How To Use iSubpathwayMiner

Chunquan Li

February 21, 2012

Contents

1 Overview

This vignette demonstrates how to easily use the iSubpathwayMiner package. The package can implement the graph-based reconstruction, analyses, and visualization of the KEGG pathways. (1) Our system provides many strategies of converting pathways to graph models (see the section 2). Ten functions related to conversion from pathways to graphs are developed. Furthermore, the combinations of these functions can get many combined conversion strategies of pathway graphs (> 20). (2) The iSubpathwayMiner can support the annotation and identification of pathways based on gene sets (see the section 3.3.1 and 3.4.1), compound sets (see the section 3.3.2), and even the combined sets of genes and compounds (see the section 3.3.3). The entire pathway and subpathway identification methods are available for these sets (see the section 3.3 and 3.4). (3) The system also supports topology-based pathway analysis of these sets (see the section 3.2). The current available topological properties contain degree, local clustering coefficient, closeness and betweenness. (4) We develop KEGG layout style of pathway graphs in R to simulate the layout of the pathway picture in KEGG website (see the section 4). In addition, our system has also provided many types of automatic layout styles. Pathway graphs can also be exported to the GML format supported by Cytoscape [Shannon *et al.*, 2003]. We firstly give several examples as follows:

The following commands can convert two metabolic pathways to graphs.

```
> #get path of the KGML files
> path<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/metabolic/ec/",sep="")
> #convert pathways to a list in R
> pList<-getPathway(path,c("ec00010.xml","ec00020.xml"))
> #convert metabolic pathways to graphs
> gmList<-getMetabolicGraph(pList)
```

The following commands visualize a pathway graph. The result is shown in Figure 1.

```
> #visualize
> plotGraph(gmList[[1]])
```

The following command gets the type of organism and identifier in the current environment variable.

```
> getOrgAndIdType()
```

```
[1] "hsa"         "ncbi-geneid"
```

The following commands annotate gene sets to the above two metabolic pathways and evaluate the enrichment significance of pathways.

```
> #To do this, let us generate an example of gene sets:
> geneList<-getExample(geneNumber=1000,compoundNumber=0)
> #see a part of the set.
> #organism:human (hsa)
> #identifier type:Entrez Gene IDs (ncbi-geneid)
> geneList[1:5]
```

```
[1] "10"    "100"   "1000"  "10000" "10005"
```

```
> #annotate the sets to pathways
> #evaluate the enrichment significance of pathways
> ann<-identifyGraph(geneList,gmList)
> #print the results to screen
> printGraph(ann)
```

```
   pathwayId                  pathwayName annComponentRatio annBgRatio
1 path:00010 Glycolysis / Gluconeogenesis           12/1000   64/21796
2 path:00020    Citrate cycle (TCA cycle)            4/1000   30/21796
        pvalue          fdr
1 2.942795e-05 5.885591e-05
2 4.678027e-02 4.678027e-02
```

2 The methods of graph-based reconstruction of pathways

The section introduces many strategies for converting pathways to different types of graphs. We firstly need to use the function `getPathway` to convert KGML files (KEGG Markup Language, `http://www.genome.jp/kegg/xml/`) of KEGG pathways to a list variable in R, which is used to store pathway data in the iSubpathwayMiner system (see the section 2.1). We can then use the function `getMetabolicGraph` or `getNonMetabolicGraph` to convert metabolic pathways or non-metabolic pathways to graphs (Figure 1 and 2). The function `getMetabolicGraph` constructs graphs based on reaction information of KGML files of pathways (see the section 2.2). The function `getNonMetabolicGraph` constructs graphs based on relation information (see the section 2.3). After using the function `getMetabolicGraph` or `getNonMetabolicGraph` to convert pathways to graphs, users can change these pathway graphs to other derivative graphs. We develop the function `getUGraph`, `mapNode`, `filterNode`, `simplifyGraph`, `mergeNode`, `getSimpleGraph`, and `expandNode` (see the section 2.4). Through these functions, many graph-based reconstruction strategies of pathways can be done such as constructing undirected graphs, organism-specific and idType-specific graphs, the metabolic graphs with enzymes (compounds) as nodes and compounds (enzymes) as edges, etc. Furthermore, the combination of these functions can also get more useful pathway graphs (see the section 2.5). For example, we can construct the directed/undirected pathway graphs of enzyme-compound (see the section 2.5.1), enzyme-enzyme (see the section 2.5.2), KO-KO (see the section 2.5.2), compound-compound (see the section 2.5.3), organism-specific gene-gene (see the section 2.5.4), etc. Most of these conversions represent current major applications [Smart *et al.*, 2008, Schreiber *et al.*, 2002, Klukas and Schreiber, 2007, Kanehisa *et al.*, 2006, Goffard and Weiller, 2007, Koyuturk *et al.*, 2004, Hung *et al.*, 2010, Xia and Wishart, 2010, Jeong *et al.*, 2000, Antonov *et al.*, 2008, Guimera and Nunes Amaral, 2005, Draghici *et al.*, 2007, Li *et al.*, 2009, Ogata *et al.*, 2000, Hung *et al.*, 2010, Barabasi and Oltvai, 2004]. The following sections will detailedly introduce the usage of the functions relative to graph-based conversion of pathways.

2.1 Convert KGML files of KEGG pathways to a list in R

The KEGG Markup Language (KGML) is an exchange format of KEGG pathway data. In a KGML file (.xml), the pathway element is a root element. The entry element stores information about nodes of the pathway, including the attribute information (id, name, type, link, and reaction), the "graphics" subelement, the "component" subelement. The relation element stores information about relationship between gene products (or between gene products and compounds). It includes the attribute information (entry1, entry2, and type), and the "subtype" subelement that specifies more detailed information about the interaction. The reaction element stores chemical reaction between a substrate and a product. It includes the attribute information (id, name, and type), the "substrate" subelement, and the "product" subelement. Detailed information is provided in `http://www.genome.jp/kegg/xml/docs/`.

In KEGG, there are two fundamental controlled vocabularies for matching genes to pathways. Enzyme commission (EC) numbers are traditionally used as an effective vocabulary for annotating genes to metabolic pathways. With the rapid development of KEGG, more and more non-metabolic pathways including genetic information processing, environmental information processing and cellular processes have been added to KEGG PATHWAY database. KEGG Orthology (KO) identifiers, which overcome limitations of enzyme nomenclature and integrate the pathway and genome information, have become a better controlled vocabulary for annotating genes to both metabolic and regulatory pathways [Kanehisa *et al.*, 2006]. Therefore, KEGG has provided the KGML files of reference metabolic pathways linked to EC identifiers, reference metabolic pathways linked to KO identifiers, and reference non-metabolic pathways linked to KO identifiers. They can be obtained from KEGG ftp site.

The function `getPathway` can convert the above KGML files to a list variable in R, which is used as pathway data in our system. The conversion only changes data structure in order to efficiently operate data in R environment. After conversion, most of original information about pathways are not ignored although data structure changed. The list that stores pathway information will be used as the input of other functions such as `getMetabolicGraph` and `getNonMetabolicGraph`. The following commands can convert KGML files of metabolic pathways to a list in R.

```
> #get path of the KGML files
> path<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/metabolic/ec/",sep="")
> #convert pathways to a list in R
> p<-getPathway(path,c("ec00010.xml","ec00020.xml"))
```

2.2 Convert metabolic pathways to graphs

2.2.1 The method to convert metabolic pathways to graphs

The function `getMetabolicGraph` can convert metabolic pathways to graphs. A result graph mainly contains three types of nodes: compounds, gene products (enzymes, KOs, or genes encoding them), and maps that represent pathways linked with the current pathway. Edges are mainly constructed from reactions. Specially, if a compound participates in a reaction as a substrate or product, a directed edge connects the compound node to the reaction node (enzymes, KOs, or genes). That is, substrates of a reaction are connected to the reaction node (enzymes, KOs, or genes) and the reaction node is connected to products. For substrates, they are directed toward the reaction node. For products, the reaction node is directed toward them. Reversible reactions have twice edges of irreversible reactions. The conversion strategy of pathway graphs has the advantage that graph algorithms and standard graph drawing techniques can be used. More importantly, almost all information can be efficiently stored in the kind of graph model. The similar strategy is also adopt by many study groups [Smart *et al.*, 2008, Klukas and Schreiber, 2007, Goffard and Weiller, 2007, Goffard and Weiller, 2007, Koyuturk *et al.*, 2004].

In addition, a compound and a linked map will be connected by an edge if they have relationships get from relation element of the KGML file. Other information such as node attribute, pathway attribute (e.g., pathway name), etc. are converted to attribute of graph.

The following commands can convert metabolic pathways to graphs.

```
> #get path of the KGML files
> path<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/metabolic/ec/",sep="")
> #convert pathways to a list in R
> p<-getPathway(path,c("ec00010.xml"))
> #convert metabolic pathways to graphs
> gm<-getMetabolicGraph(p)
```

The following commands can visualize the graph of the Glycolysis / Gluconeogenesis pathway. Figure 1 shows the result graph. In the figure, the blue rectangle nodes represent enzymes. The circle nodes represent compounds. The white rectangle nodes represent maps.

```
> #name of graph gm[[1]]
> gm[[1]]$title

[1] "Glycolysis / Gluconeogenesis"

> #visualize
> plotGraph(gm[[1]])
```

For a pathway graph, the function `summary` can print the number of nodes and edges, names of node and edge attributes, and whether the graph is directed as follows:

```
> summary(gm[[1]])

Vertices: 94
Edges: 183
Directed: TRUE
Graph attributes: name, number, org, title, image, link.
Vertex attributes: name, id, names, type, reaction, link, graphics_name, graphics_fgcolor, graphics_bg
No edge attributes.
```

The function `print` can display the information similar to the function `summary`. In addition, the function also displays edges, graph attributes, node attributes, and edge attributes. The following command prints all information of a pathway graph:

```
> print(gm[["00010"]],v=TRUE,e=TRUE,g=TRUE)
```

Because the pathway graph is usually too large, here we only display its subgraph with five nodes in order to save page space.

```
> #display a subgraph with 5 nodes.
> sgm<-subgraph(gm[[1]],V(gm[[1]])[1:5])
> print(sgm,g=TRUE,v=TRUE,e=TRUE)

Vertices: 5
Edges: 5
Directed: TRUE
Graph attributes:
```

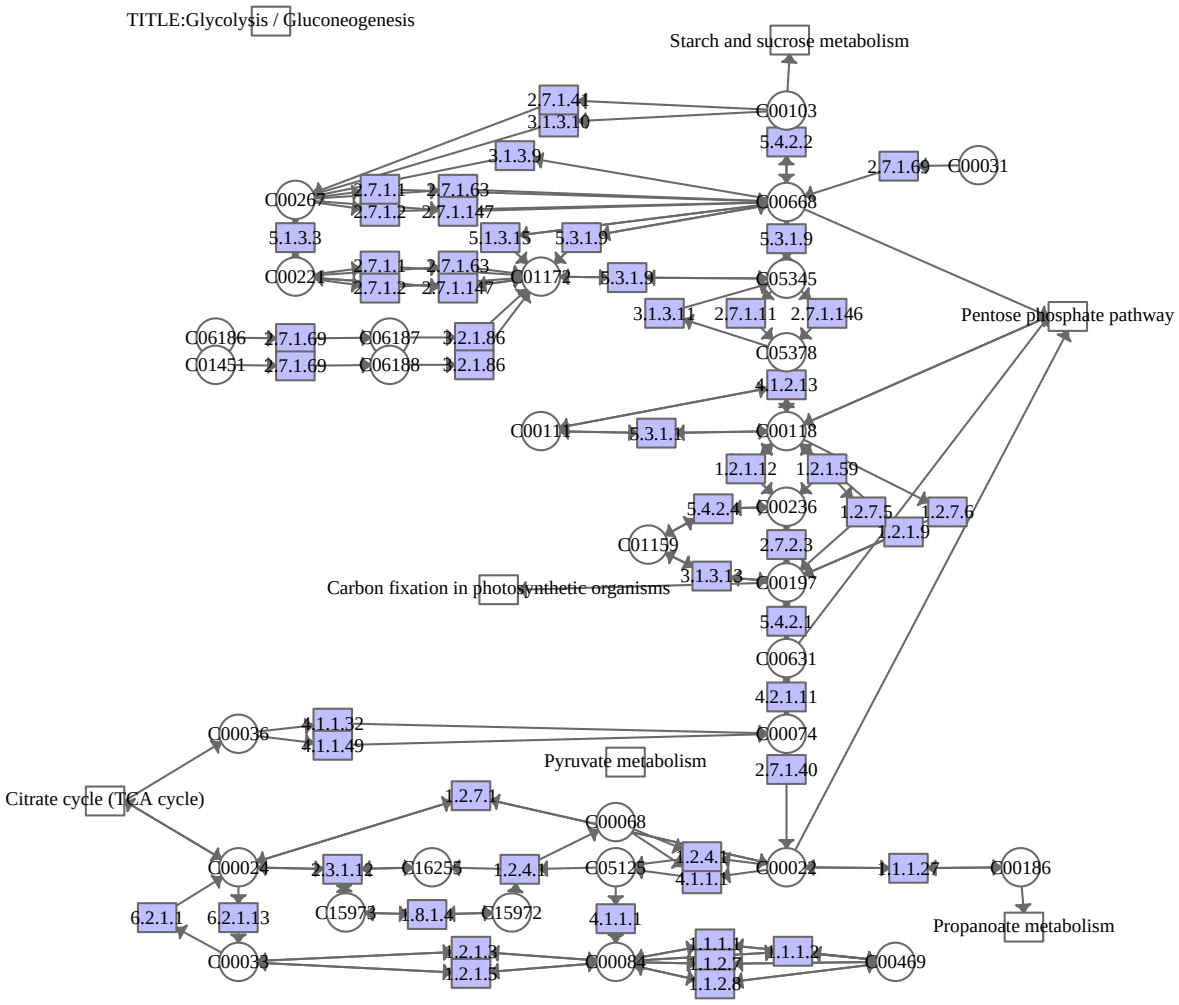

Figure 1: The Glycolysis / Gluconeogenesis pathway graph.

```
[[name]]
[1] "path:ec00010"
[[number]]
[1] "00010"
[[org]]
[1] "ec"
[[title]]
[1] "Glycolysis / Gluconeogenesis"
[[image]]
[1] "http://www.genome.jp/kegg/pathway/ec/ec00010.png"
[[link]]
[1] "http://www.genome.jp/kegg-bin/show_pathway?ec00010"
Vertex attributes:
    name id        names     type  reaction
[0]   37 37   ec:1.2.1.3   enzyme rn:R00710
[1]   38 38  ec:6.2.1.13   enzyme rn:R00229
[2]   39 39   ec:1.2.1.5   enzyme rn:R00711
[3]   40 40   cpd:C00033 compound    unknow
[4]   41 41 path:ec00030      map    unknow
                                          link             graphics_name
[0]  http://www.kegg.jp/dbget-bin/www_bget?1.2.1.3                 1.2.1.3
[1] http://www.kegg.jp/dbget-bin/www_bget?6.2.1.13                6.2.1.13
[2]  http://www.kegg.jp/dbget-bin/www_bget?1.2.1.5                 1.2.1.5
[3]   http://www.kegg.jp/dbget-bin/www_bget?C00033                  C00033
[4]  http://www.kegg.jp/dbget-bin/www_bget?ec00030 Pentose phosphate pathway
    graphics_fgcolor graphics_bgcolor  graphics_type graphics_x graphics_y
[0]          #000000          #BFBFFF      rectangle        289        943
[1]          #000000          #BFBFFF      rectangle        146        911
[2]          #000000          #BFBFFF      rectangle        289        964
[3]          #000000          #FFFFFF         circle        146        953
[4]          #000000          #FFFFFF roundrectangle        656        339
    graphics_width graphics_height graphics_coords
[0]             46              17          unknow
[1]             46              17          unknow
[2]             46              17          unknow
[3]              8               8          unknow
[4]             62             237          unknow
Edges and their attributes:

[0] '37' -> '40'
[1] '40' -> '37'
[2] '38' -> '40'
[3] '39' -> '40'
[4] '40' -> '39'
```

2.2.2 Some simple examples of operating pathway graphs

Since pathways can be converted to graphs, many analyses based on graph model are available by using the functions provided in the igraph package. For example, we can get subgraph, degree, shortest path, etc. Detailed information will be introduced in the section 3. Here, we only give some examples of operating graphs, which are very important for effectively interpreting and operating pathway graphs.

We can get the name and number of the pathway, as follows:

```
> gm[[1]]$title

[1] "Glycolysis / Gluconeogenesis"

> gm[[1]]$number

[1] "00010"
```

We can get the attribute value of a node. In all attributes, the "names" attribute is the most important. It makes us able to identify the cellular components the node includes. Its values are usually the identifiers of compound, enzyme, gene, or KO, etc. The following commands can get "names" attribute of the second node:

```
> V(gm[[1]])[2]$names

[1] "ec:6.2.1.13"
```

The result shows that the second node is the enzyme identifier. We can also use another method to get "names" attribute of the node

```
> get.vertex.attribute(gm[[1]],"names",2)

[1] "ec:6.2.1.13"
```

We can get other attributes. For example, the following command gets the "type" attribute of the second node:

```
> V(gm[[1]])[2]$type

[1] "enzyme"
```

The result shows that the second node is the enzyme.

An important application is to identify some nodes that meet the certain conditions. For example, one is likely to want to find the enzyme "ec:4.1.2.13" and "ec:1.2.1.59" in pathway graph "00010", and then calculate the shortest path between them in the graph. One may also want to identify the enzyme "ec:4.1.2.3", and then calculate its betweenness, which represents the importance of the node.

In order to do these, one firstly needs to get indexes of interesting nodes. Node indexes are used as input of most of functions in igraph package. We then use functions in the igraph package (e.g., get.shortest.paths, betweenness, etc.) to get the analysis results. The following commands get indexes of nodes with "names"="ec:4.1.2.13" and "ec:1.2.1.59" in graph "00010", then calculate shortest path of them.

```
> #get indexes of nodes
> index1<-V(gm[[1]])[V(gm[[1]])$names=="ec:4.1.2.13"]
> index2<-V(gm[[1]])[V(gm[[1]])$names=="ec:1.2.1.59"]
> #get shortest path
> shortest.path<-get.shortest.paths(gm[[1]],index1,index2)
> #display shortest path
> shortest.path

[[1]]
[1]  0 88 80
```

```
> #convert indexes to names
> V(gm[[1]])[shortest.path[[1]]]$names
```

```
[1] "ec:4.1.2.13" "cpd:C00118"  "ec:1.2.1.59"
```

Calculate betweenness of the enzyme "ec:4.1.2.3".

```
> index1<-V(gm[[1]])[V(gm[[1]])$names=="ec:4.1.2.13"]
> betweenness(gm[[1]],index1)
```

```
[1] 1756.746
```

Note that we should see node index value using the function `as.integer`. The direct display is not real node index value, but the value of the "id" attribute of nodes.

```
> #node index value
> as.integer(index1)
```

```
[1] 0
```

```
> #direct display is not real node index value.
> index1
```

```
Vertex sequence:
[1] "13"
```

```
> #it is equal to the value of the "id" attribute.
> index1$id
```

```
[1] "13"
```

2.3 Convert non-metabolic pathways to graphs

2.3.1 The default method to convert non-metabolic pathways to graphs

The function `getNonMetabolicGraph` can convert non-metabolic pathways to directed graphs. An result graph mainly contains two types of nodes: gene products (KOs) and maps that represent pathways linked with the pathway graph. Sometimes, there are several compounds in pathways such as IP3, DAG, cAMP, ca+, etc. Edges are obtained from relations. In particular, two nodes are connected by an edge if they have relationships get from relation element of the KGML file. The relation element specifies relationships between nodes. For example, the attribute PPrel represents protein-protein interaction such as binding and modification. Other information such as node attribute, pathway attribute, etc. is converted to attribute of graphs. The following commands can convert non-metabolic pathways to graphs. The result graph of the MAPK signaling pathway is shown in Figure 2.

```
> #get path
> pathn<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/non-metabolic/ko/",sep="")
> pn<-getPathway(pathn,c("ko04010.xml","ko04020.xml"))
> #Convert pathways to graphs
> gn1<-getNonMetabolicGraph(pn)
> #name of the first pathway
> gn1[[1]]$title
```

```
[1] "MAPK signaling pathway"
```

```
> #visualize
> plotGraph(gn1[[1]])
```

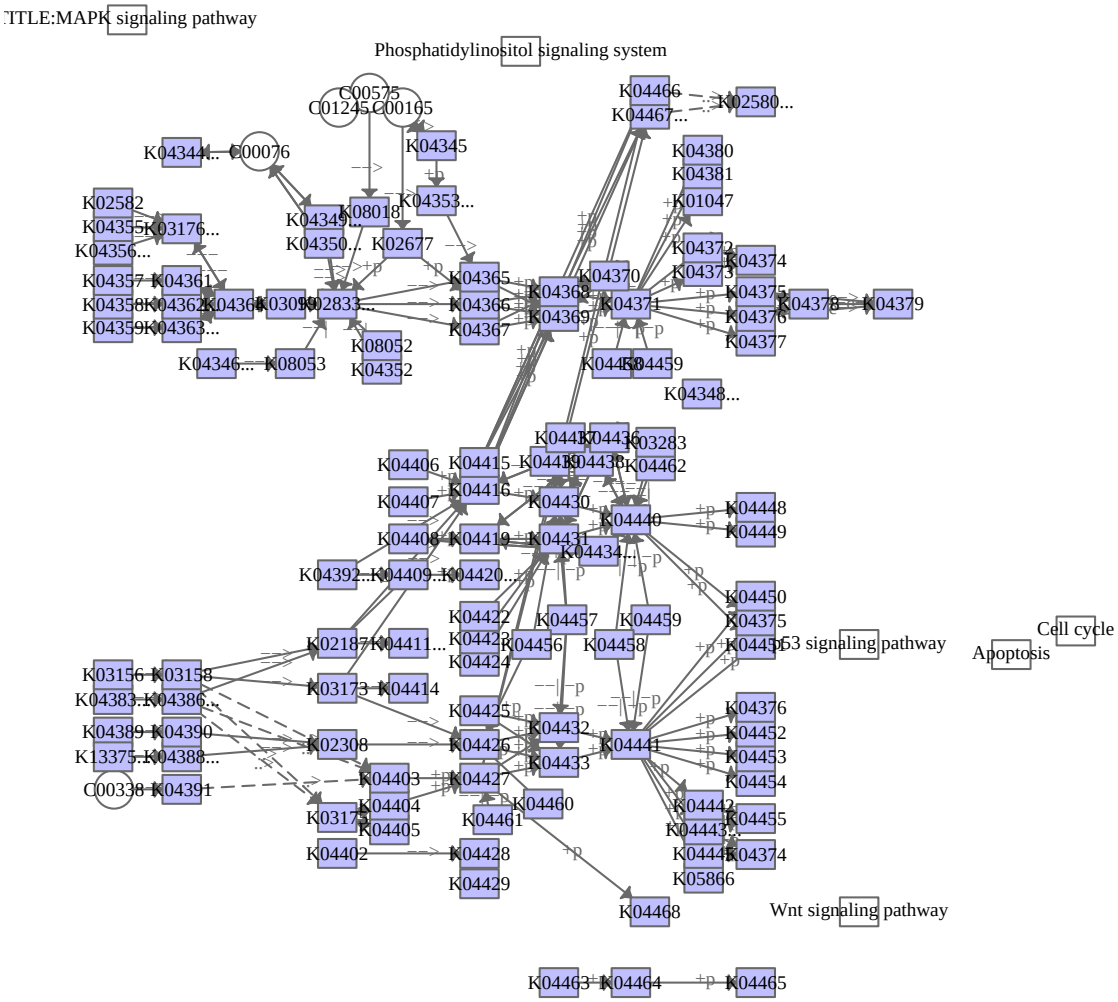

Figure 2: The MAPK signaling pathway graph with ambiguous edges as bi-directed.

2.3.2 The alternative method to convert non-metabolic pathways to graphs

In non-metabolic pathways, there are usually many different types of edges between nodes. There are four fundamental types of edges including ECrel (enzyme-enzyme relation), PPrel (protein-protein interaction), GErel (gene expression interaction) and PCrel (protein-compound interaction). Each fundamental type usually contains many subtypes such as compound, hidden compound, activation, inhibition, expression, repression, indirect effect, state change, binding/assoction, dissociation, and missing interaction. Detailed information is provided in `http://www.genome.jp/kegg/xml/docs/`.

According to these substypes, we can obtain edge direction. For example, "activation" means that protein A activates B (A–>B). However, not all types of edges have definite direction. For example, "binding/association" means that there is the binding or association relation between protein A and protein B but we don't know A–>B or B–>A. In addition, an edge is also likely to have no subtype and thus we can't know its direction. The argument `ambiguousEdgeDirection` can define direction of ambiguous edges according to subtype of edges. Users firstly define which subtype of edges are considered as ambiguous edges by setting the argument `ambiguousEdgeList`. The default ambiguous edges include "compound", "hidden compound", "state change", "binding/association", "dissociation", and "unknow". Then users can define their direction through setting the value of the argument `ambiguousEdgeDirection` as one of "single", "bi-directed" or "delete", which means to convert ambiguous edges to "–>", "<–>", or to delete these ambiguous edges. The default value is "bi-directed".

The following commands convert pathways to graphs with ambiguous edges deleted. Compared with Figure 2, some edges are deleted such as edges related with the compound "C00076" because the default ambiguous edges include "compound".

```
> #Convert pathways to graphs with ambiguous edges as deleted
> gn2<-getNonMetabolicGraph(pn,ambiguousEdgeDirection="delete")
```

2.4 Convert pathway graphs to other derivative graphs

After using the function `getMetabolicGraph` or `getNonMetabolicGraph` to convert pathways to graphs, users can change these pathway graphs to other derivative graphs. The following section will detailedly introduce the usage of the related functions.

We firstly construct metabolic pathway graphs (`gm`) and non-metabolic pathway graphs (`gn`) as examples of input data. The commands are as follows:

```
> ##get metabolic pathway graphs
> #get path of KGML files
> path<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/metabolic/ec/",sep="")
> #convert metabolic pathways to graphs
> gm<-getMetabolicGraph(getPathway(path,c("ec00010.xml")))
> #show title of pathway graphs
> sapply(gm,function(x) x$title)

                        00010
"Glycolysis / Gluconeogenesis"

> ##get non-metabolic pathway graphs
> #get path
> path1<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/non-metabolic/ko/",sep="")
> #convert non-metabolic pathways to graphs
> gn<-getNonMetabolicGraph(getPathway(path1,c("ko04010.xml")),
+ ambiguousEdgeDirection="bi-directed")
```

```
> #show title of pathway graphs
> sapply(gn,function(x) x$title)

                   04010
"MAPK signaling pathway"
```

Note that the variable `gm` is a list of metabolic pathway graphs. The variable `gn` is a list of non-metabolic pathway graphs.

2.4.1 Convert pathway graphs to undirected graphs

The function `getUGraph` can convert directed graphs to undirected graphs. The following commands can get the undirected simple pathway graph (see Figure **??** for the result graph).

```
> #get undirected pathway graphs
> g1<-getUGraph(gm,simpleGraph=TRUE)
```

2.4.2 Map current organism-specific gene identifiers to nodes in pathway graphs

The function `mapNode` can map current organism-specific gene identifiers to nodes of graphs. We can use the function `getOrgAndIdType` to know the type of organism and identifier in the current study:

```
> getOrgAndIdType()

[1] "hsa"         "ncbi-geneid"
```

The result means that the type of organism and identifier in the current study are Homo sapiens (hsa) and Entrez gene identifiers (NCBI-geneid), which is the default value of the system (see the section **??**).

The following commands use the function `mapNode` to map human gene identifiers (NCBI-geneid) to nodes in pathway graphs. We can see the value of names attribute of some nodes revised. Green rectangle nodes are those that can correspond to gene identifiers, suggesting that these nodes are enzymes that human genes can encode. White rectangle nodes are those that can't correspond to gene identifiers, indicating that they may not be enzymes which human genes can encode. Therefore, the graph can be considered as human Glycolysis / Gluconeogenesis pathway graph.

```
> #see the names attribute of nodes.
> V(gm[[1]])[1:10]$names

 [1] "ec:1.2.1.3"   "ec:6.2.1.13"  "ec:1.2.1.5"   "cpd:C00033"   "path:ec00030"
 [6] "path:ec00500" "ec:4.1.1.1"   "ec:1.1.1.2"   "ec:1.1.1.1"   "ec:4.1.1.1"

> #get the organism-specific and idType-specific graph
> g1<-mapNode(gm)
> #see the names attribute of nodes in the new graph.
> #some node names are revised as NCBI-gene IDs
> V(g1[[1]])[1:10]$names

 [1] "217 219 223 224 501"         "ec:6.2.1.13"
 [3] "218 220 221 222"             "cpd:C00033"
 [5] "path:ec00030"                "path:ec00500"
 [7] "ec:4.1.1.1"                  "10327"
 [9] "124 125 126 127 128 130 131" "ec:4.1.1.1"
```

2.4.3 Filter nodes of pathway graphs

The function `filterNode` is used to filter "not interesting" nodes. For example, it may be necessary to ignore nodes with type="map" when focusing on components such as compounds and gene products. The function will delete nodes according to the argument `nodeType` and thus related edges are also deleted.

The following commands can delete nodes whose types are "map".

```
> #We display them before nodes are filtered
> V(gn[[1]])$type
```

```
  [1] "compound" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
  [7] "compound" "compound" "compound" "compound" "ortholog" "ortholog"
 [13] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [19] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [25] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [31] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [37] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [43] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "map"
 [49] "map"      "map"      "map"      "map"      "map"      "ortholog"
 [55] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [61] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [67] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [73] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [79] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [85] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [91] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [97] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[103] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[109] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[115] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[121] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[127] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[133] "ortholog"
```

```
> #delete nodes with type="map"
> g1<-filterNode(gn,nodeType=c("map"))
> #The "map" nodes are deleted in the new graph.
> V(g1[[1]])$type
```

```
  [1] "compound" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
  [7] "compound" "compound" "compound" "compound" "ortholog" "ortholog"
 [13] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [19] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [25] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [31] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [37] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [43] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [49] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [55] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [61] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [67] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [73] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
```

```
 [79] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [85] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [91] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
 [97] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[103] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[109] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[115] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[121] "ortholog" "ortholog" "ortholog" "ortholog" "ortholog" "ortholog"
[127] "ortholog"
```

2.4.4 Simplify pathway graphs as graphs with only gene products (or only compounds) as nodes

When we focus on gene products, compounds may be not important. Similarly, gene products may be not important when focusing on metabolites (compounds). For metabolic pathway graphs, a useful approach is to get graphs with gene products (or compounds) as nodes and compounds (gene products) as edges.

The function `simplifyGraph` can convert pathways to graphs with gene products (or compounds) as nodes and compounds (or gene products) as edges. We take an example of constructing metabolic pathway graphs with enzymes as nodes and compounds as edges. Firstly, all enzymes in a pathway graph are used as nodes. For undirected, two nodes are then connected by an edge if their corresponding reactions have a common compound. For directed, two nodes are connected by an edge if their corresponding reactions have a common compound and two nodes are reachable through the compound. Finally, compound information is added into edge attribute of new graphs. Similarly, a metabolic pathway graph can be converted to a graph with compounds as nodes. Two nodes are connected by an edge if they belong to the same reaction. Enzyme information is added into edge attribute of new graphs.

The following commands construct pathway graphs with enzymes as nodes and compounds as edges.

```
> #get graphs with enzymes as nodes and compounds as edges
> g1<-simplifyGraph(gm,nodeType="geneProduct")
> #see the names attribute of three edges
> E(g1[[1]])[1:3]$names

[1] "cpd:C05378" "cpd:C00118" "cpd:C00118"
```

The following commands construct graphs with compounds as nodes and enzymes as edges.

```
> #get graphs with compounds as nodes and enzymes as edges
> g2<-simplifyGraph(gm,nodeType="compound")
> #see the names attribute of three edges
> E(g2[[1]])[1:3]$names

[1] "ec:6.2.1.1"  "ec:2.7.1.69" "ec:5.4.2.2"
```

2.4.5 Expand nodes of pathway graphs

In pathways, some nodes may have multiple components, which are considered as components of "paralogues". For example, node PDE, which is the enzyme node in Purine metabolism (ec00230), maps to two enzymes: PDE (ec:3.1.4.17) and cGMP-PDE (ec:3.1.4.35). The function `expandNode` is just used to expand those nodes with multiple components. Users can select which types of nodes are expanded using the argument `nodeType`. The default values represent that all nodes are expanded. The following commands expand nodes of non-metabolic pathway graphs:

```
> #We firstly display node number before nodes are expanded
> vcount(gn[[1]])

[1] 133

> ##expand nodes in Graphs
> g1<-expandNode(gn)
> #We can see change of node number in the new graph:
> #node number after nodes are expanded
> vcount(g1[[1]])

[1] 197
```

The argument `nodeType` can determine which types of nodes should be expanded. Expanding nodes with certain node types is also available. The following commands only expand nodes that belong to gene products.

```
> #only expand nodes with type="enzyme" or "ortholog" in graphs
> g2<-expandNode(gn,nodeType=c("ortholog","enzyme"))
```

2.4.6 Get simple pathway graphs

If a graph is simple, it does not contain loop or/and multiple edges. A loop edge is an edge where the two endpoints have the same node (vertex). Two edges are multiple edges if they have exactly the same two endpoints. If graphs are not simple, some graph-based algorithms may be not applied. We can use the function `getSimpleGraph` to get a simple graph. Note that information of multiple edges is kept in edge attribute using ";" as separator.

The function `is.simple` can check whether a graph is simple as follows:

```
> all(sapply(gm,is.simple))

[1] TRUE
```

2.4.7 Merge nodes with the same names

A pathway usually includes some nodes with the same names. For example, an enzyme may appear repeatedly in a pathway. As shown in Figure 1, the Glycolysis / Gluconeogenesis pathway contain enzymes that appear repeatedly such as 2.7.1.69, 4.1.1.1, etc. The function `mergeNode` can merge those nodes with the same names. Therefore, each node in the result graph will has unique name. The edges of the merged nodes are obtained from edges of original nodes. After nodes are merged, multiple edges or loops may appear. The argument `simpleGraph` can delete them, which will return simple graphs (see the section 2.4.6). The following commands can get the graph in which nodes with the same names are merged.

```
> #get node number before merge
> vcount(gm[[1]])

[1] 94

> #merge nodes
> g1<-mergeNode(gm,simple=FALSE)
> #get node number after merge
> vcount(g1[[1]])

[1] 83
```

2.5 The integrated application of pathway reconstruct methods

In the section, we have provided some examples for converting pathways to graphs using the combination of graph conversion functions. Through the combination of these functions, many conversion strategies of pathway graphs can be implemented.

The section introduces the 24 examples of pathway graphs. They include enzyme-compound (KO-compound) pathway graphs, enzyme-enzyme (KO-KO) pathway graphs, compound-compound pathway graphs, organism-specific gene-gene pathway graphs, etc. These examples represent current major applications [Smart *et al.*, 2008, Schreiber *et al.*, 2002, Klukas and Schreiber, 2007, Kanehisa *et al.*, 2006, Goffard and Weiller, 2007, Koyuturk *et al.*, 2004, Hung *et al.*, 2010, Xia and Wishart, 2010, Jeong *et al.*, 2000, Antonov *et al.*, 2008, Guimera and Nunes Amaral, 2005, Draghici *et al.*, 2007, Li *et al.*, 2009, Ogata *et al.*, 2000, Hung *et al.*, 2010, Barabasi and Oltvai, 2004]. In the following subsection, we will detailedly introduce the combination usage of the graph conversion functions.

2.5.1 Example 1: enzyme-compound (KO-compound) pathway graphs

For metabolic pathways, the following commands can get pathway graphs with enzymes and compounds as nodes.

```
> #get graphs with enzymes and compounds as nodes
> g1<-filterNode(gm,nodeType=c("map"))
> #visualize
> plotGraph(g1[[1]])
```

Figure 3 shows the result graph of the Glycolysis / Gluconeogenesis pathway. Compared with original pathway graph (Figure 1), the "map" nodes disappear in the new graph.

If we apply the above method to all metabolic pathways, we can get all metabolic pathway graphs with enzymes and compounds as nodes. To do it easily, we have developed the function `getMetabolicECCOGraph`. The following command can use the function to get all metabolic pathway graphs with enzymes and compounds as nodes.

```
> #get all metabolic pathway graphs with enzymes and compounds as nodes
> graphList<-getMetabolicECCOGraph()
```

The result of the function are equal to the result of the following commands:

```
> #get all metabolic pathway data
> metabolicEC<-get("metabolicEC",envir=k2ri)
> ##write the results to tab delimited file.
> graphList<-filterNode(getMetabolicGraph(metabolicEC),nodeType=c("map"))
```

The variable `metabolicEC` stores all metabolic pathway data (see the section 5). The variable `graphList` stores all metabolic pathway graphs with enzymes and compounds as nodes.

The following commands can get the corresponding undirected graphs, that is, the undirected graphs with enzymes and compounds as nodes. The function `getMetabolicECCOUGraph` can get all results.

```
> #get the undirected graphs with enzymes and compounds as nodes
> g2<-filterNode(getUGraph(gm),nodeType=c("map"))
```

The following commands can get graphs with enzymes and compounds as nodes, in which each node only contains one enzyme/compound and each enzyme/compound only appears once. The function `getMetabolicECCOEMGraph` can get all results.

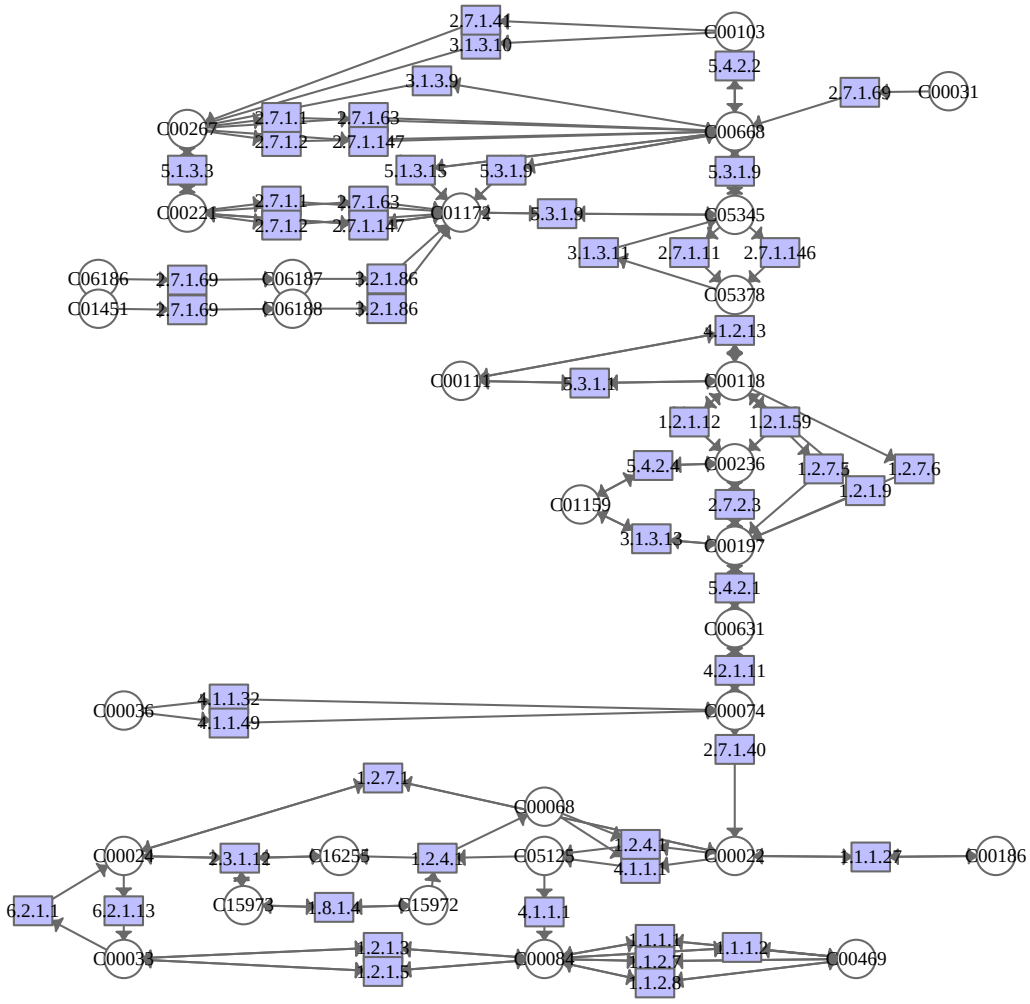

Figure 3: The Glycolysis / Gluconeogenesis pathway graph with enzymes and compounds as nodes. Compared with original pathway graph (Figure 1), the "map" nodes disappear in the new graph.

```
> #get graphs with enzymes and compounds as nodes
> #And, each node only contains one enzyme/compound and
> #each enzyme/compound only appears once in the graph.
> g3<-mergeNode(expandNode(filterNode(gm,nodeType=c("map"))))
```

The following commands can get the corresponding undirected graphs. The function `getMetabolicECCOUEMGraph` can get all results.

```
> #get the undirected graphs with enzymes and compounds as nodes
> #And, each node only contains one enzyme/compound and
> #each enzyme/compound only appears once in the graph.
> g4<-mergeNode(expandNode(filterNode(getUGraph(gm),nodeType=c("map"))))
```

For non-metabolic pathways, the following commands can get graphs with KOs and compounds as nodes. The function `getNonMetabolicKOCOGraph` can get all results.

```
> #get graphs with KOs and compounds as nodes
> g5<-filterNode(gn,nodeType=c("map"))
```

The following commands can get the undirected graphs with KOs and compounds as nodes. The function `getNonMetabolicECCOUGraph` can get all results.

```
> #get the undirected graphs with KOs and compounds as nodes
> g6<-filterNode(getUGraph(gn),nodeType=c("map"))
```

The following commands can get graphs with KOs and compounds as nodes. And, each node only contains a KO/compound and each KO/compound only appears once in the graph. The function `getNonMetabolicKOCOEMGraph` can get all results.

```
> #get graphs with KOs and compounds as nodes
> #And, each node only contains a KO/compound and
> #each KO/compound only appears once in the graph.
> g7<-mergeNode(expandNode(filterNode(gn,nodeType=c("map"))))
```

The following commands can get the corresponding undirected graphs. The function `getNonMetabolicKOCOUEMGraph` can get all results.

```
> #get the undirected graphs with KOs and compounds as nodes
> #And, each node only contains a KO/compound and
> #each KO/compound only appears once in the graph.
> g8<-mergeNode(expandNode(filterNode(getUGraph(gn),nodeType=c("map"))))
```

2.5.2 Example 2: enzyme-enzyme (KO-KO) pathway graphs

For metabolic pathways, the following commands can get graphs with enzymes as nodes and compounds as edges. The function `getMetabolicECECGraph` can get the results of all metabolic pathway graphs with enzymes as nodes and compounds as edges.

```
> #get graphs with enzymes as nodes and compounds as edges
> g1<-simplifyGraph(filterNode(gm,nodeType=c("map")),nodeType="geneProduct")
```

The following commands can get the corresponding undirected graphs, that is, the undirected graphs with enzymes as nodes and compounds as edges. The function `getMetabolicECECUGraph` can get all results.

```
> #get the undirected graphs with enzymes as nodes and compounds as edges
> g2<-simplifyGraph(filterNode(getUGraph(gm),nodeType=c("map")),nodeType="geneProduct")
```

The following commands can get graphs with enzymes as nodes and compounds as edges. And, each node contains only one enzyme and each enzyme only appears once in the graph. The graph can be treated as the enzyme-enzyme network obtained from the Glycolysis / Gluconeogenesis pathway. The function `getMetabolicECECEMGraph` can get all results.

```
> #get graphs with enzymes as nodes and compounds as edges
> #And, each node contains only one enzyme and each enzyme only appears once.
> g3<-mergeNode(expandNode(simplifyGraph(filterNode(gm,
+ nodeType=c("map")),nodeType="geneProduct")))
```

The following commands can get the corresponding undirected graphs. The function `getMetabolicECECUEMGraph` can get all results.

```
> #get undirected graphs with enzymes as nodes and compounds as edges.
> #And, each node contains only one enzyme and each enzyme only appears once.
> g4<-mergeNode(expandNode(simplifyGraph(filterNode(getUGraph(gm),
+ nodeType=c("map")),nodeType="geneProduct")))
```

For non-metabolic pathways, the following commands can get graphs with KOs as nodes. The function `getNonMetabolicKOKOGraph` can get all results.

```
> #get graphs with KOs as nodes
> g5<-simplifyGraph(filterNode(gn,nodeType=c("map")),nodeType="geneProduct")
```

The following commands can get the corresponding undirected graphs, that is, the undirected graphs with KOs as nodes. The function `getNonMetabolicKOKOUGraph` can get all results.

```
> #get the undirected graphs with KOs as nodes
> g6<-simplifyGraph(filterNode(getUGraph(gn),
+ nodeType=c("map")),nodeType="geneProduct")
```

The following commands can get graphs with KOs as nodes. And, each node contains only a KO and each KO only appears once in the graph. The function `getNonMetabolicKOKOEMGraph` can get all results.

```
> #get graphs with only KOs as nodes. And, each node contains
> #only a KO and each KO only appears once in the graph.
> g7<-mergeNode(expandNode(simplifyGraph(filterNode(gn,
+ nodeType=c("map")),nodeType="geneProduct")))
```

The following commands can get the corresponding undirected graphs. The function `getNonMetabolicKOKOUEMGraph` can get all results.

```
> #get the undirected graphs with only KOs as nodes. And, each node contains
> #only a KO and each KO only appears once in the graph.
> g8<-mergeNode(expandNode(simplifyGraph(filterNode(gn,
+ nodeType=c("map")),nodeType="geneProduct")))
```

2.5.3 Example 3: compound-compound pathway graphs

For metabolic pathways, the following commands can get graphs with compounds as nodes and enzymes as edges. The function `getMetabolicCOCOGraph` with setting the argument `type` as "EC" can get all metabolic pathway graphs with compounds as nodes and enzymes as edges.

```
> #The graph with compounds as nodes and enzymes as edges
> g1<-simplifyGraph(filterNode(gm,nodeType=c("map")),nodeType="compound")
```

The following commands can get the undirected graphs with compounds as nodes and enzymes as edges. The function `getMetabolicCOCOUGraph` with setting the argument `type` as "EC" can get all results.

```
> #The undirected graph with compounds as nodes and enzymes as edges
> g2<-simplifyGraph(filterNode(getUGraph(gm),nodeType=c("map")),nodeType="compound")
```

The following commands can get graphs with compounds as nodes and enzymes as edges. Each node only contains a compound and each compound only appears once in the graph. The function `getMetabolicCOCOEMGraph` with setting the argument `type` as "EC" can get all results.

```
> #The graph with compounds as nodes and enzymes as edges
> #Each node only contains a compound and each compound only appears once in the graph.
> g3<-mergeNode(expandNode(simplifyGraph(filterNode(gm,
+ nodeType=c("map")),nodeType="compound")))
```

The following commands can get the undirected graphs with compounds as nodes and enzymes as edges. Each node only contains a compound and each compound only appears once in the graph. The function `getMetabolicCOCOUEMGraph` with setting the argument `type` as "EC" can get all results.

```
> #The undirected graph with compounds as nodes and enzymes as edges
> #Each node only contains a compound and each compound only appears once in the graph.
> g4<-mergeNode(expandNode(simplifyGraph(filterNode(getUGraph(gm),
+ nodeType=c("map")),nodeType="compound")))
```

2.5.4 Example 4: organism-specific gene-gene pathway graphs

For metabolic pathways, the following commands can get graphs with organism-specific genes as nodes and compounds as edges. And, each node contains only a gene and each gene only appears once in the graph. The function `getMetabolicGEGEEMGraph` with setting the argument `type` as "EC" can get all metabolic pathway graphs with organism-specific genes as nodes and compounds as edges.

```
> #get graphs with organism-specific genes as nodes and compounds as edges
> g1<-mergeNode(expandNode(simplifyGraph(filterNode(mapNode(gm),
+ nodeType=c("map","enzyme")),nodeType="geneProduct")))
```

The following commands can get the corresponding undirected graphs. The function `getMetabolicGEGEEMUGraph` with setting the argument `type` as "EC" can get all results.

```
> #get the undirected graphs with organism-specific genes as nodes and compounds as edges
> g2<-mergeNode(expandNode(simplifyGraph(filterNode(mapNode(getUGraph(gm)),
+ nodeType=c("map","enzyme")),nodeType="geneProduct")))
```

For non-metabolic pathways, the following commands can get graphs with organism-specific genes as nodes and compounds as edges. Moreover, each node contains only a gene and each gene only appears once in the graph. The function `getNonMetabolicGEGEEMGraph` can get all results.

```
> #get graphs with organism-specific genes as nodes
> g3<-mergeNode(expandNode(simplifyGraph(filterNode(mapNode(gn),
+ nodeType=c("map","ortholog")),nodeType="geneProduct")))
```

The following commands can get the corresponding undirected graphs. The function `getNonMetabolicGEGEUEMGraph` can get all results.

```
> #get the undirected graphs with organism-specific genes as nodes
> g4<-mergeNode(expandNode(simplifyGraph(filterNode(mapNode(gn),
+ nodeType=c("map","ortholog")),nodeType="geneProduct")))
```

3 Methods to analyze pathway graphs

3.1 The basic analyses based on graph model

Since pathways are able to be converted to different types of graphs, many analyses based on graph model are available by using the functions provided in the igraph package. For example, we can get subgraph, degree, shortest path, etc [Csardi and Nepusz, 2006]. Here, we will give some detailed examples of operating graphs, nodes, edges, attributes. To do these, we firstly construct pathway graphs as the example graphs of the basic analyses based on graph model. The commands are as follows:

We can get metabolic pathway graphs as follows:

```
> #get path of KGML files
> path<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/metabolic/ec/",sep="")
> #convert metabolic pathways to graphs with "map" node deleted
> gmf<-filterNode(getMetabolicGraph(getPathway(path,c("ec00010.xml"))))
> #show title of pathway graphs
> sapply(gmf,function(x) x$title)

                        00010
"Glycolysis / Gluconeogenesis"

> #convert metablic pathways to graphs with enzymes as nodes and compounds as edges
> gmfs<-simplifyGraph(gmf,nodeType="geneProduct")
```

Figure 4 displays gmfs[[1]]. It is the Glycolysis / Gluconeogenesis pathway graph with enzymes as nodes and compounds as edges. The "map" nodes are deleted.

3.1.1 Node methods: degree, betweenness, local clustering coefficient, etc.

Degree (or connectivity) of a node is defined as the number of its adjacent edges [Csardi and Nepusz, 2006, Barabasi and Oltvai, 2004, Huber *et al.*, 2007]. It is a local quantitative measure of a node relative to other nodes. The following commands can get the degree of the first node in the graph.

```
> #get degree of nodes
> igraph::degree(gmfs[[1]],0)

[1] 12
```

We can see names of the first node as follows:

```
> #see name of the first node
> V(gmfs[[1]])[0]$names
```

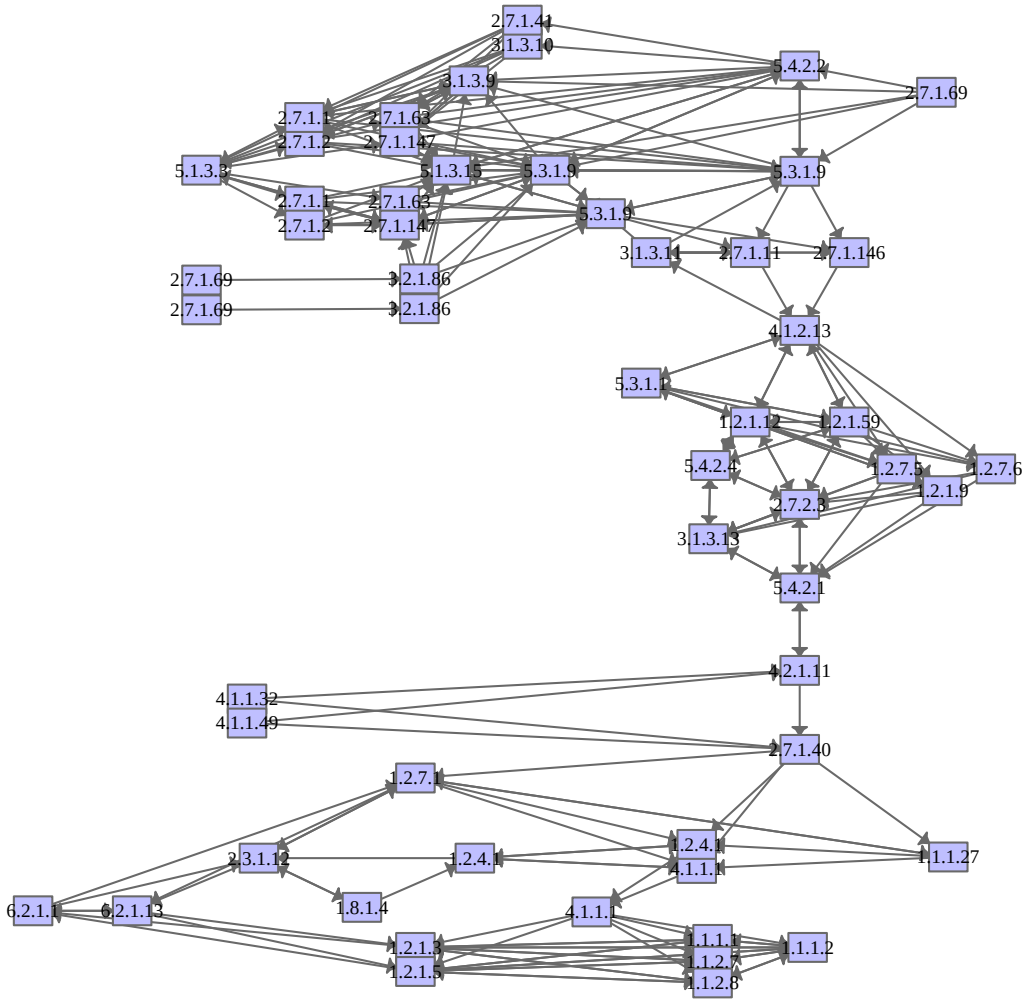

Figure 4: The Glycolysis / Gluconeogenesis pathway graph with enzymes as nodes and compounds as edges. The "map" nodes are deleted. The graph is stored in the variable gmfs[[1]].

```
[1] "ec:4.1.2.13"
```

The first node is the enzyme "ec:4.1.2.13" and is at the right-top part of Figure 4.

We can identify enzyme "ec:4.1.2.13" and get degree of a node with given names as follows:

```
> #get indexes of nodes
> index1<-V(gmfs[[1]])[V(gmfs[[1]])$names=="ec:4.1.2.13"]
> #get degree of node
> igraph::degree(gmfs[[1]],index1)
```

```
[1] 12
```

We may also want to calculate its betweeness, which is (roughly) defined by the number of shortest paths going through a node [Csardi and Nepusz, 2006, Barabasi and Oltvai, 2004, Huber *et al.*, 2007].

```
> #Calculate betweenness of enzyme "ec:4.1.2.13".
> betweenness(gmfs[[1]],index1)
```

```
[1] 960
```

The local clustering coefficient measures the probability that the adjacent nodes of a node are connected.

```
> #Calculate the clustering coefficient of enzyme "ec:4.1.2.13".
> igraph::transitivity(gmfs[[1]],type="local",vids=index1)
```

```
[1] 0.3888889
```

3.1.2 Edge method: shortest paths

The following commands can get the shortest path between the first node and the second node [Csardi and Nepusz, 2006, Barabasi and Oltvai, 2004, Huber *et al.*, 2007].

```
> #get the shortest path
> shortest.path<-get.shortest.paths(gmf[[1]],0,1,mode="out")
```

We can see name of nodes as follows:

```
> #see name of the first and second nodes
> V(gmf[[1]])[0:1]$names
```

```
[1] "ec:4.1.2.13" "ec:1.2.1.3"
```

```
> #see name of nodes in the shortest path
> V(gmf[[1]])[shortest.path[[1]]]$names
```

```
 [1] "ec:4.1.2.13" "cpd:C00118"  "ec:1.2.7.6"  "cpd:C00197"  "ec:5.4.2.1"
 [6] "cpd:C00631"  "ec:4.2.1.11" "cpd:C00074"  "ec:2.7.1.40" "cpd:C00022"
[11] "ec:4.1.1.1"  "cpd:C05125"  "ec:4.1.1.1"  "cpd:C00084"  "ec:1.2.1.3"
```

We sometimes may want to get the shortest path between two enzymes in a pathway, i.e., the shortest path between enzyme "ec:4.1.2.13" and "ec:1.2.1.3" in the Glycolysis / Gluconeogenesis pathway. To do this, we need to get indexes of interesting nodes and then use the function `get.shortest.paths` to get the result. The above strategy is usually necessary because in the igraph package, node indexes is used as input of most of functions. The following commands can calculate the shortest path between enzyme "ec:4.1.2.13" and "ec:1.2.1.3" in the Glycolysis / Gluconeogenesis pathway.

```
> #get indexes of nodes
> index1<-V(gmf[[1]])[V(gmf[[1]])$names=="ec:4.1.2.13"]
> index2<-V(gmf[[1]])[V(gmf[[1]])$names=="ec:1.2.1.3"]
> #get shortest path
> shortest.path<-get.shortest.paths(gmf[[1]],index1,index2)
> #display shortest path
> shortest.path

[[1]]
 [1]  0 81 74 52 15 44 14 51 13 57  8 58  5 60  1

> #convert indexs to names
> V(gmf[[1]])[shortest.path[[1]]]$names

 [1] "ec:4.1.2.13" "cpd:C00118"  "ec:1.2.7.6"  "cpd:C00197"  "ec:5.4.2.1"
 [6] "cpd:C00631"  "ec:4.2.1.11" "cpd:C00074"  "ec:2.7.1.40" "cpd:C00022"
[11] "ec:4.1.1.1"  "cpd:C05125"  "ec:4.1.1.1"  "cpd:C00084"  "ec:1.2.1.3"
```

3.1.3 Graph method: degree distribution, diameter, global clustering coefficient, density, module, etc.

The following command can get degree distribution of a pathway graph [Csardi and Nepusz, 2006, Barabasi and Oltvai, 2004, Huber *et al.*, 2007].

```
> #degree distribution.
> degree.distribution<-degree.distribution(gmfs[[1]])
```

The diameter of a pathway graph is the length of the longest geodesic [Csardi and Nepusz, 2006].

```
> #get diameter
> diameter(gmfs[[1]])

[1] 11
```

The following command can get the global clustering coefficient [Csardi and Nepusz, 2006].

```
> #Calculate the clustering coefficient.
> igraph::transitivity(gmfs[[1]])

[1] 0.5209302
```

The following command can get density of a pathway graph. The density of a graph is the ratio of the number of edges and the number of possible edges [Csardi and Nepusz, 2006].

```
> #Calculate the density.
> graph.density(gmfs[[1]])

[1] 0.0788961
```

The following commands can find densely connected subgraphs (modules or communities) in a pathway graph. We use walktrap community finding algorithm in the igraph package to find modules in the graph via random walks [Csardi and Nepusz, 2006]. Short random walks tend to stay in the same module.

```
> #find modules.
> wtc <- walktrap.community(gmfs[[1]])
> module<-community.to.membership(gmfs[[1]], wtc$merges, steps=53)
> module

$membership
 [1] 2 0 0 0 0 0 0 0 0 0 0 0 0 0 2 2 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 0 2 1 1 1
[39] 1 1 1 0 0 0 2 2 2 0 2 2 2 0 1 1 1 0

$csize
[1] 20 25 11
```

The result shows that three modules are found. They contain 20, 25, and 11 nodes respectively. We can also see names of nodes in the first module as follows:

```
> V(gmfs[[1]])[module$membership==0]$names

 [1] "ec:1.2.1.3"  "ec:6.2.1.13" "ec:1.2.1.5"  "ec:4.1.1.1"  "ec:1.1.1.2"
 [6] "ec:1.1.1.1"  "ec:4.1.1.1"  "ec:1.2.4.1"  "ec:1.2.4.1"  "ec:2.3.1.12"
[11] "ec:1.1.1.27" "ec:2.7.1.40" "ec:4.2.1.11" "ec:1.8.1.4"  "ec:4.1.1.32"
[16] "ec:1.1.2.7"  "ec:4.1.1.49" "ec:1.2.7.1"  "ec:6.2.1.1"  "ec:1.1.2.8"
```

This function `modularity` can calculate how modular is a given division of a graph into modules.

```
> modularity(gmfs[[1]], module$membership)

[1] 0.6122966
```

3.2 Topology-based pathway analysis of cellular component sets

The section mainly introduces topology-based pathway analysis of cellular component sets. Currently, our system can support input of three kinds of cellular component sets: gene sets, compound (metabolite) sets, and gene and compound sets at the same time. Therefore, the system can provide topology-based pathway analysis of gene sets. Topological significance of pathways can be also evaluated by the system. For example, if users input a set of interesting genes, the set can be mapped onto pathways. The topological property values can then be calculated. The topological significance of pathways can be evaluated. The available topological properties contain degree, clustering coefficient, betweenness, and closeness [Csardi and Nepusz, 2006, Barabasi and Oltvai, 2004, Huber *et al.*, 2007]. Degree of a node is the number of its adjacent edges. Local clustering coefficient quantifies the probability that the neighbours of a node are connected. Node betweenness can be calculated based on the number of shortest path passing through a given node. Closeness measures how many steps is required to access every other nodes from a given node.

3.2.1 Topology-based pathway analysis of gene sets

The function `identifyTopo` in the iSubpathwayMiner package facilitates topology-based pathway analysis of gene sets. We need to set the value of the argument `type` of the function as "gene". Moreover, we need to set the argument `propertyName` as a specific property (e.g., "degree").

To do topology-based pathway analysis of gene sets, we firstly construct a list of pathway graphs. We secondly input the interesting gene set and the list of pathway graphs to the function `identifyTopo`. The function can map interesting gene sets onto each pathway. For the mapped genes in a pathway, their topological property values can be calculated. These values can be compared with property values of all genes in the pathway. Finally, the statistical significance can be calculated using wilcoxon rank sum test.

The return value of the function `identifyTopo` is a list. Each element of the list is another list. It includes following elements: 'pathwayId', 'pathwayName', 'annComponentList', 'annComponentNumber', 'annBgComponentList', 'annBgNumber', 'ComponentNumber', 'bgNumber', 'propertyName', 'annComponentPropertyValueList', 'propertyValue', 'annBgComponentPropertyValueList', 'bgPropertyValue', 'pvalue', and 'fdr'. They correspond to pathway identifier, pathway name, the submitted components annotated to a pathway, numbers of submitted components annotated to a pathway, the background components annotated to a pathway, numbers of background components annotated to a pathway, numbers of submitted components, numbers of background components, topological property name (e.g., 'degree'), topological property values of submitted components annotated to a pathway, average topological property values of submitted components annotated to a pathway, topological property values of the background components annotated to a pathway, average topological property values of the background components annotated to a pathway, p-value of wilcoxon rank sum test for 'annComponentPropertyValueList' and 'annBgComponentPropertyValueList', and Benjamini-Hochberg fdr values. The list of results returned from the function `identifyTopo` can also be converted to `data.frame` using the function `printTopo`.

The following commands can perform topology-based pathway analysis of gene sets. The list of pathway graphs is obtained from the function `getMetabolicECECGraph`, which can get all directed metabolic pathway graphs with enzymes as nodes and compounds as edges (see the section 2.5.2).

```
> #get pathway graphs with enzymes as nodes.
> graphList<-getMetabolicECECGraph()
> #get a set of genes
> geneList<-getExample(geneNumber=1000,compoundNumber=0)
> #topology-based pathway analysis
> ann<-identifyTopo(geneList,graphList,type="gene",propertyName="degree")
> result<-printTopo(ann)
> #print a part of the result
> result[1:5,]
```

```
   pathwayId                      pathwayName annComponentRatio annBgRatio
1 path:00982 Drug metabolism - cytochrome P450           29/1000   82/21796
2 path:00380             Tryptophan metabolism           28/1000   65/21796
3 path:00562     Inositol phosphate metabolism            3/1000   55/21796
4 path:00670         One carbon pool by folate            7/1000   18/21796
5 path:00591          Linoleic acid metabolism           21/1000   42/21796
  propertyName propertyValue bgPropertyValue      pvalue       fdr
1       degree     0.5923372       0.5089431 0.006771113 0.5755446
2       degree     1.3511905       1.9128205 0.022801872 0.8365032
3       degree     6.6666667       4.2242424 0.041951602 0.8365032
4       degree    16.3809524      22.1388889 0.054268216 0.8365032
5       degree     2.6666667       4.6190476 0.054602392 0.8365032
```

The each row of the result (data.frame) is a pathway. Columns include pathwayId, pathwayName, annComponentRatio, annBgRatio, propertyName, propertyValue, bgPropertyValue, pvalue, and fdr. The `annComponentRatio` is the ratio of the annotated components. For example, 30/1000 means that 30 components in 1000 components are annotated. The `propertyValue` is average topological property value of submitted components annotated to a pathway. The `bgPropertyValue` is average topological property value of the background components annotated to a pathway. When many correlated pathways are considered, a false positive discovery rate is likely to result. Because the result is a `data.frame`, we are able to use the function `write.table` to export the result to a tab delimited file. If setting the argument `detail` as TRUE, we can also get more detailed result. For example, the topological property values of submitted genes annotated to a pathway can be exported using ";" as separator.

```
> ##write the results to tab delimited file.
> write.table(result,file="result.txt",row.names=FALSE,sep="\t")
>
> #detailed information is also outputed
> result1<-printTopo(ann,detail=TRUE)
> ##write the results to tab delimited file.
> write.table(result1,file="result1.txt",row.names=FALSE,sep="\t")
```

The result of topology-based anlysis shows that the degrees of the interesting genes in the inositol phosphate metabolism graph (path:00562) are significantly high. This suggests that these genes may play a more important role in the pathway. We can visualize the pathway using the function `plotAnnGraph`.

```
> #visualize
> plotAnnGraph("path:00562",graphList,ann)
```

The result pathway graph is shown in Figure 5. The mapped nodes, which correspond to the interesting genes, are colored red. From the figure, we can also see that degrees of these nodes are higher than the average degrees in the pathway.

The function `identifyTopo` is flexible. Users can change pathway graphs for different topological analyses. The following commands can use the function `getMetabolicGEGEUEMGraph` (see the section 2.5.4) to generate pathway graphs with genes as nodes, where each node contains only a gene and each gene only appears once. We can then use the data to analyze topological properties of gene sets in pathways. The following commands analyze local clustering coefficients of gene sets.

```
> #get undirected pathway graphs with genes as nodes.
> graphList<-getMetabolicGEGEUEMGraph(type="EC")
> #get a set of genes
> geneList<-getExample(geneNumber=1000,compoundNumber=0)
> #topology-based pathway analysis
> ann<-identifyTopo(geneList,graphList,type="gene",propertyName="clusteringCoefficient")
> result<-printTopo(ann)
> #print a part of the result
> result[1:10,c(1,3,6:8)]

    pathwayId annComponentRatio propertyValue bgPropertyValue      pvalue
1  path:00980           32/1000     0.2994552       0.4067389 0.005909343
2  path:00020            4/1000     0.4418651       0.5832804 0.014690575
3  path:00010           12/1000     0.6926918       0.5681654 0.023844855
4  path:00260           14/1000     0.8113791       0.5944147 0.026165910
5  path:00591           21/1000     0.8661994       0.9071813 0.054602392
6  path:00140           31/1000     0.7430575       0.7850713 0.061484669
7  path:00640            3/1000     0.0000000       0.3840278 0.120709223
8  path:00310            4/1000     0.3490119       0.6790014 0.131664235
9  path:00603            1/1000     0.0000000       0.7285714 0.151493992
10 path:00360            1/1000     0.4230769       0.6591660 0.152025014
```

The result shows that the local clustering coefficients of the interesting genes in the Glycolysis / Gluconeogenesis pathway (path:00562) are significantly high. This suggests that these genes tend to be in the functional module of the pathway. The local clustering coefficient measures the probability that the adjacent nodes of a node are connected [Csardi and Nepusz, 2006, Barabasi and Oltvai, 2004, Huber *et al.*, 2007].

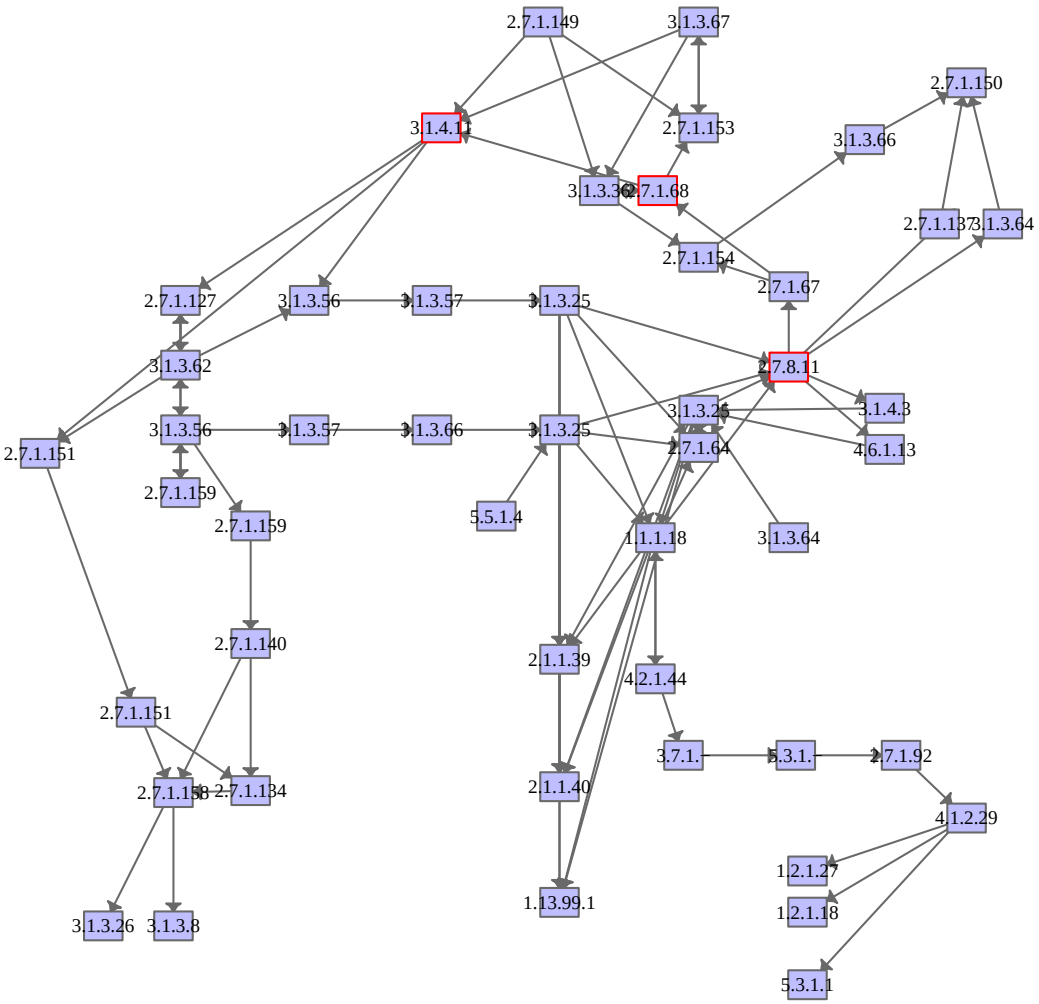

Figure 5: The inositol phosphate metabolism (path:00562) graph with enzymes as nodes and compounds as edges. The mapped nodes are colored red. We can see that degrees of these nodes are higher than the average degrees in the pathway.

3.3 Annotate cellular component sets and identify entire pathways

3.3.1 Annotate gene sets and identify entire pathways

The function `identifyGraph` in the iSubpathwayMiner package facilitates the annotation and identification of entire pathways. Firstly, we need to construct a list of pathway graphs. We then input the interesting gene set and the list of pathway graphs to the function `identifyGraph`. Through performing the function, the interesting gene set can be annotated to pathway graphs. Finally, the enrichment significance of pathways can be evaluated using hypergeometric test.

The return value of the function `identifyGraph` is a list of the annotated information. Each element of the list is another list. It includes the following elements: 'pathwayId', 'pathwayName', 'annComponentList', 'annComponentNumber', 'annBgComponentList', 'annBgNumber', 'ComponentNumber', 'bgNumber', 'pvalue', and 'fdr'. They correspond to pathway identifier, pathway name, the submitted components annotated to a pathway, numbers of submitted components annotated to a pathway, the background components annotated to a pathway, numbers of background components annotated to a pathway, numbers of submitted components, numbers of background components, p-value of the hypergeometric test, and Benjamini-Hochberg fdr values. The list of results returned from the function `identifyGraph` can also be converted to `data.frame` using the function `printGraph`.

The following commands annotate a gene set to metabolic pathways and identify significantly enriched metabolic pathways.

```
> ##Convert all metabolic pathways to graphs.
> metabolicEC<-get("metabolicEC",envir=k2ri)
> graphList<-getMetabolicGraph(metabolicEC)

> ##get a set of genes
> geneList<-getExample(geneNumber=1000)
> #annotate gene sets to pathway graphs
> #and identify significant pathway graphs
> ann<-identifyGraph(geneList,graphList)
> #convert ann to data.frame
> result<-printGraph(ann)
> #print a part of the results to screen
> result[1:10,]
```

```
    pathwayId                                  pathwayName annComponentRatio
1  path:00071                        Fatty acid metabolism           36/1000
2  path:00140                 Steroid hormone biosynthesis           31/1000
3  path:00232                          Caffeine metabolism           20/1000
4  path:00380                        Tryptophan metabolism           28/1000
5  path:00591                     Linoleic acid metabolism           21/1000
6  path:00830                           Retinol metabolism           30/1000
7  path:00980 Metabolism of xenobiotics by cytochrome P450           32/1000
8  path:00982            Drug metabolism - cytochrome P450           29/1000
9  path:00983              Drug metabolism - other enzymes           27/1000
10 path:00564               Glycerophospholipid metabolism           24/1000
   annBgRatio       pvalue          fdr
1    67/21796 0.000000e+00 0.000000e+00
2    73/21796 0.000000e+00 0.000000e+00
3    27/21796 0.000000e+00 0.000000e+00
4    65/21796 0.000000e+00 0.000000e+00
5    42/21796 0.000000e+00 0.000000e+00
6    61/21796 0.000000e+00 0.000000e+00
```

```
7    80/21796 0.000000e+00 0.000000e+00
8    82/21796 0.000000e+00 0.000000e+00
9    70/21796 0.000000e+00 0.000000e+00
10   76/21796 2.220446e-14 1.887379e-13
```

Each row of the result (data.frame) is a pathway. Its columns include pathwayId, pathwayName, annComponentRatio, annBgRatio, pvalue, and fdr. The `annComponentRatio` is the ratio of the annotated components. For example, 30/1000 means that 30 components in 1000 components are annotated to the pathway. When many correlated pathways are considered, a false positive discovery rate is likely to result. Because the result is a `data.frame`, it is able to use the function `write.table` to export the result to a tab delimited file. If setting the argument `detail` as TRUE, we can also get more detailed result. For example, the annotated components and the annotated background components can be exported using ";" as separator.

```
> ##write the annotation results to tab delimited file.
> write.table(result,file="result.txt",row.names=FALSE,sep="\t")
>
> #detailed information is also outputed
> result1<-printGraph(ann,detail=TRUE)
> ##write the annotation results to tab delimited file.
> write.table(result1,file="result1.txt",row.names=FALSE,sep="\t")
```

The following command displays a part of the return result of the function `identifyGraph`.

```
> #list of the result
> ann[1]

[[1]]
[[1]]$pathwayId
[1] "path:00071"

[[1]]$pathwayName
[1] "Fatty acid metabolism"

[[1]]$annComponentList
 [1] "10449"  "10455"  "11001"  "124"    "125"    "126"    "126129" "127"
 [9] "128"    "130"    "131"    "1374"   "1375"   "1376"   "1543"   "1544"
[17] "1545"   "1548"   "1549"   "1551"   "1553"   "1555"   "1557"   "1558"
[25] "1559"   "1562"   "1565"   "1571"   "1572"   "1573"   "1576"   "1577"
[33] "1579"   "1588"   "1632"   "1892"

[[1]]$annComponentNumber
[1] 36

[[1]]$annBgComponentList
 [1] "10449"  "10455"  "11001"  "124"    "125"    "126"    "126129" "127"
 [9] "128"    "130"    "131"    "1374"   "1375"   "1376"   "1543"   "1544"
[17] "1545"   "1548"   "1549"   "1551"   "1553"   "1555"   "1557"   "1558"
[25] "1559"   "1562"   "1565"   "1571"   "1572"   "1573"   "1576"   "1577"
[33] "1579"   "1580"   "1588"   "1632"   "1892"   "1962"   "199974" "217"
[41] "2180"   "2181"   "2182"   "219"    "223"    "224"    "23305"  "260293"
[49] "2639"   "284541" "29785"  "30"     "3028"   "3030"   "3032"   "3033"
```

```
[57] "3295"   "33"     "34"     "35"     "38"     "39"     "501"    "51"
[65] "51703"  "64816"  "8310"

[[1]]$annBgNumber
[1] 67

[[1]]$componentNumber
[1] 1000

[[1]]$bgNumber
[1] 21796

[[1]]$pvalue
[1] 0

[[1]]$fdr
[1] 0
```

The result is a list. It includes the following elements: 'pathwayId', 'pathwayName', 'annComponentList', 'annComponentNumber', 'annBgComponentList', 'annBgNumber', 'ComponentNumber', 'bgNumber', 'pvalue', and 'fdr'.

The Glycolysis / Gluconeogenesis pathway (path:00010) is significant in the analysis result of pathway. We can see the identified result of the pathway as follows:

```
> result[result[,1] %in% "path:00010",]

    pathwayId                  pathwayName annComponentRatio annBgRatio
20 path:00010 Glycolysis / Gluconeogenesis           12/1000   64/21796
         pvalue          fdr
20 2.942795e-05 0.0001250688
```

This means that the submitted interesting genes are significantly enriched to the Glycolysis / Gluconeogenesis pathway. If these genes is disease-related genes (e.g., risk genes associated with lung cancer), the Glycolysis / Gluconeogenesis pathway may be highly associted with the disease.

We can visualize the annotated pathways using the function `plotAnnGraph`. The following command displays the Glycolysis / Gluconeogenesis pathway (path:00010). The enzymes identified in the submitted genes are colored red.

```
> #visualize
> plotAnnGraph("path:00010",graphList,ann)
```

The result graph is shown in Figure 6. The red nodes in the result graph represent the enzymes which include the submitted genes.

3.3.2 Annotate compound sets and identify enire pathways

Our system can provide the annotation and identification of pathways based on compound sets. Users only need to set the value of the argument `type` of the function `identifyGraph` as "compound". We still use the above pathway graphs. We then input the interesting compound set and the list of pathway graphs to the function `identifyGraph`. Through performing the function `identifyGraph`, the interesting gene set can be annotated to pathway graphs. Finally, the enrichment significance of pathways can be evaluated using hypergeometric test. The following commands can annotate a compound set and identify statistically significantly enriched metabolic pathways.

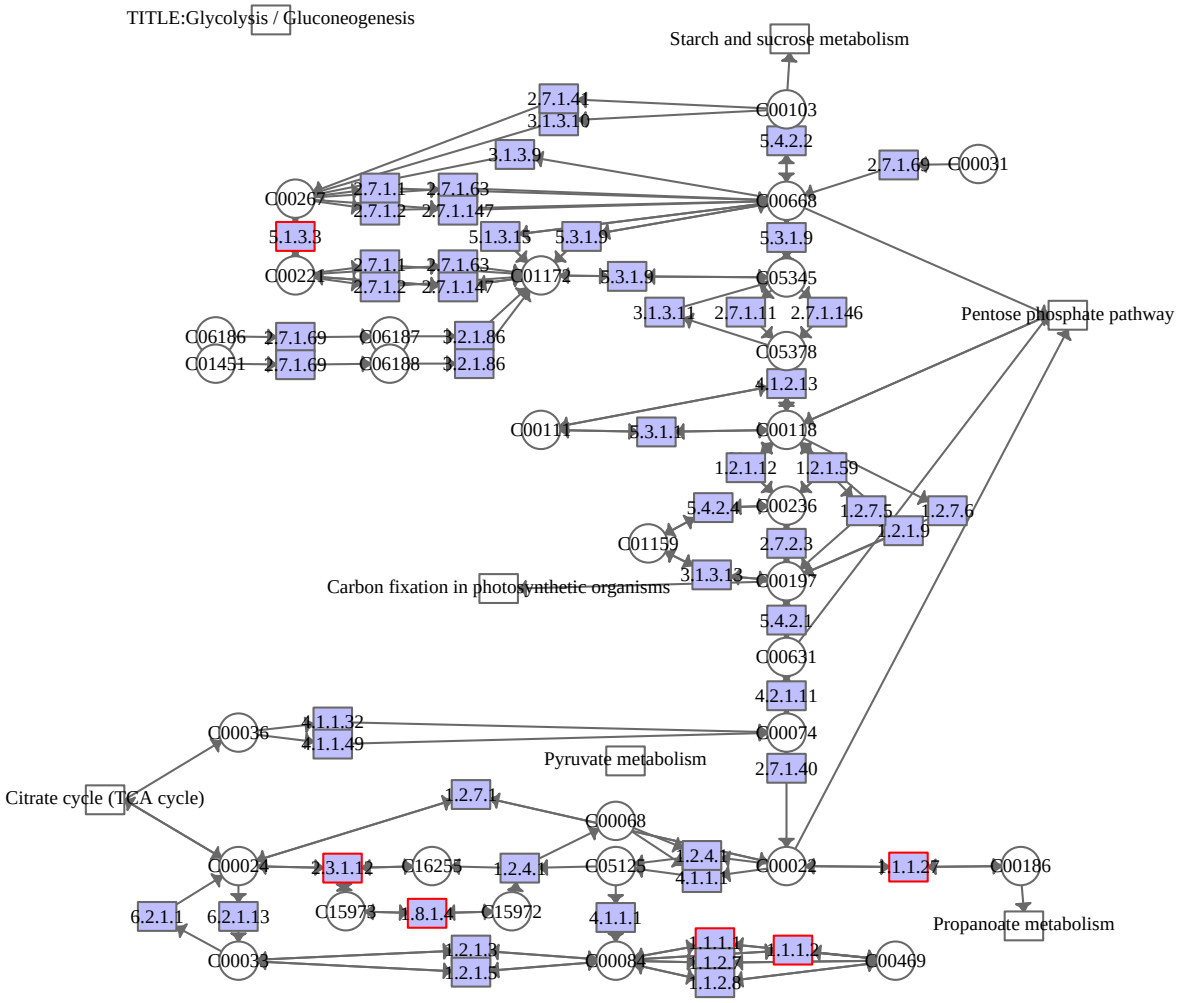

Figure 6: The Glycolysis / Gluconeogenesis pathway (path:00010). The enzymes identified in the submitted genes are colored red.

```
> #get a set of compounds
> compoundList<-getExample(geneNumber=0,compoundNumber=100)
> #annotate compound sets and identify significant pathways
> ann<-identifyGraph(compoundList,graphList,type="compound")
> result<-printGraph(ann)
> #display a part of the result
> result[1:5,c(1,3,4,5)]

   pathwayId annComponentRatio annBgRatio       pvalue
1 path:00190            11/100   16/14931 0.000000e+00
2 path:00230            17/100   92/14931 0.000000e+00
3 path:00970            14/100   53/14931 0.000000e+00
4 path:00250             9/100   24/14931 2.253753e-14
5 path:00270            11/100   56/14931 8.026912e-14
```

3.3.3 Annotate compound and gene sets and identify entire pathways

If users have not only interesting gene sets but also interesting compound sets, then users can annotate them at the same time and identify significant entire pathways. To do this, we need to set the argument `type` of the function `identifyGraph` as "gene_compound". We input the interesting compound set and the list of pathway graphs to the function `identifyGraph`. Through performing the function `identifyGraph`, the interesting gene and compound set can be annotated to pathway graphs. Finally, the enrichment significance of pathways can be evaluated using hypergeometric test. The following commands can annotate a combined set of genes and compounds and identify statistically significantly enriched metabolic pathways.

```
> #get a set of compounds and genes
> componentList<-getExample(geneNumber=1000,compoundNumber=100)
> #annotate gene and compound sets to metabolic graphs
> #and identify significant graphs
> ann<-identifyGraph(componentList,graphList,type="gene_compound")
> result<-printGraph(ann)
> #display a part of results
> result[1:5,c(1,3,4,5)]

   pathwayId annComponentRatio annBgRatio pvalue
1 path:00071           39/1100  117/36727      0
2 path:00190           32/1100  146/36727      0
3 path:00230           44/1100  243/36727      0
4 path:00232           21/1100   48/36727      0
5 path:00240           32/1100  153/36727      0
```

3.3.4 Other examples

The function `identifyGraph` is flexible in input of pathway data. We can change pathway data for different analyses. For example, we can use reference pathways linked to KO identifiers to support the identification of not only metabolic pathways but also non-metabolic pathways. The following commands annotate a gene set and identify significantly enriched metabolic and non-metabolic pathways:

```
> ##Convert all metabolic pathways to graphs.
> metabolicKO<-get("metabolicKO",envir=k2ri)
> gm<-getMetabolicGraph(metabolicKO)
> ##Convert all non-metabolic pathways to graphs,
```

```
> nonMetabolicKO<-get("nonMetabolicKO",envir=k2ri)
> gn<-getNonMetabolicGraph(nonMetabolicKO)
> graphList<-c(gm,gn)
> ##get a set of genes
> geneList<-getExample(geneNumber=1000,compoundNumber=0)
> #annotate gene sets and identify significant pathways
> ann<-identifyGraph(geneList,graphList,type="gene")
> result<-printGraph(ann)
> #display part of results
> result[1:5,c(1,3,4,5)]
```

```
   pathwayId annComponentRatio annBgRatio pvalue
1 path:00830           29/1000   65/21796      0
2 path:00980           26/1000   71/21796      0
3 path:04080           66/1000  272/21796      0
4 path:04142           35/1000  117/21796      0
5 path:04740           76/1000  384/21796      0
```

The result includes both metabolic pathways and non-metabolic pathways.

Note that for metabolic pathways, the results of pathway analyses based on KO may be slightly different from that based on EC. We suggest users to use reference pathways linked to KO identifiers to analyze metabolic pathways because KEGG uses KO to annotate genes to pathways. In this vignette, many examples of pathway analyses use reference pathways linked to EC identifiers because enzymes may be more easily understood by users. The following commands can annotate a gene set and identify significantly enriched metabolic pathways by using KO metabolic pathways:

```
> ##Convert all metabolic pathways to graphs.
> metabolicKO<-get("metabolicKO",envir=k2ri)
> graphList<-getMetabolicGraph(metabolicKO)
> ##get a set of genes
> geneList<-getExample(geneNumber=1000,compoundNumber=0)
> #annotate gene sets and identify significant pathways
> ann<-identifyGraph(geneList,graphList)
> result<-printGraph(ann)
> #display part of results
> result[1:10,c(1,3,4,5)]
```

```
    pathwayId annComponentRatio annBgRatio       pvalue
1  path:00830           29/1000   65/21796 0.000000e+00
2  path:00980           26/1000   71/21796 0.000000e+00
3  path:00982           24/1000   73/21796 7.993606e-15
4  path:00564           24/1000   79/21796 5.873080e-14
5  path:00071           16/1000   42/21796 1.851408e-11
6  path:00140           18/1000   56/21796 2.838807e-11
7  path:00561           16/1000   49/21796 2.749426e-10
8  path:00240           22/1000   99/21796 5.635397e-10
9  path:00190           25/1000  132/21796 1.388163e-09
10 path:00591           12/1000   29/21796 2.051083e-09
```

3.4 The k-cliques method to identify subpathways

The section mainly introduces the annotation and identification of subpathways. We developed the k-cliques subpathway identification method [Li *et al.*, 2009] according to pathway structure data provided

by KEGG.

3.4.1 Annotate gene sets and identify subpathways

Users can annotate the interesting gene sets and identify significantly enriched subpathways. Firstly, we need to construct a list of the undirected pathway graphs with enzymes as nodes. Enzymes in a graph are connected by an edge if their corresponding reactions have a common compound. Secondly, we use the function `getKcSubiGraph` to mine subpathways with the parameter k. We then input the interesting gene set and the list of subpathways to the function `identifyGraph`. Through performing the function, the interesting gene set can be annotated to subpathways. Finally, the enrichment significance of pathways can be evaluated using hypergeometric test.

The following commands can annotate gene sets and identify statistically significantly enriched metabolic subpathways based on the k-cliques method. The list of pathway graphs is obtained from the function `getMetabolicECECUGraph`, which can get all undirected metabolic pathway graphs with enzymes as nodes and compounds as edges (see the section 2.5.2).

```
> ##identify metabolic subpathways based on gene sets
> #get the enzyme-enzyme pathway graphs
> graphList<-getMetabolicECECUGraph()
> #get all 4-clique subgraphs
> subGraphList<-getKcSubiGraph(k=4,graphList)
> #get a set of genes
> geneList<-getExample(geneNumber=1000,compoundNumber=0)
> #annotate gene sets to subpathways
> #and identify significant graphs
> ann<-identifyGraph(geneList,subGraphList,type="gene")
> result<-printGraph(ann)
> #display a part of results
> result[1:15,c(1,3,4,5)]

       pathwayId annComponentRatio annBgRatio pvalue
1   path:00071_8           27/1000   38/21796      0
2   path:00140_5           25/1000   36/21796      0
3   path:00140_6           25/1000   43/21796      0
4   path:00140_7           25/1000   43/21796      0
5   path:00140_8           24/1000   41/21796      0
6   path:00140_9           25/1000   43/21796      0
7  path:00140_10           28/1000   64/21796      0
8  path:00140_19           27/1000   63/21796      0
9  path:00140_20           27/1000   63/21796      0
10 path:00140_21           27/1000   63/21796      0
11  path:00232_1           20/1000   27/21796      0
12  path:00232_2           20/1000   27/21796      0
13  path:00380_5           24/1000   40/21796      0
14  path:00591_1           21/1000   42/21796      0
15  path:00830_1           30/1000   61/21796      0
```

We find that the subpathway "path:00010_3", which is a subpathway of the Glycolysis / Gluconeogenesis pathway, is statistically significant. We can see the identified result of the subpathway as follows:

```
> result[result[,1] %in% "path:00010_3",]
```

```
      pathwayId                   pathwayName annComponentRatio annBgRatio
72 path:00010_3 Glycolysis / Gluconeogenesis           11/1000   36/21796
         pvalue          fdr
72 3.747776e-07 3.523951e-06
```

The following commands can display the subpathway.

```
> plotAnnGraph("path:00010_3",subGraphList,ann)
```

The result is shown in Figure 7. The nodes identified in the submitted genes are colored red.

4 Visualize a pathway graph

We provide the function `plotGraph` for visualization of a pathway graph. The function can display a pathway graph using varieties of layout styles. The default is the KEGG style. We implement it by using detailed information about pathway map obtained from KGML files, which are converted to attributes of the corresponding graph, including graphics_x, graphics_y, graphics_name, graphics_type, names, type, etc. The function is developed based on the function `plot.igraph` in the igraph and the function `plot`. Therefore, most of functions in `plot.igraph` and `plot` are also available for the `plotGraph`. We will detailedly describe how to efficiently use the function. The following command is a simple usage for the function to visualize pathway graphs with the KEGG style.

We firstly generate a pathway graph.

```
> path<-paste(system.file(package="iSubpathwayMiner"),
+ "/localdata/kgml/metabolic/ec/",sep="")
> gm<-getMetabolicGraph(getPathway(path,c("ec00010.xml")))
```

We can use `plotGraph` to visualize the pathway graph as follows:

```
> #visualize
> plotGraph(gm[[1]])
```

The result graph is shown in Figure 8. The default layout style of the function is the KEGG style.

4.1 Change node label of the pathway graph

We can change node labels into the gene identifiers of the current organism as follows:

```
> plotGraph(gm[[1]],vertex.label=getNodeLabel(gm[[1]],
+ type="currentId",displayNumber=1))
```

4.2 The basic commands to visualize a pathway graph with custom style

We can display a pathway graph with different styles by using some basic commands. For example, we can set a color vector and then use it to change color of each node frame. Figure **??** shows an example of changing the certain enzyme node as red frame. The commands are as follows:

```
> #add red frame to the enzyme "ec:4.1.2.13"
> vertex.frame.color<-ifelse(V(gm[[1]])$names=="ec:4.1.2.13","red","dimgray")
> vertex.frame.color
```

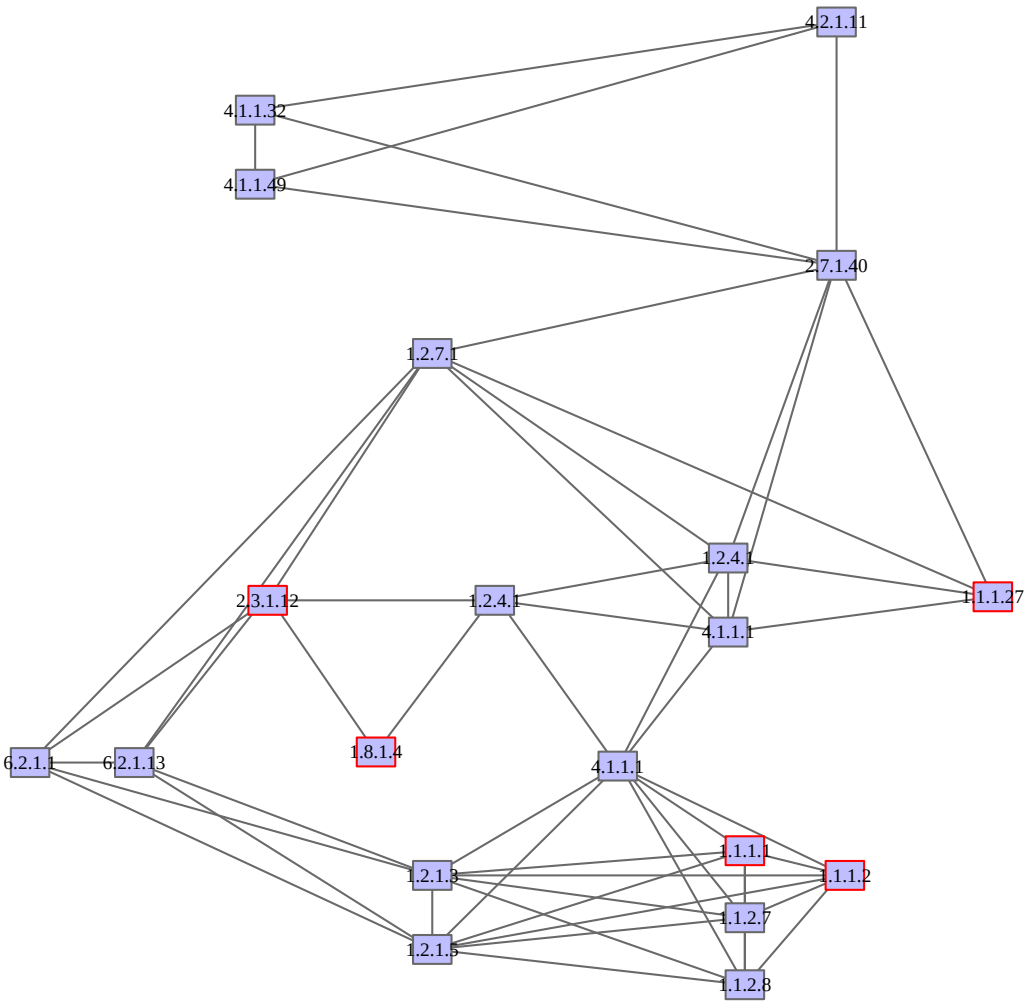

Figure 7: A significant subpathway of the Glycolysis / Gluconeogenesis pathway. The subpathway is constructed based on the k-clique method. In the subpathway, the distance between any two nodes is no greater than 4. The nodes identified in the submitted genes are colored red.

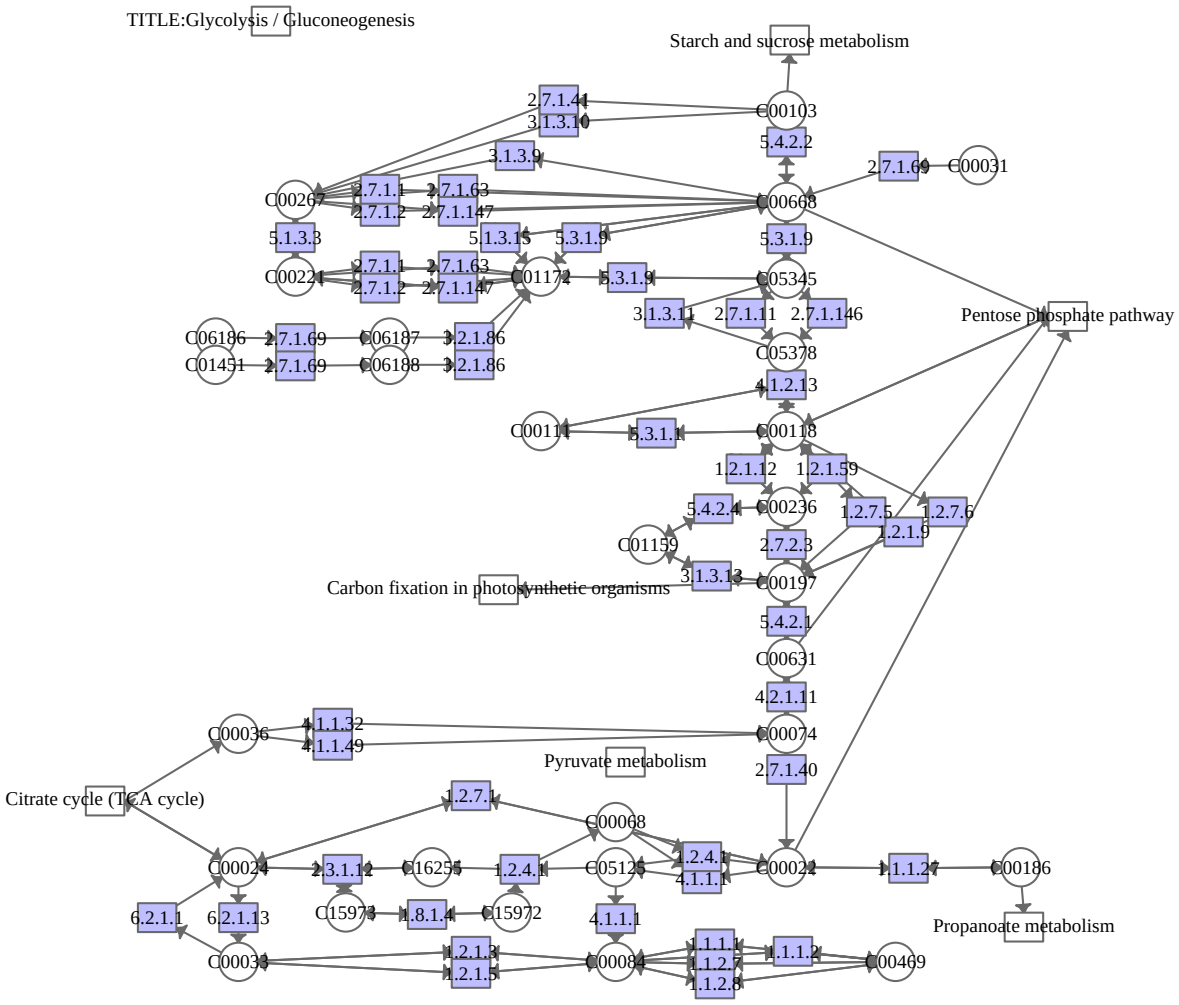

Figure 8: The Glycolysis / Gluconeogenesis pathway graph with the KEGG style

```
 [1] "red"     "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
 [8] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[15] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[22] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[29] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[36] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[43] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[50] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[57] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[64] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[71] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[78] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[85] "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray" "dimgray"
[92] "dimgray" "dimgray" "dimgray"
```

```
> #display new graph
> plotGraph(gm[[1]],vertex.frame.color=vertex.frame.color)
```

Operations to change other settings are similar to the example. In order to change styles of a graph, we only need to get and change the value of vectors related to styles and then transfer them to the function `plotGraph`. Detailed information can be provided in the help of the function `plot.igraph` in the igraph package and the function `plot` in the graphics package. Here, we only provide some examples of setting some styles for interpreting the usages of the function `plotGraph`. For instance, we can change node color, size, label font, x-y coordinates, etc. Figure 9 shows the results and the corresponding commands as follows:

```
> #add green label to the comound "cpd:C00111"
> vertex.label.color<-ifelse(V(gm[[1]])$names=="cpd:C00111","green","dimgray")
> #change node color
> vertex.color<-sapply(V(gm[[1]])$type,function(x) if(x=="enzyme"){"pink"}
+ else if(x=="compound"){"yellow"} else{"white"})
> #change node size
> size<-ifelse(V(gm[[1]])$graphics_name=="Starch and sucrose metabolism",20,8)
> #change a compound label
> #font size
> vertex.label.cex<-ifelse(V(gm[[1]])$names=="cpd:C00036",1.0,0.6)
> #italic
> vertex.label.font<-ifelse(V(gm[[1]])$names=="cpd:C00036",3,1)
> #change y coordinate of an enzyme
> layout<-getLayout(gm[[1]])
> index<-V(gm[[1]])[V(gm[[1]])$names=="ec:4.1.1.32"]
> layout[index+1,2]<-layout[index+1,2]+50
> #display the new graph
> plotGraph(gm[[1]],vertex.frame.color=vertex.frame.color,
+ vertex.label.color=vertex.label.color,vertex.color=vertex.color,
+ vertex.size=size,vertex.size2=size,vertex.label.cex=vertex.label.cex,
+ vertex.label.font=vertex.label.font,layout=layout)
```

4.3 The layout style of a pathway graph in R

The argument `layout` of the function `plotGraph` is used to determine the placement of the nodes for drawing a graph. There are mainly two methods to determine the placement of the nodes for drawing a

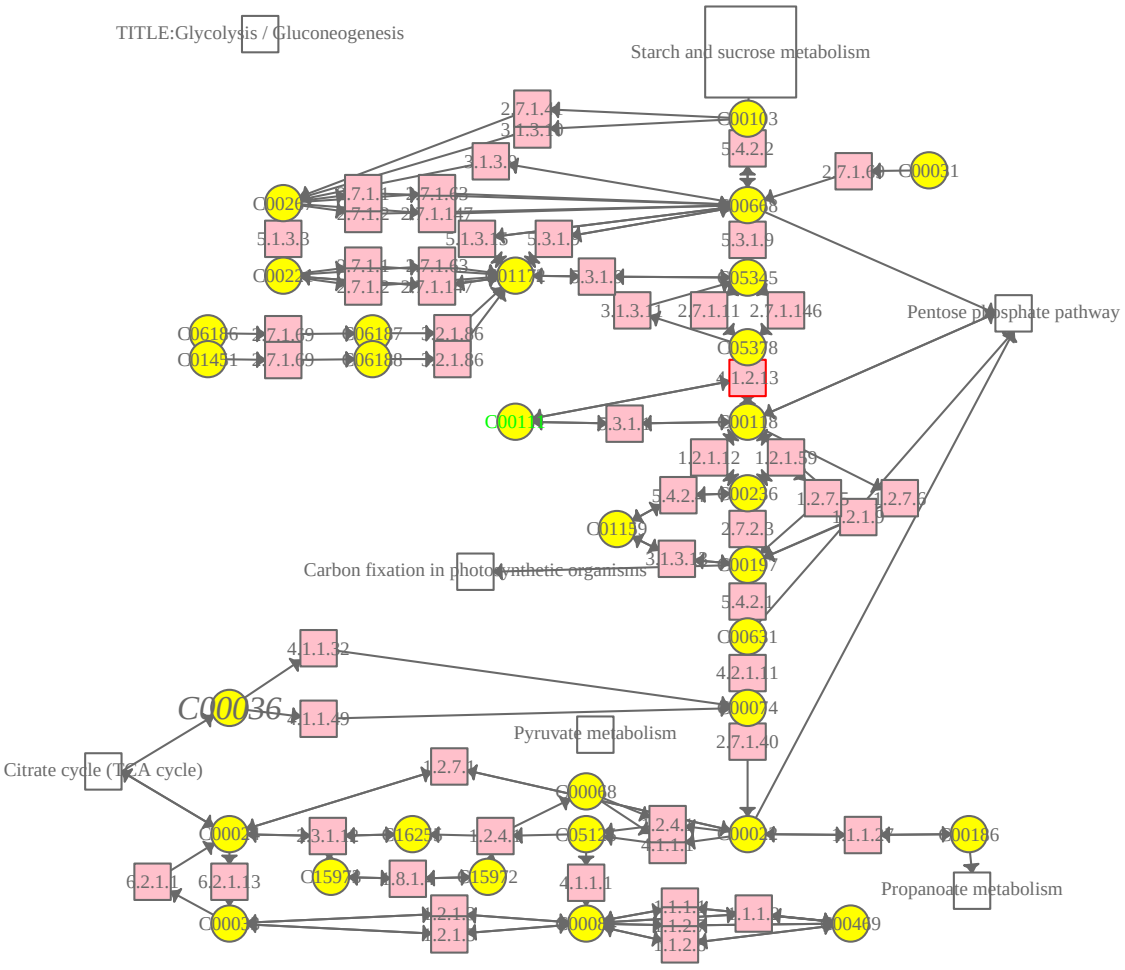

Figure 9: The new graph after changing some setting of visualization

pathway graph: the KEGG layout style and `layout` provided in the function `plot.igraph` of the igraph package. The default layout is the KEGG layout style, for which the coordinates of nodes in KEGG pathway maps is used to determine the placement of the nodes for drawing a graph. Therefore, the returned figure by the function can be very similar to the KEGG pathway graph. Figure 8 displays a pathway graph with the KEGG layout style.

The layout styles provided in igraph include `layout.random`, `layout.circle`, `layout.sphere`, `layout.sphere`, `layout.fruchterman.reingold`, `layout.kamada.kawai`, `layout.spring`, `layout.lgl`, `layout.fruchterman.reingold.g` `layout.graphopt`, `layout.mds`, `layout.svd`, `layout.norm`, `layout.drl`, and `layout.reingold.tilford`. For example, as shown in Figure 10, the `layout.random` places the nodes randomly. The `layout.circle` (e.g., Figure 11) places the nodes on an unit circle.

The following command displays a pathway graph using `layout.random` style.

```
> plotGraph(gm[[1]],layout=layout.random)
```

The result is shown in Figure 10.

The following command displays a pathway graph using `layout.circle` style.

```
> plotGraph(gm[[1]],layout=layout.circle)
```

The result is shown in Figure 11.

4.4 Visualize the result graph of pathway analyses

We can use the function `plotAnnGraph` to visualize the result graph of a pathway analysis (e.g., most of result graphs in the section 3). We take an example of visualizing a metabolic pathway, which is obtained from the annotation and identification method of entire pathways based on gene sets.

The following commands annotate a gene set to metabolic pathways and identify significantly enriched metabolic pathways.

```
> ##Convert all metabolic pathways to graphs.
> metabolicEC<-get("metabolicEC",envir=k2ri)
> graphList<-getMetabolicGraph(metabolicEC)
> ##get a set of genes
> geneList<-getExample(geneNumber=1000)
> #annotate gene sets to pathway graphs
> #and identify significant pathway graphs
> ann<-identifyGraph(geneList,graphList)
```

The following command displays the Glycolysis / Gluconeogenesis pathway (path:00010). Users need to input pathway identifier, a list of pathway graphs, and the result variable `ann` of pathway analysis.

```
> #visualize
> plotAnnGraph("path:00010",graphList,ann)
```

The result graph is shown in Figure 12. The red nodes in the result graph represent the enzymes which include the submitted genes. In fact, the function `plotAnnGraph` can obtain the annotated genes from the variable `ann` , match the genes to the given pathway, and display the pathway with the annotated genes colored red.

We can also use the function `plotAnnGraph` to visualize pathways not only in R but also in KEGG web site. The annotated genes are also colored red in KEGG maps.

```
> #visualize
> plotAnnGraph("path:00010",graphList,ann,gotoKEGG=TRUE)
```

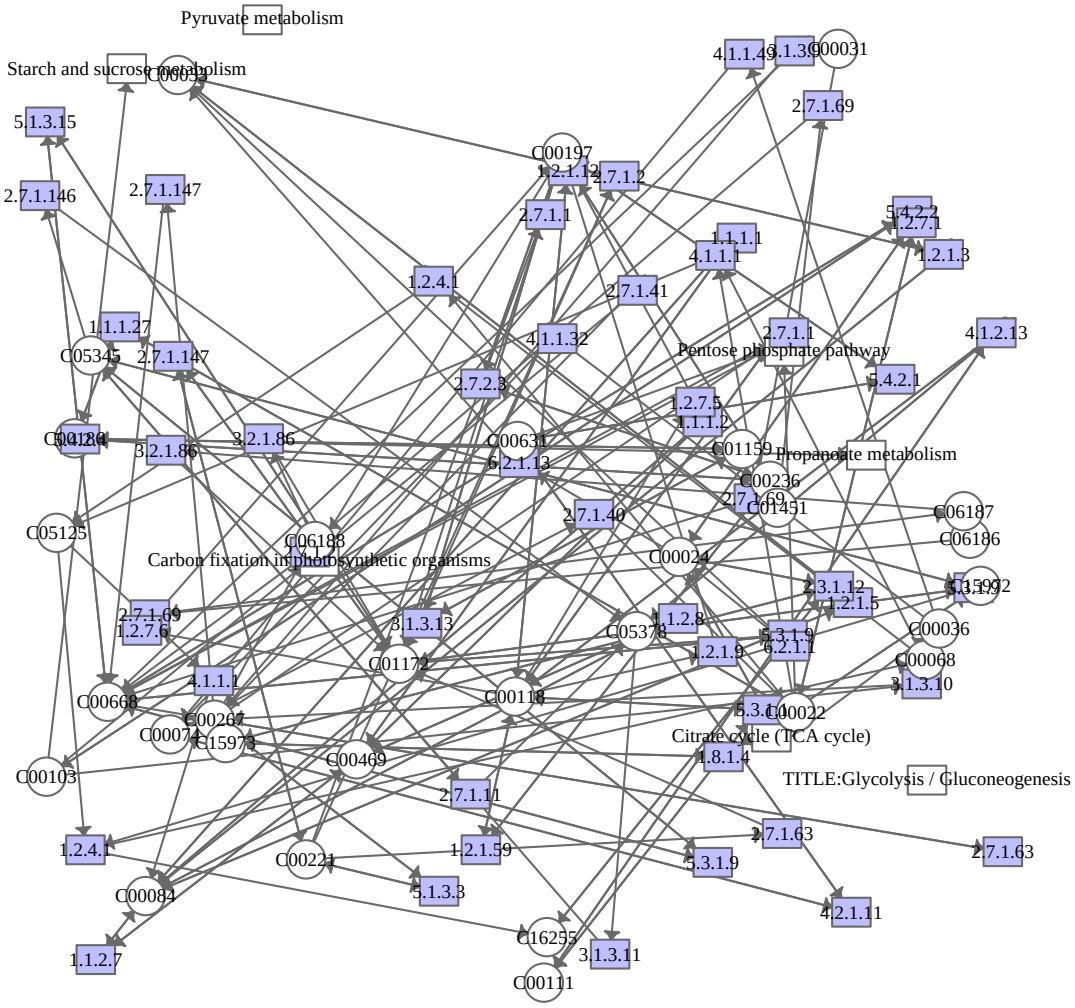

Figure 10: The pathway graph with the `layout.random` style

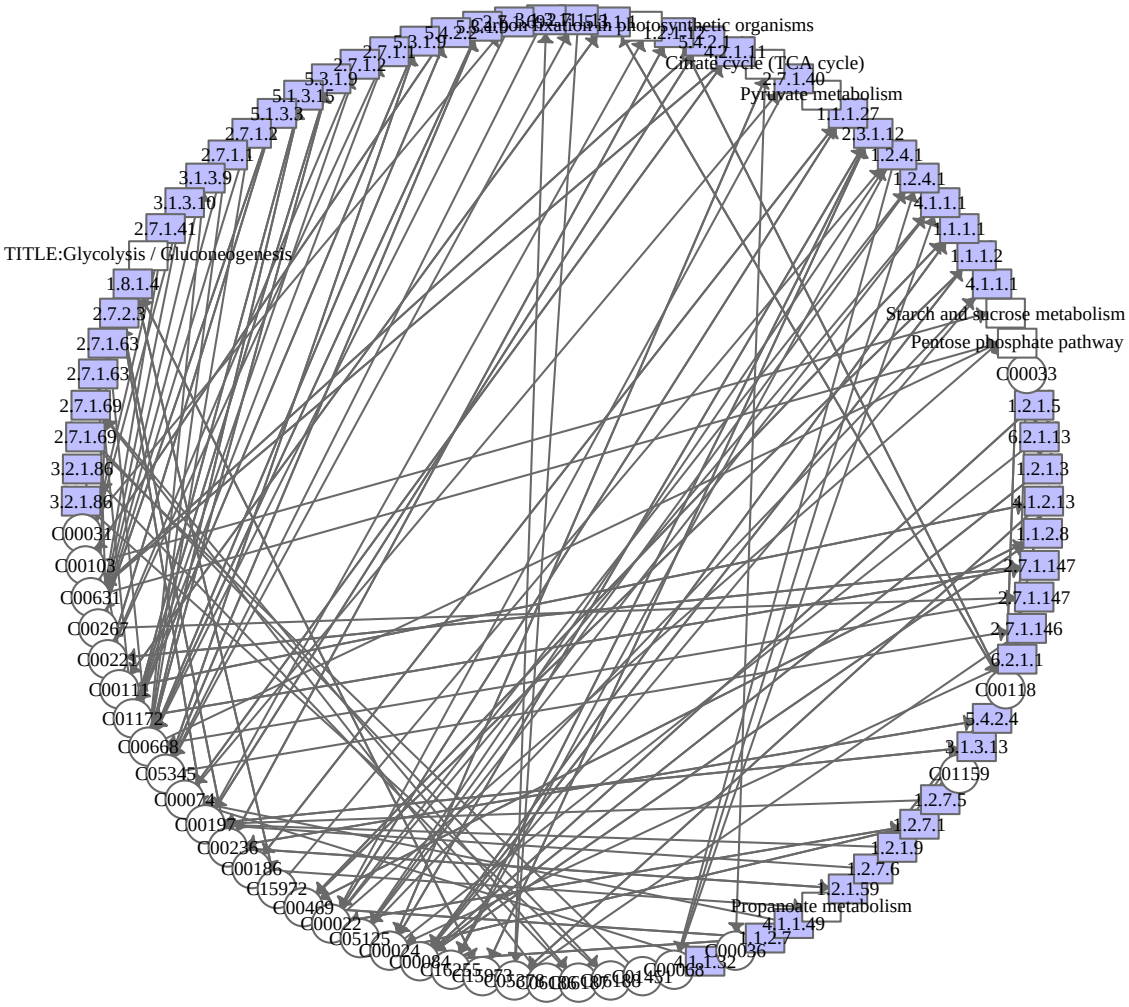

Figure 11: The pathway graph with the `layout.circle` style

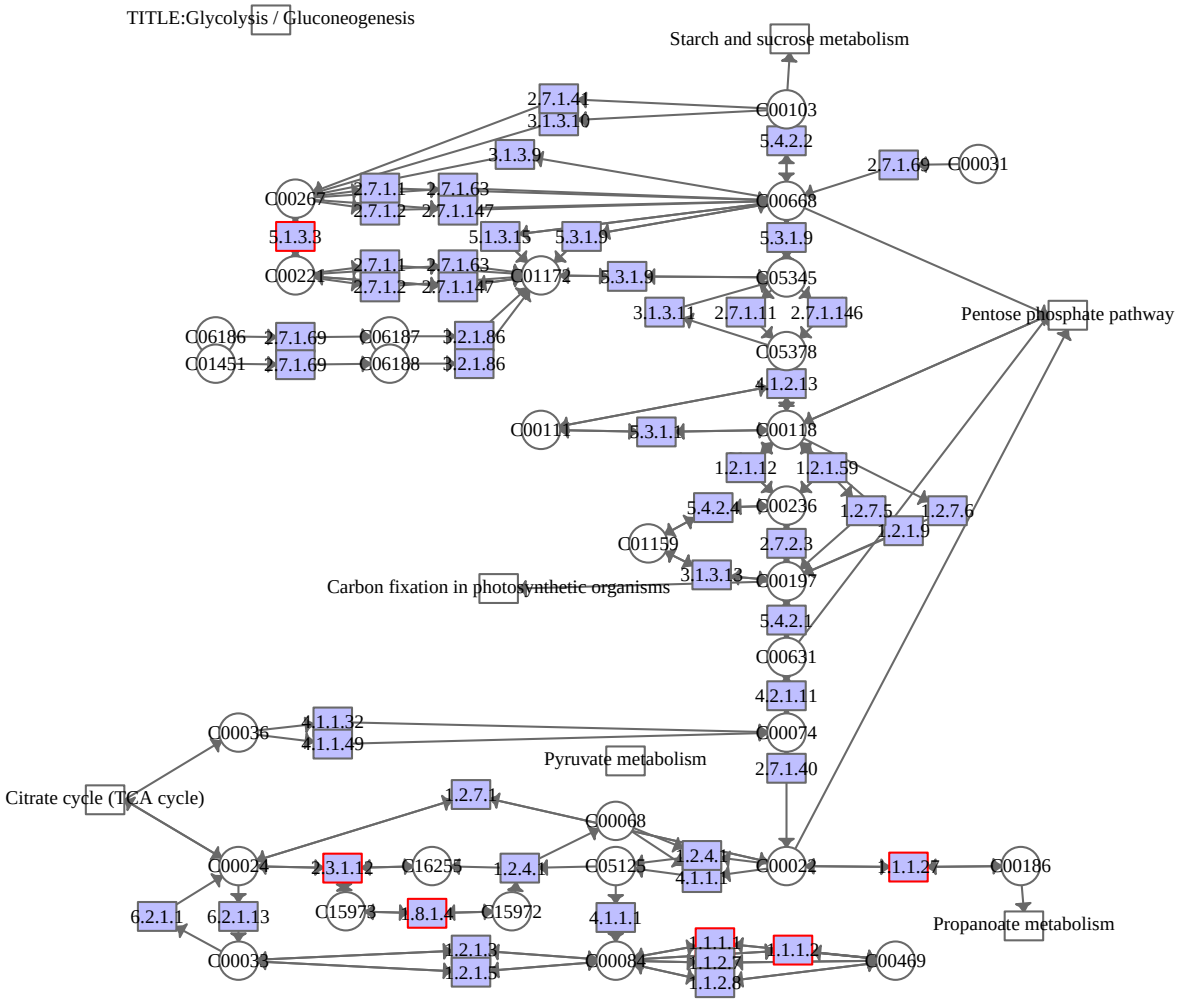

Figure 12: The Glycolysis / Gluconeogenesis pathway (path:00010). The enzymes identified in the submitted genes are colored red.

4.5 Export a pathway graph

The function `write.graph` can export a pathway graph to foreign file formats. The following command exports a metabolic pathway graph to the GML format `http://www.infosun.fim.uni-passau.de/Graphlet/GML/`. The format is supported by Cytoscape software [Shannon *et al.*, 2003] that provides more advanced visualization facilities `http://www.cytoscape.org`.

```
> write.graph(gm[[1]], "ec00010.txt", "gml")
```

5 Data management

The environment variable `k2ri`, which is used as the database of the system, stores many data relative to pathway analyses. We can use the function `ls` to see the environment variable and use `ls(k2ri)` to see data in it. These data include `gene2ec`, `gene2ko`, `metabolicEC`, `metabolicKO`, `nonMetabolicKO`, etc. For example, the variable `gene2ec` stores relation betweeen genes and enzymes in the current organism (e.g., relation between human genes and enzymes). The variable `metabolicEC` stores reference metabolic pathways linked to EC identifiers. The variable `metabolicKO` stores reference metabolic pathways linked to KO identifiers. The variable `nonMetabolicKO` stores reference non-metabolic pathways linked to KO identifiers.

```
> ##data in environment variable k2ri
> ls(k2ri)
```

```
 [1] "compbackground" "compound"       "gene2ec"        "gene2ko"
 [5] "gene2path"      "gene2symbol"    "genebackground" "keggGene2gene"
 [9] "metabolicEC"    "metabolicKO"    "nonMetabolicKO" "orgAndIdType"
[13] "taxonomy"
```

We can obtain these data in the environment variable `k2ri` using the function `get`. The following command gets reference metabolic pathways linked to EC identifiers in the variable `metabolicEC` in R.

```
> #get all metabolic pathway data
> metabolicEC<-get("metabolicEC",envir=k2ri)
```

The section will introduce the functions relative to the data management of the environment variable `k2ri`.

5.1 Set or update the current organism and the type of gene identifier

When using the pathway analysis functions of iSubpathwayMiner, users need to know the type of organism and identifier in the current study. Users can check the type of organism and identifier in the current study through the function `getOrgAndIdType`:

```
> getOrgAndIdType()
```

```
[1] "hsa"         "ncbi-geneid"
```

The result means that the type of organism and identifier in the current study are Homo sapiens and Entrez gene identifiers, which is the default value of the system. Users should ensure that the organism and gene identifiers in the expected study accord with the return value of the function `getOrgAndIdType`. If the result is different from the type of your genes, you need to change them through some functions, e.g., `updateOrgAndIdType` and `loadK2ri`.

The function `updateOrgAndIdType` can download data relative to organism and gene identifiers, and then treat and store them in the environment variable `k2ri`. The following command can set the type of organism and identifier in the current study as Saccharomyces cerevisiae and sgd identifier in Saccharomyces Genome Database.

```
> path<-paste(system.file(package="iSubpathwayMiner"),"/localdata",sep="")
> updateOrgAndIdType("sce","sgd-sce",path)
```

The function `updateCompound` is able to update the variable `compound` in the environment variable `k2ri`. The function `updateTaxonomy` is able to update the variable `taxonomy` in the environment variable `k2ri`. The variable stores information about organism name and the three- or four-letter KEGG organism code.

Through these functions, iSubpathwayMiner can support multiple species in KEGG and different gene identifiers (KEGG compound, Entrez Gene IDs, gene official symbol, NCBI-gi IDs, UniProt IDs, PDB IDs, etc.). It can also provide the most up-to-date pathway analysis results for users.

5.2 Update pathway data

The function `updatePathway` can update pathways in the environment variable `k2ri` from KEGG ftp site. The function `importPathway` can construct the pathway variable metabolicEC, metabolicKO, and nonMetabolicKO from local system. Firstly, users need to download KGML pathway files from KEGG ftp site.

5.3 Load and save the environment variable of the system

Through the above functions, data in the environment variable of the system can be updated. The system provides two functions (`saveK2ri` and `loadK2ri`) to easily save and load the new environment variable. The following command is used to save the environment variable of Saccharomyces cerevisiae.

```
> saveK2ri("sce_sgd-sce.rda")
```

When one needs to use the environment variables of Saccharomyces cerevisiae next time, one can use the function `loadK2ri` to load the last environment variable. The following command is used to load the environment variables of Saccharomyces cerevisiae.

```
> loadK2ri("sce_sgd-sce.rda")
```

6 Session Info

The script runs within the following session:

```
R version 2.14.1 (2011-12-22)
Platform: i386-pc-mingw32/i386 (32-bit)

locale:
[1] LC_COLLATE=C
[2] LC_CTYPE=Chinese_People's Republic of China.936
[3] LC_MONETARY=Chinese_People's Republic of China.936
[4] LC_NUMERIC=C
[5] LC_TIME=Chinese_People's Republic of China.936

attached base packages:
[1] stats     graphics  grDevices utils     datasets  methods   base

other attached packages:
[1] iSubpathwayMiner_2.0 XML_3.4-0.2          igraph_0.5.5-4
[4] RBGL_1.30.1          graph_1.32.0

loaded via a namespace (and not attached):
[1] tools_2.14.1
```

References

[Antonov *et al.*, 2008] Antonov, A.V., et al. (2008) Kegg Spider: Interpretation of Genomics Data in the Context of the Global Gene Metabolic Network. Genome Biol, 9, R179.

[Barabasi and Oltvai, 2004] Barabasi, A.L. and Oltvai, Z.N. (2004) Network Biology: Understanding the Cell's Functional Organization. Nat Rev Genet, 5, 101-113.

[Csardi and Nepusz, 2006] Csardi, G. and Nepusz, T. (2006) The igraph software package for complex network research. InterJournal, Complex Systems, 1695.

[Draghici *et al.*, 2007] Draghici, S., et al. (2007) A Systems Biology Approach for Pathway Level Analysis. Genome Res, 17, 1537-1545.

[Gentleman *et al.*, 2004] Gentleman, R.C., et al. (2004) Bioconductor: Open Software Development for Computational Biology and Bioinformatics. Genome Biol, 5, R80.

[Goffard and Weiller, 2007] Goffard, N. and Weiller, G. (2007) Pathexpress: A Web-Based Tool to Identify Relevant Pathways in Gene Expression Data. Nucleic Acids Res, 35, W176-181.

[Guimera and Nunes Amaral, 2005] Guimera, R. and Nunes Amaral, L.A. (2005) Functional Cartography of Complex Metabolic Networks. Nature, 433, 895-900.

[Huber *et al.*, 2007] Huber, W., et al. (2007) Graphs in Molecular Biology. BMC Bioinformatics, 8 Suppl 6, S8.

[Hung *et al.*, 2010] Hung, J.H., et al. (2010) Identification of Functional Modules That Correlate with Phenotypic Difference: The Influence of Network Topology. Genome Biol, 11, R23.

[Jeong *et al.*, 2000] Jeong, H., et al. (2000) The Large-Scale Organization of Metabolic Networks. Nature, 407, 651-654.

[Kanehisa *et al.*, 2006] Kanehisa, M., et al. (2006) From Genomics to Chemical Genomics: New Developments in Kegg. Nucleic Acids Res, 34, D354-357.

[Klukas and Schreiber, 2007] Klukas, C. and Schreiber, F. (2007) Dynamic Exploration and Editing of Kegg Pathway Diagrams. Bioinformatics, 23, 344-350.

[Koyuturk *et al.*, 2004] Koyuturk, M., et al. (2004) An Efficient Algorithm for Detecting Frequent Subgraphs in Biological Networks. Bioinformatics, 20 Suppl 1, i200-207.

[Li *et al.*, 2009] Li, C., et al. (2009) Subpathwayminer: A Software Package for Flexible Identification of Pathways. Nucleic Acids Res, 37, e131.

[Ogata *et al.*, 2000] Ogata, H., et al. (2000) A Heuristic Graph Comparison Algorithm and Its Application to Detect Functionally Related Enzyme Clusters. Nucleic Acids Res, 28, 4021-4028.

[Schreiber *et al.*, 2002] Schreiber, F. (2002) High Quality Visualization of Biochemical Pathways in Biopath. In Silico Biol, 2, 59-73.

[Shannon *et al.*, 2003] Shannon, P., et al. (2003) Cytoscape: A Software Environment for Integrated Models of Biomolecular Interaction Networks. Genome Res, 13, 2498-2504.

[Smart *et al.*, 2008] Smart, A.G., et al. (2008) Cascading Failure and Robustness in Metabolic Networks. Proc Natl Acad Sci U S A, 105, 13223-13228.

[Strimmer, 2008] Strimmer, K. (2008) fdrtool: a versatile R package for estimating local and tail area-based false discovery rates. Bioinformatics, 24, 1461-1462.

[Team , 2004] Team, R.D.C. (2008) R: A Language and Environment for Statistical Computing. R Foundation Statistical Computing.

[Xia and Wishart, 2010] Xia, J. and Wishart, D.S. (2010) Metpa: A Web-Based Metabolomics Tool for Pathway Analysis and Visualization. Bioinformatics, 26, 2342-2344.

[Zhang and Wiemann, 2009] Zhang, J.D. and Wiemann, S. (2009) Kegggraph: A Graph Approach to Kegg Pathway in R and Bioconductor. Bioinformatics, 25, 1470-1471.