



Week 5 notes:

- Projects: some ideas
- Journal club
 - signups via pull request
 - Feedback forms
 - "Benchmarking"



Project ideas

- n.b.: 50% of mark
- As always, reproducing analyses from a paper or designing your own simulation to evaluate some methods is always a possibility
- Not urgent; I will put pressure on in a few weeks
- “Consulting” type possibilities:
 - Comparing fixed effects and mixed effects models for the paired comparison problem
 - Time course differential gene expression



33 signed up

| | | | | | |
|------------|-----------|--|---|--|--|
| 22.10.2018 | Mark | limma + friends | linear model simulation + design matrices | Averaged gene expressions for regression (AS, LB, MK) | Detection and accurate false discovery rate control of differentially methylated regions from whole genome bisulfite sequencing (DT, HP) |
| 29.10.2018 | Charlotte | hands-on session #1: RNA-seq | FASTQC/Salmon/etc. | Capturing Heterogeneity in Gene Expression Studies by Surrogate Variable Analysis (MS, CR) | X |
| 05.11.2018 | Mark | edgeR+friends 1 | basic edgeR/voom | Overcoming systematic errors caused by log-transformation of normalized single-cell RNA sequencing data (RB, RG) | |
| 12.11.2018 | Mark | edgeR+friends 2 | GLM/DEXSeq | A general and flexible method for signal extraction from single-cell RNA-seq data (AL, VL) | Integrating single-cell transcriptomic data across different conditions, technologies, and species (PV, FN, ES) |
| 19.11.2018 | Mark | single-cell dim. reduction + clustering; FDR | conquer | Normalization of RNA-seq data using factor analysis of control genes or samples (RM, JD, CV) | Diffusion maps for high-dimensional single-cell analysis of differentiation data (SP, GK) |
| 26.11.2018 | Lukas | hands-on session #2: cytometry | cytof null comparison | Epigenome-wide association studies without the need for cell-type composition (RL, SG) | X |
| 03.12.2018 | Hubert | classification | MLInterfaces | Predicting cell types in single cell mass cytometry data (CM, SS) | Bayesian Trees for Automated Cytometry Data Analysis (CB, XC) |
| 10.12.2018 | Mark | loose ends: HMM, EM, robustness | segmentation, peak finding | Differential expression analysis for sequence count data (AA, PS) | Visualizing Data using t-SNE (MJT, TB, MP) |
| 17.12.2018 | Mark | hands-on session #3: single-cell RNA-seq | full scRNA-seq pipeline | tba (SB,ST) | X |



Journal Club procedure

- Your job during journal clubs: give the presenters some constructive feedback
- n.b.: Please put your GitHub names on the first slide of journal club and/or add them to the Slack on the day.

STA426 Journal Club Feedback Form

Please give your classmates some constructive feedback on their presentations. Note that several of the questions were taken from https://neurograd.ucsf.edu/sites/neurograd.ucsf.edu/files/journal_club_evaluation_form.pdf

What is my github username? *

Short answer text

What are the github usernames of the presenters? *

Long answer text

What is the topic of today's Journal Club? (i.e., not the title, but a quick description of what the method is/does) *

Long answer text

Paper Selection (importance, interest): *

- ☐ Poor
- ☐ Good
- ☐ Very Good
- ☐ Outstanding



From the feed: “Over-optimism” + Terry’s IMS Bulletin

We will see a lot of methods in this course – **how do we evaluate what works well in practice ?**

<http://bulletin.imstat.org/2012/11/terences-stuff-does-it-work-in-practice/>

BIOINFORMATICS ORIGINAL PAPER

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Gene expression

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Over-optimism in bioinformatics: an illustration

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EDITORIAL

Ten Simple Rules for Reducing Overoptimistic Reporting in Methodological Computational Research

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“if the improvement of a quantitative criterion such as the error rate is the main contribution of a paper, the superiority of new algorithms should always be demonstrated on independent validation data.”



In class exercise + discussion

- (5 minutes) Read the excerpt from “Terence’s Stuff” column
- (5-10 minutes; discuss with your neighbour) Answer the following 4 questions:
 1. How do we tell what works in practice?
 2. What problems arise using simulated (synthetic) data?
 3. What problems arise using real data?
 4. What are positive/negative controls?
- Discuss
- If simulation: what metrics could/should we use?
- **n.b. include this method comparison context in your Journal Club talks**