

## In this issue

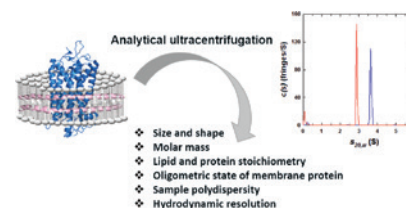
Sayaka Inagaki and Rodolfo Ghirlando

### Nanodisc characterization by analytical ultracentrifugation

DOI 10.1515/ntrev-2016-0082  
Nanotechnol Rev 2017; 6(1): 3–14

**Review:** Analytical ultracentrifugation can characterize nanodisc preparations, reporting on their size, shape, molar mass, and polydispersity in a straightforward manner without sample modification.

**Keywords:** analytical ultracentrifugation; lipid; membrane protein; nanodisc.



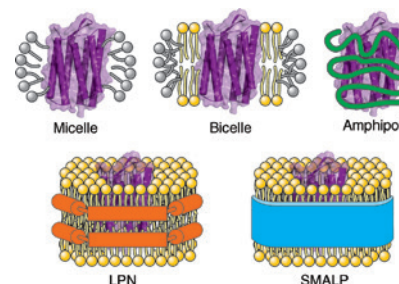
Konstantin S. Mineev and Kirill D. Nadezhdin

### Membrane mimetics for solution NMR studies of membrane proteins

DOI 10.1515/ntrev-2016-0074  
Nanotechnol Rev 2017; 6(1): 15–32

**Review:** Review on the properties, history of implementation, and peculiarities of membrane mimetics that are conventionally used in solution NMR studies of membrane proteins.

**Keywords:** bicelles; membrane mimetics; micelles; nanodiscs; NMR.



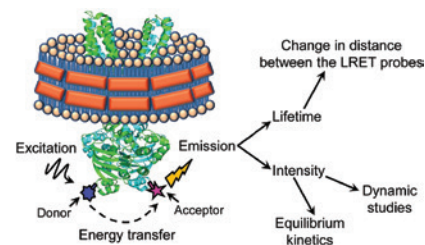
Maria E. Zoghbi and Guillermo A. Altenberg

### Membrane protein reconstitution in nanodiscs for luminescence spectroscopy studies

DOI 10.1515/ntrev-2016-0078  
Nanotechnol Rev 2017; 6(1): 33–46

**Review:** Nanodiscs are ideal platforms for spectroscopic studies of membrane proteins in a near-physiologic environment.

**Keywords:** ATP-binding cassette; LRET; luminescence resonance energy transfer; MsbA; multidrug resistance.



Kushal Sejwal, Mohamed Chami,  
Paul Baumgartner, Julia Kowal,  
Shirley A. Müller and Henning  
Stahlberg

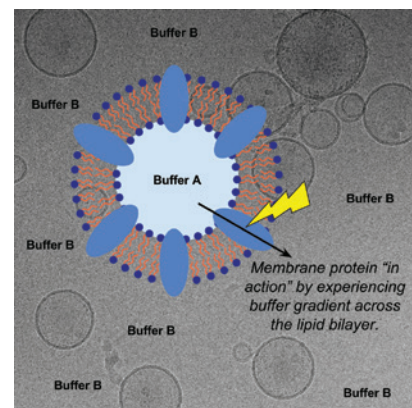
**Proteoliposomes – a system to study  
membrane proteins under buffer  
gradients by cryo-EM**

DOI 10.1515/ntrev-2016-0081

Nanotechnol Rev 2017; 6(1): 57–74

**Review:** Proteoliposomes are lipidic vesicles with reconstituted membrane proteins in their lipid bilayer. These proteoliposomes can be imaged by cryo-electron microscopy, as shown in this figure, allowing the structural determination of the membrane proteins in the membranes. Preparing proteoliposomes in buffer A and exchanging the outer buffer to buffer B allow the setting up of a buffer gradient across the membranes, thereby establishing a salt, pH, or ion concentration gradient, or even electric voltage.

**Keywords:** buffer gradient; cryo-electron microscopy; image processing; membrane proteins; proteoliposomes.



Elka R. Georgieva

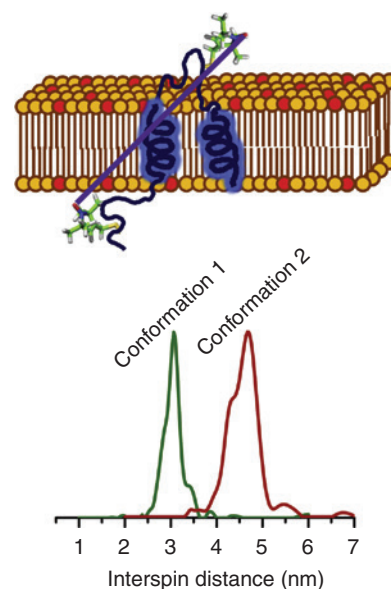
**Nanoscale lipid membrane mimetics  
in spin-labeling and electron para-  
magnetic resonance spectroscopy  
studies of protein structure and  
function**

DOI 10.1515/ntrev-2016-0080

Nanotechnol Rev 2017; 6(1): 75–92

**Review:** Described are applications of phospholipid-based nanosized membrane mimetics, such as liposomes, bicelles and nanodiscs, in conjunction with the powerful biophysical technique electron paramagnetic resonance in study of protein structure and function.

**Keywords:** bicelles; electron paramagnetic resonance spectroscopy; liposomes; membrane protein structure and function; nanodiscs.



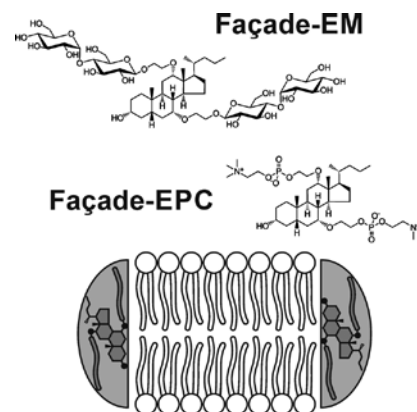
Konstantin S. Mineev, Kirill D. Nadezhdin, Sergey A. Goncharuk and Alexander S. Arseniev

**Façade detergents as bicelle rim-forming agents for solution NMR spectroscopy**

DOI 10.1515/ntrev-2016-0069  
Nanotechnol Rev 2017; 6(1): 93–103

**Research highlight:** Façade-EM and Façade-EPC were implemented as rim-forming agents for small isotropic bicelles for structural studies by solution NMR spectroscopy.

**Keywords:** bicelles; Façade detergents; membrane protein; NMR spectroscopy.



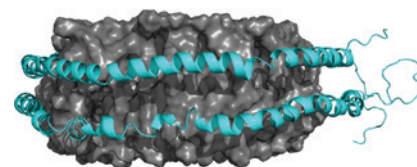
Cédric Eichmann, Stefan Bibow and Roland Riek

**$\alpha$ -Synuclein lipoprotein nanoparticles**

DOI 10.1515/ntrev-2016-0062  
Nanotechnol Rev 2017; 6(1): 105–110

**Review:** The protein  $\alpha$ -synuclein plays a pivotal role in Parkinson's disease. It shares a high sequence similarity with apolipoproteins, and in the presence of phospholipids  $\alpha$ -synuclein can form apolipoprotein nanodisc-type particles by a disorder-to-helix structural change.

**Keywords:** lipoprotein; nanodiscs; NMR; Parkinson's disease;  $\alpha$ -synuclein.

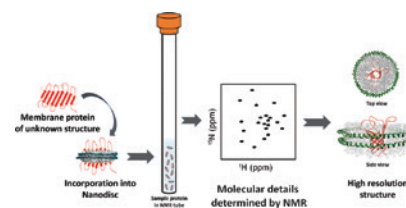


Robbins Puthenveetil, Khiem Nguyen and Olga Vinogradova  
**Nanodiscs and solution NMR: preparation, application and challenges**

DOI 10.1515/ntrev-2016-0076  
Nanotechnol Rev 2017; 6(1): 111–125

**Review:** Structure determination of membrane proteins in nanodiscs using solution NMR.

**Keywords:** beta barrel; integrin; membrane proteins; nanodisc; nanoscale phospholipid bilayers; saposin-A; SMALP; solution NMR; styrene maleic acid; transmembrane.



Svetla Stoilova-McPhie

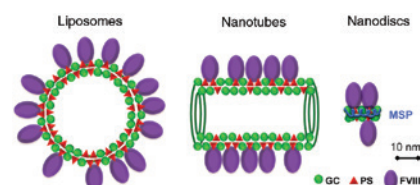
**Lipid nanotechnologies for structural studies of membrane-associated clotting proteins by cryo-electron microscopy**

DOI 10.1515/ntrev-2016-0066

Nanotechnol Rev 2017; 6(1): 127–137

**Review:** Blood coagulation factor VIII (FVIII) binds specifically to phosphatidylserine (PS) containing lipid membranes. Liposomes are single bilayer lipid vesicles. Nanotubes are single bilayer lipid nanotubes self-assembled from galactosylceramide lipids (GC). Nanodiscs are lipid bilayer patches held together by membrane scaffolding proteins (MSPs). The macromolecules and nanotechnologies are drawn to scale.

**Keywords:** blood coagulation factors; cryo-electron microscopy; lipid nanotechnologies; macromolecular structure; membrane-associated proteins.



Kirill Grushin, Mark Andrew White and Svetla Stoilova-McPhie  
**Reversible stacking of lipid nanodiscs for structural studies of clotting factors**

DOI 10.1515/ntrev-2016-0073

Nanotechnol Rev 2017; 6(1): 139–148

**Research highlight:** Electron micrograph of negatively stained Y-branched and side-attached ND stacks formed after a 60-min incubation in the presence of 10 mM  $\text{CaCl}_2$ . The insets show short ND stacks of two and three ND formed at less than 5 mM  $\text{CaCl}_2$ . The number of ND stacks per micrograph and the distribution of the ND stacks by lengths at different  $\text{CaCl}_2$  concentrations are shown on the graph.

**Keywords:** cryo-electron microscopy; lipid nanodiscs; phospholipids; stacking of nanodiscs.

