Using the RCircos Package

Hongen Zhang, Ph.D. Genetics Branch, Center for Cancer Research, National Cancer Institute, NIH

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1 Introduction

The RCircos package provides a set of graphic functions which implement basic Circos 2D track plot [1] for visualizing similarities and differences of genome structure and positional relationships between genomic intervals. The package is implemented with R graphics package that comes with R base installation and aimed to reduce the complexity of usage and increase the flexibility in integrating into other R pipelines of genomic data processing.

Currently, following graphic functions are provided:

- Chromosome ideogram plots for human, mouse, and rat
- Data plots include:
 - heatmap
 - histogram
 - lines
 - scatterplot
 - tiles
- Plot items for further decoration include:
 - connectors
 - links (lines and ribbons)
 - text (gene) labels

After successful installation of RCircos, one needs to load the library to get started using it:

> library(RCircos)

2 Input Data Format

RCircos takes the input data in the form of a data frame that could be an object returned from read.table() or generated with other pipelines in the current R session. The first three columns of the data frame, except for input to the link plot, must be genomic position information in the order of chromosome names, chromosome start, and chromosome end positions.

- > data(RCircos.Histogram.Data)
- > head(RCircos.Histogram.Data)

```
Chromosome chromStart chromEnd
                                      Data
1
        chr1
               45000000 49999999 0.070859
2
        chr1
               55000000 59999999 0.300460
3
               60000000 64999999 0.125421
        chr1
4
               70000000 74999999 0.158156
        chr1
5
        chr1
               75000000 79999999 0.163540
6
               80000000 84999999 0.342921
        chr1
```

For gene labels and heatmap plots, the gene/probe names must be provided in the fourth column. For other plots, this column could be optional.

```
> data(RCircos.Heatmap.Data)
```

> head(RCircos.Heatmap.Data)

```
Chromosome chromStart chromEnd GeneName
                                             X786.0
                                                          A498 A549.ATCC
                                                                             ACHN
        chr1
                  934341
                            935552
                                       HES4 6.75781
                                                      7.38773
                                                                 6.47890 6.05517
1
2
                  948846
                            949919
                                      ISG15 7.56297 10.49590
                                                                 5.89893 7.58095
        chr1
3
        chr1
                 1138887
                           1142089
                                   TNFRSF18 4.69775
                                                      4.55593
                                                                 4.38970 4.50064
4
        chr1
                 1270657
                           1284492
                                       DVL1 7.76886
                                                      7.52194
                                                                 6.87125 7.03517
5
                 1288070
                           1293915
                                      MXRA8 4.49805
                                                      4.72032
                                                                 4.62207 4.58575
        chr1
6
                          1624243 SLC35E2B 8.73104
        chr1
                 1592938
                                                      8.10229
                                                                 8.36599 9.04116
    BT.549 CAKI.1
   8.85062 7.00307
1
2 12.08470 7.81459
   4.47525 4.47721
   7.65386 7.69733
5
   5.66389 4.93499
   9.24175 9.89727
```

Different from other plot data, the input data for link line plot has only paired genomic position information for each row in the order of chromosome name A, chromStart A, chromEnd A, chromosome name B, chromStart B, and chromEnd B.

```
> data(RCircos.Link.Data)
> head(RCircos.Link.Data)
```

	${\tt Chromosome}$	${\tt chromStart}$	${\tt chromEnd}$	${\tt Chromosome.1}$	${\tt chromStart.1}$	${\tt chromEnd.1}$
1	chr1	8284703	8285399	chr1	8285752	8286389
2	chr1	85980143	85980624	chr7	123161313	123161687
3	chr1	118069850	118070319	chr1	118070329	118070689
4	chr1	167077258	167077658	chr1	169764630	169764965
5	chr1	171671272	171671550	chr1	179790879	179791292
6	chr1	174333479	174333875	chr6	101861516	101861840

Note: RCircos will convert the input data to RCircos plot data but it does not provide functionality for general data processing. If the data frame does not have genomic position information, you have to add the information to the data frame before passing it to RCircos functions. Sample datasets are included in the package for demo purpose and they could be easily explored with data() method.

Starting from version 1.1.2, user can append to input data a column of plot color names with header of "PlotColor" to control the colors for each data point except of heatmap plot.

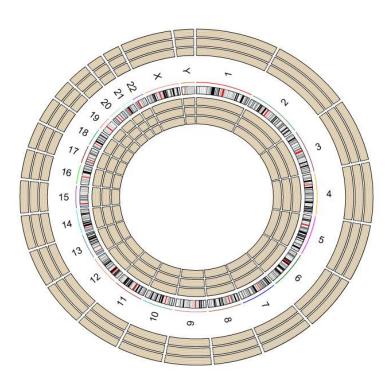
3 Plot Track Layout

RCircos follows the same algorithm of Circos plot and arranges data plots in tracks. A track could be placed either inside or outside of chromosome ideogram

and the detailed position for a track could be easily manipulated by changing of the track width and track numbers.

The figure below shows a human chromosome ideogram plus three empty tracks arranged in both inside and outside of chromosome ideogram.

R Circos Layout Demo



4 Getting Started: Initialize RCircos core components first

4.1 Initialize RCircos core components

The first step of making RCircos plot is to initialize RCircos core components. To setup RCircos core components, user needs load the chromosome ideogram data into current R session. The RCircos package have three build-in datasets for human, mouse, and rat chromosome ideograms which can be loaded with data() command. Ideogram data in text files with same format can also be

loaded with read.table() function in R.

```
> data(UCSC.HG19.Human.CytoBandIdeogram);
> head(UCSC.HG19.Human.CytoBandIdeogram);
  Chromosome ChromStart ChromEnd
                                    Band
                                           Stain
1
        chr1
                       0
                         2300000 p36.33
                                            gneg
2
        chr1
                2300000 5400000 p36.32 gpos25
3
                5400000 7200000 p36.31
        chr1
                                            gneg
                7200000 9200000 p36.23 gpos25
4
        chr1
5
        chr1
                9200000 12700000 p36.22
                                            gneg
6
                12700000 16200000 p36.21 gpos50
        chr1
> data(UCSC.Mouse.GRCm38.CytoBandIdeogram);
> head(UCSC.Mouse.GRCm38.CytoBandIdeogram);
  Chromosome ChromStart ChromEnd Band
                                          Stain
        chr1
                       0 8840440
1
                                   qA1 gpos100
2
        chr1
                8840440 12278390
                                   qA2
                                           gneg
3
        chr1
                12278390 20136559
                                   gA3
                                        gpos33
4
        chr1
               20136559 22101102
                                   qA4
                                           gneg
5
               22101102 30941543
        chr1
                                   qA5 gpos100
6
        chr1
               30941543 43219933
                                    qΒ
                                           gneg
> data(UCSC.Baylor.3.4.Rat.cytoBandIdeogram);
> head(UCSC.Baylor.3.4.Rat.cytoBandIdeogram);
  Chromosome ChromStart ChromEnd Band Stain
1
        chr1
                       0 10142096
                                   p13
                                        gneg
2
        chr1
               10142096 24272657
                                   p12
                                        gvar
3
               24272657 38517175
        chr1
                                   p11
                                        gneg
4
        chr1
               38517175 48659271
                                   q11
                                         gpos
5
        chr1
               48659271 69741157
                                   q12
                                        gneg
6
        chr1
               69741157 90025350
                                   q21
                                        gpos
```

After the chromosome ideogram data is loaded, RCircos core components can be initialized with function of RCircos.Set.Core.Components(). This function needs four arguments:

cytoinfo the chromosome ideogram data loaded

chr.exclude chromosomes should be excluded from ideogram, e.g., chr.exclude <- c("chrX", "chrY");. If it is set to NULL, no chromosome will be excluded.</p>

tracks.inside how many tracks will be plotted inside chromosome ideogram

tracks.outside how many data tracks will be plotted outside chromosome ideogram

Following code initialize RCircos core components with all human chromosome ideogram and 10 data track space inside of chromosome ideogram.

```
> chr.exclude <- NULL;
> cyto.info <- UCSC.HG19.Human.CytoBandIdeogram;
> tracks.inside <- 10;
> tracks.outside <- 0;
> RCircos.Set.Core.Components(cyto.info, chr.exclude,
+ tracks.inside, tracks.outside);
```

RCircos use three core components to perform data transformation and data plot:

RCircos cytoband data RCircos cytoband data is derived from the input chromosome ideogram data. Except of the chromosome name, start and end positions, band name and stain intensity for each band, chromosome highlight colors, band colors, band length in base pairs and chromosome units as well as the relative location on the circular layout are also included. These data are used to calculate the plot location of each genomic data.

RCircos plot positions RCircos plot positions are x and y coordinates for a circular line of radius 1.0 and the total number of points for the circular line are decided by the total number of chromosome units. One chromosome units is a plot point which covers a defined number of base pairs and total units for chromosome ideogram include units of each band plus chromosome padding area, both of them are defined in the list of plot parameters.

RCircos plot parameters RCircos plot parameters are only one core components open to users. With the get and reset methods, users can modify the parameters for updating other two core components. Following are RCircos plot parameters and their default values:

radiu.len The radius of a circular line which serves as baseline for calculation of plot items, default: 1

chr.ideog.pos Radius of chromosome ideogram position, default: 1.1

highlight.pos Radius of chromosome ideogram highlights, default: 1.25

chr.name.po Radius of chromosome name position, default: 1.4

plot.radius Radius of plot area, default: 1.5

track.in.start Radius of start position of the first track inside of chromosome idogram, default: 1.05

track.out.start Radius of start position of the first track outside of chromosome idogram, default: 1.6

chrom.width Width of chromosomes of the ideogram, default: 0.1 **track.padding** Width of padding between two plot tracks, default: 0.02

- **track.height** Height of data plot track, default: 0.1 (Note: Parameters above are all relative to the radius.len and will be updated automatically when reset plot parameters with new radiu.len).
- base.per.unit Number of base pairs a chromosome unit (a plot point) will cover, default: 3000
- chrom.paddings Width of padding between two chromosomes in chromosome unit, default: 3000 (Note: chrom.paddings is binded to base.per.unit. It will be automatically updated if the base.per.unit is changed, unless be set to zero).
- heatmap.width Width of heatmap cells in chromosome unit, default: 1000
- hist.width Width of histogram column in chromosome unit, default: 1000
- color.map Color scales for heatmap plot, default: BlueWhiteRed
- text.size Character size (same as cex in R graphics package) for text plot, default: 0.4
- highlight.width Line type (same as lty in R graphics package) for chromosome highlight, default: 2
- **point.type** Point type (same as pch in R graphics package) for scatter plot, default: "."
- **point.size** Point size (same as cex in R graphics package) for scatter plot, default: 1

hist.color Color for histgram plot, default: red

tile.color Color for tile plot, default: black

track.background Color of track background, default: wheat

grid.line.color Color for track grids, default: gray

Bezier.point Total number of points for a link(Bezier) line default: 1000

max.layers Maximum number of layers for tile plot, default: 5

sub.tracks Number of sub tracks in a data track, default: 5

The core components are stored in RCircos session and each component is supplied with one Get method for advanced usage. In addition, simply call the function RCircos.List.Parameters() will list all current plot parameters.

```
> rcircos.params <- RCircos.Get.Plot.Parameters();
> rcircos.cyto <- RCircos.Get.Plot.Ideogram();
> rcircos.position <- RCircos.Get.Plot.Positions();
> RCircos.List.Parameters()
```

4.2 Modifying RCircos core components

Among the three RCircos core components, RCircos cytoband data and RCircos plot positions are calculated based on plot parameter setting. Users can modify RCircos core components by changing plot parameters. Once the plot parameter(s) is changed, call and pass the new parameters to the function of RCircos.Reset.Plot.Parameters(), other two components will be checked for update.

```
> rcircos.params <- RCircos.Get.Plot.Parameters();
> rcircos.params$base.per.unit <- 30000;
> RCircos.Reset.Plot.Parameters(rcircos.params);
> RCircos.List.Parameters();
```

5 Making a Plot with RCircos

Plotting with RCircos is a stepwise process. First, an initialization step is needed. Then, tracks and other aspects of the plot are added sequentially. The result is available after the plot has been entirely constructed. The next subsections walk through the process in detail.

5.1 Initialize Graphic Device

RCircos provides a set of graphic plot functions but does not handle graphic devices. To make RCircos plots, a graphic device has to be opened first. Currently, RCircos works with files supported by R graphics package such as tiff, png, pdf images as well as GUI windows. For example, to make a pdf file with RCircos plot image:

```
> out.file <- "RCircosDemoHumanGenome.pdf";
> pdf(file=out.file, height=8, width=8, compress=TRUE);
> RCircos.Set.Plot.Area();
```

Note: RCircos.Set.Plot.Area() will setup plot area base on total number of tracks inside and outside of chromosome ideogram. User can also setup plot area by summit the R plot commands for user defined plot area, for example:

```
> par(mai=c(0.25, 0.25, 0.25, 0.25));
> plot.new();
> plot.window(c(-2.5,2.5), c(-2.5, 2.5));
```

Note: After everything is done, the graphic device needs to be closed with dev.off().

5.2 Plot Chromosome Ideogram

For RCircos plot, a common first step is to draw chromosome ideograms and label chromosomes with names and highlights. After the RCircos core components were initialized and graphic device was open, simply call the function of RCircos.Chromosome.Ideogram.Plot() will add the chromosome ideogram to the current plot.

> RCircos.Chromosome.Ideogram.Plot();

5.3 Gene Labels and connectors on RCircos Plot

Due to the resolution issues, only limited number of gene names can be labeled. For best visualization, cex should be no less than 0.4 when draw gene labels. When cex is set to 0.4, width of character will be 5000 chromosome units when each unit covers 3000 base pairs. If the gene name list supplied is too long, it will be truncated to fit the chromosome length. Also the long gene name will span more than one track so one or more tracks may be needed to skip for next track.

Connectors are used to mark a genomic position with their names or variant status. Currently, RCircos only provide connector plot between genes and their genomic positions. The following code draw connectors on the first track inside chromosome ideogram and plot gene names on the next track.

```
> data(RCircos.Gene.Label.Data);
> name.col <- 4;
> side <- "in";
> track.num <- 1;
> RCircos.Gene.Connector.Plot(RCircos.Gene.Label.Data,
+ track.num, side);
> track.num <- 2;
> RCircos.Gene.Name.Plot(RCircos.Gene.Label.Data,
+ name.col,track.num, side);
```

5.4 Heatmap, Histogram, Line, Scatter, and Tile Plot

Heatmap, histogram, line, scatter, and tile plot with RCircos require that the first three columns of input data are genomic position information in the order of chromosome name, start, and end position. RCircos provides one function for each type of plots and each function will draw one data track. User can simply call each function with appropriate arguments such as plot location (which track and which side of chromosome ideogram). No more data processing needed.

The code below will draw data tracks of heatmap, scatter, line, histogram, and tile plots.

```
> data(RCircos.Heatmap.Data);
> data.col <- 6;</pre>
```

```
> track.num <- 5;
> side <- "in";
> RCircos.Heatmap.Plot(RCircos.Heatmap.Data, data.col,
                           track.num, side);
> data(RCircos.Scatter.Data);
> data.col <- 5;
> track.num <- 6;</pre>
> side <- "in";
> by.fold <- 1;
> RCircos.Scatter.Plot(RCircos.Scatter.Data, data.col,
                           track.num, side, by.fold);
> data(RCircos.Line.Data);
> data.col <- 5;
> track.num <- 7;
> side <- "in";
> RCircos.Line.Plot(RCircos.Line.Data, data.col,
                           track.num, side);
> data(RCircos.Histogram.Data);
> data.col <- 4;
> track.num <- 8;
> side <- "in";</pre>
> RCircos.Histogram.Plot(RCircos.Histogram.Data,
                           data.col, track.num, side);
> data(RCircos.Tile.Data);
> track.num <- 9;
> side <- "in";
> RCircos.Tile.Plot(RCircos.Tile.Data, track.num, side);
```

5.5 Links: A Special Plot

Links presents relationship of two genomic positions and it is always the last track inside chromosome ideogram. Different from other data plots, input data for links plot is a data frame with paired genomic positions in the order of chromosome, start, and end position for each one genomic position. RCircos supports two types of link plot: lines and ribbons. Link lines are used for presenting relationship of two small genomic regions and ribbons are plotted for bigger genomic regions. Colors for links between chromosomes or same chromosomes could be modified by defining by chromosome=TRUE (or FALSE).

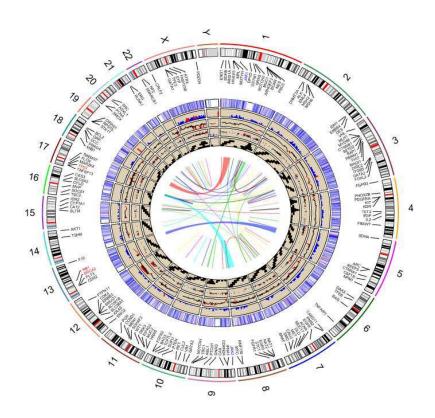
Following code draw link lines and ribbons in the center of plot area.

```
> data(RCircos.Link.Data);
> track.num <- 11;</pre>
```

```
> RCircos.Link.Plot(RCircos.Link.Data, track.num, TRUE);
> data(RCircos.Ribbon.Data);
> RCircos.Ribbon.Plot(ribbon.data=RCircos.Ribbon.Data,
+ track.num=11, by.chromosome=FALSE, twist=FALSE);
> dev.off();
```

Run all code above will generate an image like below.

RCircos Demo with Human Genome



6 More Information

Several demo samples are included in the package. Simply run following demos to see how the RCircos works for simple and complex RCircos plot.

```
> library(RCircos);
> demo("RCircos.Demo.Human");
> demo("RCircos.Demo.Mouse.And.Rat");
```

7 sessionInfo

```
> sessionInfo()
R version 3.2.2 (2015-08-14)
Platform: x86_64-pc-linux-gnu (64-bit)
Running under: Red Hat Enterprise Linux Server release 6.7 (Santiago)
locale:
[1] C
attached base packages:
[1] stats graphics grDevices utils datasets methods base
other attached packages:
[1] RCircos_1.1.3
loaded via a namespace (and not attached):
[1] tools_3.2.2
```

References

[1] Krzywinski, Martin I and Schein, Jacqueline E and Birol, Inanc and Connors, Joseph and Gascoyne, Randy and Horsman, Doug and Jones, Steven J and Marra, Marco A. Circos: An information aesthetic for comparative genomics. *Genome Research*, 2009.