

## **Laboratory of Transcriptional Regulation**

Development and evolution, Pax genes, Wnt/\(\beta\)-catenin signalling, Tcf/Lef

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We are interested in development and evolution of development [evo-devo]. Our focus is on the role of transcription factors and signalling cascades, especially on the role of the Wnt/ $\beta$ -catenin signalling pathway and transcription factors of Pax and Tcf/Lef families. A combination of gain-of-function (transgenic) and loss-of-function (conditional knock-outs) approaches is used to study mammalian development using laboratory mouse as a model organism. In addition, cell culture approaches are used to study the role of the Tcf/Lef family during induced pluripotency. Our second main interest in the Laboratory is evolution of animal development. Several model systems including amphioxus, platynereis, fish and cnidarians are used in the laboratory to study various aspects of evo-devo.

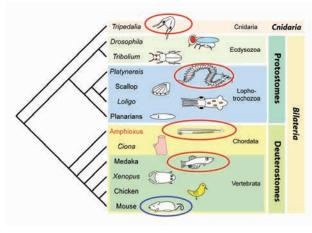


Fig. 1. Main model organisms used in the Laboratory of Transcriptional Regulation.

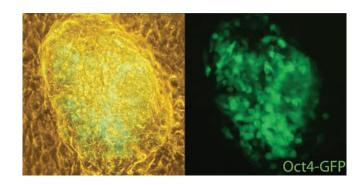
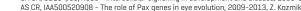


Fig. 2. Induction of pluripotent cells using PiggyBac-OSKM transposon. Activation of endogenous Oct4 is visualized by Oct4-GFP reporter.





GA CR, GCP305/10/J064 - Reconstructing urbilaterian photoreceptors: comparative study between Branchiostoma (Chordata) and Platynereis (Annelida), 2010-2013, Z. Kozmik

GA CR, GAP305/11/2198 - Genetics of mammalian eye development, 2011-2014, Z. Kozmik

Ministry of Education, Youth and Sports of the Czech Republic, LK11214 LK-NÁVRAT - Genetic regulation of embryonic development of the brain and the eye, 2012-2016, O. Machoň

Ministry of Education, Youth and Sports of the Czech Republic, LH12047 LH-KONTAKT II - The role of alternative splicing in evolution of vertebrate body plan, 2012-2015, Z. Kozmik

GA CR, GAP305/12/2042 – The role of transcription factors Tcf in the induced pluripotency and during neurogenesis, 2012-2015, 0. Machoň

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5. Waaler J, Machon O, von Kries JP, Wilson SR, Lundenes E, Wedlich D, Gradl D, Paulsen JE, Machonova O, Dembinski JL, Dinh H, Krauss S. Novel synthetic antagonists of canonical Wnt signaling inhibit colorectal cancer cell growth. Cancer Res 2011 71[1]:197-205





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<sup>4.</sup> Waaler J, Machon Q, Tumova L, Dinh H, Korinek V, Wilson SR, Paulsen JE, Pedersen NM, Eide TJ, Machonova O, Gradl D, Voronkov A, von Kries JP, Krauss S. A novel tankyrase inhibitor decreases canonical Wnt signaling in colon carcinoma cells and reduces tumor growth in conditional APC mutant mice. Cancer Res 2012 72[11]: 2822-2832.



## From the left:

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Not in the picture:
Ondřej Machoň, PhD / Research Fellow · Michal Koc, PhD / Postdoctoral Fellow (until 2012) · Jana Smolíková, PhD / Postdoctoral Fellow (until 2012) · Daniela Gurská, MSc / PhD Student (until 2012)