



# Laboratory of Genomics and Bioinformatics

Genome analysis, cancer genomics, next-generation sequencing, transcriptome analysis

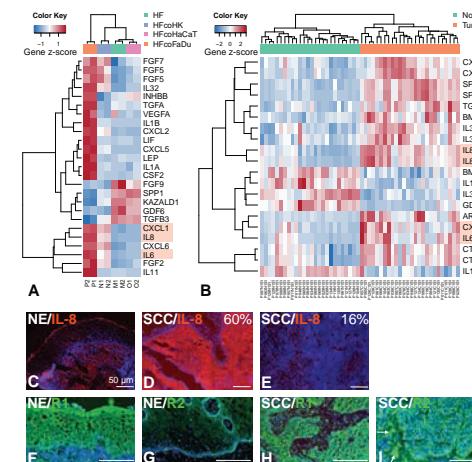
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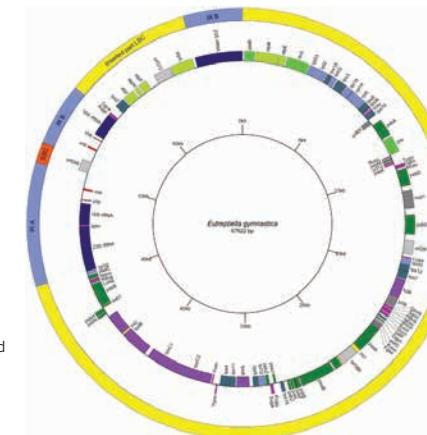
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Genome sequences are the ultimate source for phylogenomics. To understand the evolution of eukaryotes and the developmental processes that they regulate, it is necessary to analyse their genomes and transcriptomes. We sequenced transcriptomes of two multicellular eukaryotes, both of phylum Radiata, sweet water medusa *Craspedacusta sowerbyi* and cubozoan *Tripedalia cystophora*. The expression of selected developmental genes was characterized *in situ*. Single-cell eukaryotes [protists] with their branching close to the root of the evolutionary tree are the best candidates for genome studies. The availability of the genomic sequences will allow inferences to be made about the gene complement of the common eukaryotic ancestor. The main interest is also focused on endosymbiotic origin of two emblematic organelles of the eukaryotic cell, the mitochondrion and the plastid. Using next-generation sequencing platforms we characterize genomes and transcriptomes of many protist species, namely *Diplonema papillatum*, *Mastigamoeba balamuthi*, *Andalucia godoyi* and *Malawimonas*. Adding genome sequences from diverse protists to currently available eukaryotic genomes enables us to deduce, with a much higher accuracy, details of many steps and processes of the evolution of the eukaryotic cell. A second major topic of our group is directed towards cancer genomics. Using the Illumina microarray technique, we study intracellular interactions in malignant melanoma and identify markers

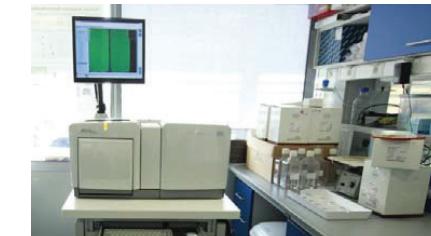
specific for different cancers. We also analyse the endogenous retroviral elements in human that are associated with cancers and explore the anticancer effects of statins.



**Fig. 1.** Expression changes of genes coding for paracrine factors in human fibroblasts cultured with the epithelial cells [A], comparison of tumour and normal tissues of clinical samples of squamous cell carcinoma [B], and expression of selected genes [IL8, CXCL1 and CXCL2] in human tissues visualized by immunofluorescence [C-I].



**Fig. 2.:** Map of the plastid genome of *Etreptiella gymistica*.



**Fig. 3.** Next-generation sequencer GS FLX (Roche).

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Not in the picture:

Jana Šáchová MSc / PhD Student [maternity leave]