RETISTRUCT manual

David C. Sterratt

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Chapter 1

Installation

1.1 Install the necessary system packages

1.1.1 Ubuntu Linux

Follow the instructions at https://cran.r-project.org/bin/linux/ubuntu/ to obtain the latest stable version of R. For example, for Ubuntu 16.04 (Xenial) to download R from the Bristol UK mirror, enter the following in a terminal window:

```
sudo apt-key adv --keyserver keyserver.ubuntu.com --recv-keys E084DAB9 sudo apt-add-repository deb https://https://www.stats.bris.ac.uk/R/bin/linux/ubuntu xenial/sudo apt-get install r-base r-base-dev libgtk2.0-dev libgl1-mesa-dev libglu1-mesa-dev
```

1.1.2 Fedora Linux

NOTE: these instructions have not been tested since at least 2015. As root,

```
yum install R-core
yum install R-devel
yum install gtk2
yum install gtk2-devel
yum install mesa-libGL-devel
yum install mesa-libGLU-devel
```

1.1.3 Mac OS X

First install R for Mac OS X, available from http://www.r-project.org/. Under Mac OS X Leopard and Lion, this should be sufficient.

Under Mac OS X 10.8 (Snow Leopard),

it may be necessary to install the following external packages:

- Xcode (free from App Store)
- Xquartz (http://xquartz.macosforge.org/)
- GTK (http://r.research.att.com/libs/GTK_2.18.5-X11.pkg)

Under Mac OS X 10.9 (Mavericks) or Mac OS X 10.10 (Yosemite),

See this issue on Github: https://github.com/davidcsterratt/retistruct/issues/4

1.1.4 MS/Windows

First install R for MS/Windows, available from http://www.r-project.org/.

1.2 Install the core RETISTRUCT package

- 1. Start R
- 2. Type:

```
install.packages("retistruct")
```

The first time this runs, it should create a personal directory for R packages, and it will take a few minutes to install some required packages.

3. Test that it is working as follows (from within R)

```
library(retistruct)
demo("retistruct.method")
```

If all is well, you should be prompted to press **Return** a few times and Figure 1 from the paper [1] will appear (drawing the 3D renditions takes some time). There will be reports of warnings; they are to do with printing the figures to files. You can view them by typing warnings().

4. A further test produces Figure 2:

```
demo("low.high")
```

5. A further (quite long-running) test produces Figure 6:

```
demo("figure6")
```

1.3 Install the RETISTRUCT GUI

If the previous step works:

1. Type:

```
retistruct()
```

- 2. Some more packages will be downloaded this may take a minute or so.
- 3. If you are using MS/Windows, there may be an error message saying that libatk-1.0-0.dll is missing from your computer. Click on **OK**. Another message will appear, asking if you want to install GTK+. Agree to this. After this has finished, quit R (without saving the workspace image) and restart.
- 4. If all works, an interface window should appear. A number of sets of demonstration data are available from the **Demo** menu item.

Chapter 2

Running RETISTRUCT

To start the program, start R. At the R prompt type:

library(retistruct)
retistruct()

A window should appear.

2.1 Preparing and opening the files for a retina

There are a number of types of information associated with a flat-mount retina that RETISTRUCT can process:

- The coordinates of the outline of the flattened retina
- An image of the flat-mount retina (optional)
- The coordinates of labelled data points within the flat-mount retina (optional)
- The coordinates of labelled counts of data within the flat-mount retina (optional)
- The scale of the coordinates provided in the other files (optional)

To import this information into RETISTRUCT, for each retina, a directory should be created containing files with the above information, as will be described below. At present there are three formats of directory that RETISTRUCT can read. Most users will probably find the the IMAGEJ ROI format most convenient.

2.2 Reading files using the IMAGEJ ROI format

This format allows you to load images of retinae whose outlines have been marked up in IMAGEJ. To create the image and outline files for this format:

- 1. Create a directory to save the files created to.
- 2. Open up IMAGEJ (or FIJI).
- 3. Use **File** \rightarrow **Open** to open the image.
- 4. Use **Image**→**Scale** to down sample so that the resolution is less than 1000x1000. (This is not crucial, but will speed up things later.)
- 5. Save the down-sampled version of the image to the file name image.png in the directory created in Step 1.

- 6. Use the Polygon Tool () to mark the edge of the retina. According to the IMAGEJ manual : "To create the selection, click repeatedly with the mouse to create line segments. When finished, click in the small box at the starting point (or double-click), and ImageJ automatically draws the last segment.
 - The points that define a polygon selection can be moved or deleted, and new points can be added. To delete a point, click on it with the alt key down. To add a point, click on an existing point with the shift key down."
- 7. Open the ROI manager by selecting **Analyze**→**Tools**→**ROI Manager**.
- 8. Click on the **Add** [t] button.
- 9. Click on the More → Save button. In the Save selection... box that appears, enter the file name outline.roi and make sure this file is saved to the same directory as the image.png file, i.e. the directory created in Step 1.
- 10. Optionally mark up the optic disc in the same way as the outline. Save this ROI to a file called od.roi in the same directory as image.png.
- 11. Open RETISTRUCT.
- 12. Click on the **Open** icon and select the directory containing image.png and outline.roi. The retinal image should now appear, with the outline shown. By default the outline is in black. If this isn't visible against the image, press the **Properties** button in the interface (or select **Edit** → **Properties**), and change the **Outline colour**.

2.2.1 Specifying the scale

Optionally, the scale of the image can be specified by providing a file scale.csv in the same directory as the image and ROI files. This file should contain two lines, the first with the heading Scale and the second with the length of the side of a pixel in micrometres. For example:

"Scale"

2.2.2 Reading in data points

In general, the coordinates of data points are read in from a csv file called datapoints.csv. The two cells of the first line of the file contain the name of the group of data points and the colour that these should be displayed in Retistruct. Marking up the points from the image in IMAGEJ will ensure that the coordinates are in the same system as those used for the image and the outline. To do this:

- 1. Open the image used for marking up the outline (image.png) in IMAGEJ.
- 2. Threshold the image (**Image** \rightarrow **Adjust** \rightarrow **Threshold...** so that the labelled points are visible.
- 3. Select **Analyze**→**Set Measurements...** and make sure that **Centroid** is checked
- Select Analyze → Analyze Particles. Make sure that Display results and Clear results are checked. Click on OK.
- 5. A window entitled **Results** should appear. This contains the X and Y coordinates of the detected points. In this window select **File**→**Save as...** and save the file as datapoints.csv.
- 6. Open this file with either EXCEL or OPENOFFICE CALC and remove all the columns apart from the X and Y columns. Replace the X with the name of the data set and the Y with the colour it should be displayed in RETISTRUCT.
- 7. Save datapoints.csv in the retina's directory, along with outline.roi (and image.png if an image is desired).

¹http://rsbweb.nih.gov/ij/docs/tools.html

2.2.3 Reading in data counts

In general, the coordinates of data counts (counts of data at X, Y positions) are read in from a csv file called datacounts.csv.

The two cells of the first line of the file contain the name of the group of data counts and the colour that these should be displayed in RETISTRUCT.

Each subsequent row has three numbers: The first two columns contain the X and Y coordinates at which the count was measured, and the third column contains the count at that location. It is important to ensure that the coordinates are in the same system as those used for the image and the outline.

Save datapoints.csv in the retina's directory, along with outline.roi (and image.png if an image is desired).

2.3 Reading in files using other formats

2.3.1 CSV format

This is the same as the IMAGEJ ROI format, except that the outline is contained in the two columns of a file called outline.csv. Each column should have a heading, e.g.:

```
"X", "Y"
1,5
10,76,
```

The optic disc outline can be supplied in a similar file called od.csv.

2.3.2 IDT format

This is the format used in Ian Thompson's lab (see Appendix A). These files are contained in a directory. To open the files corresponding to a retina, click on the open file icon, and navigate to the directory containing the SYS and MAP files. On opening this directory, the retinal outline should appear in the RETISTRUCT window.

2.4 Editing the retinal mark-up

After a set of files is opened, the **Edit** tab in the RETISTRUCT window will be open. Before the retina can be reconstructed, you need to mark-up tears and landmarks on the retina using the following controls:

Add tear To add a tear, click on this button, then click on three points in turn which define a tear. The order in which the points are added does not matter. Tears contained within a tear can be marked up, but tears cannot cross over one another.

Move Point To move one of the points defining a tear, click on this button, then click on the point which you desire to move, then click on the point to which it should be moved.

Remove tear To remove a tear, click on this button, then click on the apex of the tear (marked in cyan on the plot)

Mark nasal To mark the nasal pole, click on this button, then click on the point which is the nasal pole. If the nasal or dorsal pole has already been marked, the marker is removed from the existing location. The nasal pole should not be in a tear. If the nasal tear is placed within a tear, no error is reported at this stage, but it will be reported later.

Mark dorsal As above, except for the dorsal pole.

Mark OD To mark the optic disc, click on the structure marked in orange which you think is the OD. Once clicked on, the structure should become blue.

Phio This determines the latitude of the rim of the reconstructed hemisphere. In mouse, it depends on the age of the animal [1]. **Ideally you will have measured or estimated this value from an intact retina.** The default value of 0 may lead to reconstructions that are not as accurate as they could be, since the template that RETISTRUCT is trying to morph the flat-mount retina onto is not the same as the original shape.

2.5 Editing metadata

You can also edit the following metadata:

Data/Flip DV Flip the DV axis to compensate for microscope orientation. Affects display of retinae.

Eye Specify whether the eye is Right or Left. Affects display of retinae.

2.6 Saving the markup and metadata

To save the markup, click on the "Save" button in the toolbar. This saves various markup files to the directory containing the data files. This saved data can be used to reconstruct the retina using a batch process (Section 3.3).

2.7 Reconstructing the retina

To reconstruct the retina, click on the "Reconstruct retina" button. This causes RETISTRUCT to perform a (sometimes lengthy) sequence of operations:

Stitching Links between corresponding points on parts of the retinal outline contained in tears are made.

Triangulation A triangular mesh is placed over the flattened retina

Initial projection to sphere The mesh is projected roughly onto a sphere

Optimisation The locations of the mesh points on the hemisphere is adjusted so as to minimise a weighted sum of the squared differences between the lengths of links in the mesh on the hemisphere and on the flattened retina, whilst ensuring that as few triangles as possible are flipped.

At the end of the reconstruction process, a polar plot appears next to the flattened retina. By default the location of the cuts and tears in the polar coordinates can be seen.

2.8 Assessing the quality of the reconstruction

It is important to assess the quality of the reconstruction. The choice of where to mark up tears can affect this adversely, as can the rim angle.

One diagnostic is to view the strain in each mesh link by clicking on the **Strain** checkbox in the **Edit** tab. When **Strain** is shown, the flat plot shows the strain in the links of the mesh using a colour code from blue (compressed) to red (expanded). The polar plot is replaced by a scatter plot of the length of links in the reconstructed object versus the length on the flattened object. The flat plot of a good reconstruction will look mostly green, and in the scatter plot the points will lie close to the black line.

2.9 Selecting viewing options

To select viewing options click on the **View** tab. A number of options are now shown, grouped into a number of sections:

2.9.1 Show

In the **Show** section there are checkboxes that allow you to show various types of information:

Markup Locations of tears and the dorsal or nasal pole

Stitch Locations of how the algorithm has stitched tears (only visible after the reconstruction step)

Grid Lines of latitude and longitude projected back onto the flattened retina (only visible after the reconstruction step) and grid lines are shown on the reconstructed retina

Landmarks Landmarks such as the optic disc

Points Locations of data points, such as the locations of beads of dye, which have been imported into RETISTRUCT as described in Section 2.2.2

Point means Locations of the Karcher mean (mean in spherical coordinates) of each group of data points

Point contours Kernel Density Estimate contours of the data points. By default, contours at 5%, 25%, 50%, 75% and 95% of the maximum density are shown. This can be changed by a command like this at the R prompt:

```
options(contour.levels=c(5, 25, 50))
```

This command would cause the 5%, 25% and 50% contour lines to be displayed.

Counts Counts (for example of cells), represented by a number at the location at which the count was made, which have been imported into RETISTRUCT as described in Section 2.2.3

Count contours Kernel Regression contours of the data counts. By default, contours at 5%, 25%, 50%, 75% and 95% of the maximum density are shown. This can be changed by a command, as shown under "Point contours".

2.9.2 IDs

If you included a datapoints.csv and/or a datacounts.csv file (see Sections 2.2.2 and 2.2.3) the **IDs** section will contain checkboxes with names corresponding to the names of the groups of points and the names of the groups of counts specified in those files. If you have marked the optic disc, it will also contain an **OD** checkbox. Check or uncheck these boxes to show or hide the corresponding points and counts.

2.9.3 Projection

The reconstructed data can be viewed in a number of projections, just as there are various ways that the surface of the globe is projected to a 2D image. Because the geometry of the surface of a sphere is fundamentally different from a flat sheet of paper, no projection is a perfect representation; all have advantages and disadvantages.

Note: Coordinate systems

The co-ordinates of a point on the reconstructed sphere can be described in two ways:

Latitude and longitude Here latitude and longitude are defined analogously to the standard geographical coordinate system. We imagine the retina lying with its optic pole at the south pole, and with its widest point lying on the equator. The latitude of the optic pole is -90° and the latitude of point on the widest point of the retina (its "equator") is 0° . We denote the latitude φ and the longitude λ .

Colatitude and **longitude** Here we imagine the retina in the same orientation with respect to the earth, but we rather than latitude, we use colatitude measured from the south pole, i.e. the south pole has a colatitude of 0° and the widest point (the "equator") has a colatitude of 90° . We denote colatitude by ψ and it is related to latitude by $\psi = \varphi + 90^{\circ}$.

Currently the options are:

Azimuthal Equidistant The default projection. This is a polar plot in which the radial distance is proportional to the colatitude (latitude measured from the "South Pole"; see note on coordinate systems), and angle from the horizontal is equal to the longitude on the original retina. Points equidistant on line of longitude are also equidistant in this projection. It is not an area-preserving projection: it makes regions closer to the rim look bigger than they actually are relative to areas close to the south pole.

Projection	Latitude & longitude	Colatitude & longitude
Azimuthal Equidistant	$ ho=90^\circ+arphi$, $\lambda=\lambda$	$\rho = \psi, \lambda = \lambda$
Azimuthal Equal Area	$ ho = \sqrt{2(1+\sin\varphi)}$, $\lambda = \lambda$	$ \rho = \sqrt{2(1-\cos\psi)}, \lambda = \lambda $
Azimuthal Conformal	$ \rho = \tan{(45^{\circ} + \varphi/2)}, \lambda = \lambda $	$ \rho = \tan(\psi/2), \lambda = \lambda $
Sinusoidal	$x = (\lambda - \lambda_0) \cos \varphi, y = \varphi$	
Orthographic	$x = \cos\varphi\sin(\lambda - \lambda_0)$	
	$y = \cos \varphi_0 \sin \varphi - \sin \varphi_0 \cos \varphi \cos(\lambda - \lambda_0)$	

Table 2.1: Projections. For the three azimuthal projections, the polar coordinates (ρ, λ) are given in terms of the latitude and longitude (φ, λ) or colatitude and longitude (ψ, λ) on the original sphere. For the sinusoidal and orthographic projection the (x, y) coordinates in the plane are given. For these projections (φ_0, λ_0) is the latitude and longitude at the centre of the projection.

Azimuthal Equal Area In this projection, also known as a Lambert projection, points that are equidistant on lines of longitude on the spherical retina are not equidistant: points close to the rim are mapped to be closer together. This is done so that area is preserved in the sense that equal areas on the sphere project to equal areas on the plane of projection. It is thus an area-preserving projection.

Azimuthal Conformal In this projection, also known as a Wulff projection, angles are preserved. This is done at the expense of expanding the representation of areas close to the rim.

Sinusoidal The sinusoidal projection projects the entire globe onto the plane and preserves area. The user selects the longitude at the centre of the projection in the **Az** box in the **Projection centre** section of the window. The sinusoidal projection is helpful for data transformed into visuotopic space (see later).

Orthographic The orthographic projection gives a perspective view of one side of the globe. The projection is centred on a latitude and longitude specified by the values in degrees in the **El** and **Az** boxes in the **Projection centre** section.

Note: Convention on longitude of nasal, dorsal, temporal and ventral poles

By convention, the polar plots are viewed as though the animal is facing towards the observer. This means that when plotting a retina from a right eye, the nasal pole on the right and N, D, T, V are in anticlockwise order; for a retina from a left eye, nasal is on the left and N, D, T, V are in clockwise order. The longitude of a point was defined so that 0° is always at the right of the plot and 90° at top. This means that for a right eye the poles correspond to the longitudes as follows: N, 0° ; D, 90° ; T, 180° ; V, 270° . For a left eye the longitudes of the nasal and temporal poles are interchanged: N, 180° ; D, 90° , T, 0° ; V, 270° .

2.9.4 Transform

The Transform option allows the reconstructed retina to be projected into visual space. The options are

None (default) No transformation is made.

Invert The image is inverted as though by a pinhole lens, a crude approximation to the optics of the eye. This allows visual space to be mapped onto the retina via the optical system of the eye. This mapping depends on the angle that a ray makes with the optic axis (the **Axis direction**) which we define as being oriented at azimuth (**Az**) and elevation (**El**).

Invert to hemisphere Similar to **Invert**, except that the projection is compressed or expanded so that the field of view is 180°.

There are more details of the transformation in the supplemental information of [1].

2.10 Exporting the plots

The **Bitmap** and **PDF** buttons above the flatmount view and the reconstructed view export the image show to a PNG file format or an PDF respectively. To change the width of the image in pixels, select **Properties** and set the **Maximum width of projection** to the desired number of pixels. **Warning:** setting a high number of pixels will create a high-resolution image, but can also take some minutes.

2.11 Saving the reconstruction

Note: Saving reconstructions in Retistruct v0.6.0

The ability to save reconstructions is currently disabled in v0.6.0, but it is planned to enable it in v0.6.1.

To save the markup, click on the "Save" button in the toolbar. This saves various files to the directory containing the data files. When the files in the directory are opened again, all the markup information (cut locations, location of nasal point and rim angle) is loaded, and the reconstruction will also appear, unless there has been a major upgrade of the software, in which case the retina will need to be reconstructed using the "Reconstruct retina" button.

Chapter 3

Further topics

3.1 Accessing reconstruction data from R

It can be useful to have access to the data underlying a reconstruction directly, for example to allow statistical analysis. To load saved date into R, type the following into R:

```
r <- retistruct.read.recdata("/path/to/reconstruction/directory")</pre>
```

The resulting object r contains various fields, which can be accessed using the R \$ operator, e.g.

```
r$Dss$green
```

gives a list of the spherical co-ordinates of green labelled data points. See Table 3.1 for a list of the most useful quantities. Other quantities (for example the locations of the mesh points in the flat and spherical retina) are also available. The documentation for ReconstructedOutline, ReconstructedDataset, getDss and getSss in the user manual has more information about this.

3.2 Reading reconstruction data into MATLAB

By default when a reconstruction is saved, a subset of the data is stored in a file called r.mat in the same directory as the raw data and the markup. To import this data into MATLAB, cd into that directory, and type:

```
clear
load r.mat
```

This puts a number of variables into the workspace, as shown in Table 3.1.

To produce a polar plot of data points, try the following code:

```
polar(Dss.green(:,2), Dss.green(:,1)*180/pi+90, '.g')
hold on
polar(Dss.red(:,2), Dss.red(:,1)*180/pi +90, '.r')
hold off
```

The radial axis indicates the latitude in degrees measured from the retinal pole.

In the matlab subdirectory of the distribution, there are some scripts to produce polar polar plots, including the locations of tears and landmarks. To create PDF plots of all the retinae in a directory, try:

```
makefigures('retinae', 'output_directory')
```

There are also some embryonic scripts to create polar plots: $plot_datapoints_polararea.m$ and polararea.m.

1:0	Th. 1. Co. 1 6.0 - 1	
phi0	ressed in degrees	
Dss	_	ons of labelled cell bodies in spherical coordinates.
		ne latitude of each point, measured in radians. The
		e longitude of each point, measured in radians.
DssMean	Location of Karcher mean	of green-labelled cell bodies in spherical coordin-
	ates. The first column conta	ins the latitude of each point, measured in radians.
	The second column contain	s the longitude of each point, measured in radians.
	A value of -2147483648 co	rresponds to NA in R.
DssHullarea,	Structure containing area o	f convex hull of points on sphere. The convex hull
		awn around data points. A value of -2147483648
	corresponds to NA in R.	•
Tss	-	lement contains spherical coordinates (in the same
	latitude-longitude format a	
Sss	_	lement contains spherical coordinates (in the same
	latitude-longitude format a	
KDE		nation about Kernel Density Estimates of the loca-
ומו	tions of cell bodies.	nation about refrict benoity Estimates of the focu
	green_flevels	Contour heights, determined by finding heights
	810011101015	that exclude a certain fraction of the probability.
		For example, the 95% contour is excludes 95% of
		the probability mass, and it should enclose about
		5% of the points.
	green_labels	
	green_rapers	Contour labels. These give the label (e.g. 5, 25, 50,
		where these are the percentages above) of each
		contour. Note that there may be more than one
		contour at the same level, so this vector may con-
		tain more elements than flevels. The first ele-
		ment of green_labels labels the contour whose
		coordinates are specified in green_contours1,
		the second element of green_labels relates to
		green_contours2 and so on.
	green_tot_contour_areas	The total area in square degrees enclosed by each
		contour. This is a matrix with the first column
		giving the contour label (see above) and the next
		column giving the area.
	${ t green_kappa}$	The concentration parameter of the Fisher density
		determined by the kernel fitting algorithm.
	green_h	A pseudo-bandwidth parameter, the inverse of
		the square root of kappa. Units of degrees.
	green_maxs_phi	Lattitude of maximum point of kernel estimate.
	${\tt green_maxs_lambda}$	Longitude of maximum point of kernel estimate.
	green_g_xs green_g_ys	Kernel density estimates on standard polar
	green_g_f	grid. This can be plotted in MATLAB us-
		ing the command contour(KDE.green_g_xs,
		KDE.green_g_ys, KDE.green_g_f).
	green_gpa_xs	Kernel density estimates on area-preserving po-
	green_gpa_ys	lar grid. Plotted in MATLAB as above.
	green_gpa_f	
	green_contours1,	Coordinates of contours. See green_labels
	green_contours2	above for more explanation.
	green_contour_areas1,	Area contained within each individual contour.
	_	See green_labels above for more explanation.
KR		Regression estimates of the density of points, de-
1116		a points. All fields correspond to KDE above.
	Tivea from the grouped dat	m points. I'm neids correspond to the above.

Table 3.1: Variables exported in the ${\tt r.mat}$ file.

3.3 Running a batch of reconstructions

The RETISTRUCT library can be used to reconstruct a batch of retinae which have been marked up. Suppose that the directory retinae contains a directory tree in which there are data directories containing the raw outline, data point and image files and the saved markup files. In order to perform the reconstructions, we create a new directory retinae/reconstructions, and run the following sequence of commands in R:

```
R
> library(retistruct)
> retistruct.batch(tldir='retinae', outputdir='retinae/reconstructions')
```

This command will go through the retinae directory, looking for valid data directories. If it finds one, it sets about trying to reconstruct the retina. As it reconstructing each retina, it writes to log file in retinae/reconstructions. Once the reconstruction is complete, it saves a number of plots in this directory in PDF format. It also adds a line to a summary log file in retinae/reconstructions called retistruct-batch.csv. This file contains a number of columns:

Dataset The directory of the data set

Return The return value from the process

Result A summary of the result, including if any errors were returned

E The total error of the optimised reconstruction

El The error due to purely to the lengths of links in the optimised reconstruction

nflip The number of flipped triangles

EOD The distance of the Optic Disc from the inferred centre of the retina, in degrees. If the OD has not been marked up, this is NA.

To export the reconstruction data in a directory hierarchy in which retistruct.batch() has been run, run the following sequence of commands in R:

```
R
> library(retistruct)
> retistruct.batch.export.matlab(tldir='retinae')
```

Appendix A

The IDT Data format

The data for each retina is stored in a separate directory. Within each directory there are two files:

- SYS.SYS A table in SYSTAT format containing the coordinates of the red, green and doubly labelled cell bodies, and counts of labelled cell bodies within each grid box. The column headings shown in Table A.1. Each row of the table contains information only on a subset of the data, e.g. the coordinates of a red-labelled cell.
- ALU.MAP A text file containing the coordinates of the map outline. The file comprises a number of sections, each starting with a single number, which is the number of lines to read in the next section. These lines have two numbers each, the *x* and *y* coordinates of a vertex of the map outline.

FOR EACH BOUNDARY							
MAPNUM	id number of boundary						
MINLAT	min latitude						
MAXLAT	max latitude						
MINLON	min longitude						
MAXLON	max longitude						
LABLAT	latitude of label						
LABLON	longitude of label						
FOR EACH CELL							
XRED	<i>x</i> -coordinate if cell labelled red but not doubly						
YRED	<i>y</i> -coordinate if cell labelled red but not doubly						
XGREEN	<i>x</i> -coordinate if cell labelled green but not doubly						
YGREEN	<i>y</i> -coordinate if cell labelled green but not doubly						
XDOUBLE	<i>x</i> -coordinate if cell labelled doubly						
YDOUBLE	<i>y</i> -coordinate if cell labelled doubly						
XGRID	sample box cell is in						
YGRID	sample box cell is in						
PERIM	perimeter of cell						
AREA	area of cell						
	ONE PER GRID BOX						
GRIDX	grid location of centre of sample box						
GRIDY	grid location of centre of sample box						
XGRIDCOO	<i>x</i> -coordinate of centre of sample box						
YGRIDC00	<i>y</i> -coordinate of centre of sample box						
BOXSIZEX	size (half width) of sample box in <i>x</i> -direction						
BOXSIZEY	size (half width) of sample box in <i>y</i> -direction						
COMPLETE	whether counting of sample has been completed						
TOTALCEL	total number of cells in this box						
TOTALRED	total number of red-only cells in this box						
TOTALGRE	total number of green-only cells in this box						
TOTALDOU	total number of double cells in this box						
MEANPERI	average perimeter of all cells						
MEANAREA	average area of all cells						

Table A.1: Column headings of the SYS.SYS file.

Bibliography

[1] Sterratt DC, Lyngholm D, Willshaw DJ, Thompson ID (2013) Standard anatomical and visual space for the mouse retina: computational reconstruction and transformation of flattened retinae with the Retistruct package PLoS Comp. Biol. 9