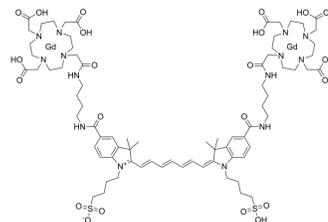


Gd-NMC-3

| | |
|--------------------|---|
| Cat. No.: | HY-150979 |
| CAS No.: | 2678579-76-3 |
| Molecular Formula: | $C_{77}H_{116}Gd_2N_{14}O_{22}S_2$ |
| Molecular Weight: | 1968.46 |
| Target: | Fluorescent Dye |
| Pathway: | Others |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | | | | | | | | | |
|--------------------|---|---------------|---|----------------|------------------|------------------|---|---------|--|
| Description | Gd-NMC-3 is a near-infrared fluorescence/magnetic resonance (NIRF/MR) bimodal imaging probe. Gd-NMC-3 shows high resolution and sensitivity in tumor imaging with good biocompatibility, indicating huge application potential ^[1] . | | | | | | | | |
| In Vitro | <p>Gd-NMC-3 shows the maximum excitation wavelength and emission wavelength are 755 and 792 nm, respectively. Both wavelengths are located in the near-infrared region^[1].</p> <p>Gd-NMC-3 acts as a bimodal imaging molecule, can be accumulated in tumor sites^[1].</p> <p>Gd-NMC-3 (50 μM; 24 h) can be internalized into cancer cells by OATPs and NTCP, indicating an excellent specificity to tumor tissues^[1].</p> <p>Gd-NMC-3 (6.25-800 μM, 24 h) exhibits significant fluorescence accumulation (with the optimal concentration of 100, 200 μM) and reasonable relaxation property (11.64 M/m/s) in tumors^[1].</p> <p>Gd-NMC-3 (6.25-100 μM, 48 h) displays low cytotoxicity and good biocompatibility^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HepG2 and LM3: human hepatocarcinoma cell line; L02: human hepatocyte cell line</td> </tr> <tr> <td>Concentration:</td> <td>6.25-100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Resulted more than 90% cell viability maintained after 48 h.</td> </tr> </table> | Cell Line: | HepG2 and LM3: human hepatocarcinoma cell line; L02: human hepatocyte cell line | Concentration: | 6.25-100 μ M | Incubation Time: | 48 hours | Result: | Resulted more than 90% cell viability maintained after 48 h. |
| Cell Line: | HepG2 and LM3: human hepatocarcinoma cell line; L02: human hepatocyte cell line | | | | | | | | |
| Concentration: | 6.25-100 μ M | | | | | | | | |
| Incubation Time: | 48 hours | | | | | | | | |
| Result: | Resulted more than 90% cell viability maintained after 48 h. | | | | | | | | |
| In Vivo | <p>Gd-NMC-3 (20 mg/kg; i.v.) holds an excellent tumor targeting ability, shows high resolution and sensitivity and provides real-time visual navigation in LM3 orthotopic and subcutaneous tumor models to guide the resection of tumors^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>HepG2 subcutaneous xenograft mice^[1]</td> </tr> <tr> <td>Dosage:</td> <td>20 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intravenous injection; 1.5 h later dissected tumors</td> </tr> <tr> <td>Result:</td> <td>Accumulated in the tumor after injection and produced stronger fluorescence intensity in</td> </tr> </table> | Animal Model: | HepG2 subcutaneous xenograft mice ^[1] | Dosage: | 20 mg/kg | Administration: | Intravenous injection; 1.5 h later dissected tumors | Result: | Accumulated in the tumor after injection and produced stronger fluorescence intensity in |
| Animal Model: | HepG2 subcutaneous xenograft mice ^[1] | | | | | | | | |
| Dosage: | 20 mg/kg | | | | | | | | |
| Administration: | Intravenous injection; 1.5 h later dissected tumors | | | | | | | | |
| Result: | Accumulated in the tumor after injection and produced stronger fluorescence intensity in | | | | | | | | |

tumor tissues.

Remained fluorescence signal longer than 1.5 h, and provided high-resolution images of the tumor tissues with a SNR of 4.32.

Animal Model: LM3 orthotopic mice^[1]

Dosage: 20 mg/kg

Administration: Intravenous injection

Result: Decreased gradually the fluorescence intensity in LM3 orthotopic liver tumors after administration, whereas tumor-to-skin fluorescence ratios increased due to high accumulation and low clearance in tumor tissues.

REFERENCES

[1]. Li Q, et al. Tumor-Targeting NIRF/MR Dual-Modal Molecular Imaging Probe for Surgery Navigation. Anal Chem. 2022 Aug 3.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA