研究简报

¹⁸F-FLT的制备及其microPET显像

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摘要 [摘要] 本文制备了增殖显像剂18F-FLT,考察其稳定性及研究其在肿瘤模型鼠的microPET显像。本文以3-N-t-叔丁氧羰基-1-[5'-O-(4,4'-二甲氧基三苯甲基)-2'-脱氧-3'-O-(4-硝基苯磺酰基-β-1)-苏戊呋喃糖]胸腺嘧啶脱氧核苷(N-BOC-FLT)为标记前体进行氟代亲核置换反应,用HPLC检测放射化学纯度(RCP),进行稳定性研究和正常小鼠体内分布试验和肿瘤模型鼠microPET显像;研究结果显示RCP〉95%,6h内稳定,正常小鼠体内分布显示,在60min时,肾,脾,肠摄取较多,心,肝,肺,膀胱摄取次之;肿瘤模型鼠microPET显像能够清晰地观察到接种部位的放射性浓聚。

关键词 制备,18F-FLT,肿瘤,microPET

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Synthesis and MicroPET image of 18F-FLT

Abstract Abstract Synthesis, stability study and MicroPET image of proliferation imaging agent 3'-deoxy-3'-[18F] fluorothymidine (18F-FLT) were reported. Nucleophilic substitution of fluo ro replacement reaction was proceeded with N-BOC-FLT as labeling precursor. Radiochemical purity(RCP)was determined by high pressure liquid chromatography(HPLC). Biodistribufion was performed in mice. RCP determined by HPLC was over 95% and were stable within 6 h. Biodistribufion studies in mice showed that the uptake of 18F-FLT in kidney, spleen and intestine was higher than that of 18F-FLT in heart, liver, lung and bladder at 60min postinjection. MicroPET image of tumor in nude mice bearing tumor xenografts was clear.

Key words Synthesis, 18F-FLT, tumor, MicroPET

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