Effective targeted therapy for drug-resistant infection by ICAM-1 antibody-conjugated TPGS modified β -Ga₂O₃:Cr³⁺ nanoparticles

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Figure S1. Morphology observation of I-TPGS/Ga₂O_{3.}

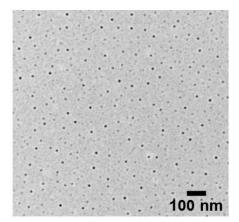
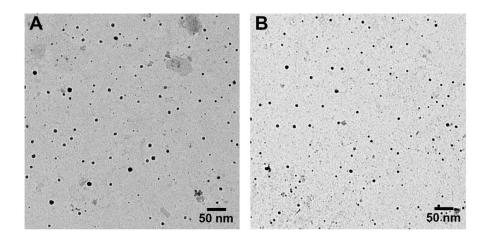
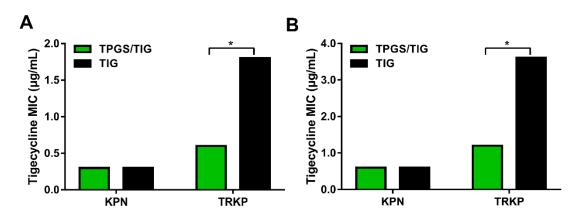


Figure S2. Morphology observation of I-TPGS/ Ga_2O_3 /TIG.



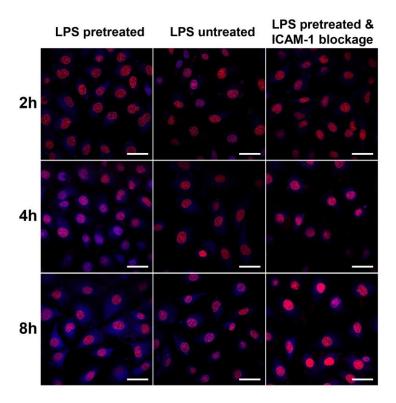
(A) TEM image of I-TPGS/ Ga_2O_3 /TIG before the pH 5.5 drug release study; (B) TEM image of I-TPGS/ Ga_2O_3 /TIG after the pH 5.5 drug release study.

Figure S3. *In vitro* antibacterial studies of TPGS/TIG and control samples on KPN and TRKP.



(**A**) MIC susceptibility semiquantitative profiles of TPGS/TIG and TIG against KPN and TRKP using microplate broth dilution method (n=3). (**B**) MBC susceptibility semiquantitative profiles of TPGS/TIG and TIG against KPN and TRKP (n=3). *p<0.05.

Figure S4. Cellular uptake study.



The cellular fluorescent uptake images of I-TPGS incubated with HUEVC for 2 h, 4 h and 8 h in the presence or absence of LPS pretreatment and ICAM1 blockage, respectively. HUEVC were labelled with DAPI (red), and the blue fluorescence signal indicated the excited blue fluorescence from β -Ga₂O₃:Cr³⁺. The bar is 40 μ m.

Figure S5. Body weights of the acute KPN-infected and TRKP-infected pneumonia mice within 5 days after treatments.

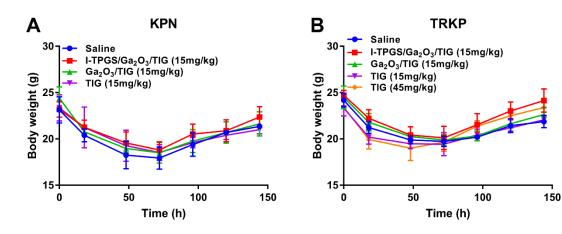


Table S1. Haematology parameters of mice in control group and I-TPGS/Ga₂O₃/TIG treated group (mean \pm SD, n = 6).

Blood chemistry parameters	Control group	Experimental group
WBC (10 ⁹ /L)	3.70 ± 1.19	4.17 ± 1.23
Lymph (10 ⁹ /L)	3.15 ± 0.45	3.27 ± 0.74
Mon (10 ⁹ /L)	0.10 ± 0.07	0.10 ± 0.08
Gran (10 ⁹ /L)	0.85 ± 0.23	0.80 ± 0.20
RBC $(10^{12}/L)$	7.60 ± 0.12	7.79 ± 1.24
HGB (g/L)	131.33 ± 2.12	134.00 ± 11.38
HCT (%)	42.87 ± 2.55	42.20 ± 6.97

^{*} The white blood cell (WBC) count, lymphocytes (Lymph) count, monocytes (Mon) count, neutrophilic granulocyte (Gran) count, red blood cell (RBC) count and haemoglobin (HGB) of the whole blood were determined.