

Features of Bio Membrane Structure and Cytoskeleton Organization

Kayla Kumawat^{*}

Department of Ecology and Evolutionary Biology, Monash University, Melbourne, Australia

DESCRIPTION

A biological membrane, sometimes called a cell membrane or a bio membrane, is a selectively permeable membrane that separates a cell's interior from its environment or creates intracellular compartments by serving as a wall between various cell regions. Biological membranes for eukaryotic cell membranes are made up of a phospholipid bilayer containing embedding, integral, and peripheral proteins that are used for communication as well as chemical and ion transport. For physiological function, proteins must be able to rotate and distribute laterally in a fluid matrix to the majority of lipid in a cell membrane. By possessing an annular lipid shell comprised of lipid on their surface, proteins are adapted to the high membrane fluidity of the lipid bilayer. The cell membranes are distinct from the isolating tissues formed up of layers of cells, such as mucous membranes, basement membranes, and serous membranes. One fascinating issue in membrane biophysics is the existence of lipid micro domains inside the plasma membrane known as rafts, which are thought to be essential for its complex activity. Recent experimental work has shown that biological membranes are composed of floating regions with different lipid and protein compositions rather than being laterally homogeneous. Contrary to popular belief, the present understanding of how a biological membrane is organised structurally still strongly relies on the fluid-mosaic model of a fluid-lipid bilayer put forward by Singer and Nicholson in 1972. It is still unclear exactly how these domains' exact composition and functional duties are determined at the molecular level. Furthermore, in addition to the random motion predicted by the fluid mosaic model, a number of lateral transport mechanisms exist for a variety of membrane proteins. Certain lipid mixtures, including cholesterol, have been found to include micro domains that show coexisting liquid phases when particular temperature, lateral pressure, and composition conditions are fulfilled. A few hundred lipid diameters (100 nm-200 nm) to one micrometre are the range of sizes of the membrane domains in this case.

The biological membrane system where the existence of lateral domains has now been definitively proven is the plasma membrane of mammalian cells. Raft domains in living cells appear to be quite small and most likely diverse. This may shed light on why they have managed to elude direct microscopic observation. It was feasible to demonstrate indirectly that small rafts exist by using single-particle tracking of the thermal position fluctuation, which showed that raft-associated membrane proteins are permanently attached to a small, cholesteroldependent lipid assembly of around 50 nm in diameter.

Lipids

The biological membrane is composed of lipids with hydrophilic heads and hydrophobic tails. The hydrophobic tails, which are hydrocarbon tails, are essential for identifying the cell because of their length and saturation. Lipid rafts are created when lipid species and proteins collect in membrane domains. They facilitate the division of membrane components into distinct zones that are involved in certain processes, such as signal transduction. Red blood cells, or erythrocytes, have unique lipid structures. The bilayer of red blood cells is composed of cholesterol and phospholipids in an equal weight ratio. The erythrocyte membrane plays a crucial role in blood coagulation. A part of the bilayer of red blood cells is phosphatidylserine. This usually takes place on the cytoplasmic side of the membrane. It is sent to the outer membrane to be used while blood clotting.

Proteins

Proteins of many types are present in phospholipid bilayers. These membrane proteins play a number of activities and exhibit a wide range of characteristics while catalysing various chemical processes. Integral proteins that bridge membranes have different domains on each side. Integral proteins are difficult to remove because they are firmly bonded to the lipid bilayer. They won't split naturally; only a chemical process that breaks the

Correspondence to: Kayla Kumawat, Department of Ecology and Evolutionary Biology, Monash University, Melbourne, Australia, E-mail: Kaylakumawat@gmail.com

Received: 22-Feb-2023, Manuscript No. BEG-23-20314; Editor assigned: 24-Feb-2023, PreQC No. BEG-23-20314 (PQ); Reviewed: 10-Mar-2023, QC No. BEG-23-20314; Revised: 17-Mar-2023, Manuscript No. BEG-23-20314 (R); Published: 27-Mar-2023, DOI: 10.35248/2167-7662.23.11.199

Citation: Kumawat K (2023) Features of Bio Membrane Structure and Cytoskeleton Organization J Bio Energetics. 11:199.

Copyright: © 2023 Kumawat K. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

membrane would do that. Peripheral proteins are different from integral proteins in that they have weaker ties to the surface of the bilayer and can easily split from the membrane. Peripheral proteins, which are only located on one side of a membrane, are what generate membrane asymmetry.

Oligosaccharides

Sugar is a component of oligosaccharide polymers. They may establish covalent bonds with proteins or lipids to create glycoproteins or glycolipids in the membrane. Membranes include glycolipids, or lipid molecules that contain sugar. The sugar groups of glycolipids are exposed at the bilayer's cell surface and are capable of forming hydrogen bonds. Glycolipids offer the clearest example of asymmetry in the lipid bilayer. One of the many communication-related functions that glycolipids do in the biological membrane is cell-cell adhesion, which is just one of the many duties they perform. Glycoproteins are examples of integral proteins. They are essential for the defence and reaction of the immune system.