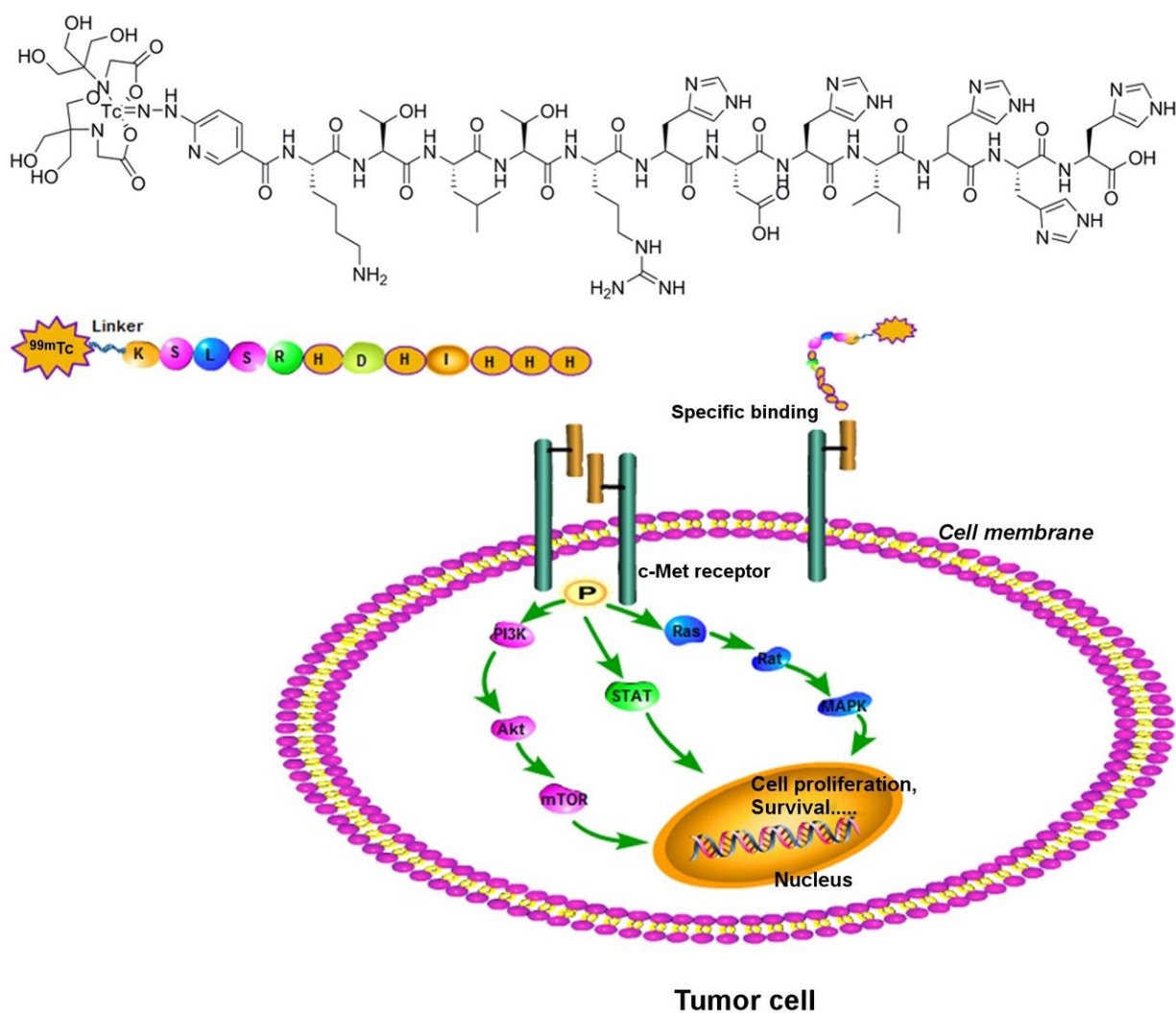


Novel tracer developed for precision targeting of non-small cell lung cancer

November 6 2018



The schematic diagram illustrates the process of radiochemical synthesis and molecular structure of 99mTc-HYNIC-cMBP (top) and the specific binding of the tracer to c-Met receptor (bottom). Credit: Zhaoguo Han, Yadi Xiao, Kai

Wang, Ji Yan, Zunyu Xiao, Fang Fang, Zhongnan Jin, Yang Liu and Baozhong Shen, Harbin Medical University, Harbin, Heilongjiang, China

Researchers have developed a new nuclear medicine tracer that could improve diagnosis and treatment of non-small cell lung cancer. Research published in the November issue of *The Journal of Nuclear Medicine* found that the new tracer, ^{99m}Tc -HYNIC-cMBP, produced clearer images in less time than currently used tracers and was more rapidly eliminated from the body, reducing radiation exposure.

The c-Met receptor—a promising therapeutic target for this disease—is a tyrosine kinase receptor that promotes the growth of cancer [cells](#), particularly in NSCLC. When c-Met receptors are elevated, cells may be resistant to EGFR-targeted therapy. However, with precise targeting of the c-Met receptor, physicians may be able to more effectively treat this disease.

In the study, the novel radiotracer was developed and cultured in two NSCLC cell lines, H1993 (high c-Met expression) and H1299 (no c-Met expression), and the activity and uptake of the ^{99m}Tc -HYNIC-cMBP tracer was assessed. These [tumor](#) cells were then transplanted into mice, and in vivo tumor specificity was measured by SPECT at various intervals after injection of the tracer. Blocking assays, biodistribution and autoradiography also were conducted to determine the tracer's specificity.

The authors found the preparation of the ^{99m}Tc -HYNIC-cMBP imaging agent to be simple, providing a high yield of the tracer with few associated costs. Biodistribution and autoradiography showed significantly higher accumulation of ^{99m}Tc -HYNICcMBP in H1993 tumors than in H1299 tumors. In addition, the H1993 tumors were

clearly visualized after only a half hour in SPECT images, while the H1299 tumors were not observed at any time. The tracer was also cleared rapidly, which is favorable for reducing exposure to radiation, background noise and long delays between treatment and imaging readout.

"To the best of our knowledge, there are no other c-Met-targeted imaging studies in NSCLC using peptide-based radiotracer to date," noted Baozhong Shen, MD, Ph.D., professor of medical imaging and nuclear medicine and the first president of the Fourth Hospital of Harbin Medical University. "Our research results confirmed the feasibility of detecting c-Met expression in NSCLC with the novel SPECT tracer. Upon further optimization, the SPECT tracer may be translated into clinical use for screening and monitoring therapeutic response in NSCLC patients."

More information: Zhaoguo Han et al, Development of a SPECT Tracer to Image c-Met Expression in a Xenograft Model of Non–Small Cell Lung Cancer, *Journal of Nuclear Medicine* (2018). DOI: [10.2967/jnumed.117.206730](https://doi.org/10.2967/jnumed.117.206730)

Provided by Society of Nuclear Medicine and Molecular Imaging

Citation: Novel tracer developed for precision targeting of non-small cell lung cancer (2018, November 6) retrieved 6 May 2024 from <https://medicalxpress.com/news/2018-11-tracer-precision-non-small-cell-lung.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.
