Organ Biodistribution of Radiolabelled δγ T Cells Following Liposomal Alendronate Administration in Different Mice Tumour Models

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SUPPLEMENTARY FIGURES



Figure S1: *In vivo* Biodistribution of radiolabelled [¹¹¹In]L-ALD in A375P β 6 tumours, after single dose administration *via* tail vein injection in NSG mice. NSG mice were inoculated with the A375P β 6 cell line to form subcutaneous (SC), intraperitoneal (IP) or pseudo-metastatic lung tumours. Mice were i.v. injected with [¹¹¹In]L-ALD at a dose of 2 µmol lipid/mouse. After 24 h the mice were sacrificed and the amount of liposomes was quantified by gamma counting. (A) Organ biodistribution of [¹¹¹In]L-ALD expressed as per cent injection dose organ (%ID). (B) SC-tumour and IP-tumours uptake of [¹¹¹In]L-ALD expressed as %ID. (C) Tumour-bearing lung and healthy lung uptake of [¹¹¹In]L-ALD expressed as %ID. Data was expressed as mean ± SD (n=4)



Figure S2: *In vivo* biodistribution of radiolabelled $\gamma\delta$ T cells in A375P $\beta\delta$ tumour bearing NSG mice, after single dose administration *via* tail vein injection. NSG mice were inoculated with luciferase-expressing A375P $\beta\delta$ cell line to form (A) subcutaneous (SC), (B) intraperitoneal (IP) or (C) pseudo-metastatic lung tumours. Mice were i.v. injected with [¹¹¹In] $\gamma\delta$ T cells at a dose of 5 x 10⁶ $\gamma\delta$ T cells/mouse. Mice were pre-treated with 0.5 µmol ALD or L-ALD, 24 h prior to injection of $\gamma\delta$ T cells. After 24 h the mice were sacrificed and the amount of $\gamma\delta$ T cells was quantified by gamma counting. Results are expressed as percentage injection dose (%ID) per organ. Data was expressed as mean ± SD (n=4).