The mice were group-housed in ventilated cages

Abbreviations used in this paper: ARE, anti-oxidant response element; Asbt, apical sodium-dependent bile acid transporter; CDCA, chenodeoxycholic acid; cRNA, complementary RNA; FGF, fibroblast growth factor; FXR, farnesoid X receptor; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; GFP, green fluorescence protein; GSH, reduced glutathione; GSSG, oxidized glutathione; Ibabp, ileal bile acid binding protein; mRNA, messenger RNA; NEC, necrotizing enterocolitis; Nox, reduced nicotinamide adenine dinucleotide phosphate oxidases; Nrf2, nuclear factor (erythroid-derived 2)-like 2; Ost, organic solute transporter; PBS, phosphate-buffered saline; ROS, reactive oxygen species; TNF, tumor necrosis factor; TUNEL, terminal deoxyuridine triphosphate nick-end labeling; WT, wild type.

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(Super Mouse 750 Microisolator System; Lab Products, Seaford, DE) containing bedding (1/8" Bed-O-Cobbs; Andersons Lab Bedding Products, Maumee, OH) in the same temperature- (22°C) and light/dark cycle- (12-h; 6 AM to 6 PM) controlled room of the animal facility to minimize environmental differences. The breeding mice were maintained in cages with standard bedding (1/8" Bed-O-Cobbs) and pulp cotton fiber nesting material (Nestlets; Ancare, Bellmore, NY), fed ad libitum rodent breeder Chow (21% of calories as fat; PicoLab Diet 20 no. 5058; catalog no. 0007689, LabDiet, St. Louis, MO), and offspring were weaned at postnatal day 20. The higher fat content breeder chow was fed to the dams because previous studies noted small postnatal growth deficits in Osta−/− and Asbt−/− mice.29

Adult nonbreeding mice were maintained on rodent Chow (13% of calories as fat; PicoLab Diet 20 no. 5053, catalog no. 0007686, LabDiet). For the ontogeny study, WT and Osta−/− breeders were closely monitored for pregnancy status. Offspring were collected within 4 hours of the end of the dark phase on postnatal days 5, 10, 15, 20, and 30. The sex of the pups at day 5 was confirmed by genotyping for the presence of the Y chromosome as described.30 Body weight, small intestinal length, and segment wet weight were recorded at necropsy from 2 to 4 litters (n ≥ 7 mice) per genotype and time point. For the ontogeny study, the small intestine was divided into 3 equal-length segments for histology and analysis of gene expression or 5 equal-length segments for bile acid analysis; segments were flushed with ice-cold phosphate-buffered saline (PBS) to remove luminal contents before weighing and analysis. For histologic analysis, the small intestinal segments were fixed for paraffin embedding. For measurements of bile acid content or gene and protein expression, the tissues were flash-frozen in liquid nitrogen and stored at -80°C before analysis. For analysis of the Asbt-Osta double-null mice, male Osta−/−Asbt−/− and corresponding control mice were analyzed at 10 days and at 8–10 weeks of age. The adult mice were fasted for approximately 4 hours at the end of the dark phase before being euthanized to isolate tissues. For the adult Osta−/−Asbt−/− and control mice, the small intestine was subdivided into 5 equal-length segments and flushed with ice-cold PBS to remove luminal contents before being fixed for histologic analysis or flash-frozen in liquid nitrogen and stored at -80°C for measurements of gene expression.8,10

**Histologic and Immunohistochemical Analysis**

The intestinal segments were flushed with PBS to remove luminal contents, fixed overnight in 10% neutral formalin (Sigma-Aldrich), and stored in 60% ethanol until processed for histology. Each segment was cut into longitudinal strips, stacked, and encased in 2% agarose before being embedded in paraffin and processed by Children's Healthcare of Atlanta Pathology Services. Histologic sections (5 μm) were cut and stained with H&E or used for immunohistochemical or immunofluorescence analysis. Microscopy was performed at the Emory University Integrated Cellular Imaging Core. The average villus heights of the proximal and midintestinal segments were measured for Osta−/− and WT mice at postnatal days 5, 10, 15, 20, and 30. At least 20 well-oriented, full-length, villus units per segment per mouse were measured; quantitative analysis of the digitally acquired images was performed using ImageJ software (National Institutes of Health, Bethesda, MD).30 Because of the aberrant morphology of the Osta−/− mice, quantitative measurements of crypt height were not performed for distal (ileal) segment at postnatal days 5, 10, 15, 20, and 30.

For detection of phosphohistone H3, a marker of cell proliferation, paraffin-embedded ileal sections underwent a heat-induced epitope retrieval procedure and were stained using a rabbit anti-phosphohistone H3 (Ser10) primary antibody (catalog no. 9701S, lot 13; Cell Signaling Technology, Danvers, MA) and Alexa Fluor 488-conjugated goat anti-rabbit antibody (catalog no. A-11034, lot 1670152; Thermo Fisher Scientific, Waltham, MA). For detection of apoptosis, terminal deoxynucleotidyl transferase-mediated fluorescein-deoxyuridine triphosphate nick-end labeling was performed using an in situ Cell Death Detection kit (catalog no. 11684795910, lot 10711900; Sigma-Aldrich) as described by the manufacturer. At least 3 mice were included for each experimental group or condition, and 6–8 field views at a magnification of 20× were taken for each ileal segment and analyzed for both phosphohistone H3 and terminal deoxynucleotidyl transferase-mediated deoxyuridine triphosphate nick-end labeling (TUNEL)-positive cells. Because individual crypt-villus structures were difficult to discern for Osta−/− mice owing to the altered morphology, comparisons were made using the number of phosphohistone H3 or TUNEL-positive cells counted per unit of intestinal length. For detection of mouse ileal bile acid binding protein (Ibabp) (Fabp6), paraffin-embedded ileal sections underwent a heat-induced epitope retrieval procedure and were stained using rabbit anti-fatty acid binding protein 6 primary antibody (catalog no. ab91184, lot GR245572-3; Abcam, Cambridge, MA) and Alexa Fluor 488-conjugated goat anti-rabbit antibody (catalog no. A-11034, lot 1670152; Thermo Fisher Scientific).

**Bile Acid and Glutathione Measurements**

Feces were collected from single-housed adult mice over a 72-hour period to measure the total bile acid content by enzymatic assay.7 To measure ileal tissue-associated bile acids, the small intestine was divided into 5 equal segments, flushed with PBS, flash-frozen, and ground under liquid nitrogen using a mortar and pestle. After addition of [14C]cholic acid (PerkinElmer Life Sciences, Boston, MA) to monitor bile acid recovery, aliquots of ileal tissue were extracted in ethanol and the bile acids were quantified by enzymatic assay as described.31 To measure ileal tissue-associated reduced and oxidized glutathione levels, the small intestine was divided into 5 equal segments, flushed with PBS, and ileum (segment 5) was used to isolate mucosal cells that were frozen and stored at -80°C. After thawing, the samples were extracted using perchloric acid and chemically modified for analysis of dansyl derivatives by high-performance liquid chromatography using a fluorescence detector (Children's Healthcare of Atlanta and Emory University’s Pediatric Biomarkers Core, Atlanta, GA).8,10
Microarray and Measurements of Messenger RNA Expression

Total RNA was extracted from frozen tissue using TRIzol reagent (Invitrogen, Carlsbad, CA). For microarray analyses, ileal total RNA samples were purified further using an RNeasy MinElute Cleanup Kit (Qiagen, Hilden, Germany), followed by quality assessment using an Agilent 2100 bioanalyzer. Samples with RNA integrity number values > 8.0 were carried forward for complementary RNA (cRNA) synthesis and hybridization to GeneAtlas MG-430 PM Array Strips (Affymetrix, Santa Clara, CA) following the manufacturer’s recommended protocol. Briefly, approximately 250 ng of purified total RNA was reverse-transcribed and biotin-labeled to produce biotinylated cRNA targets according to the standard Affymetrix GeneAtlas 3’-IVT Express labeling protocol (GeneAtlas 3’ IVT Expression Kit User Manual, P/N 702833 revision 4; Affymetrix). After fragmentation, 6 μg of biotinylated cRNA was hybridized for 16 hours at 45°C with the Affymetrix GeneAtlas Mouse MG-430 PM Array Strips. Strips were washed and stained using the GeneAtlas Fluidics Station according to standard Affymetrix operating procedures (GeneAtlas System User’s Guide; P/N 08-0306 revision A January 2010; Affymetrix) and subsequently scanned using the GeneAtlas Imager system. Fluidics control, scan control, and data collection were performed using the GeneAtlas Instrument Control Software version 1.0.5.267 (Affymetrix). The raw data generated were normalized using the robust multi-array average method.34 Ontology and pathway analysis was performed using the Database for Annotation, Visualization and Integrated Discovery (DAVID).35 All microarray analyses were performed by the Wake Forest School of Medicine Cancer Genomics Shared Resource Core (Winston-Salem, NC). Quantitative real-time polymerase chain reaction analysis was performed and messenger RNA (mRNA) expression levels were calculated based on the delta-delta threshold cycle method; values are means of triplicate determinations and expression was normalized using cyclophilin.34 The primer sequences are provided in Table 1.

Measurements of Protein Expression

Small intestinal extracts were prepared, subjected to sodium dodecyl sulfate–polyacrylamide gel electrophoresis using a 4%–20% gradient (Tris-Glycine Midi Gel; Invitrogen), and analyzed by immunoblotting.36 Blots were stripped before re-probing with antibodies to the different bile acid transport proteins or re-probing with antibody to glyceraldehyde-3-phosphate dehydrogenase (GAPDH) to normalize for protein load. Protein expression was quantified by densitometry using a Microtek (Hsinchu, Taiwan) ScanMaker i900 and FujiFilm (Tokyo, Japan) Multiguage 3 software, and expression data were normalized to levels of the GAPDH loading control. Sources of the antibodies used in the study were as follows: anti-mouse Asbt,8 anti-mouse Osta and anti-mouse Ostβ,8 anti-mouse Ibabp (anti-fatty acid binding protein 6, catalog no. ab91184, lot GR245572-3; Abcam), anti-GAPDH (catalog no. MA5-15738; Thermo Fisher Scientific), horseradish-peroxidase–conjugated goat anti-rabbit antibody (catalog no. A9169, lot 015M48581; Sigma-Aldrich).

Table 1. List of Primer Sequences Used for Real-Time PCR Analysis

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Asbt</td>
<td>5’ TGGTCTCTTCCCTGCTGACTAGCTGCTT</td>
<td>5’ TTGCTCTGATTCCATGGTTCCAAAGCT</td>
</tr>
<tr>
<td>Cyclophilin</td>
<td>5’ TCTTCTCATACGACCCAGGTGATTCTT</td>
<td>5’ CCCATCAAGATGTTGCACTGCTCAGGAG</td>
</tr>
<tr>
<td>Cyp7a1</td>
<td>5’ AGCACTAAACCCGGTCTAATATGCT</td>
<td>5’ GTCCTGATATGCAAGCATCTTACAGGAG</td>
</tr>
<tr>
<td>Cyp8B1</td>
<td>5’ GCTTCAGTGTAGCTCCGGTCTT</td>
<td>5’ GATCTCTTGCGCCACGTGTCTGAGAG</td>
</tr>
<tr>
<td>Diet1</td>
<td>5’ CACTCCAGTGAATGTTGCTCGAAT</td>
<td>5’ CGAAGTTGCAGGTTAAGAAGGAGC</td>
</tr>
<tr>
<td>FGF15</td>
<td>5’ GAGGACCCAGAACGGAGGATTTGTCGT</td>
<td>5’ ACGTCCTGATGACATCTTACAGGAG</td>
</tr>
<tr>
<td>GSTa1</td>
<td>5’ GGCAGAATGGAGGTGATGCTGA</td>
<td>5’ TCCAAATTCGGGACTGCTGCTGAGAG</td>
</tr>
<tr>
<td>GSTa3</td>
<td>5’ AGGAAAGACGCTTTTAAACAGGAGAA</td>
<td>5’ CCATCAAGGTGATTGCACTGCTGAGAG</td>
</tr>
<tr>
<td>GSTa4</td>
<td>5’ CCCCTGATCATGCGCTGCTTGCTT</td>
<td>5’ GAATGGTGCTGATCCCTTCTGCTGAC</td>
</tr>
<tr>
<td>GSTu1m</td>
<td>5’ GCAGCTCATCGTCGTCTGCTGCTGCT</td>
<td>5’ CATTTCCTAGGATGTTGCTGCTGCTG</td>
</tr>
<tr>
<td>GSTu3</td>
<td>5’ CGCCGATCAGCTGCTGCTGCTGCTGCT</td>
<td>5’ TGGCCAGAATCTGACTGCTGAGAGG</td>
</tr>
<tr>
<td>Ibabp</td>
<td>5’ CAAGGCTCGCTGGAATGGATGGGAGAG</td>
<td>5’ CCCGACAGCGCTTACAAGGAGG</td>
</tr>
<tr>
<td>Ibt1</td>
<td>5’ TGAATGAAAAGGACCGAACCCCTGCTT</td>
<td>5’ TCTTCTTGTTGGATTCTGCTGCTGAG</td>
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<tr>
<td>Nox1</td>
<td>5’ CGTCCCGGAGACCTGCTGATCAAGGAG</td>
<td>5’ GGATGGACCCCACTCCGGCGCCACCAAC</td>
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<tr>
<td>Nqo1</td>
<td>5’ AGCTTCTGCTGCTGCTGCTGCTGCTGCT</td>
<td>5’ AGTACAAATCGGATCCTGCTGCTGCTG</td>
</tr>
<tr>
<td>Nrf2</td>
<td>5’ CATGAGAATGGAGGATGCTGCTGCTGCT</td>
<td>5’ CCTCAAGAATGCGATCAACCTGCTG</td>
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<tr>
<td>OSTa</td>
<td>5’ TACAGAAACCCCGCTTTGCGCTGCTG</td>
<td>5’ CGAGAATCCAGAGCCTGCTGCTGCT</td>
</tr>
<tr>
<td>OSTb</td>
<td>5’ GTTATTGTCGTCGAAATGGGAGAGAGAG</td>
<td>5’ TGGTTCTGGGCGCGATCTGCTGCTG</td>
</tr>
<tr>
<td>SHP</td>
<td>5’ CATCCGCTTACCCGATGCTGCTGCTGCTG</td>
<td>5’ AGCTTCTGCTGCTGCTGCTGCTGCTG</td>
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<tr>
<td>S10C3</td>
<td>5’ AATAGCGGACTGGCTGCTGCTGCTGCTG</td>
<td>5’ TTGGTTCTGGGCGGACCTGCTGCTG</td>
</tr>
<tr>
<td>TNFa</td>
<td>5’ TCTTCTGATTCCCTGCTGCTGCTGCTGCTG</td>
<td>5’ GGCTTGGGATCCAGAATCTGCTGCTG</td>
</tr>
</tbody>
</table>

GST, glutathione-S-transferase.
and horseradish-peroxidase–conjugated goat anti-mouse antibody (catalog no. sc-2005, lot K1915; Santa Cruz Biotechnology, Dallas, TX).

**Drosophila Bile Acid Resistance and Staining Assays**

The WT and indicated *Drosophila* lines were obtained from previously described sources and maintained on standard media at 25°C. Unless otherwise indicated, the WT strain w1118 was used for all assays. Whole-animal cytoprotection in *Drosophila* was measured in response to dietary bile acids. Groups of 30–50 adult WT *Drosophila* (age, 5 days) were starved for 3 hours and then transferred to a vial containing a 2 × 2 cm piece of Whatman filter paper (Sigma-Aldrich) soaked with a solution of 5% sucrose containing the indicated concentration of bile acid. Survivors were scored for up to 7 days or until 100% lethality. The percentage of surviving flies was recorded and compared by log-rank Martel–Cox test. For detection of nuclear factor (erythroid-derived 2)-like 2 (Nrf2)/anti-oxidant response element (ARE)-dependent green fluorescent protein (GFP) expression, 5-day adult GSTD1-gfp *Drosophila* were transferred to vials containing 5% sucrose plus 25 mmol/L chenodeoxycholate (CDCA) or to vials containing 5% sucrose plus 25 mmol/L N,N,N′-dimethyl-4,4′-bipyridinium dichloride (Paraquat) as a positive control to induce oxidative stress. After 30 minutes, the intestinal tract was removed, fixed in 4% paraformaldehyde, and fluorescence was detected by confocal microscopy. For detection of reactive oxygen species (ROS) generation, 5-day adult WT flies were transferred to vials containing 5% sucrose or 5% sucrose plus 25 mmol/L CDCA. After 12 hours, the intestinal tract was removed and incubated with 100 μmol/L ROSstar 550 hydrocyanine ROS-sensitive reagent (catalog no. PN926-20000; LI-COR, Lincoln, NE) in Schneider’s *Drosophila* media at room temperature. The intestine then was washed, fixed in 4% paraformaldehyde, and the fluorescence was visualized by confocal microscopy. Representative fluorescent images of 4 independent measurements are shown.

**Statistical Analyses**

Mean values ± SEM are shown unless otherwise indicated. The data were evaluated for statistically significant differences using the Mann–Whitney test, the 2-tailed Student *t* test, or by analysis of variance (Tukey–Kramer honestly significant difference) (GraphPad Prism; Mountain View, CA). Differences were considered statistically significant at *P* < .05.

**Access to the Data**

All authors had access to the study data and reviewed and approved the final manuscript.

**Results**

**Ontogeny of the Morphologic Changes in Small Intestine**

As previously reported,8–10 *O斯塔−/−* mice backcrossed onto the C57BL/6J background are viable, fertile, and similar to WT littermates in terms of survival and gross appearance. The body weights of male or female *O斯塔−/−* mice were similar to WT mice at days 5, 10, and 15, but reduced at days 20 and 30 (Figure 1A). Small intestinal length and weight were increased as early as day 5 in male and female *O斯塔−/−* mice compared with WT mice, and remained increased at days 10, 15, 20, and 30 (Figure 1B and C). The increase in mass per unit length was particularly evident for the distal small intestine (ileum) (Figure 1D). Analysis of the tissue morphology and histology showed little change in the proximal (duodenum) or midintestinal (jejunum) regions (data not shown). The villus heights were similar between WT and *O斯塔−/−* mice in the proximal and midintestinal segments at postnatal days 5–15, with evidence of potential compensatory increases in villus height in jejunum at days 20 and 30 (Figure 2). In contrast, the ileal morphology and histology were altered dramatically in *O斯塔−/−* mice (Figure 3). Compared with WT mice, the distal small intestine of *O斯塔−/−* mice showed evidence of mucosal injury with features of regeneration. Specifically, *O斯塔−/−* mice showed villous blunting that ranged from mild to severe, which was prominent at early ages of postnatal development (days 5, 10, and 15). The epithelium lining the villi showed regenerative changes including cellular pseudostratification and loss of nuclear polarity, particularly within the transit-amplifying zone and crypt neck (Figure 3B). In addition, the epithelium on the tips of villi in *O斯塔−/−* mice showed altered apical borders and mucin synthesis. Although the tips of villi of WT mice are lined by absorptive enterocytes with an intact brush border and occasional mucin-producing goblet cells, the villous tips of *O斯塔−/−* mice showed loss of the brush border and replacement by markedly increased numbers of mucus-secreting cells (Figure 3B). Consistent with increased rates of cell turnover, numerous apoptotic epithelial cells also were present near the villous tips as well as in the crypt regions. The lamina propria of *O斯塔−/−* mice also contained mildly increased numbers of lymphocytes and macrophages with occasional neutrophils, eosinophils, and plasma cells. To confirm the histologic impression of increased crypt mitotic rate and the number of apoptotic cells in the ileum of *O斯塔−/−* mice, cell proliferation and apoptosis were assessed using phosphohistone H3 and TUNEL staining, respectively. As early as postnatal day 5, there was an increase in the numbers of proliferating (phosphohistone H3-positive) cells in ileum of *O斯塔−/−* compared with WT mice (Figure 4), consistent with the increased number of mitotic bodies noted in the histopathology assessment. There was also a trend toward an increased number of TUNEL-positive nuclei in the ileal villus structures of *O斯塔−/−* vs WT mice (Figure 4). In agreement with the histologic absence of an active inflammatory response, the mRNA expression for proinflammatory genes tumor necrosis factor α (TNFα) and interleukin 1β was not increased consistently in the ileal tissue of *O斯塔−/−* mice (Figure 5). There was a trend toward higher expression of TNFα at early postnatal time points, with a significant increase at day 10 in males, but this was followed by significant reductions. In contrast to the markedly altered ileal tissue morphology, the liver weights and liver histology were similar in male and female WT and *O斯塔−/−* at postnatal days.
Figure 2. Intestinal morphology of proximal small intestine and jejunum in male and female WT and Ostα−/− mice. Quantitative morphometric analysis of small intestine of WT and Ostα−/− mice. The small intestine of mice of the indicated age and genotype was divided into 3 equal segments. H&E-stained sections from segments 1 and 2 were used to measure the villus height in 20 well-oriented, high-power fields per mouse. Data are expressed as the means (n = 3–5 mice per group). Mean values ± SEM are shown. Significant differences between genotypes for that age were as follows: *P < .05.

Expression of Bile Acid Transporters and FXR Target Genes in Distal Small Intestine

The intestinal morphologic changes in Ostα−/− mice are evident before the normal induction of ileal concentrative bile acid uptake, which occurs between postnatal days 17 and 21 in WT mice.26,41 To determine if this temporal relationship is altered in Ostα−/− mice, ileal bile acid transporter expression was examined in male (Figure 6) and female (Figure 7) WT and Ostα−/− mice. In WT male and female mice, the mRNA and protein for Ostα-Ostβ were expressed abundantly in distal small intestine at postnatal day 5, and increased 3- to 5-fold by day 20. In Ostα−/− mice, mRNA and protein for the Ostα subunit were undetectable (Figures 6A and 7A). For the Ostβ subunit (Figures 6B and 7B), the mRNA expression was increased in Ostα−/− mice, but levels of Ostβ protein were reduced in the absence of its partner protein Ostα, as previously reported.8,9,42 In WT mice, ileal Asbt mRNA and protein were almost undetectable at postnatal days 5 and 10, and their levels dramatically increased between postnatal days 15 and 20. By contrast, Asbt was expressed prematurely in Ostα−/− mice, with ileal mRNA levels increased 12- to 28-fold in days 5 and 10 male and female Ostα−/− as compared with WT mice (Figures 6C and 7C). This is associated with increased Asbt protein expression. After day 15, ileal Asbt mRNA and protein levels were lower in Ostα−/− vs WT mice, in agreement with previous findings that ileal Asbt expression is reduced in adult (age, >8 wk) Ostα−/− mice.10 An increase in Asbt expression in the absence of Ostα-Ostβ-mediated bile acid export is predicted to increase the intracellular burden of bile acids in ileal enterocytes and increase expression of target genes for the bile acid–activated nuclear receptor FXR. Consistent with an increased intracellular bile acid load, expression of the FXR target genes Ostβ (Figures 6B and 7B), the cytosolic bile acid binding protein Ibabp (Fabp6) (Figures 6D and 7D), Shp (Nr0b2) (Figures 6E and 7E), and the sodium-sulfate co-transporter (Slc13a1) (Figures 6F and 7F) were induced in male and female Ostα−/− as compared with WT mice at postnatal days 5–15. This was particularly evident for Ibabp, whose mRNA and protein levels were increased more than 100-fold at postnatal days 5 and 10. By immunofluorescence, Ibabp protein was readily detected in ileal enterocytes of Ostα−/−, but not WT mice at postnatal days 10 and 15 (Figure 6G). In agreement with the increased expression of FXR target genes, the ileal tissue bile acid content also was increased in 10-day-old Ostα−/− vs WT mice (Figure 6H).

Molecular Mechanisms

Cellular accumulation of bile acids is associated with injury in a variety of diseases or disorders. The cytotoxic effects of bile acids have been attributed to several molecular mechanisms, including detergent-associated membrane damage, disruption of mitochondrial membrane potential, enhanced generation of ROS, direct activation of cell death receptors such as CD95/Fas and TNF-related apoptosis-inducing ligand receptor 2, and induction of an inflammatory response.15,22,43,44 To determine if similar mechanisms may be engaged in Ostα−/− mice, microarray analysis of ileal gene expression was performed for adult male WT and Ostα−/− mice at approximately 56 days of age (GEO series accession number: GSE99579). This analysis identified 244 differentially expressed genes (regulated more than 2-fold,
$P < .05$); the list included 54 and 190 genes that were increased and decreased, respectively. The induced genes are shown in the heat map in Figure 8A. Ontogeny analysis performed for differentially expressed genes showed that inactivation of Ostα induced ileal gene expression for pathways related to oxidation-reduction, cell proliferation, and glutathione metabolism. In particular, there was increased expression of oxidant-responsive genes, with
Nrf2/ARE target genes among the most highly induced. The Nrf2 system responds to oxidative stress and is a major regulator of cytoprotective pathways. In addition to the glutathione-S-transferase genes, Nrf2/ARE target genes induced in Osta−/− mice included those involved in glutathione biosynthesis, such as glutamate-cysteine ligase catalytic subunit (increased 1.7-fold), glutamate-cysteine ligase modifier (increased 1.4-fold), and glutathione synthetase (increased 1.6-fold), and those involved in glutathione metabolism, such as glutathione reductase (increased 2.2-fold) and glutathione peroxidase 2 (increased 1.5-fold). To determine how these gene expression changes may relate to the ileal glutathione redox status, the levels of reduced glutathione (GSH) and oxidized glutathione (GSSG) were

Figure 3. (continued)
measured in ileal mucosa from adult Ostα-/- and WT mice (n = 4/group). GSH levels (means ± SEM, 2.09 ± 0.99 vs 6.34 ± 1.06 nmol/mg protein; P < .05) and the GSH-to-GSSG ratio (2.6 ± 1.2 vs 11.6 ± 3.0; P < .05), but not levels of GSSG (1.02 ± 0.27 vs 0.62 ± 0.1.0 nmol/mg protein; P = .22) were reduced significantly in the ileal mucosa of Ostα-/- vs WT mice, respectively.

To determine if there are similar gene expression changes in the neonatal mice, mRNA expression of the Nrf2 target genes Gsta1, Gsta4, and Nqo1 was measured in male and female WT and Ostα-/- mice (Figure 8B). Consistent with the microarray analysis, expression of these genes was induced significantly in Ostα-/- mice vs WT mice as early as postnatal day 5. Because the ROS-generating enzyme reduced nicotinamide adenine dinucleotide phosphate oxidase 1 (Nox1) is expressed by the ileal epithelium and potentially involved in the redox-signaling associated with epithelial repair,39,46-48 Nox1 mRNA expression also was

![Figure 4. Ileal proliferation and apoptosis.](image)

**Figure 4. Ileal proliferation and apoptosis.** Quantitation of the number of phosphohistone H3 and TUNEL-positive cells per unit area. Mice from 2 to 4 litters were included in the analysis for each genotype and postnatal age group (n = 5–8 per group). Mean values ± SEM are shown. Significant differences between genotypes for that age were as follows: *P < .05, **P < .01, and ***P < .001.

![Figure 5. Expression of proinflammatory genes in ileum of male and female WT and Ostα-/- mice.](image)

**Figure 5. Expression of proinflammatory genes in ileum of male and female WT and Ostα-/- mice.** RNA was isolated from the distal small intestine of individual mice (n = 4–5 per group) and used for real-time PCR analysis. The mRNA expression was normalized using cyclophilin. Mean values ± SEM are shown. Significant differences between genotypes for that age and sex were as follows: *P < .05, **P < 0.01.
Figure 6. Expression of bile acid transporters and FXR target genes in distal small intestine of male WT and Ostα⁻/⁻ mice. Expression of mRNA and protein for bile acid transport–related genes: (A) Ostα, (B) Ostβ, (C) Asbt, and (D) Ibabp. Expression of mRNA for FXR target genes: (E) Shp and (F) Slc13a1. For mRNA expression measurements, RNA was isolated from the distal small intestine of individual male mice (n = 4–5 per group) and used for real-time PCR analysis. The mRNA expression was normalized using cyclophilin. For protein expression, extracts were prepared from distal intestine from individual male mice. Equal amounts of protein from 3 to 5 mice per group were pooled and duplicate samples were subjected to immunoblotting analysis. Blots were stripped and re-probed using antibodies to GAPDH as a loading control. (G) Immunostaining of transverse sections of distal small intestine for Ibabp. Original magnification: 20×. Scale bar: 100 μm. (H) Ileal bile acid content of day 10 male WT and Ostα⁻/⁻ mice. Bile acids were extracted from terminal ileum (last 20% of the small intestine) of 10-day-old male WT or Ostα⁻/⁻ mice and quantified by enzymatic assay. The bile acid content is expressed as mass per whole ileum, mass per unit length of ileum, and mass per unit weight of ileum. For mRNA expression, bars indicate means ± SEM. Significant differences between genotypes for that age were as follows: *P < .05, **P < .01, and ***P < .001, ****P < .0001.
measured and shown to be increased significantly in neonatal Ostα−/− mice (Figure 8C).

**Nrf2 Protects Against Bile Acid Cytotoxicity**

The results suggest that bile acids are inducing oxidative injury that is countered by expression of Nrf2-regulated cytoprotective genes. The findings are in agreement with a prior in vitro study in Caco-2 cells that showed bile acids such as deoxycholic acid and lithocholic acid induced Nrf2-ARE target gene expression and Nrf2-knockdown cells were more susceptible to lithocholic acid-induced toxicity.49 To directly test if intestinal Nrf2 is protective against bile acid–induced cytotoxicity in vivo, the model organism Drosophila was used. Components of the Nrf2 signaling pathway are conserved and fully developed in Drosophila.
Figure 7. Ontogeny of distal intestinal expression of bile acid transporters and FXR target genes in female WT and Ostα−/− mice. Expression of mRNA and protein for bile acid transport–related genes: (A) Ostα, (B) Ostβ, (C) Asbt, and (D) Ibabp. Expression of mRNA for FXR target genes: (E) Shp and (F) Slc13a1. RNA was isolated from the distal small intestine of individual female mice (n = 4–5 per group) and used for real-time PCR analysis. The mRNA expression was normalized using cyclophilin. Mean values ± SEM are shown. Significant differences between genotypes for that age were as follows: *P < .05, **P < .01, ***P < .001, and ****P < .0001. For protein expression, extracts were prepared from distal intestine from individual mice. Equal amounts of protein from 3 to 5 mice per group were pooled and duplicate samples were subjected to immunoblotting analysis. Blots were re-probed using antibodies to GAPDH as a load control.
where Nrf2 functions in oxidative stress tolerance, and in ROS-induced cytoprotective mechanisms in the gut. Although invertebrates lack bile acids, we reasoned that the ability of the Nrf2 pathway to respond to bile acids as an environmental insult may be evolutionarily conserved in Drosophila. We first examined the sensitivity of adult Drosophila to bile acid feeding. The concentration of bile acids selected for these studies were based on levels measured in the intestinal lumen of human beings and experimental animal models, which are in the millimolar range. However, because the intestine in Drosophila is covered by a protective membrane (the peritrophic matrix) and likely lacks transporters for membrane-impermeable conjugated bile acids, the flies were fed unconjugated hydrophobic bile acids, which

Figure 8. Ileal expression of Nrf2 ARE target genes. (A) Microarray analysis of ileum in adult male WT and Ostα⁻/⁻ mice. Differentially expressed genes whose expression was induced (2-fold change; \( P < .05; n = 4 \)) in Ostα⁻/⁻ vs WT mice are shown as a heat map. The blue line indicates the Z score for each gene. (B) Expression of mRNA for Nrf2 target genes in male and female WT and Ostα⁻/⁻ mice. (C) Expression of Nox1 mRNA in ileum of male and female WT and Ostα⁻/⁻ mice. RNA was isolated from the distal small intestine of individual mice at the indicated ages (n = 4–5 per group) and used for real-time PCR analysis. The mRNA expression was normalized using cyclophilin. Mean values ± SEM are shown. Significant differences between genotypes for that age and sex were as follows: * \( P < .05 \), ** \( P < .01 \), *** \( P < .001 \), and **** \( P < .0001 \).
Figure 8. (continued)
would not require a specific transporter for enterocyte uptake. Despite the presence of the peritrophic membrane, we observe reduced survival of adult WT Drosophila-fed CDCA at concentrations as low as 6.25 mmol/L, and almost 100% lethality within 100 hours at 25 mmol/L (data not shown). As shown in Figure 9A, feeding the bile acids CDCA or deoxycholic acid to the flies induced death within 4 days. To examine the potential cytoprotective role of intestinal Nrf2 (known as Cap 'n' collar isoform-c in Drosophila), the effect of bile acid feeding on survival of midgut enterocyte-specific Cap 'n' collar isoform-c-deficient Drosophila was determined. Reduction of enterocyte Nrf2 expression significantly

Figure 9. Bile acid cytotoxicity and protective role of intestinal Nrf2 (Cap 'n' collar isoform-c [cncC]) in Drosophila. (A) Relative survival of 5-day-old adult Drosophila in response to feeding CDCA or deoxycholic acid (DCA). Log-rank test for CDCA or DCA vs sucrose, $P < .001$, $n = 30–44$ per group. (B) Relative survival of 5-day-old adult Drosophila in which the levels of cncC (UAS-CncCIR) are diminished; log-rank test for myoA-GAL4 vs myoA-GAL4, UAS-cncCIR, $P < .0001$, $n = 50$ per group. (C) Detection of ARE-dependent GFP expression in the midgut of gstD-gfp adult Drosophila fed sucrose, sucrose plus chenodeoxycholic acid, or sucrose plus paraquat. (D) ROS generation in midgut after ingestion of sucrose or sucrose plus 25 mmol/L CDCA. ROS were detected by oxidation of the hydrocyanine ROS-sensitive dye, Hydro-Cy3. Images within a panel were taken at the same confocal settings: original magnification, $20\times$. 
Asbt−/−Ost−/− mice. (A) Fecal bile acid excretion. (B) Ileal expression of Asbt and FGF15 mRNA. (C) Hepatic expression of Cyp7a1 and Cyp8b1 in adult male mice for the indicated genotypes. The mRNA expression (n = 5) was determined by real-time PCR analysis (in triplicate) and normalized using cyclophilin. The mRNA expression is expressed relative to WT (set at 100%). Male mice (age, 8 wk) were included in the analysis (n = 5 per genotype). Mean values ± SEM are shown. Different lowercase letters indicate significant differences (P < .05) between genotypes. BW, body weight.

**Figure 10. Bile acid metabolism and small intestinal morphologic changes in adult male WT, Asbt−/−, Ost−/−, and Asbt−/−Ost−/− mice.**

Inactivation of the Asbt Prevents Ileal Injury in Ostα−/− Mice

The findings suggest that continued Asbt activity in the absence of the basolateral bile acid transporter Ostα-Ostβ leads to ileal accumulation of cytotoxic levels of bile acids early in life, which induces an oxidative stress–related injury and restitution response. To directly test this hypothesis, the Asbt null allele was crossed into Ostα−/− mice. In mice, the Asbt accounts for the vast majority of bile acid absorption from the gut lumen, with negligible contributions by other mechanisms. As such, if enterocyte bile acid uptake is required for the intestinal injury/restitution phenotype, inactivation of the Asbt should be protective. The Ostα−/−Asbt−/− mice are viable and fertile; crosses between heterozygous mice produced the predicted Mendelian distribution of WT and mutant genotypes. The adult Ostα−/−, Asbt−/−, and Ostα−/−Asbt−/− mice were indistinguishable from WT mice in terms of survival and gross appearance. To confirm that inactivation of the Asbt was sufficient to block intestinal bile acid uptake, fecal bile acid excretion and expression of hepatic and intestinal genes involved in bile acid metabolism were measured. As shown in Figure 10A, fecal bile acid excretion was similar in WT and Ostα−/− mice, and inactivation of the Asbt in Ostα−/− mice increased fecal bile acid excretion almost 6-fold to levels observed in Asbt−/− mice. In ileum, expression of the Asbt was reduced in Ostα−/− mice and undetectable in Asbt−/− and Asbt−/−Ost−/− mice. In agreement with the reduced bile acid uptake in the absence of the Asbt, expression of ileal FGF15 mRNA was reduced dramatically in Ostα−/−Asbt−/− and Asbt−/− mice to almost undetectable levels (Figure 10B). The intestinal bile acid malabsorption was associated with significant increases in the mRNA expression for hepatic Cyp7a1 and Cyp8b1, critical enzymes for bile acid synthesis (Figure 10C).

Because Ostα-Ostβ shows a gradient of expression along the cephalocaudal axis of the small intestine with highest expression levels in ileum, the segmental intestinal mass was examined in adult mice (age, 2 mo) for the different genotypes. For distal ileum (segment 5), the intestinal weight per unit length was increased significantly (approximately 45%) in male Ostα−/− mice (Figure 11A), and restored to near WT levels in Ostα−/−Asbt−/− mice. Histologically, the ileum of postnatal day 10 and adult (age, 8 wk) Ostα−/−Asbt−/− mice was almost indistinguishable from WT
mice, showing only mild epithelial reactive changes (Figure 11B). Furthermore, transcript levels of Nrf2 target genes Gsta1, Gsta3, Gsta4, Gstmu1, and Gstmu3 were increased from 3- to 8-fold in adult male Ostα−/− mice vs WT mice, but reduced to WT levels in the Ostα−/−Asbt−/− mice (Figure 11C). Together, these findings provide compelling evidence in support of the hypothesis that continued expression of the Asbt in the absence of Ostα-Ostβ leads to ileal bile acid stasis and injury. In that event, the Nrf2 pathway is one component of the adaptive response that functions to protect the ileal enterocyte from bile acid-induced cytotoxicity.

**Discussion**

The major finding of this study was that Ostα-Ostβ is required to protect the terminal ileum from bile acid-induced injury. We propose that inactivation of Ostα-Ostβ combined with continued Asbt expression results in bile acid accumulation, injury, and morphologic changes starting in early postnatal development. In a similar fashion, it has been postulated that hepatic Ostα-Ostβ plays a role in protecting against bile acid accumulation in hepatocytes under cholestatic conditions.39,57,58 In Ostα−/− mice, enterocyte bile acid stasis is associated with significant histologic mucosal alterations, including villous blunting, crypt hyperplasia, increased proliferation, apoptosis, and an increase in the number of mucin-secreting cells at the villous tips. These changes occur in the absence of histologic evidence of necrosis or an active inflammatory response. Despite a mild increase in TNFα mRNA levels early in postnatal development, ileal mRNA levels of TNFα and interleukin 1β are lower than WT mice by day 30, in agreement with our previous studies of adult Ostα−/− mice.10 The mechanisms responsible for the lower ileal cytokine mRNA levels in the postweaning Ostα−/− mice are unclear, but may be related to persistent increases in Nrf2 and antioxidant activity. For example, previous studies have
shown that administration of L-cysteine can increase intestinal Nrf2 activity, reduce intestinal cytokine levels, and suppress lipopolysaccharide-induced intestinal inflammation in a piglet model. In addition, intestinal levels of proinflammatory cytokines were found to be higher in Nrf2−/− vs WT mice after traumatic brain injury.

In WT mice or rats, ileal Asbt expression and active bile acid uptake remains low in early postnatal development, increasing after day 15. In contrast, there is an early induction of Asbt expression in Ostα−/− mice, correlating with increased expression of bile acid–activated FXR target genes, including Shp and Ibabp. Although premature ileal Asbt expression had been noted previously in mice and rats under pathophysiologic conditions such as experimental induction of necrotizing enterocolitis (NEC) or formula feeding, the underlying cause was not identified. Our attempts to interrogate the pathways/factors known to regulate the Asbt failed to identify a mechanism for the altered ileal Asbt temporal pattern in Ostα−/− mice. Among those factors that regulate Asbt expression are FXR, GATA4, tristetraprolin (TTR)/Human antigen R (HuR), and caudal (CDx) transcription factors 1 and 2. FXR is activated early in postnatal development in Ostα−/− mice, but is unlikely to be involved because the ileal morphologic changes still are present in Ostα−/−Fxr+/− mice. GATA4 and TTP/HuR are important regulators of ASBT gene transcription and RNA stability, respectively, in small intestine, and CDX1/2 are important transcriptional regulators of many intestinal genes. However, the ileal mRNA expression of these factors was similar in male and female WT and Ostα−/− mice at the early postnatal ages examined (data not shown), arguing against a direct role. Nrf2 has not been reported to directly regulate the Asbt, but Nrf2−/− mice have reduced levels of ileal Asbt expression and bile acid absorption. That observation raises the possibility that Nrf2 may contribute to the early postnatal increase in Asbt expression in Ostα−/− mice. However, it also should be noted that the relative increase in Asbt expression is transient and ileal Asbt mRNA and protein expression is reduced in postweaning adult Ostα−/− mice compared with age-matched WT male or female mice. Finally, these studies have equated ASBT expression with bile acid uptake because the ASBT appears to have a relatively narrow specificity and no endogenous substrates other than bile acids have yet been identified. Although the protection against ileal injury observed in Asbt−/−Ostα−/− mice supports the hypothesis that the phenotype is secondary to bile acid trapping, it is possible that ASBT transports another solute besides bile acids or its activity is coupled to another cellular pathway, which is responsible for the phenotype.

The cellular and molecular mechanisms of hepatocyte injury caused by retention of hydrophobic bile acids in cholestatic disease have been the subject of intensive study. In contrast, less is known regarding the mechanisms and signaling pathways underlying the cytotoxic effects of bile acids in other cell types and in extrahepatic tissues. This is particularly true for ileal enterocytes, which by virtue of their role in the enterohepatic circulation are exposed to a high daily flux of bile acids. In the 10-day-old Ostα−/− mice, ileal tissue bile acid levels were increased vs WT mice, and ranged from approximately 1.5 to 3 μmol/g of tissue. A limitation of these measurements of tissue-associated bile acids (which are primarily in the mucosa vs muscularis layers) is that it does not distinguish between the extracellular and intracellular compartments or directly measure the bile acid concentrations within the enterocytes. However, the ileal tissue bile acid levels in the 10-day-old Ostα−/− mice are appreciable and in the range of that observed in liver of bile duct–ligated mice. In the gut, bile acids such as deoxycholate have been shown to induce apoptosis in colonocytes through mechanisms involving protein kinase C delta and phospholipase A2, possibly by altering plasma membrane organization. Indeed, a myriad of pathways have been implicated in bile acid–induced cytotoxicity, including altering mitochondrial outer membrane structure, inducing mitochondrial membrane permeability transition, activating Nox, and signaling via epidermal growth factor receptor and CD95/Fas. One of the most important and relevant pathways for this study is bile acid–induced generation of ROS and oxidative stress, secondary to bile acid interaction with plasma membrane enzymes such as Nox and interaction with the mitochondria. Low to moderate levels of ROS maintain tissue homeostasis and promote cell proliferation and survival, but higher ROS levels can induce cell damage and death. In response, cells engage an evolutionarily conserved system, the Nrf2 pathway, to induce expression of a regulon of genes involved in xenobiotic and ROS detoxification, including those involved in glutathione synthesis and utilization, thioredoxin production and utilization, and quinone detoxification. In vitro studies of cells, including intestine-derived Caco-2 cells, have shown that bile acids can induce ROS and increase expression of Nrf2–activated genes. However, activation of Nrf2 by bile acids in enterocytes has not been shown in vivo. In this study, we show that blocking ileal enterocyte bile acid export in Ostα−/− mice leads to significant induction of Nrf2/ARE cytoprotective gene expression that correlates with the ileal morphologic changes. Tissue ROS levels were not directly measured in this study, but ileal levels of reduced glutathione were found to be decreased in Ostα−/− mice. In conjunction with the changes in gene expression, these findings suggest that induction of the glutamate-cysteine ligase/glutathione synthetase/GSH axis and expression of glutathione reductase, glutathione peroxidase 2, and glutathione-S-transferase may be quenching ROS or otherwise neutralizing free radicals to protect the epithelium and promote restitution. Exploration of this mechanism in intestine of the reductionist Drosophila model directly showed increased generation of ROS, activation of Nrf2–ARE signaling, and reduced survival of midgut enterocyte-specific Nrf2-deficient flies in response to CDCA feeding, supporting the concept of enterocyte bile acid–induced oxidative stress and injury. However, in addition to a pathophysiologic role in promoting cell damage, ROS also has a physiologic role in stimulating cell proliferation and tissue restitution. In particular, there is compelling evidence that ROS generation by Nox1 in the gut epithelium stimulates enterocyte or colonocyte proliferation and is required for restitution and wound repair after mechanical injury or chemical-induced colitis. In that regard, the significant early induction of Nox1 expression may have a role in promoting ileal mucosal repair in...
Ostα/− mice. Although the adaptive mechanisms are not sufficient to restore the ileal morphology back to that of the WT mice, it should be noted that the Ostα/− mice survive and continue to thrive. The persistence of the ileal morphologic changes may be because the hypothesized offending agent, bile acids, continuously are synthesized and secreted into the small intestine over the life of the mouse. As a result, an unremitting but controlled cycle of epithelial injury and restitution ensues. A schematic summarizing the findings and proposed mechanisms underlying the intestinal changes associated with loss of Ostα is shown in Figure 12.

In theory, enterocyte injury secondary to imbalanced expression of the ASBT and OSTα-OSTb may be involved in human disease. For example, this mechanism may contribute to the pathogenesis of NEC because ASBT expression has been reported to be higher in premature infants diagnosed with NEC vs premature infants with non-NEC diagnoses.61 Furthermore, in rat and mouse models of NEC, ileal damage correlated with increased Asbt expression and intracellular bile acid accumulation, and administration of an ASBT inhibitor or inactivation of the Asbt gene was protective.23,61 By comparison with the NEC models, the intestinal phenotype is relatively mild in Ostα/− mice, although the pups are born full-term and dam-fed, raising the possibility that the neonatal Ostα/− mice may be particularly susceptible to additional stressors such as hypoxia or formula feeding. Outside of inflammatory bowel disease, ileal atrophy is not often seen without accompanying duodenojejunal changes,98 but cases of apparent primary ileal villous atrophy and bile acid malabsorption of unknown etiology have been reported.94–96 Although the relationship of these disorders and the intestinal bile acid transporters remains to be determined, it should be noted that an inherited OSTb (SLC51B) deficiency recently was identified in a family with 2 affected brothers. These pediatric patients presented with congenital diarrhea, fatsoluble vitamin deficiency, and increased liver serum chemistries.97 Although a cursory endoscopic examination showed no obvious ileal abnormalities in the older affected sibling, additional studies will be required to determine the mechanisms underlying the patients’ diarrhea and apparent steatorrhea. Their clinical presentation may be owing solely to bile acid malabsorption and reduced luminal bile acid concentrations, similar to patients with ASBT mutations.5,6 Alternatively, changes in the ultrastructure of the ileal mucosa secondary to imbalanced ASBT and OSTα-OSTb expression also may contribute to the phenotype.

In conclusion, we have provided data indicating that in addition to its role in maintaining the enterohepatic circulation of bile acids, Ostα-Ostb appears to play a major role in protecting the ileal epithelium against bile acid accumulation and bile acid–induced injury. Inactivation of Ostα resulted in increased expression of FXR target genes as well as villous blunting, cell apoptosis, and oxidative stress in early postnatal development. There is also an early robust induction of Nrf2/anti-oxidant and cytoprotective genes, which may be involved in the subsequent healing and restitution response.

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Author contributions
Courtney Ferrebee, Paul Dawson, Brian Robinson, and Rheinallt Jones were responsible for the study concept and design; Courtney Ferrebee, Jianing Li, Jamie Haywood, Kimberly Pachura, Brian Robinson, Anuradha Rao, and Paul Dawson acquired data; Courtney Ferrebee, Jianing Li, Jamie Haywood, Anuradha Rao, Brian Robinson, Rheinallt Jones, Benjamin Hinrichs, and Paul Dawson were responsible for the analysis and interpretation of data; Courtney Ferrebee, Rheinallt Jones, and Paul Dawson drafted the manuscript; Rheinallt Jones and Paul Dawson obtained funding; and Paul Dawson was responsible for study supervision.

Conflicts of interest
The authors disclose no conflicts.

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