

# Transferable skills training: how to make a poster

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# Preamble

**Disclaimer:** these are just my views, and hopefully will make you think about ways to present your work.

## Why give a poster?

- Posters are often regarded as second prize for presenting your work at a conference.
- Conference organizers need many attendees; people come normally to present their work. Don't assume acceptance of your poster means it has undergone standard peer review process.
- Posters give more opportunity for interaction, and longer time to talk. (e.g. SFN talks are 10 minutes; posters are one hour).
- Good forum to show your ideas and meet people.
- Think of it as an advert. (Display one too if you are looking for a job, or if your group is hiring).

# Examples: old style



# Examples: current style

## Mechanisms underlying the formation of beta retinal ganglion cell mosaics

Stephen J. Egle<sup>1</sup>, Peter J. Diggle<sup>2</sup>, John B. Troy<sup>3</sup>

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### Introduction

Beta retinal ganglion cells (RGCs) are labelled ON-cones or OFF-cones, depending on their response to light (Fig. 1). Cell bodies of each type form a non-explicit pattern, termed "retinal mosaics". We do not yet know how the mosaics of ON- and OFF-cone cells emerge during development.

- A population of indistinguishable beta cells may divide into two typologically different types through bistatic interactions, possibly mediated by activity.
- The two types of cell may develop independently of each other.
- Previous statistical approaches based on testing for statistical dependence between ON and OFF cells. This is not a statistical evidence when both types of neuron are located in the same layer, since the constraint that two neurons cannot occupy the same (x, y) location does not independence a priori.

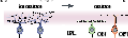


Figure 1: Development of stratification in beta RGCs (drawing from Wang & Chavira, 2002). Stratification reflects functional class.

**Approach:** we fit models of the joint spatial pattern which require the constraint that no two neurons can be separated by less than their soma diameter. If model replicates real maps without equating bistatic interactions, this might suggest bistatic interactions do not occur during development.



Figure 2: Real (WBL; Watzel et al., 1991) and simulated RGC mosaics.

### Methods

- $d_{\text{min}}$  model (Gelli-Bozza et al., 1997) adapted to bistatic case (Fig. 3). Size of histotypic exclusion zone derives from a bivariate distribution (mean  $\mu$  and  $\sigma$ ); histotypic exclusion zone fixed at soma diameter.
- Model parameters used to find best fit:  $d_{\text{min}}$  (M23) and  $d_{\text{diff}}$  (see: L. 1) — mean (total) number of cells within diameter of a cell. 1. function are constant versions of DRP; Rodick, 1991).
- regularity index — mean of  $d$  of the distance to nearest neighbour.
- function of  $1^{\text{st}}, 2^{\text{nd}}, 3^{\text{rd}}$  or all nearest neighbours of opposite type.

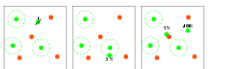


Figure 3: Bivariate  $d_{\text{min}}$  model. On- and off-cone cells are initially located randomly throughout the array. All cells are then moved within the array according to the following procedure. A cell is selected (1) and repositioned randomly (e.g. at 4) avoiding histotypic exclusion zones (dotted circles); (2) and smaller histotypic zones (solid red circles, which are cell bodies of opposite type); (3). One sweep consists of moving all cells in the array once. Cells are moved for many sweeps to allow the pattern to stabilize.

### Results

Best fits could be replicated by bivariate  $d_{\text{min}}$  model (Table 4; Fig. 2, 4, 5). DRP is equivalent DRP for each function.

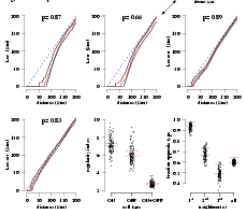


Figure 4: Results for field M62. Red lines indicate experimental data; black lines indicate envelope from 99 simulations. Dotted blue lines indicate the expectation of L for a Poisson pattern. In inset charts, each black dot indicates one simulation, and dotted black line indicates median.

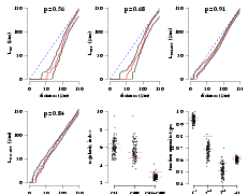


Figure 5: Results for field WBL (same format as Fig. 4).

Field	#ON	#OFF	$d_{\text{min}}$	$d_{\text{diff}}$	RMSE
WBL1	65	70	1.16 ± 0.20 $\mu\text{m}$	1.30 ± 0.25 $\mu\text{m}$	0 $\mu\text{m}$
M621	74	82	1.00 ± 0.13 $\mu\text{m}$	0.96 ± 0.15 $\mu\text{m}$	1.5 $\mu\text{m}$

Table 4: Best-fit parameters of the  $d_{\text{min}}$  model to the two datasets.  $d_{\text{min}}$  and  $d_{\text{diff}}$  — mean  $\pm$  s.d. of histotypic exclusion zones; soma diameter of histotypic exclusion zone.

### Conclusions

- Real RGC maps can be simulated with limited interaction between the two mosaics. Histotypic interactions are limited to preventing axonal overlap.
- Continuous general principle that mosaics are functionally independent of each other (Reckart et al., 2000).
- Previous model suggested axial dependency between two mosaics (Chan & Troy, 2000); may be by-product of model simplification.
- Functional implications of independence in arrays?
- Can cone mosaic coexist with axial maps (imposing developmental processes, such as cell death). Limited data sets (n=5). Interactions between dendritic confinement and axon positioning unknown.

**Acknowledgements:** Wellcome Trust (SE) and NREY0669 (JT). Thanks to Haini Wang for providing comp. WBLs.



POSTER SESSIONS

## Preparation before the day

1. Make sure you know the poster board size and orientation (portrait/landscape).
2. Aim to finish a few days before you depart for the meeting. Poster making takes time. For your first poster, I suggest allowing three days.
3. Do a dry-run with your supervisor (e.g. the “3–5 minute talk”) before printing final version.
4. Check what materials will be available for hanging poster (pins/velcro).
5. Beware of relying on using a laptop to show multimedia; power sockets and tables are rare.
6. At large meetings, check what other posters will be shown in the same session as yours, and arrange cover.
7. Find a poster tube and **label it!**

## Planning the content

- A poster is not a journal article; focus on core figures and results.
- Use an A4 version of poster as a handout; use separate handouts for details of method.
- Posters full of text are ignored!
- Layout:

Top-left Problem and aims of the poster.

Bottom-right Summary of what you've shown.

In-between Method, results, FIGURES.

- Figures are the key: design them first (with good legends) and give minimal supporting text.
- Do not include abstract (submitted previously to the conference) or references. People will find them.
- Provide acknowledgements and contact info, but keep them small!

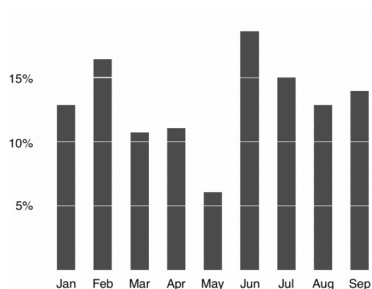
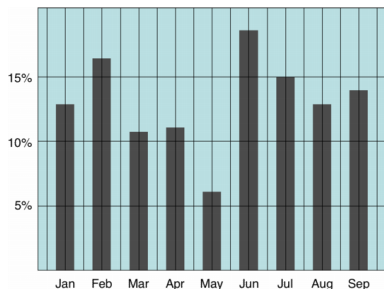


## Effective figures: Edward Tufte

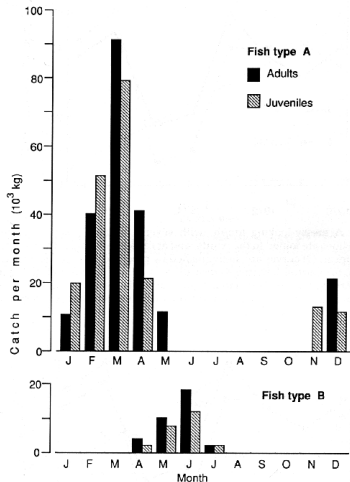
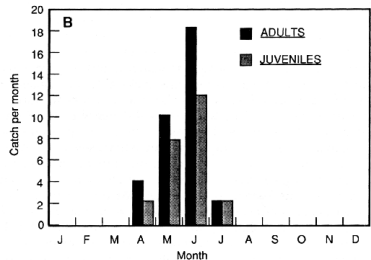
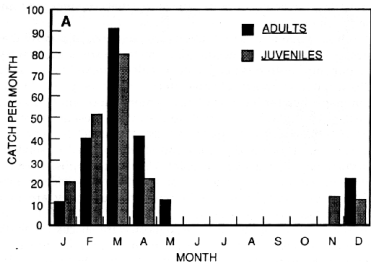
Keep the “Data to ink ratio” high. Remove chartjunk.

<http://www.edwardtufte.com/tufte/>

Sequence: <http://www.tbray.org/ongoing/data-ink/di1>



# Improving figures: another example (O'Connor)



Source: Maeve O'Connor (1996). Writing successfully in science. Chapman & Hall. (Figure 4.1, 4.2)

## Top ten worst graphs

See [http://www.biostat.wisc.edu/~kbroman/topten\\_worstgraphs/](http://www.biostat.wisc.edu/~kbroman/topten_worstgraphs/)  
“What’s worse than one piechart?”

## What happens on the day

- Hang your poster up early (you *might* get a good spot):



- Find out how long you are expected to be at your poster: don't miss the allotted session.
- Offer to talk people through your work.
- Prepare a 3–5 min. summary of your poster. Don't just read the poster.
- Hang envelope for handouts (+ to collect business cards).

## Mechanics of poster making

- Mount several A4 sheets; easy to carry.
- Use A0/A1 poster. Nicer, but hard to carry, and requires specific printing.
- Print shops: PANDIS, Anatomy, Engineering. (DAMTP students can get P/O from John Turner, to avoid VAT.)
- Since Summer 2010 CMS has an A0 printer:  
<http://www.damtp.cam.ac.uk/internal/computing/printing/poster.html>
- Most print shops can handle rescaling, but ensure your aspect ratio is the same as the poster board.
- What software?
  1. Illustrator / Inkscape
  2. Powerpoint (?)
  3.  $\text{\LaTeX}$  ...
- Note: I am a fan of  $\text{\LaTeX}$  but posters require much re-arrangement of material, so you might find GUIs more suitable.

# Using $\text{\LaTeX}$ for making posters

<http://www.damtp.cam.ac.uk/user/eglen/damtp/cuposter> allows you to make poster in a simple 3 col format.

## Mechanisms underlying the formation of beta retinal ganglion cell mosaics

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<sup>1</sup>University of Cambridge and <sup>2</sup>University of Edinburgh, UK; <sup>3</sup>University of Lancaster, UK; <sup>4</sup>Northwestern University, USA.



### Introduction

Retinal ganglion cells (RGCs) are lateral Offspring or OFF cells depending on their response to light (Fig. 1). Cell bodies of each type form a non-overlapping pattern, termed 'lateral mosaics'. The two patterns have the same size and are separated by a distance equal to the diameter of the RGCs.

A population of RGCs that are lateral OFF cells may arise via two independent mechanisms through the same process, possibly mediated by one of:

- 1) The two types of RGC arise independently of each other.
- 2) The two types of RGC arise from the same progenitor, but the two types of RGC are formed in the same place, with the same time and space constraints.



Figure 1: Development of RGCs in the retina. RGCs are formed from a single progenitor cell (Eglen et al., 2002). The diagram shows the formation of lateral RGC mosaics.

Figure 2: Final RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.

Figure 3: Final RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.

### Methods

1. **Fig. 1**—see text for details. The diagram shows the formation of lateral RGC mosaics.
2. **Fig. 2**—see text for details. The diagram shows the formation of lateral RGC mosaics.
3. **Fig. 3**—see text for details. The diagram shows the formation of lateral RGC mosaics.



Figure 4: RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.



Figure 5: RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.

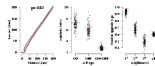


Figure 6: RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.

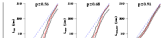


Figure 7: RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.

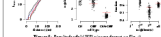


Figure 8: RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.



Figure 9: RGC mosaics. The diagram shows the formation of lateral RGC mosaics. The RGCs are shown in different colors (red, green, blue) to represent different RGC types.

But I recommend you now try the Beamer poster style:

<http://www-i6.informatik.rwth-aachen.de/~drew/latexbeamerposter.php>

# Summary

- Leave lots of time to prepare.
- Work on figures first.
- Prepare an effective 3–5 minute summary.
- Questions?

## Further reading

- Edward Tufte. The visual display of quantitative information.
- Maeve O'Connor. Writing successfully in science.
- “Poster perfect”  
<http://the-scientist.com/2011/09/01/poster-perfect/>
- <http://colinpurrington.com/tips/academic/posterdesign>
  
- Image credits: Images taken from NCSU and Swarthmore sites;  
SfN poster session image from  
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