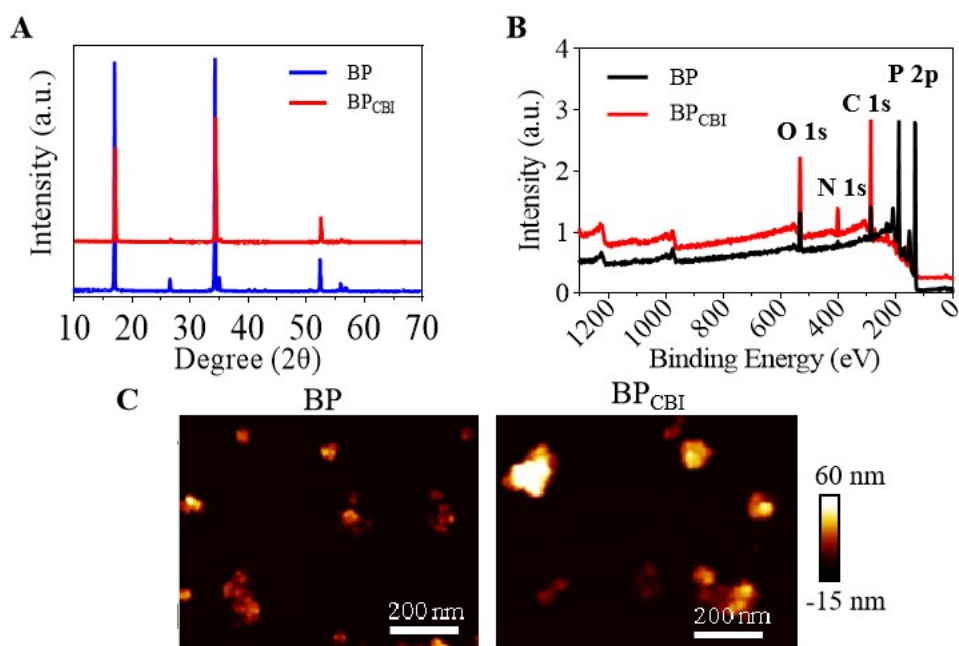


**Table S1 Drug loading content (DLC, wt%) and loading efficiency (DLE, %) of CI and BI into M@BP determined by HPLC**

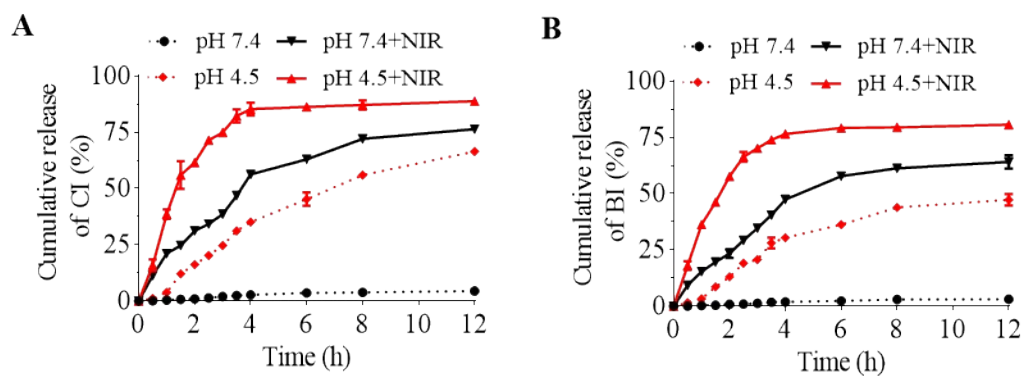
Nanosystem	Drug	DLC (wt %)		DLE (%)
		Theory	Determined	
M@BP <sub>CBI</sub>	CI	10	9.29	96.90
	BI	10	9.44	98.40
	CI/BI	10	9.21/9.35	96.19/98.31

**Figure S1**



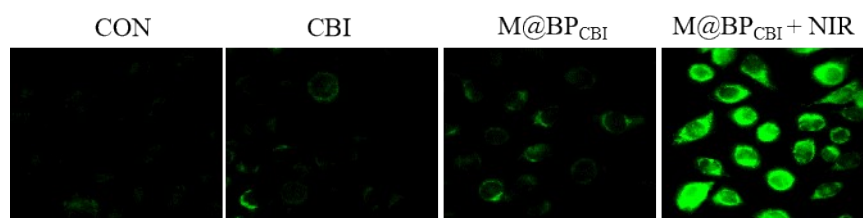
The characterization of BP and CBI-loaded BP with XRD assay (A), XPS spectra (B) and AFM (C), respectively

**Figure S2**



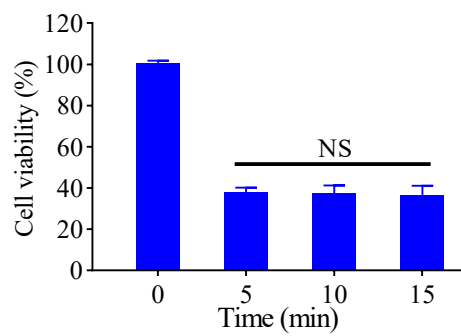
The cumulative release of CI (A) and BI (B) under different pH conditions (pH7.4 and 4.5) treated with or without NIR irradiation was determined by HPLC

**Figure S3**



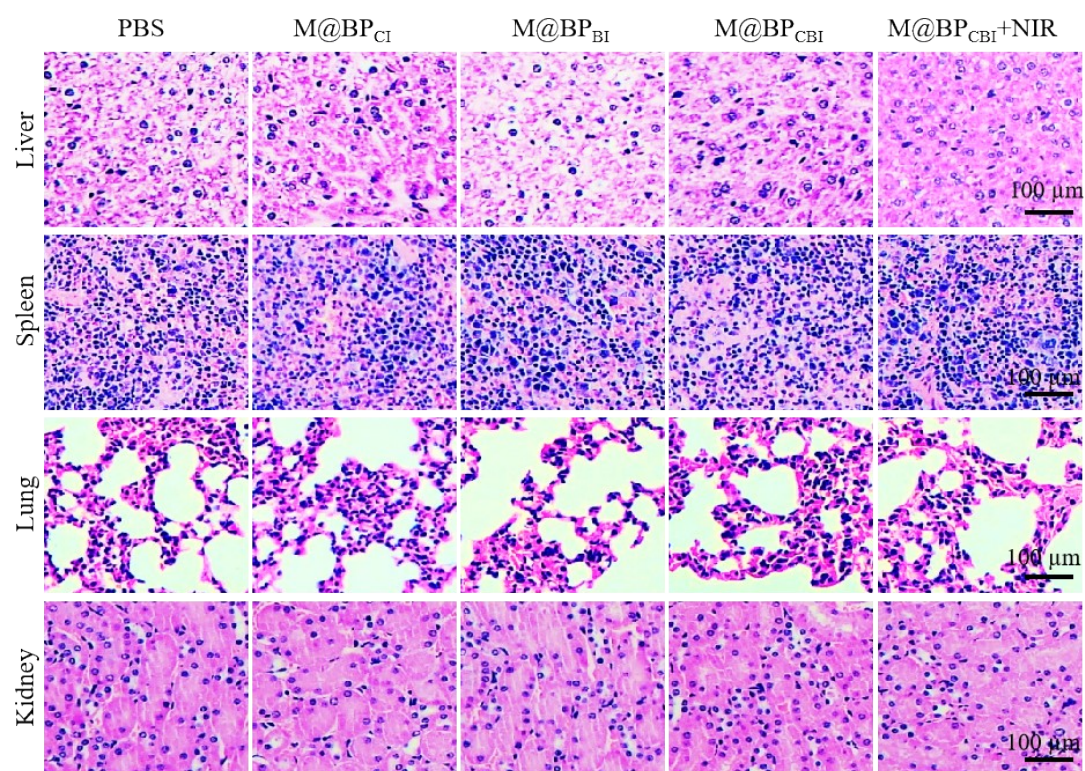
MFC cells were treated with different formulations (CI: 9.8  $\mu\text{M}$ , BI: 10  $\mu\text{M}$ , BP: 45  $\mu\text{g mL}^{-1}$ ) for 6h followed with or without NIR irradiation (NIR irradiation: 1.5  $\text{W cm}^{-2}$ ) for 10 min, incubated with ROS probe of DCFH-DA for 30 min, and then examined under fluorescence microscope

**Figure S4**



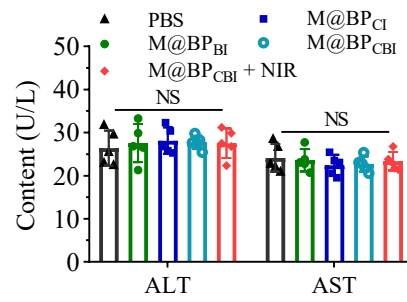
The CI and BI were incubated at 50 °C for different time, then added to MFC cells (Concentration: 10  $\mu$ M) for 48 h and examined by MTT assay. NS indicated non-significant difference

**Figure S5**



H&E staining of the major organs separated from different groups at the end of experiments

**Figure S6**



Aspartate aminotransferase (AST) and alanine aminotransferase (ALT) measurement in mice with different treatments for liver function assay at the end of experiments. NS indicated non-significant difference

Raw data

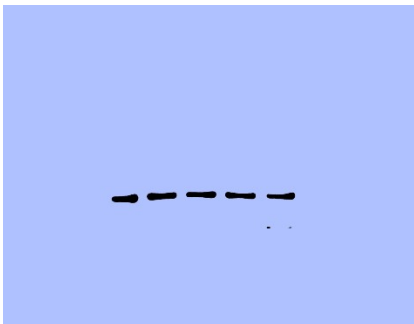
Pol II pSer2



Pol II



CDK9



BRD4

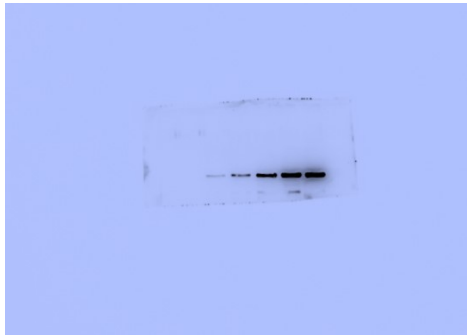




MYC



Cleaved Casp.3



GAPDH

