

Supporting Information

**Comparison of penetration depth in mouse brain in vivo through
3PF imaging labelled by AIE nano particles and THG imaging
within the 1700 nm window**

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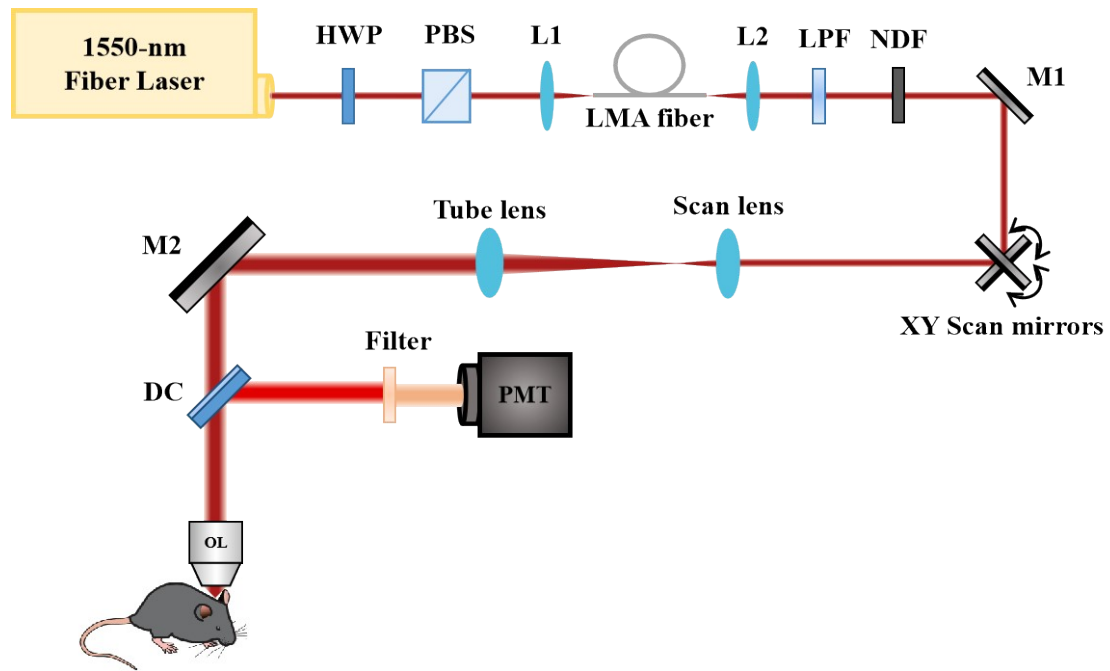


Fig. S1 The experimental setup of 3PM imaging system. HWP: half-wave plate; PBS: polarization beam splitter; L1 and L2: focusing and collimating lens; LPF: long-pass filter; NDF: neutral density filter; M1 and M2: silver-coated mirror; DC: dichroic mirror; Filter: band-pass filter; PMT: GaAsP photo-multiplier tube; OL: objective lens.

Table S1 In vivo mouse deep-brain imaging of different fluorescence indicator with 1700nm window

Wavelength (nm)	Materials	MPM	Concentration	Depth (μm)	Repetition rate (MHz)	Ref.
1600	MTTCM NPs	3PM	1 mM	840	6	*
1610	BONAPs	3PM	50 mg/kg	1680	1	[1]
1617	ICG	2PM	2 mM	2000	1	[2]
1620	SR101	3PM	3.3 mg/ml	1340	1	[3]
1660	MTTCM NPs	3PM	2 mM	1900	1	[4]
1665	Texas red	3PM	700 μM	1650	1	[5]
1665	Qtracker655	3PM	2 μM	2100	1	[6]
1665	DPNA-NZ	3PM	2 mM	1700	1	[7]
1665	DPCZ-BT	3PM	2 mM	1860	1	[8]
1700	DCTBT	2PM	2 mM	2180	1	[9]
1700	MTTCM NPs	3PM	1 mM	890	6	*
1720	OEFT NPs	3PM	550 μM	1696	1	[10]
1800	MTTCM NPs	3PM	1 mM	810	6	*

* Experimental parameters from this article

Note: The repetition rate of the laser used in this experiment was 6 MHz, while all other experiments used a repetition rate of 1 MHz. Because of the higher repetition rate used in this experiment, the pulse energy is lower and the imaging depth is smaller.

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