A Novel RGD-Containing Cyclic Peptide for use in Cancer Imaging and as a Targeted-Therapy Ligand

Tech ID: 20851 / UC Case 2010-162-0

ABSTRACT

Integrin plays a key role in the angiogenesis and metastasis of human tumors. avß3 integrin binding ligands have value in cancer diagnostic imaging and targeted therapy. The RGD motif binds to several integrins, including avß3, aIßb3, avß5, and aßß1. It is known that amino acids lateral to RGD affect RGD binding specificity to different integrins. Researchers at the University of California Davis have discovered a novel RGD-containing peptide useful in cancer imaging and as a targeted-therapy ligand.

FULL DESCRIPTION

The compound has the ability to bind to target avß3 integrin on tumor cells and neovasculatures and can be used in tumor diagnostic imaging and therapy. This compound outperforms existing commonly used RGD ligands in both targeting efficacy and lower non-specific binding to normal organ tissues. The compound can be easily functionalized to conjugate imaging payload without decreasing binding strength.

- The compound bound avß3 integrin as evidenced by stained glioblastoma U-87 MG cells and melanoma A375M cells. The binding of the compound to said cells was blocked by anti-avß3 antibody.
- The compound specifically targeted tumor xenografts on nude mouse models. Six hours after injecting a biotinylated form of the compound coupled to Streptavidin-Cy5.5 into tail vein of nude mice implanted with U-87 MG xenograft, imaging displayed preferential uptake of biotinylated compound in the tumor. Experimental design was replicated for nude mice implanted with A375M xenograft, with similar results.
- Tumor uptake was higher and liver uptake was lower in mouse studies, compared to three commonly used RGD ligands.

APPLICATIONS

- Targeting reagent against avß3 integrin.
- Payload carrier (such as radioisotype or drug) for tumor radioimaging, radiotherapy, and targeting chemotherapy.

FEATURES/BENEFITS

- The compound shows remarkable positive binding with avß3; very weak cross-reaction with aIßb3; and no binding with a1, a2, a3, a4, a5, aß, and aß9.
- In contrast to other RGD-containing peptides, the incorporation of payloads (labeling groups, etc.) at the C-terminus of this novel compound does not affect the binding affinity of the ligand to the integrin.
- Tumor uptake is higher and liver uptake is lower, compared to other existing RGD ligands.

RELATED MATERIALS

- Lam, et al. 2010. The Use of One-Bead One-Compound Combinatorial Library Technology to Discover High-Affinity avß3 Integrin and Cancer Targeting Arginine-Glycine-Aspartic Acid Ligands with a Built-in Handle. Mol Cancer Ther. October 2010 9:2714-2723; Published Online First September 21, 2010; doi:10.1158/1535-7163.MCT-10-0308 - 09/21/2010

PATENT STATUS

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RELATED TECHNOLOGIES

- Ligands for Alpha-4-Beta-1 Integrin
- Three-Dimensional Cell Adhesion Matrix

ADDITIONAL TECHNOLOGIES BY THESE INVENTORS

- Ligands for Alpha-4-Beta-1 Integrin
- Three-Dimensional Cell Adhesion Matrix
- Functional Illumination in Living Cells
- Nanoparticles for Drug Delivery, Tissue Targeting and Imaging Analysis
- Nanoporphyrin Nanoparticles for Combination Phototherapy and Drug Delivery to Infantile Hemangiomas
- Early Detection of Ovarian Cancer Using Markers to Short Chain Carbohydrates
- PVA Nanocarrier System for Controlled Drug Delivery
- Site-Specific Ligation and Compound Conjugation to Existing Antibodies

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Ligands for Improved Angiogenesis and Endothelialization of Blood Contacting Devices
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DM1-Conjugated Nanotherapeutic Agent for Cancer Treatment
Engineered Biomaterial to Prevent Endothelial Inflammation