

# GRAPH Interaction from pathway Topological Environment

Gabriele Sales (gabriele.sales@unipd.it)  
Enrica Calura (enrica.calura@gmail.com)  
Chiara Romualdi (chiara.romualdi@unipd.it)

October 13, 2015

## Contents

---

|          |  |          |
|----------|--|----------|
| <b>1</b> | <b>Introduction</b>                                | <b>1</b> |
| <b>2</b> | <b>Pathway topology conversion to gene network</b> | <b>1</b> |
| 2.1      | Pathway definition . . . . .                       | 2        |
| 2.2      | Nodes with multiple elements . . . . .             | 2        |
| 2.3      | Compound mediated interactions . . . . .           | 2        |
| 2.4      | Relation attributes . . . . .                      | 3        |
| 2.5      | Pathway functions recovery . . . . .               | 3        |
| <b>3</b> | <b>Graph</b>                                       | <b>4</b> |
| <b>4</b> | <b>Identifiers</b>                                 | <b>5</b> |
| <b>5</b> | <b>Cytoscape Plot</b>                              | <b>6</b> |
| <b>6</b> | <b>Topological pathway analysis</b>                | <b>6</b> |
| 6.1      | SPIA . . . . .                                     | 6        |
| 6.2      | DEGraph . . . . .                                  | 7        |
| 6.3      | topologyGSA . . . . .                              | 7        |
| 6.4      | clipper . . . . .                                  | 8        |

## 1 Introduction

---

*graphite* (GRAPH Interaction from pathway Topological Environment) an R package built to i) provide networks from six databases (KEGG, [1]; Biocarta, [www.biocarta.com](http://www.biocarta.com); Reactome, [2]; NCI/Nature Pathway Interaction Database, [3]; HumanCyc, [10]; Panther, [11]) and, since the version 1.14, from 14 species; ii) discriminate between different types of gene groups; iii) propagate pathway connections through chemical compounds; iv) allow the selection of edge attributes and the conversion of nodes identifiers to EntrezGene IDs and HUGO Symbols; finally, v) to run *SPIA*, *DEGraph*, *ClipPER* and *topologyGSA* analyses directly on *graphite* networks.

## 2 Pathway topology conversion to gene network

---

In order to gather curated information about human pathways, we have collected data from the four public databases that have emerged as reference points for the systems biology community. The KEGG database has been in development by Kanehisa Laboratories since 1995, and is now a prominent reference knowledge base for integration and interpretation

of large-scale molecular data sets generated by genome sequencing and other high-throughput experimental technologies. KEGG is the only pathway database not in biopax format, they use the KGML format. Reactome, backed by the EBI, is one of the most complete repository; it is frequently updated and provides a semantically rich description of each pathway [2]. KEGG Pathways (KGML format) provides maps for both signaling and metabolic pathways [1]. BioCarta ([www.biocarta.com](http://www.biocarta.com)) is a developer, supplier and distributor company of reagents and assays for biopharmaceutical and academic research. Through an "open source" approach, this community-fed forum constantly integrates emerging proteomic information from the scientific community. It also catalogs and summarizes important resources providing information for over 120,000 genes. BioCarta pathway data in biopax format are available through NCI website. NCI (NCI/Nature Pathway Interaction Database [3]) is a highly-structured, curated collection of information about known biomolecular interactions and key cellular processes assembled into signaling pathways. This was a collaborative project between the NCI and Nature Publishing Group (NPG). Finally, Panther [11] data are a comprehensive, curated database of pathways, protein families, trees, subfamilies and functions available at <http://pantherdb.org> backed by the University of Southern California. HumanCyc is part of the BioCyc database collection of pathways [10].

## 2.1 Pathway definition

KEGG database provides separate xml files, one for each pathway, thus, a pathway is defined by all the reactions defined within each file. For the other databases, we identify a pathway every time in the BioPax format is defined the xml tag pathway.

## 2.2 Nodes with multiple elements

Within a pathway nodes often correspond to multiple gene products. These can be divided into protein complexes (proteins linked by protein-protein interactions) and the groups containing alternative members (genes generally with similar biochemical functions). Thus, when considering signal propagation these groups should be considered differently; the first (hereafter group AND) should be expanded into a clique (all proteins connected to the others), while the second (hereafter group OR) should be expanded without connection among the contained elements.

In the KGML format there are two ways of defining nodes with multiple elements: protein complexes (group AND defined by entry type="group") and group with alternative members (group OR defined by entry type="gene"). In the BioPax format only one type of group is allowed: protein complexes (group AND) with the owl tag complex. However, it often happened that protein tag contains multiple xref pointing to alternative elements of the process (group OR).

## 2.3 Compound mediated interactions

Compound mediated interactions are interactions for which a compound acts as a bridge between two elements. Since chemical compounds are generally not measured with high-throughput technology, they should be removed from the network. However, the trivial elimination of the compounds, without signal propagation, will strongly bias the topology interrupting the signals that pass through them. If element  $A$  is linked to compound  $c$  and compound  $c$  is linked to element  $B$ , thus elements  $A$  should be linked to elements  $B$ .

Within the KGML format there are two different ways of describing a compound mediated interaction: i) direct interaction type="PPrel" ( $A$  interacts whit  $B$  through compound  $c$ ) and ii) indirect one type="PCrel" ( $A$  interacts whit compound  $c$  and  $c$  interacts whit  $B$ ). In the BioPax owl language, on the other hand, only a indirect way of defining compound mediated signals is available.

Since signal propagation reconstruction is crucial for topological gene set analysis we decide to include additional rules for the propagation reconstructing a connection between two genes connected through a series of chemical compounds. Not all compound are considered for the propagation because some of them, such as Hydrogen, H<sub>2</sub>O, ATP, ADP etc., are highly frequent in map descriptions and the signal propagation through them would lead to degenerate too long chains of compounds. The compounds not considered for propagation are not characteristic of a specific reaction but secondary substrates/products widely shared among different processes.

## 2.4 Relation attributes

*graphite* allows the user to see the single/multiple relation types that characterized an edge. The type of edges have been conserved as much as possible to those annotated in the data formats. Some new types have been introduced due to topological conversion needs.

## 2.5 Pathway functions recovery

Human pathway data are natively distributed with the package. But, since the version 1.14.0, non-human pathway data are also available and can be downloaded using *pathways*

*pathways* requires the name of the specie of interest and the name of the pathway database name as follow:

```
> humanReactome <- pathways("hsapiens", "reactome")
> names(humanReactome)[1:10]

[1] "2-LTR circle formation"
[2] "3' -UTR-mediated translational regulation"
[3] "Abacavir metabolism"
[4] "Abacavir transmembrane transport"
[5] "Abacavir transport and metabolism"
[6] "Abasic sugar-phosphate removal via the single-nucleotide replacement pathway"
[7] "ABC-family proteins mediated transport"
[8] "ABC transporters in lipid homeostasis"
[9] "Abortive elongation of HIV-1 transcript in the absence of Tat"
[10] "Acetylcholine Binding And Downstream Events"

> p <- humanReactome[["ABC-family proteins mediated transport"]]
> p

"ABC-family proteins mediated transport" pathway
Native ID      = R-HSA-382556
Database       = Reactome
Species        = hsapiens
Type of identifiers = UNIPROT
Number of nodes = 21
Number of edges = 40
Retrieved on   = 07-10-2015
```

A pathway database is a list of pathways. We can access to a pathway through its name, as above, or through its position in the list, as follow:

```
> p <- humanReactome[[21]]
> p@title

[1] "Activation of APC/C and APC/C:Cdc20 mediated degradation of mitotic proteins"
```

In the pathway, nodes represent genes/proteins and the edges represent the functional relationships:

```
> head(nodes(p))

[1] "A5LHX3" "000231" "000232" "000233" "000487" "000762"
```

Edges can be characterized by multiple functional relationships:

```
> head(edges(p))

      src  dest direction      type
1 A5LHX3 000231 undirected Binding
2 A5LHX3 000487 undirected Binding
```

```

3 A5LHX3 000762 directed Control(Out: ACTIVATION of BiochemicalReaction)
4 A5LHX3 043684 directed Control(Out: ACTIVATION of BiochemicalReaction)
5 A5LHX3 060566 directed Control(Out: ACTIVATION of BiochemicalReaction)
6 A5LHX3 P06493 directed Control(Out: ACTIVATION of BiochemicalReaction)

```

This same steps can be used to access the KEGG, Biocarta, Panther, HumanCyc and NCI databases for Homo sapiens pathways (through the kegg, biocarta, panther, humancyc and nci lists, respectively), but not all the databases are present for all the species. To know the pathway data available the user can use the `pathwayDatabases`.

```

> pathwayDatabases()

      species database
1    athaliana   kegg
2    athaliana reactome
3     btaurus   kegg
4     btaurus reactome
5    celegans reactome
6   cfamiliaris   kegg
7   cfamiliaris reactome
8 dmelanogaster   kegg
9 dmelanogaster reactome
10     drerio   kegg
11     drerio reactome
12      ecoli   kegg
13    ggallus   kegg
14    ggallus reactome
15   hsapiens biocarta
16   hsapiens humancyc
17   hsapiens   kegg
18   hsapiens   nci
19   hsapiens panther
20   hsapiens reactome
21  mmusculus   kegg
22  mmusculus reactome
23 rnorvegicus   kegg
24 rnorvegicus reactome
25 scerevisiae   kegg
26 scerevisiae reactome
27   sscrofa   kegg
28   sscrofa reactome
29    xlaevis   kegg

```

### 3 Graph

---

The function `pathwayGraph` builds a *graphNEL* object from a pathway `p`:

```

> g <- pathwayGraph(p)
> g

```

A *graphNEL* graph with directed edges

Number of Nodes = 73

Number of Edges = 3758

```

> edgeData(g)[1]

```

```
$`A5LHX3|000231`
$`A5LHX3|000231`$weight
[1] 1

$`A5LHX3|000231`$edgeType
[1] "Binding"
```

## 4 Identifiers

---

Gene annotations databases are widely used as public repositories of biological information. Our current knowledge on biological elements is spread out over a number of databases (such as: Entrez Gene , RefSeq, backed by the NCBI <http://www.ncbi.nlm.nih.gov/>, UniProt, ENSEMBL backed by the EBI <http://www.ebi.ac.uk/> to name just a few), specialised on a subset of specific biological entities (for instance, UniProt focuses on proteins while Entrez Gene focuses on genes). Key identifiers (IDs) in the internal structure of each database uniquely represent biological entities, thus biological entities can be identified by homogeneous IDs according to the selected database they refer to. Due to their different origins and specificity, switching from an ID to another is possible but not trivial: there could be either no correspondence between them or many-to-many relations. For detailed information about IDs, their structures and differences please consult those resources.

The function `converterIdentifiers` allows the user to convert the pathway IDs into different types, to fit the user needs. This mapping process, however, may lead to the loss of some nodes (not all identifiers may be recognized) and has an impact on the topology of the network (one ID may correspond to multiple IDs in another annotation or vice versa). We based the function of ID conversion through the species-specific Bioconductor [AnnotationDbi](#), such as [org.Hs.eg.db](#). Since the version 1.14, all the conversion supported in the annotation packages can be used. `converterIdentifiers` needs a list of pathways or a single pathway and a string describing the type of the identifier as provided by an Annotation package (for example, "ENTREZID"), the values "entrez", "symbol" remains for backward compatibility.

```
> pSymbol <- convertIdentifiers(p, "SYMBOL")
> pSymbol

"Activation of APC/C and APC/C:Cdc20 mediated degradation of mitotic proteins" pathway
Native ID           = R-HSA-176814
Database            = Reactome
Species             = hsapiens
Type of identifiers = SYMBOL
Number of nodes     = 73
Number of edges     = 2933
Retrieved on       = 07-10-2015

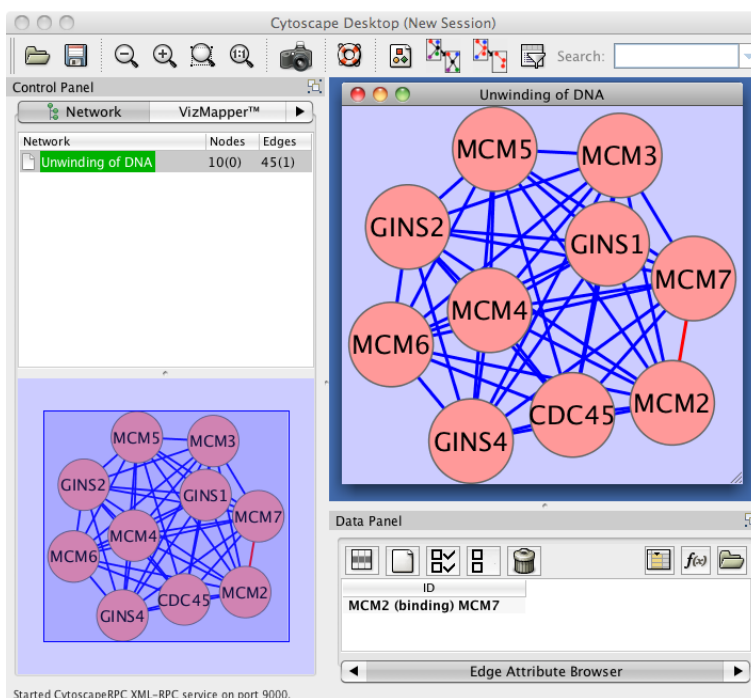
> head(nodes(pSymbol))
[1] "PSMD5" "PSMA1" "PSMA5" "PSMB9" "PTTG1" "PSMD1"

> reactomeSymbol <- convertIdentifiers(humanReactome[1:5], "SYMBOL")
```

## 5 Cytoscape Plot

Several pathways have a huge number of nodes and edges, thus there is the need of an efficient system of visualization. To this end *graphite* uses the *Rcytoscape* package to export the network to Cytoscape. Cytoscape is a Java based software specifically built to manage biological network complexity and for this reason it is widely used by the biological community. Unfortunately, since *Rcytoscape* use CytoscapeRCP plugin, which is not available for the new Cytoscape 3.0 version, the use of *cytoscapePlot* is currently limited to Cytoscape 2.8 and lower versions.

```
> cytoscapePlot(convertIdentifiers(reactome$`Unwinding of DNA`, "symbol"))
```



## 6 Topological pathway analysis

*graphite* gives access to three types of topological pathway analyses recently proposed. For more details on the results obtained by these analyses see the corresponding R packages.

### 6.1 SPIA

The analysis with *SPIA* requires the conversion of the networks in a suitable format. This conversion is performed by the function *prepareSPIA* that must be executed before the analysis command *runSPIA*. The *SPIA* data will be saved in the current working directory; every time you change it, you should also re-run *prepareSPIA*. Edges not included in *SPIA* have been coerced into the admitted *SPIA* types. Compound mediated interactions annotated in *graphite* with "indirect" type are mapped into the *SPIA* edge type "indirect effect" by default set to zero. To use the signal propagated through compounds type 1 in "indirect effect".

```
> library(SPIA)
> data(colorectalcaner)
> library(hgu133plus2.db)
> x <- hgu133plus2ENTREZID
> top$ENTREZ <- unlist(as.list(x[top$ID]))
```

```

> top <- top[!is.na(top$ENTREZ), ]
> top <- top[!duplicated(top$ENTREZ), ]
> tg1 <- top[top$adj.P.Val < 0.05, ]
> DE_Colorectal = tg1$logFC
> names(DE_Colorectal) <- as.vector(tg1$ENTREZ)
> ALL_Colorectal <- top$ENTREZ
> prepareSPIA(biocarta[1:2], "biocartaEx")
> res <- runSPIA(de=DE_Colorectal, all=ALL_Colorectal, "biocartaEx")

```

Done pathway 1 : acetylation and deacetylation ..

Done pathway 2 : actions of nitric oxide in the..

```
> res
```

|   | Name   | pSize    | NDE      | pNDE       | tA         | pPERT |
|---|--|----------|----------|------------|------------|-------|
| 1 | acetylation and deacetylation of rela in nucleus | 5        | 3        | 0.06107832 | -0.2757872 | 0.919 |
|   | pG   | pGFdr    | pGFWER   | Status     |            |       |
| 1 | 0.217792   | 0.217792 | 0.217792 | Inhibited  |            |       |

For more details see the [SPIA](#) package [5, 6, 4].

## 6.2 DEGraph

[DEGraph](#) implements recent hypothesis testing methods which directly assess whether a particular gene network is differentially expressed between two conditions.

In [graphite](#), [DEGraph](#) has a dedicated function, called `runDEGraph`, which performs the analysis on a single pathway or on a pathway list.

```

> library(DEGraph)
> data("Loi2008_DEGraphVignette")
> biocarta <- pathways("hsapiens", "biocarta")
> p <- convertIdentifiers(biocarta[["actions of nitric oxide in the heart"]], "ENTREZID")
> res <- runDEGraph(p, exprLoi2008, classLoi2008)
> res$`1`

```

\$p.value

|  | T2        | T2 (1 Fourier components) |
|--|-----------|---------------------------|
|  | 0.6106982 | 0.3221514                 |

\$graph

A graphNEL graph with directed edges

Number of Nodes = 2

Number of Edges = 1

\$k

[1] 1

The function `runDEGraph`, which easily performs the analysis on the entire pathway database, provides as result a list of two elements: a list with the results of the pathway analyses and the list of generated errors.

For more details see the [DEGraph](#) package [7].

## 6.3 topologyGSA

[topologyGSA](#) uses graphical models to test the pathway components and to highlight those involved in its deregulation.

In *graphite*, *topologyGSA* has a dedicated function, `runTopologyGSA`, which performs the analysis on a single pathway or on a pathway list.

```
> library(topologyGSA)
> data(examples)
> kegg <- pathways("hsapiens", "kegg")
> p <- convertIdentifiers(kegg[["Fc epsilon RI signaling pathway"]], "SYMBOL")
> runTopologyGSA(p, "var", y1, y2, 0.05)
```

Pathway Variance Test

data: exp1, exp2 and g

lambda = 26.02199, df = 10, p-value = 0.003710726, equal variances: TRUE

The function `runTopologyGSA`, which easily performs the analysis on the entire pathway database, provides as result a list of two elements: a list with the results of the pathway analyses and the list of generated errors.

For more details see the *topologyGSA* package [8].

## 6.4 clipper

*clipper* is a package for topological gene set analysis. It implements a two-step empirical approach based on the exploitation of graph decomposition into a junction tree to reconstruct the most relevant signal path. In the first step clipper selects significant pathways according to statistical tests on the means and the concentration matrices of the graphs derived from pathway topologies. Then, it "clips" the whole pathway identifying the signal paths having the greatest association with a specific phenotype.

In *graphite*, *clipper* has a dedicated function, `runClipper`, which performs the analysis on a single pathway or on a pathway list.

```
> library(clipper)
> library(ALL)
> kegg <- pathways("hsapiens", "kegg")
> paths <- convertIdentifiers(kegg[1:5], "ENTREZID")
> genes <- unlist(lapply(paths, nodes))
> data(ALL)
> all <- as.matrix(exprs(ALL[1:length(genes),1:20]))
> classes <- c(rep(1,10), rep(2,10))
> rownames(all) <- genes
> clipped <- runClipper(paths, all, classes, "mean", pathThr=0.1)
> resClip <- do.call(rbind,clipped$results)
> resClip[,1:5]
```

|  | startIdx | endIdx | maxIdx | length | maxScore          |
|--|----------|--------|--------|--------|-------------------|
| Acute myeloid leukemia.1;14            | 1        | 14     | 1      | 9      | 0.339081447250642 |
| Acute myeloid leukemia.1;30            | 1        | 30     | 1      | 16     | 0.244501437839259 |
| Adherens junction                      | 1        | 12     | 2      | 6      | 2.48839241597474  |
| Adipocytokine signaling pathway.1;12   | 1        | 12     | 1      | 12     | 0.575646273248511 |
| Adipocytokine signaling pathway.1;22   | 1        | 22     | 1      | 10     | 0.342950408053159 |
| Adipocytokine signaling pathway.1;23   | 1        | 23     | 1      | 3      | 1.10611558349724  |
| Adipocytokine signaling pathway.1;36   | 1        | 36     | 2      | 12     | 1.20281011203905  |
| Adipocytokine signaling pathway.1;39   | 1        | 39     | 1      | 6      | 0.744328933622543 |
| Adrenergic signaling in cardiomyocytes | 1        | 6      | 3      | 5      | 5.93217779586003  |

The function `runClipper`, which easily performs the analysis on the entire pathway database, provides as result a list of two elements: the list with the results of the pathway analyses and the list of generated errors.



For more details see the [clipper](#) package [9].

## References

---

- [1] Ogata H, Goto S, Sato K, Fujibuchi W, Bono H, Kanehisa M. KEGG: Kyoto Encyclopedia of Genes and Genomes. *Nucleic Acids Res.* 1999 Jan 1;27(1):29-34.
- [2] Matthews L, Gopinath G, Gillespie M, Caudy M, Croft D, de Bono B, Garapati P, Hemish J, Hermjakob H, Jassal B, Kanapin A, Lewis S, Mahajan S, May B, Schmidt E, Vastrik I, Wu G, Birney E, Stein L, D'Eustachio P. Reactome knowledgebase of human biological pathways and processes. *Nucleic Acids Res.* 2009 Jan;37(Database issue):D619-22. Epub 2008 Nov 3.
- [3] Schaefer CF, Anthony K, Krupa S, Buchoff J, Day M, Hannay T, Buetow KH. PID: the Pathway Interaction Database. *Nucleic Acids Res.* 2009 Jan;37(Database issue):D674-9. Epub 2008 Oct 2.
- [4] Draghici, S., Khatri, P., Tarca, A.L., Amin, K., Done, A., Voichita, C., Georgescu, C., Romero, R. A systems biology approach for pathway level analysis. *Genome Research*, 17, 2007.
- [5] Tarca AL, Draghici S, Khatri P, Hassan SS, Mittal P, Kim JS, Kim CJ, Kusanovic JP, Romero R. A novel signaling pathway impact analysis. *Bioinformatics.* 2009 Jan 1;25(1):75-82.
- [6] Adi L. Tarca, Sorin Draghici, Purvesh Khatri, et. al. A Signaling Pathway Impact Analysis for Microarray Experiments. *Bioinformatics*, 2009, 25(1):75-82.
- [7] L. Jacob, P. Neuviat, and S. Dudoit. Gains in power from structured two-sample tests of means on graphs. Technical Report arXiv:q-bio/1009.5173v1, arXiv, 2010.
- [8] Massa MS, Chiogna M, Romualdi C. Gene set analysis exploiting the topology of a pathway. *BMC System Biol.* 2010 Sep 1;4:121.
- [9] Martini P, Sales G, Massa MS, Chiogna M, Romualdi C. Along signal paths: an empirical gene set approach exploiting pathway topology. *Nucleic Acids Res.* 2013 Jan 7;41(1):e19. doi: 10.1093/nar/gks866. Epub 2012 Sep 21.
- [10] Caspi R, Altman T, Dale JM, Dreher K, Fulcher CA, Gilham F, Kaipa P, Karthikeyan AS, Kothari A, Krummenacker M, Latendresse M, Mueller LA, Paley S, Popescu L, Pujar A, Shearer AG, Zhang P, Karp PD. The MetaCyc database of metabolic pathways and enzymes and the BioCyc collection of pathway/genome databases. *Nucleic Acids Research* 38:D473-9 2010.
- [11] PANTHER in 2013: modeling the evolution of gene function, and other gene attributes, in the context of phylogenetic trees. Huaiyu Mi, Anushya Muruganujan and Paul D. Thomas *Nucl. Acids Res.* (2012) doi: 10.1093/nar/gks1118