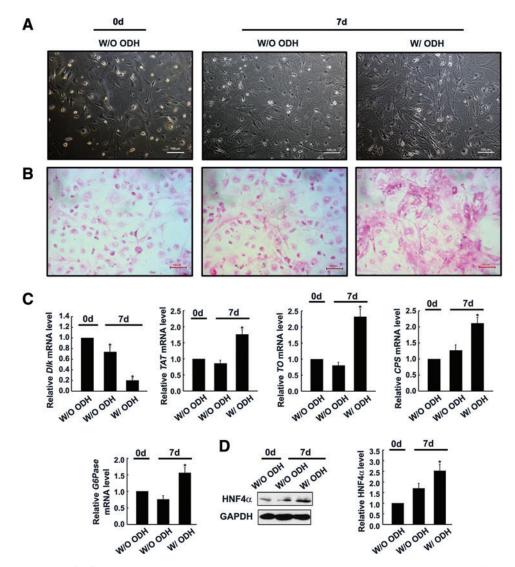
Supplementary Data



SUPPLEMENTARY FIG. S1. The capability of hepatoblasts to develop into mature hepatocytes. (**A**) The morphology under phase-contrast microscopy of the hepatoblasts with or without ODH induction for 7 days is presented. (**B**) Glycogen contents in cultured hepatoblasts. The hepatoblasts were cultured for 7 days in the absence or presence of ODH. Then, the intracellular glycogen accumulation was analyzed by staining the cells with the PAS reagent. Glycogen is shown in *magenta*. Scale bar = 100 μm. (**C**) Expression of hepatocyte marker genes in cultures supplemented with ODH. After 7 days of induction, the expression levels of immature hepatocyte markers (*DLK*) and mature hepatocyte markers (*TAT*, *TO*, *CPS*, and *G6Pase*) were evaluated by qRT-PCR. The results are the means \pm SDs (n=5). *P<0.05 compared with the values at day 0 without ODH. (**D**) Cells were treated with ODH for 7 days. At the end of the incubation, the cells were lysed, and 100 μg of protein extract was loaded and separated by SDS/PAGE. Western blotting was performed using antibodies against HNF4α. The relative densities of the HNF4α bands were normalized to GAPDH. The values are expressed as the means \pm SDs of four independent experiments.