## **Supporting Information**

## Tumor-targeting Polycaprolactone nanoparticles with co-delivery of paclitaxel and IR780 for combinational therapy of drug-resistant ovarian cancer

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Figure S1. TEM images of (A) PCL and (B) PCL/IR780-PTX nanoparticles.



**Figure S2**. The temperature of 0.5 mg/mL PCL-LHRH/IR780-PTX nanoparticle suspension as a function of irradiation time under different light dosage.



**Figure S3**. Fluorescent intensity of endothelial cells cultured with Nile red-labelled PCL NPs or PCL-LHRH NPs, as a function of incubation time.



**Figure S4**. Plasma ID% of IR780 in tumor bearing mice as a function of time after administration of free IR780, PCL/IR780 and IR780-LHRH/IR780, respectively.



**Figure S5**. Photos of ST30 tumors harvested from mice after sacrifice. Circles indicate that there is no tumor left.



**Figure S6**. The density of proliferative (Ki67 positive) cells. **\*\*** indicates significant difference at p<0.01 level. (I) control group, (II) PTX + Light group, (III) PCL-LHRH/IR780-PTX NPs, (IV) PCL-LHRH/IR780 + Light, (V) PCL/IR780-PTX + Light, and (VI) PCL-LHRH/IR780-PTX + Light, respectively (n = 5).



**Figure S7**. Blood test results of mice received (I) PCL-LHRH NPs and (II) PCL-LHRH/IR780-PTX NPs (n=4). The blood was collected and tested after 1 day after i.v. injection of nanoparticles. The dosages of PCL-LHRH NPs and IR780 were 2 mg and 148  $\mu$ g for each mouse, respectively.