# PETER BRAZDA

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EDUCATION		
University of Debrecen, Debrecen, Hungary Ph.D., Cell Biology and Biophysics		9/2005 - 3/2015
University of Debrecen, Debrecen, Hungary M.Sc., Biotechnology		9/1999 - 7/2005

### RESEARCH EXPERIENCE

### Princess Máxima Center for Pediatric Oncology (PMC), Utrecht, The Netherlands

Postdoctoral research with Professor Henk Stunnenberg

6/2019 – present

- Project 1: Single cell transcriptomics of Pheochromocytoma
- Project 2: (Single cell) genomics in pediatric AML relapse
- Project 3: (Single cell) genomics of next-generation CAR-T cells

## Radboud University, Institute for Molecular Life Sciences (RIMLS), Nijmegen, The Netherlands

Postdoctoral research with Professor Henk Stunnenberg

6/2017 - 6/2019

- Project 1: Transcriptome profiling in Pheochromocytoma at single cell level
- Project 2: Transcriptomic analysis of early innate immune events after pertussis booster vaccination

### Delft University of Technology, Department of Bionanoscience, Delft, The Netherlands

Postdoctoral research with Professor Nynke Dekker

4/2015 - 5/2017

- Project 1: Accessory helicase (UvrD) action in *E. coli* replication-transcription conflicts with single molecule microscopy
- Project 2: The dynamics of *E. coli* replication termination (Tus) with super-resolution microscopy and large population genomics

# German Cancer Research Center (DKFZ), Biophysics of Macromolecules Division, Heidelberg, Germany

Guest researcher with Professor Jorg Langowski

4/2013 - 12/2014

• Several visits to test and carry out FCS, FCCS (fluorescence (cross-)correlation spectroscopy) and also single-plane illumination microscopy (SPIM-FCS) measurements on the setup designed and built by the group.

# **University of Debrecen, Department of Biochemistry and Molecular Biology, Debrecen, Hungary** *Graduate research* with Professor Laszlo Nagy and Dr Gyorgy Vamosi 9/2005 – 3/2015

- Thesis: Determination of dynamic properties of nuclear receptors
- Project 1: The mobility of retinoic acid receptor (RAR) during activation by fluorescence correlation spectroscopy (FCS)
- Project 2: The dynamic properties of retinoic-x receptor (RXR) during activation, chromatin association and coregulator binding as seen by single-plane illumination FCS (SPIM-FCS) and large population genomics (ChIP-seq)
- Project 3: The ligand-dependent dimerization (homo- and heterodimer) of RXR investigated by flow cytometry and single cell imaging

*Undergraduate research* with Professor Laszlo Nagy

12/2002 - 9/2005

- Thesis: Investigation of a novel retinoid-x receptor antagonist ligand (Best presentation in Natural Sciences at the National Conference of Scientific Students' Association (OTDK))
- Project 1: The study of a novel ligand on the dimerization- and transactivation ability of RXR with transient transfection assay

**PUBLICATIONS:** https://scholar.google.com/citations?user=vXZEPpEAAAAJ&hl=hu&oi=ao

#### LEADERSHIP EXPERIENCE

- International Genetically Engineered Machines (iGEM), tutor for the Debrecen team 2010-Boston, 2011-Amsterdam
- Researchers' Night, organizer at the University of Debrecen, 2008, 2009, 2011, 2012
- South-East European (SEE) Science Festival, organizer of local events, 2013
- Association of Biologist Students of the University of Debrecen, president, 2007

### **TEACHING ACTIVITIES**

- <u>Practical</u> in Biochemistry and Molecular Biology for medical students (Uni. of Debr.)
- Seminar in Molecular Biology for medical students. (Uni. of Debr.)
- Co-tutory support of 2 undergraduate students and support for 7 more students (Uni. of Debr.)
- Co-tutory support of 2 undergraduate students (TU Delft, RIMLS)

#### RESEARCH-RELATED SKILLS

RNA: isolation, RNA sequencing (bulk, single cell: CELseq2, 10x), scATACseq, RT-qPCR

DNA: isolation, molecular cloning, mutagenesis, chromatin IP

NGS: performing sequencing runs on Illumina (MiniSeq, Novaseq) platforms

**cell culturing**: tissue disintegration, transfection, cell-line related tasks, mammalian two-hybrid assay, bacterial strain engineering (recombination, transduction), processing of primary tissue samples

**analysis**: flow cytometry, confocal microscopy (imaging, FCS (fluorescence (cross)correlation spectroscopy, FRAP, FRET), fluorescence microscopy (wide field, super-resolution), single cell RNAseq data analysis, Rlanguage