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Corrigendum: Effects of *Poria cocos* extract on metabolic dysfunction-associated fatty liver disease via the FXR/PPAR α -SREBPs pathway

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KEYWORDS

MAFLD (metabolic-associated fatty liver disease), *Poria cocos* (Schw.) Wolf., bile acid metabolism, FXR/PPAR α -SREBP pathway, lipid homeostasis, UPLC Q-TOF/MS

A Corrigendum on

Effects of *Poria cocos* extract on metabolic dysfunction-associated fatty liver disease via the FXR/PPAR α -SREBPs pathway

by He J, Yang Y, Zhang F, Li Y, Li X, Pu X, He X, Zhang M, Yang X, Yu Q, Qi Y, Li X and Yu J (2022). *Front. Pharmacol.* 13:1007274. doi: 10.3389/fphar.2022.1007274

In the published article, there was an error in the legend for **Figure 2** as published. In the legend for **Figure 2**, “(N) Brown adipose tissue (BAT).” is a duplicate and needs to be deleted because the BAT is explained in the legend for **Figure 2I**. The corrected legend appears below.

“FIGURE 2 | EPC ameliorated MAFLD in rats. (A) Body weight (BW). (B) BW gain. (C–E) Organ wet weight. (F) Inguinal white adipose tissue (iWAT). (G) Perirenal white adipose tissue (pWAT). (H) Epididymis white adipose tissue (eWAT). (I) Brown adipose tissue (BAT). (J) iWAT/BW ratio. (K) pWAT/BW ratio; (L) eWAT/BW ratio. (M) BAT/BW ratio. (N) Representative rat liver images of hematoxylin and eosin (H and E) and Oil Red O staining per group (X200). (O) Representative iWAT, pWAT, eWAT, BAT. One-way analysis of variance (ANOVA) was conducted for the group comparison. $n = 8$, data are presented as mean \pm SEM. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ vs. MOD group. EPC, *P. cocos* ethanol extract; CON, normal diet control group; MOD, high-fat diet group; FC, Fenofibrate capsules; EPC-L, low-dose *P. cocos* ethanol extract; EPC-H, high-dose *P. cocos* ethanol extract.]”

Furthermore, there was an error in **Figure 6P** as published. The authors apologize for uploading the ERK protein image in **Figure 6** incorrectly, with image of p-JNK, in this article. Furthermore, P-ERK should be p-ERK in **Figure 6P**. The corrected **Figure 6** appears below.

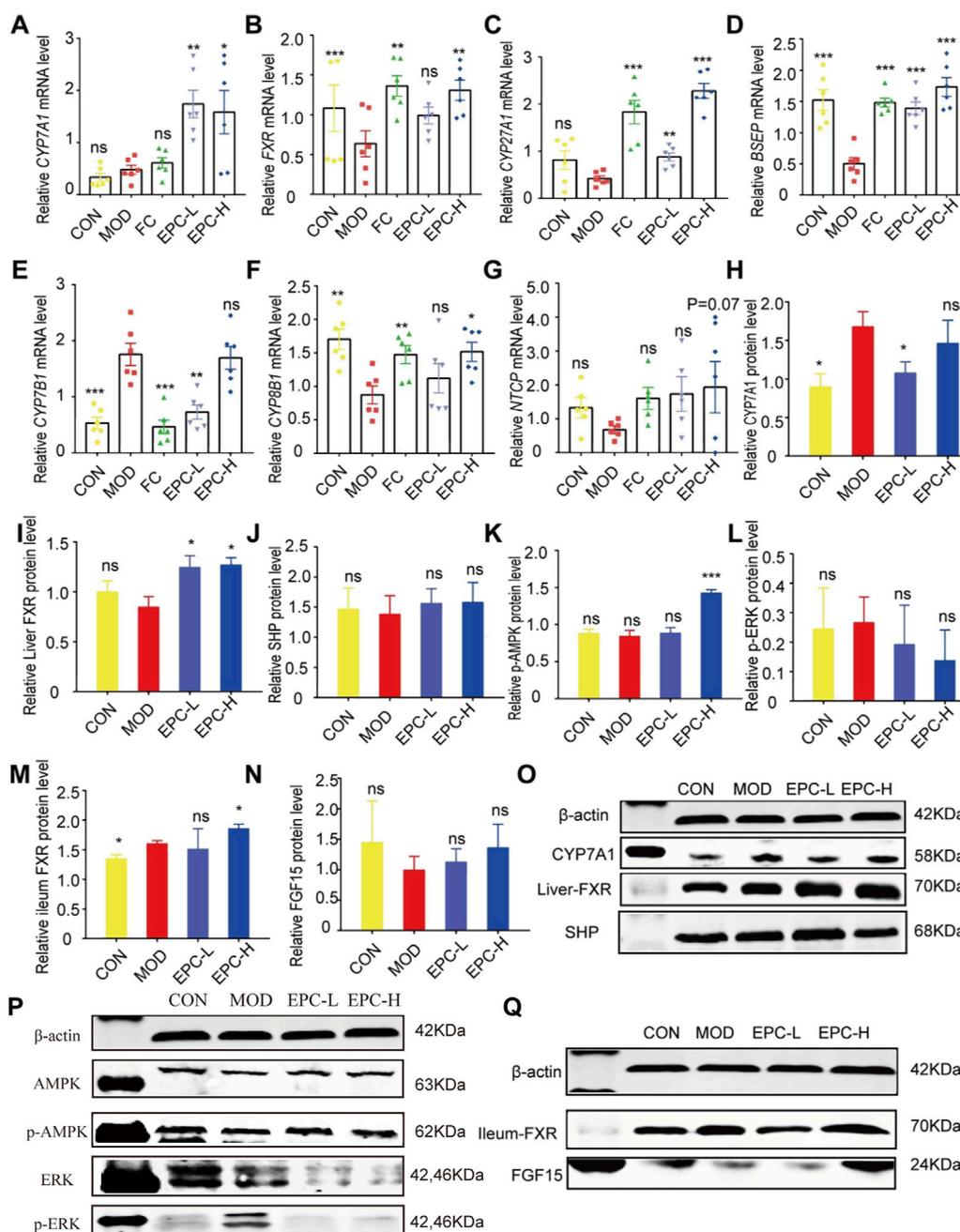


FIGURE 6

EPC ameliorated MAFLD formation in rats by regulating BA metabolism. (A–G) Relative expression of CYP7A1, FXR, CYP27A1, BSEP, CYP7B1, CYP8B1, NTCP mRNA in liver, n = 6; (H–L) Relative expression of protein CYP7A1, FXR, SHP, p-AMPK, and p-ERK in the liver, n = 4; (M–N) Relative expression of protein FXR and FGF15 in the ileum, n = 4. (O–P) Representative immunoblotting images of CYP7A1, FXR, SHP, p-AMPK, and p-ERK in the liver. (Q) Representative immunoblotting images of FXR and FGF15 in the ileum. Data are presented as mean ± SEM. One-way analysis of variance (ANOVA) was conducted for the group comparison. **p* < 0.05, ***p* < 0.01, ****p* < 0.001 vs. MOD group. CYP7A1, cholesterol 7 α -hydroxylase; FXR, farnesoid X receptor; CYP27A1, sterol 27-hydroxylase; BSEP, bile salt export protein; CYP7B1, oxysterol 7 α -hydroxylase; CYP8B1, sterol 12 α hydroxylase; NTCP, Na + -taurocholate co-transporting polypeptides; SHP, small heterodimer partner; AMPK, 5'-AMP-activated protein kinase; ERK, Extracellular signal-regulated kinase.

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