

## عنوان مقاله:

Evaluation of 99m Tc-MccJ25 peptide analog in mice bearing B16F10 melanoma tumor as a diagnostic radiotracer

## محل انتشار:

مجله پزشکی هسته ای و زیست شناسی آسیا اقیانوسیه, دوره 7, شماره 2 (سال: 1398)

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## خلاصه مقاله:

**Objective(s):** Despite recent advances in treatment modalities, cancer remains a major source of morbidity and mortality throughout the world. Currently, the development of sensitive and specific molecular imaging probes for early diagnosis of cancer is still a problematic challenge. Previous studies have been shown that some of the antimicrobial peptides (AMPs) exhibit a broad spectrum of cytotoxic activity against cancerous cells in addition to their antimicrobial activities. MicrocinJ25 (MccJ25) is an antimicrobial peptide that is produced by *Escherichia coli* (E. coli) strain. The aim of this study was to investigate the potential of a new peptide radiopharmaceutical derived from MccJ25 for diagnosis of melanoma tumor bearing C57BL/6 mice. **Methods:** A 14 amino acid analog of MccJ25 was labeled with technetium-99m (99mTc) through hydrazinonicotinamide (HYNIC) chelator and tricine as coligand. In vivo tumor uptake and tissue distribution were evaluated. The in vivo biodistribution studies were determined in C57BL/6 mice bearing B16F10 tumor. **Results:** The amount of non-peptide related 99mTc-impurities that measured by thin layer chromatography (TLC) did not exceed 5% of the total radioactivity. The in vitro binding to B16F10 cells was  $30.73 \pm 0.9\%$  after 1 h incubation at 37°C, and saturation binding experiments showed good affinity for radio-complex ( $K_d=47.98 \pm 6.25$  nM). The melanoma tumor was clearly visible up 1 h post-injection by gamma camera imaging. **Conclusion:** The results showed that 99mTc-labeled peptide could be a promising candidate as a targeting radiopharmaceutical for melanoma tumor imaging in mice.

## کلمات کلیدی:

99m Tc-MccJ25, Radiotracer, B16F10, Tumor

## لینک ثابت مقاله در پایگاه سیویلیکا:

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