- **Supplementary Materials:**
- 2 Fig. S1. Representative images for (A) normal hepatocyte, (B) steatosis, (C) Hyperplasia, (D)
- 3 Dysplasia, and (**E**,**F**) HCC in transgenic zebrafish.

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- 5 **Fig. S2.** Pathways analysis for the differential expressed genes in [HBx,src], and [HBx,src,p53⁻¹]
- 6 /+] transgenic zebrafish following oligo-fucoidan treatment. (A) Enriched pathways of oligo-
- 7 fucoidan upregulated genes. (**B**) Enriched pathways of oligo-fucoidan downregulated genes..

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- 9 Fig. S3. Upstream regulator analysis using IPA for the differential expressed genes in
- 10 [HBx,src], and [$HBx,src,p53^{-/+}$] transgenic zebrafish following oligo-fucoidan treatment. (A)
- 11 Clustering of the upstream regulators identified by IPA for the [HBx, src, p53^{-/+}]-DIO transgenic
- fish compare to NOR (DIO-batch2), [HBx,src]-DIO+OF compare to DIO (DIO-batch1), and
- 13 [HBx,src,p53^{-/+}]-DIO+OF compare to DIO (DIO-batch2). (**B-E**) The upstream regulators
- predicted by IPA including MYCN, KRAS, TGFB1 and STK1 which were predicted induced
- by DIO (as shown in orange), and repressed by OF (as shown in blue) together they either
- upregulates the downstream genes (as shown in red) or inhibit the downstream genes (as shown
- in green) in [HBx, src, $p53^{-/+}$] transgenic fish.

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