

OptiPrep™ Mini-Review MS09

Resolution of soluble cytosolic proteins from membrane vesicles and organelles: a bibliography

There are three Application Sheets listed in the Application Sheet Index under “Protein localization (membrane *versus* cytosol)” which describe different gradient strategies; they can be found on the OptiPrep™ Applications flash drive or accessed via the following website www.axis-shield-density-gradient-media.com (click on “Methodology”, then “Organelles and subcellular membranes” and follow the links from the Index):

- ◆ Discontinuous gradient: Application Sheet S35
- ◆ Self-generated gradient: Application Sheet S36
- ◆ A special strategy for rapid resolution of protein complexes and cytosol: Application Sheet S37
- ◆ Note that the resolution of mammalian cell exosomes and other microvesicles from soluble proteins is covered in Mini-Review MS17 and the similar resolution of bacterial and fungal microvesicles in Mini-Review MS16.

The reference list, which follows, includes principally papers describing the separation of membranes and soluble (cytosolic) proteins (**Section 1**); it is divided alphabetically into source material (**cell or tissue type**). It includes both mammalian and non-mammalian sources and in each of the 25 sections, references are listed alphabetically according to first author.

Some papers report the study of previously prepared subcellular membranes to determine the distribution of a particular protein between the soluble fraction and the organelle(s). Others papers describe the separation of vesicles either budded from the cells or obtained from permeabilized cells. These are listed in **Section 2**. In some cases gradients also resolve lipid droplets.

Because a significant number of papers use the methodology for the study of virus processing, these may be listed both in the main cell/tissue references (**Section 1**) and in **Section 3**, which is devoted solely to virus processing.

- ◆ To facilitate identification of references of scientific interest **key words in titles are highlighted in light blue**.

1. Cells or tissues

1.1. Algae

Wood, C.R. and Rosenbaum, J.L. (2014) *Proteins of the ciliary axoneme are found on cytoplasmic membrane vesicles during growth of cilia* Curr. Biol., **24**, 1114-1120

1.2. Bacteria

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Valent, Q.A., Scotti, P.A., High, S., de Gier, J-W.L., et al (1998) *The Escherichia coli SRP and SecB targeting pathways converge at the translocon* EMBO J., **17**, 2504-2512

1.3. Brain

Ding, T.T., Lee, S-J., Rochet, J-C. and Lansbury, Jr., P.T. (2002) *Annular α -synuclein protofibrils are produced when spherical protofibrils are incubated in solution or bound to brain-derived membranes* Biochemistry, **41**, 10209-10217

Wang, X., Bowers, S.L., Wang, F., Pu, X-a., et al (2009) *Cytoplasmic prion protein induces forebrain neurotoxicity* Biochim. Biophys. Acta **1792**, 555–563

1.4. Carcinoma cells/tissues

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Jorgensen, I., Bednar, M.M., Amin, V., Davis, B.K., et al (2011) *The Chlamydia protease CPAF regulates host and bacterial proteins to maintain pathogen vacuole integrity and promote virulence* Cell Host Microbe, **10**, 21–32

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Lee, S.M., Olzmann, J.A., Chin, L-S. and Li, L. (2011) *Mutations associated with Charcot–Marie–Tooth disease cause SIMPLE protein mislocalization and degradation by the proteasome and aggresome–autophagy pathways* J. Cell Sci., **124**, 3319–3331

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Mira, E., Lacalle, R.A., Buesa, J.M., Gonzalez de Buitrago, G., et al (2004) *Secreted MMP9 promotes angiogenesis more efficiently than constitutive active MMP9 bound to the tumor cell surface* J. Cell Sci., **117**, 1847-1856

Murillo, A., Vera-Estrella, R., Barkla, B.J., Méndez, E. and Arias, C.F. (2015) *Identification of host cell factors associated with astrovirus replication in Caco-2 cells* J. Virol., **89**, 10359–10370

Salman, E.D., He, D., Runge-Morris, M., Kocarek, T.A., et al (2011) *Site-directed mutagenesis of human cytosolic sulfotransferase (SULT) 2B1b to phospho-mimetic Ser348Asp results in an isoform with increased catalytic activity* J. Steroid Biochem. Mol. Biol., **127**, 315– 323

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1.5. CHO cells

Lin, C-C., Love, H.D., Gushue, J.N., Bergeron, J.J.M., et al (1999) *ER/Golgi intermediates acquire Golgi enzymes by Bredelfin A – sensitive retrograde transport in vitro* J. Cell. Biol., **147**, 1457-1472

Love, H.D., Lin, C.C., Short, C.S. and Ostermann, J. (1998) *Isolation of functional Golgi-derived vesicles with a possible role in retrograde transport* J. Cell Biol., **140**, 541-551

1.6. COS cells

Dicu, A.O., Topham, M.K., Ottaway, L. Epanand, R.M. (2007) *Role of the hydrophobic segment of diacylglycerol kinase ϵ* Biochemistry, **46**, 6109-6117

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1.13. Kidney proximal tubule cells (incl. LLC-PK1)

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Nürnberg, J., Bacallao, R.L. and Phillips, C.P. (2002) *Inversin forms a complex with catenins and N-cadherin in polarized epithelial cells* Mol. Biol. Cell, **13**, 3096-3106

1.14. Liver

Nakatsuka, A., Wada, J., Iseda, I., Teshigawara, S., et al (2012) *Vaspin is an adipokine ameliorating ER stress in obesity as a ligand for cell-surface GRP78/MTJ-1 complex* Diabetes, **61**, 2823–2832

1.15. MDCK cells

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1.17. Mouse embryo fibroblasts

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1.20 Pheochromocytoma (PC12) cells

Thayanidhi, N., Liang, Y., Hasegawa, H., Nycz, D.C., et al (2012) *R-SNARE [ykt6](#) resides in membrane-associated protease-resistant protein particles and modulates [cell cycle progression](#) when over-expressed* Biol. Cell, **104**, 397-417

1.21 Plant cells

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1.22 Squid axons

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Du, L-L. and Novick, P. (2001) *Yeast [Rab GTPase-activating protein Gyp1p](#) localizes to the [Golgi apparatus](#) and is a negative regulator of [Ypt1p](#)* Mol. Biol. Cell, **12**, 1215-1226

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Ge, W., Chew, T.G., Wachtler, V., Naqvi, S.N., et al (2005) *The novel [fission yeast protein Pal1p](#) interacts with [Hip1-related Sla2p/End4p](#) and is involved in cellular morphogenesis* Mol. Biol. Cell, **16**, 4124-4138

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- Sciskala, B.** and Kölling, R. (2013) *Interaction maps of the Saccharomyces cerevisiae ESCRT-III protein Snf7* Eukaryot. Cell, **12**, 1538-1546
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2. Subcellular membranes

2.1 Golgi membranes

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2.2 Lysosomes/endosomes

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2.3 Vesicles (budded and from permeabilized cells)

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3. Virus processing

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