

Integrins are cell surface proteins that bind components of the extracellular matrix and play a critical role in facilitating tumor growth and metastasis. Many solid tumors express a variety of these integrins, making integrins a prime target for anti-cancer therapy. PIP-LYTAC molecules work by recognizing a variety of integrins, including $\alpha 1$, $\alpha 3$, $\alpha 5$, $\alpha 6$, and $\beta 1$ integrins, on the cell surface. The molecule can then link the engaged integrin to a lysosome targeting receptor and trigger its internalization and degradation. Removing integrins from cancer cells could slow tumor progression and prevent metastasis. The PIP-LYTAC can also be tailored to target organ- or tissue-specific lysosomal targeting receptors, allowing this molecule to be functionalized for specific tumor types. Integrins are also involved in viral infections, making PIP-LYTACs a potential antiviral therapy.

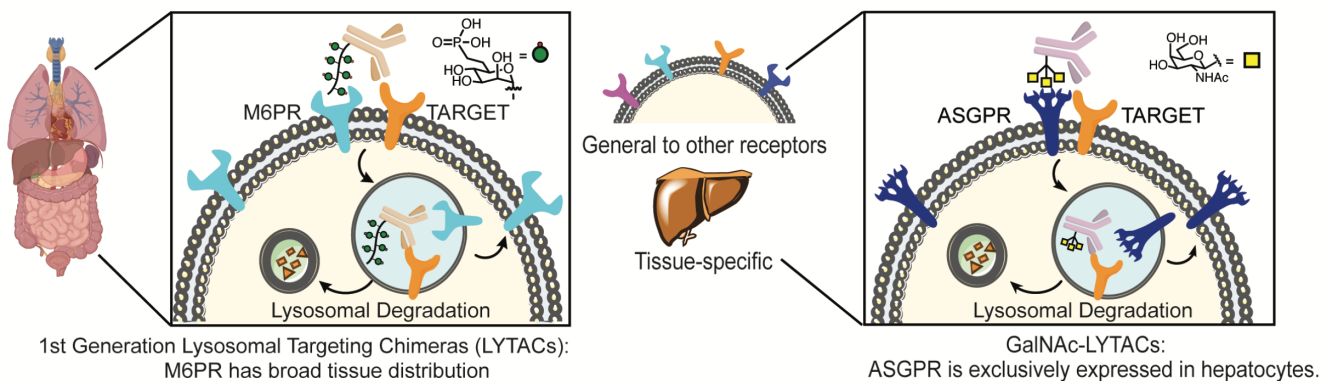


Figure: LYTACs can hijack the asialoglycoprotein receptor (ASPGR) for targeted and cell-specific protein degradation. LYTACs targeting M6PR can degrade broadly expressed targets while LYTACs targeting ASPGR can degrade targets specifically on hepatocytes. Courtesy of Carolyn Bertozzi and Jennifer Cochran.

Related Technology: [Stanford Docket 06-356: "Integrin Binding Peptides Based On Knottin Scaffolds"](#)

Stanford Technology 18-179: "Lysosome targeting chimeras (LYTACs)"

Applications

- Cancer therapeutic for integrin-overexpressing cancers
- Antiviral therapy

Advantages

- Bifunctional: single protein can both target the integrin and trigger the degradation pathway
- Can be tailored for tissue- or organ-specific targeting
- Alternative to antibody approach

Publications

- Ahn, G., Banik, S., Miller, C.L., et al. ["Lysosome Targeting Chimeras \(LYTACs\) That Engage a Liver-Specific Asialoglycoprotein Receptor for Targeted Protein Degradation."](#) ChemRxiv. Preprint. (2020).

Patents

- Published Application: [20230211001](#)

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