Bile acid homeostasis controls CAR signaling pathways in mouse testis through FXRalpha.

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Fig. S1. a/ Testicular mRNA accumulation of *Lcgr*; *Fshr*; *Nanog*, *Oct3/4*, *G9a*, *Ccna1 and Smad6*, normalized to β -actin mRNA levels in whole testes of C57BL/6J mice treated with busulfan (20 mg/kg, one injection IP) at T0, 4, or 8 weeks (n=8 per group). *Denotes significant difference from the T0 time point; #denotes significant difference from the 4-week time point (P < 0.05). **b**/ Analysis of the relative liver and testicular levels of *Cyp7a1*, *Cyp8b1*, *Hsd3b7*, *Cyp27a1* and *Cyp7b1*. The relative levels are represented as delta-Ct versus β -actin. **c**/ Testicular concentration of LCA in testis of Wt and Fxr α -/- males fed a control or BA-diet for 2 weeks (n=5 per group).

In all panels, data are expressed as means \pm standard error of the mean. Statistical analysis: *P < 0.05.



Fig. S2. a/ Testis weight normalized to body weight in wild-type C57Bl6 males treated 2 weeks with PCN (50mg/kg/day) or vehicle (DMSO) (n=6 to 7 per group). **b**/ Spermatozoa count in the head of the epidydimis of C57Bl6 males treated 2 weeks with PCN (50mg/kg/day) or vehicle (DMSO) (n=6 to 7 per group). **c**/ Quantification of TUNEL analyses in C57Bl6 males treated 2 weeks with PCN (50mg/kg/day) or vehicle (DMSO), the number of tubules with TUNEL-positive cells is indicated as the number of positive tubes per 100 seminiferous tubules (n = 6-7). **d**/ Testicular mRNA accumulation of *Shp, Stra8 and Cyp2b10* normalized to β -actin mRNA levels in C57Bl6 males treated 2 weeks with PCN (50mg/kg/day) or vehicle (DMSO). Vehicle-treated mice were arbitrarily fixed at 100% (n=6 to 7 per group). **e**/ Liver mRNA accumulation of *Mrp3 and Cyp2b10* normalized to β -actin mRNA levels in C57Bl6 males treated 2 weeks with PCN (50mg/kg/day) or vehicle (DMSO). Vehicle-treated mice were arbitrarily fixed at 100% (n=6 to 7 per group). **e**/ Liver mRNA accumulation of *Mrp3 and Cyp2b10* normalized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA accumulation of *Mrp3 and Cyp2b10* normalized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA levels in C57Bl6 males treated 100% (n=6 to 7 per group). **e**/ Liver mRNA accumulation of *Mrp3 and Cyp2b10* normalized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to β -actin mRNA levels in C57Bl6 males treated 2 mellized to 7 per group).

In all panels, data are expressed as means \pm standard error of the mean. Statistical analysis: *P < 0.05.



Fig. S3. Liver mRNA expression of *Sult2a1*, *Cyp2b10* and *Cyp3a11* normalized to β -actin mRNA levels in wild-type and Fxr α -/- adult mice fed a control or BA-diet (n = 6 -10 per group).

Data are expressed as means \pm standard error of the mean. Statistical analysis: **P* < 0.05.

Table S1

FW	ACACAGACAGACCCATACTGG
REV	TCACCTGTATGTGAGCGAAGG
FW	ACCTGCACCTGAGTTCTTGG
REV	CAATCACATTCTGTGTGCCC
FW	GTGGTGCTGATCCCCTATGT
REV	CTTGCCAGGAAATCATCCAG
FW	ACCAGCCGCAAGCTAAAGACTCAT
REV	TCAGACTGGCATTTACCCACTCT
FW	ATGATGACACCGACGATCAGG
REV	TCGGTACTGCTGACTCTGTCC
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