

| Known Property | microglia selective probe |
|--|-------------------------------|
| Application | Immunofluorescence |
| Cell selectivity mechanism: MOLD (Ugt1a7c) | |
| Storage | ① Delivery: Room Temperature |
| | ② Dried compound: 4 °C or -20 |

(3) Compound solution: 4 °C or -20 °C





General Use Guide

Suck solution is essential Suck solution is essential Working concentrations for specific applications should be determined by the investigator. It is recommended to use up the buffer diluted solution within one day Tr precipitated out from buffer solution It is recommended to use up the buffer diluted solution within one day. The compound may be decomposed or Jalize With

°C

OH

Molecular Weight

354.16 (C₁₉H₁₇BF₂N₂O₂)

 $\lambda_{ex} / \lambda_{em}$

568 / 600 nm

CDr20 works both in vitro cell culture and on brain slice. In mouse embryo, CDr20 penetrated into brain tissue and live microglia was fluorescently stained. Through iv injection, CDr20 was introduced to Alzheimer mouse brain and the microglia could be in situ imaged by two photon fluorescence microscope.



Identification of **CDr20** as a microglia-specific fluorescent probe. A) Left: Experimental setup. Right: Related probes: CDr10 Chemical structure of CDr20 with the structure-activity relationships. B) Densitometry of nonstained or CDr20-stainedCsf1r–EGFP brain cells. C) Superimposed images of CDr20 live-cell labeling and the indicated immunolabeling in primary cultured glial cells. Scale bars=100 μ m.

Reference

1. Visualizing Microglia with a Fluorescence Turn-On Ugt1a7c Substrate, Kim, B.; Fukuda, M.; Lee, J. Y.; Su, D.; Sanu, S.; Silvin, A.; Khoo, A. T. T.; Kwon, T.; Liu, X.; Chi, W.; Liu, X.; Choi, S.; Wan, D. S. Y.; Park, S. J.; Kim, J. S.; Ginhoux, F.; Je, H. S.*; Chang, Y. T.* Angew. Chem. Int. Ed. Engl. 2019, 58, 7972-7976.