

## PUBLICATIONS

### Emerging bioinformatics approaches for analysis of NGS-derived coding and non-coding RNAs in neurodegenerative diseases

Guffanti A., Simchovitz A. & Soreq A.  
Frontiers in Cellular Neuroscience 8 Article 89 (2014)

### BayMeth: improved DNA methylation quantification for affinity capture sequencing data using a flexible Bayesian approach

Riebler A., Menigatti M., Song, J. Z., Statham A.L., Stirzaker C., Mahmud N., Mein C. A., Clark S. J. & Robinson M. D. Genome Biology 15(2):R35 (2014)

### h5vc: scalable nucleotide tallies with HDF5

Pyl P. T., Gehring J., Fischer B. & Huber W. Bioinformatics  
Published online 5 Feb (2014)

### Cell-to-cell expression variability followed by signal reinforcement progressively segregates early mouse lineages

Ohnishi Y., Huber W., Tsumura A., Kang M., Xenopoulos P., Kurimoto K., Oleś A.K., Araúzo-Bravo M. J., Saitou M., Hadjantonakis A-K. & Hiiragi T.  
Nature Cell Biology 16:27-37 (2014)

### Improved variational Bayes inference for transcript expression estimation

Papastamoulis P., Hensman J., Glaus P. & Rattray M.  
Statistical Applications in Genetics and Molecular Biology 13(2):203-216 (2014)

### Accounting for technical noise in single-cell RNA-seq experiments

Brennecke P., Anders S., Kim J.K., Kołodziejczyk A.A., Zhang X., Proserpio V., Baving B., Benes V., Teichmann S.A, Marioni J.C, & Heisler M.G.  
Nature Methods 10:1093-1095 (2013)

### Hierarchical Bayesian modelling of gene expression time series across irregularly sampled replicates and clusters

Hensman J., Lawrence N. D. & Rattray M.  
BMC Bioinformatics 14:252 (2013)

### Count-based differential expression analysis of RNA sequencing data using R and Bioconductor

Anders S., McCarthy D. J., Chen Y., Okoniewski M., Smyth G. K., Huber W. & Robinson M. D.  
Nature Protocols 8:1765-1786 (2013)

### Drift and conservation of differential exon usage across tissues in primate species

Reyes A., Anders S., Weatheritt R.J., Gibson T. J., Steinmetz L. M., & Huber W.  
PNAS 110(38):15377-15382 (2013)

## The RADIANT Consortium

### Principal Investigator

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RADIANT is a research project funded by the 7th Framework Programme of the European Commission. The project started in December 2012 and brings together ten project partners from five European countries. The aim of the RADIANT project is to develop new statistical analysis tools for solving open problems in next generation sequencing (NGS) data analysis.

NGS is a rapidly evolving family of technologies with many applications, including genetics of both rare and common diseases, understanding disease mechanism progression. RADIANT aims to provide an integrated computational framework for NGS data analysis that is robust and user-friendly, and provide tools to benchmark experimental protocols and statistical methods. The project will establish training materials and an extensive training programme for the rapid dissemination of these new tools to the biomedical community.



<http://radiant-project.eu>

## PROJECT OUTPUT

RADIANT stands for Rapid Development and Distribution of Statistical Tools for High-Throughput Sequencing.

The project consists of work packages around three themes: Analysis, Integration and Dissemination.

### Analysis

WP1 Improved Read Mapping

WP2 Genomic Variations

WP3 Counts and Regions

WP4 Methylation Data Analysis

WP5 Transcriptome Expression

### Integration

WP6 Genome, Epigenome, Transcriptome

WP7 Systems Biology of Gene Regulation

WP8 Visualisation

### Dissemination

WP9 Scientific Publishing

WP10 Benchmarking

WP11 Training

For more information about the project visit the RADIANT website:  
<http://radiant-project.eu>

## SOFTWARE

### DESeq2

Differential gene expression analysis based on the negative binomial distribution

Estimate variance-mean dependence in count data from high-throughput sequencing assays and test for differential expression based on a model using the negative binomial distribution.

Authors: Love M., Anders S., & Huber W.

### BiocStyle

Provides standard formatting styles for Bioconductor documents.

Authors: Morgan M., Oleś A. & Huber, W.

### Repitools

Epigenomic tools for the analysis of enrichment-based epigenomic data. Features include summarization and visualization of epigenomic data across promoters according to gene expression context, finding regions of differential methylation/binding and BayMeth for quantifying methylation.

Authors: Robinson M., Strbenac D., Statham A., & Riebler A.

## Meetings/Training

### Computational statistics for genome biology, Brixen, Italy

RADIANT has been involved this intensive week-long course on current approaches in statistical and computational analysis of large-scale experiments. Members of the RADIANT team delivered lectures on the 2013 course, and will be doing so again on the course in June, 2014.

### RADIANT Workshop at ECCB'14, Strasbourg, France 6-7 September 2014

Analysis of differential isoform usage by RNA-seq: statistical methodologies and open software workshop. Bringing together leading researchers working on computational tools for exploring differential isoform usage to explore different approaches to method benchmarking.

### Genomic Workshop at MASAMB 2014

RADIANT sponsored workshop on mathematical and computational aspects of dealing with big data at the MASAMB (Mathematical and Statistical Aspects of Molecular Biology) workshop, April 2014.

### RADIANT 2nd General Meeting, January 2014

RADIANT held its 2nd General Meeting at the Institute Marie Curie, Paris on 9-10 January. Participants included members of the RADIANT consortium, Sasa Jenko, RADIANT EC Project Officer, members of the Scientific Advisory Board, and Peter Fraser, invited speaker from The Babraham Institute Cambridge.

### European Bioconductor Developers Meeting, December 2013

2013 European Bioconductor Developers Meeting, hosted by The University of Cambridge. The meeting was aimed at bioinformaticians, programmers and software engineers who contribute to the Bioconductor project, or are interested in developing packages for Bioconductor.

### SIG Frontiers in Somatic Variant Calling, Workshop at the EMBL Cancer Genomics Conference, November 2013

RADIANT sponsored workshop bringing together computational biologists directly involved in designing, benchmarking and applying somatic variant calling.