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Putar Video Sex Luna Maya Arielcom Putar Video Sex Luna Maya Arielcom Targeting cell-specific proteins or molecular markers to cell membranes has generally been achieved by attaching the protein or marker to a lipid or lipid-based molecule, and is commonly referred to as “lipid targeting.” In certain circumstances, however, it is desirable to attach a protein or other molecular marker directly to the surface of a cell membrane. This may be desirable, for example, to provide cell markers for identification of specific cell types, and/or to allow pharmacologic intervention. For lipid targeting to be useful in achieving the above objectives, it is necessary for the lipid to be able to traverse the plasma membrane of the target cell, and for the lipid to be able to be cleaved or degraded at a rate that does not endanger the viability of the target cell. Proteins can be covalently attached to cell membranes by various means. This process is referred to as “protein targeting.” For example, a protein can be modified with hydrophilic polymers, such as polyethylene glycol (PEG), that “mask” the hydrophobic protein so that it is unlikely to interact with the lipid membrane. Alternatively, a synthetic vector can be used to attach the protein to a lipid by chemical means. Lipid targeting, however, has generally been more difficult and problematic. Previously, known lipid targeting systems have focused on the use of synthetic membrane penetration molecules (e.g., amphipathic peptides) and have not addressed the issue of lipids that are biologically compatible. More recently, lipid-based targeting has been achieved using cationic peptides and proteins (to form a so-called “cationic peptide lipoplex”). These compounds typically are constructed of a cationic lipid (e.g., DOTAP, DOTMA, DOTGA, or DOTMA) and a peptide or protein (e.g., a polylysine or a cationized polyanion such as poly-L-lysine). The principal purpose of the cationic lipid is to condense the peptide or protein into a charged lipid phase, thereby enabling the peptide or protein to traverse the cell membrane. However, certain difficulties and limitations exist with the use of cationic peptide-lipid formulations. For example, the cationic lipid and peptide/protein must have complementary charges, which generally limits the 595f342e71

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