Name: Nullin Divecha (Professor in cellular signalling)

Education		
1982-1985	Bsc	Biochemistry Manchester University.
1985-1988	Ph.D	Biochemistry Sheffield University.

Positions held.

1988-1993	Postdoctoral fellow	Prof Rf.Irvine	The Babraham Institute
1993-1997	Babraham Fellow	Group leader	The Babraham Institute
1997-2003	Junior Group leader		The Netherlands Cancer Institute
2003-2007	Senior Group Leader		The Netherlands Cancer Institute
2007-2014	Senior Group Leader		The Paterson institute for cancer Res
2014-2016	Associate Professor.		Southampton University
2015-2016	EU Marie Curie fellow		INGM Milan
2016-	Professor in cell signalli	ing	University of Southampton

Prizes and awards and positions:

The Babraham Institute

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Positions of Esteem

2016-	Theme	lead for	molecular	and	cellular	Biosciences.
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- 2016- Senior member of the Research Strategy group in Biological Sciences
- 2018- Co-lead Cell biology in life sciences drive.
- 2021 member of UOS implementation of BBRSC initiatives

Conferences organised.

2007	Vice Chair	Gordon conference on nuclear signalling
2009	Co-Chair	Gordon conference on nuclear signalling
2011	Co-chair	Keystone Symposia: Inositide Signaling and
		PtdIns3P Kinases

Invited international Conference seminars (2007-).

- 2007 Gordon conference on nuclear signalling
- 2007 Babraham Institute Signal transduction meeting

- 2007 GRC "Signal Transduction within the Nucleus, CA, USA; Janilia Conferences, WashingtonDC, USA
- 2009 Gordon conference on nuclear signalling
- 2009 FEBS Lipid signalling in disease Ortona, Italy
- 2009 Mt.St. Odile 34th Symposium Inositide metabolism, France
- 2010 Japanese biochemical society meeting BM2010 (Tokyo) and DGK satellite meeting (Kobe)
- 2010 Advances in enzyme regulation (Bologna)
- 2011 Gordon conference on nuclear signalling, Ventura
- 2011 Keystone symposia Inositide signalling (USA)
- 2012 Faseb: Lipid signalling, Vermont
- 2012 Advances in Biological regulation (Bologna)
- 2012 FEBS Lipid Signalling (Vico Equensa, italy)
- 2012 Inositide signalling in health and disease (Coorg Bangalore India)
- 2014 European Symposium on Hormones and Cell Regulation (mt st Odile Strasbourg)
- 2015 ASBMB (Boston USA)
- 2015 Biochemical society lipid signalling meeting UK
- 2015 Advances in biological regulation: Bologna
- 2015 Chromatin Signalling Keynote speaker: London chromatin club
- 2016 FASEB: phospholipid Signaling in Cancer, neurodegeneration and cardiovascular
- 2017 Special ABR symposium: Ulsan korea
- 2019 Advances in biological regulation: Bologna

Publications.

- 1. Andrews, D.M., et al., *Identification and optimization of a novel series of selective PIP5K inhibitors.* **Bioorg Med Chem**, 2021. **54**: p. 116557.
- 2. Poli, A., et al., *PIP4Ks impact on PI3K, FOXP3, and UHRF1 signaling and modulate human regulatory T cell proliferation and immunosuppressive activity.* **Proc Natl Acad Sci** U S A, 2021. **118**(31).
- 3. Ratti, S., et al., "Modulating Phosphoinositide Profiles as a Roadmap for Treatment in Acute Myeloid Leukemia". Front Oncol, 2021. 11: p. 678824.
- 4. Poli, A., et al., *Exploring the controversial role of PI3K signalling in CD4*(+) *regulatory T (T-Reg) cells.* Adv Biol Regul, 2020. 76: p. 100722.
- 5. Bowler, E.H., et al., *Deep proteomic analysis of Dnmt1 mutant/hypomorphic colorectal cancer cells reveals dysregulation of epithelial-mesenchymal transition and subcellular re-localization of Beta-Catenin.* Epigenetics, 2020. 15(1-2): p. 107-121.
- 6. Bowler, E.H., et al., *Proteomic Analysis of Azacitidine-Induced Degradation Profiles Identifies Multiple Chromatin and Epigenetic Regulators Including Uhrf1 and Dnmt1 as Sensitive to Azacitidine.* **J Proteome Res**, 2019. **18**(3): p. 1032-1042.
- 7. Poli, A., et al., *Phosphatidylinositol 5 Phosphate (PI5P): From Behind the Scenes to the Front (Nuclear) Stage.* Int J Mol Sci, 2019. 20(9).
- 8. Fiume, R., et al., *Nuclear Phosphoinositides: Their Regulation and Roles in Nuclear Functions*. Int J Mol Sci, 2019. 20(12).
- 9. Lundquist, M.R., et al., *Phosphatidylinositol-5-Phosphate 4-Kinases Regulate Cellular Lipid Metabolism By Facilitating Autophagy*. **Mol Cell**, 2018. **70**(3): p. 531-544 e9.
- Stijf-Bultsma, Y., et al., *The basal transcription complex component TAF3 transduces changes in nuclear phosphoinositides into transcriptional output*. Mol Cell, 2015. 58(3): p. 453-67.
- Gelato, K.A., et al., Accessibility of different histone H3-binding domains of UHRF1 is allosterically regulated by phosphatidylinositol 5-phosphate. Mol Cell, 2014. 54(6): p. 905-19.