

Technical Information

NpFlamma[®] HGC series

Overview

NpFlamma[®] HGC series is a fluorescent dye incorporated chitosan based amphiphilic nanoparticles that enables to selectively detect tumor cells. Chitosan nanoparticles can selectively accumulate in cancer tissues due to high permeability for new blood vessels in cancer tissues. Chitosan particles display low toxicity along with absence of noticeable side effect *in vivo*, yet they exhibit a long half-life, high stability and aqueous solubility. Thus, NpFlamma[®] HGC series is an ideal fluorescence agent for *in vivo* imaging of angiography and tumor progression. The hydrophobic nature of NpFlamma[®] HGC series can embed hydrophobic materials, thus they can be utilized as a selective carrier for hydrophobic cancer drugs such as doxorubicin and paclitaxel, etc.

Table 1. NpFlamma[®] HGC series list

Cat. No.	Product name	Ex (nm)	Em (nm)	Common filter set	Excitation source
PNC1201	NpFlamma [®] HGC 648	648	675	Cy 5	594, 633 nm
PNC1401	NpFlamma [®] HGC 675	675	698	Су 5.5	633, 680 nm
PNC1301	NpFlamma [®] HGC 749	750	782	Су 7	680 nm
PNC1601	NpFlamma [®] HGC 774	777	802	Су 7.5	785 nm
PNC1501	NpFlamma [®] HGC ICG	785	821	Су 7.5	785 nm

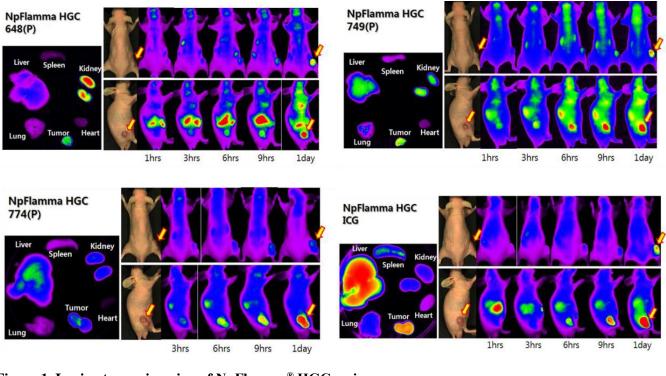
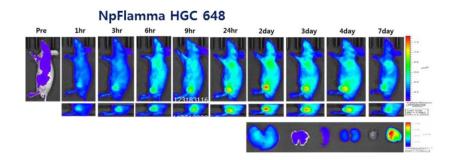
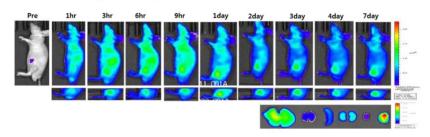
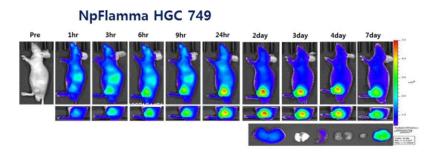


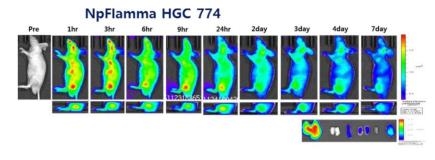
Figure 1. In vivo tumor imaging of NpFlamma[®] HGC series



NpFlamma HGC 675







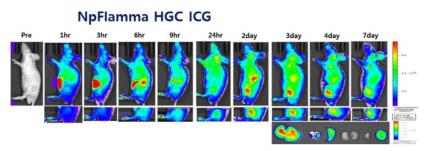
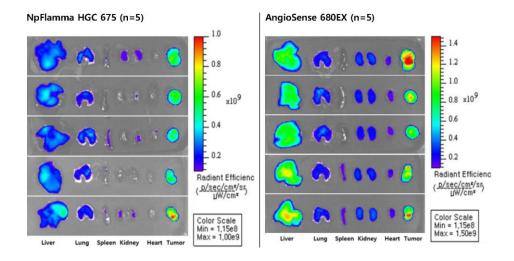


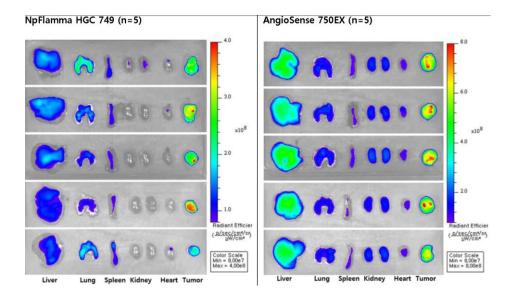
Figure 2. In vivo imaging of five NpFlamma[®] **HGC series over seven days** Fluorescence from cancer cell continued after 7 days injection, but fluorescence from other organs disappeared.

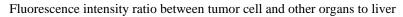


Fluorescence intensity ratio between tumor cell and other organs to liver

	Liver	Lung	Spleen	Kidney	Heart	Tumor	
1	1	0.940	-	0.509	-	1.812	
2	1	0.669	-			1.747	
3	1	0.944				1.650	
4	1	0.657				1.509	
5	1	0.613	-			1.998	

ור		Liver	Lung	Spleen	Kidney	Heart	Tumor	
11	1	1	0.325	-	0.406	0.226	1.634	
11	2	1	0.431	-	0.368	0.237	1.329	
11	3	1	0.358	-	0.324	0.203	1.022	
11	4	1	0.276	0.158	0.330	0.214	1.058	
11	5	1	0.305	0.168	0.373	0.209	1.072	





Liver	Lung	Spleen	Kidney	Heart	Tumor			Liver	Lung	Spleen	Kidney	Heart	Tumor
1	1.632		-		2.271		1	1	0.357	-	0.388	-	1.311
1	0.879				2.292		2	1	0.398	-	0.410		1.626
1	0.834	-	-	-	2.171		3	1	0.319	-	0.372	-	1.358
1	0.682	-	-	-	2.762		4	1	0.360	-	0.405	-	1.400
1	1.476	-	-	-	1.955		5	1	0.396	-	0.451		1.389
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Figure 3. Comparison of ex-vivo imaging of NpFlamma[®] **HGC and another imaging agent** NpFlamma[®] HGC series displays higher tumor cell accumulation than that of AngioSense

In vivo Imaging Protocol

General

- Prepare the fluorescent probe solution by adding DW or PBS to the NpFlamma^{\circ} HGC series powder and vortex the mixture (1does = 120 µg per 100 µL)
- Since fluorescent substances are unstable under light, they should be stored in the dark.
- Mouse fur may cause scattering or absorption of excitation of light during optical imaging process. Use nude mouse or remove the mouse fur in advance.
- It is recommended to use 31 G syringe needle.
- Prepare 5 week-old male Balb/c-nude mouse.

Typical procedure for mouse model tumor imaging with NpFlamma® HGC

- Inject SCC7 cell line (1x10⁶ per 0.1 mL) into subcutaneous of Balb/c-nude mouse.
- When the volume of tumor cell reaches to 60~80 mm³, take the zero time image of each subject.
- Inject NpFlamma® HGC series (120 µg per 100 µL) intravenously to mouse.
- The optimal interval for fluorescence imaging is 1 h, 3 h, 6 h, 9 h, and 24 h after injection.
- After take the 24 h imaging, extract major organs (liver, lung, spleen, kidney, heart) and tumor cell, and perform the ex-vivo imaging process.

Custom Labeling Service

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Nucleic acid



Peptide/Protein



Antibody



Small molecules /Polymer

Technical Support

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