

# STRINGdb Package Vignette

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## 1 INTRODUCTION

STRING (<https://www.string-db.org>) is a database of known and predicted protein-protein interactions. The interactions include direct (physical) and indirect (functional) associations. The database contains information from numerous sources, including experimental repositories, computational prediction methods and public text collections. Each interaction is associated with a combined confidence score that integrates the various evidences. We currently cover over 24 millions proteins from 5090 organisms.

As you will learn in this guide, the STRING database can be usefull to add meaning to list of genes (e.g. the best hits coming out from a screen or the most differentially expressed genes coming out from a Microarray/RNAseq experiment.)

We provide the STRINGdb R package in order to facilitate our users in accessing the STRING database from R. In this guide we explain, with examples, most of the package's features and functionalities.

In the STRINGdb R package we use the new ReferenceClasses of R (search for "ReferenceClasses" in the R documentation.). Besides we make use of the iGraph package (<http://igraph.sourceforge.net>) as a data structure to represent our protein-protein interaction network.

To begin, you should first know the NCBI taxonomy identifiers of the organism on which you have performed the experiment (e.g. 9606 for Human, 10090 for mouse). If you don't know that, you can search the NCBI Taxonomy (<http://www.ncbi.nlm.nih.gov/taxonomy>) or start looking at our species table (that you can also use to verify that your organism is represented in the STRING database).

Hence, if your species is not Human (i.e. our default species), you can find it and their taxonomy identifiers on STRING webpage under the 'organisms' section ([https://string-db.org/cgi/input.pl?input\\_page\\_active\\_form=org](https://string-db.org/cgi/input.pl?input_page_active_form=org)) or download the full list in the download section of STRING website.

```
> library(STRINGdb)
> string_db <- STRINGdb$new( version="11.5", species=9606,
+                             score_threshold=200, network_type="functional", input_directory="")
```

```
WARNING: Only 'full' and 'physical' network types are valid. Setting to the network type to 'full' ST
```

As it has been shown in the above commands, you start instantiating the STRINGdb reference class. In the constructor of the class you can also define the STRING version to be used and a threshold for the combined scores of the interactions, such that any interaction below that threshold is not loaded in the object (by default the score threshold is set to 400).

You can also specify the network type "functional" for full functional STRING network or "physical" for physical subnetwork, which link only the proteins which share a physical complex.

Besides, if you specify a local directory to the parameter input-directory, the database files will be downloaded into this directory and most of the methods can be used off-line. Otherwise, the database files will be saved and cached in a temporary directory that will be cleaned automatically when the R session is closed.

For a better understanding of the package two other commands can be useful:

```
> STRINGdb$methods()           # To list all the methods available.

[1] ".objectPackage"           ".objectParent"
[3] "add_diff_exp_color"       "add_proteins_description"
[5] "benchmark_ppi"            "benchmark_ppi_pathway_view"
[7] "callSuper"                "copy"
[9] "enrichment_heatmap"       "export"
[11] "field"                    "getClass"
[13] "getRefClass"              "get_aliases"
[15] "get_annotations"          "get_bioc_graph"
[17] "get_clusters"             "get_enrichment"
[19] "get_graph"                "get_homologs"
[21] "get_homologs_besthits"     "get_homology_graph"
[23] "get_interactions"          "get_link"
[25] "get_neighbors"            "get_paralogs"
[27] "get_pathways_benchmarking_blackList" "get_png"
[29] "get_ppi_enrichment"        "get_ppi_enrichment_full"
[31] "get_proteins"              "get_pubmed"
[33] "get_pubmed_interaction"    "get_subnetwork"
[35] "get_summary"              "get_term_proteins"
[37] "import"                   "initFields"
[39] "initialize"               "load"
[41] "load_all"                 "map"
[43] "mp"                       "plot_network"
[45] "plot_ppi_enrichment"       "post_payload"
[47] "ppi_enrichment"            "remove_homologous_interactions"
[49] "set_background"           "show"
[51] "show#envRefClass"         "trace"
[53] "untrace"                  "usingMethods"
```

```
> STRINGdb$help("get_graph")  # To visualize their documentation.
```

Call:

```
$get_graph()
```

#### Description:

Return an igraph object with the entire STRING network.  
We invite the user to use the functions of the iGraph package to conveniently search/analyze the network.

#### References:

Csardi G, Nepusz T: The igraph software package for complex network research, InterJournal, Complex Systems 1695. 2006.  
<http://igraph.sf.net>

#### See Also:

In order to simplify the most common tasks, we do also provide convenient functions that wrap some iGraph functions.

```
get_interactions(string_ids) # returns the interactions in between the input proteins
get_neighbors(string_ids)    # Get the neighborhoods of a protein (or of a vector of proteins).
get_subnetwork(string_ids)   # returns a subgraph from the given input proteins
```

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For all the methods that we are going to explain below, you can always use the help function in order to get additional information/parameters with respect to those explained in this guide.

As an example, we use the analyzed data of a microarray study taken from GEO (Gene Expression Omnibus, GSE9008). This study investigates the activity of Resveratrol, a natural phytoestrogen found in red wine and a variety of plants, in A549 lung cancer cells. Microarray gene expression profiling after 48 hours exposure to Resveratrol has been performed and compared to a control composed by A549 lung cancer cells treated only with ethanol. This data is already analyzed for differential expression using the limma package: the genes are sorted by fdr corrected pvalues and the log fold change of the differential expression is also reported in the table.

```
> data(diff_exp_example1)
> head(diff_exp_example1)
```

	pvalue	logFC	gene
1	0.0001018	3.333461	VSTM2L
2	0.0001392	3.822383	TBC1D2
3	0.0001720	3.306056	LENG9
4	0.0001739	3.024605	TMEM27
5	0.0001990	3.854414	L0C100506014
6	0.0002393	3.082052	TSPAN1

As a first step, we map the gene names to the STRING database identifiers using the "map" method. In this particular example, we map from gene HUGO names, but our mapping function supports several other common identifiers (e.g. Entrez GeneID, ENSEMBL proteins, RefSeq transcripts ... etc.).

The map function adds an additional column with STRING identifiers to the dataframe that is passed as first parameter.

```
> example1_mapped <- string_db$map( diff_exp_example1, "gene", removeUnmappedRows = TRUE )
```

Warning: we couldn't map to STRING 15% of your identifiers

As you may have noticed, the previous command prints a warning showing the number of genes that we failed to map. In this particular example, we cannot map all the probes of the microarray that refer to position of the chromosome that are not assigned to a real gene (i.e. all the LOC genes). If we remove all these LOC genes before the mapping we obtain a much lower percentage of unmapped genes (i.e. < 6 %).

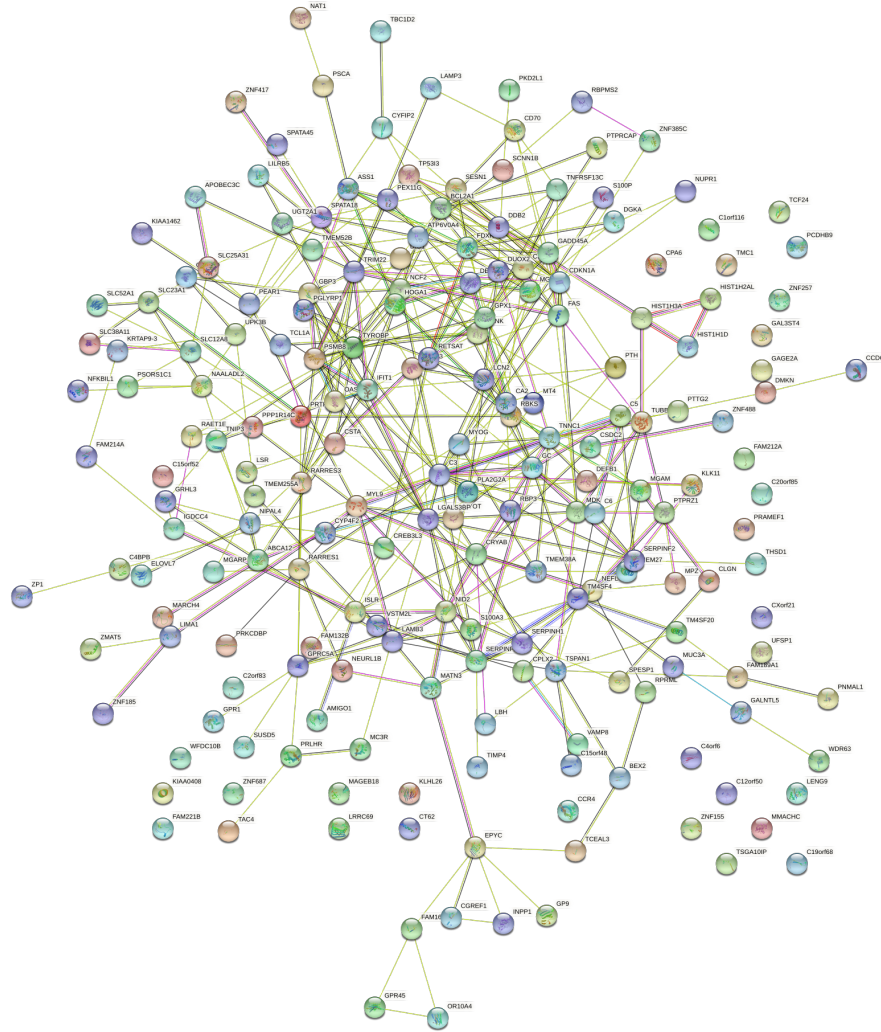
If you set to FALSE the "removeUnmappedRows" parameter, than the rows which corresponds to unmapped genes are left and you can manually inspect them.

Finally, we extract the most significant 200 genes and we produce an image of the STRING network for those. The image shows clearly the genes and how they are possibly functionally related. On the top of the plot, we insert a pvalue that represents the probability that you can expect such an equal or greater number of interactions by chance.

```
> hits <- example1_mapped$STRING_id[1:200]
```

```
> string_db$plot_network( hits )
```

proteins: 200  
interactions: 382  
expected interactions: 229 (p-value: 0)



## 2 PAYLOAD MECHANISM

This R library provides the ability to interact with the STRING payload mechanism. The payload appears as an additional colored "halo" around the bubbles.

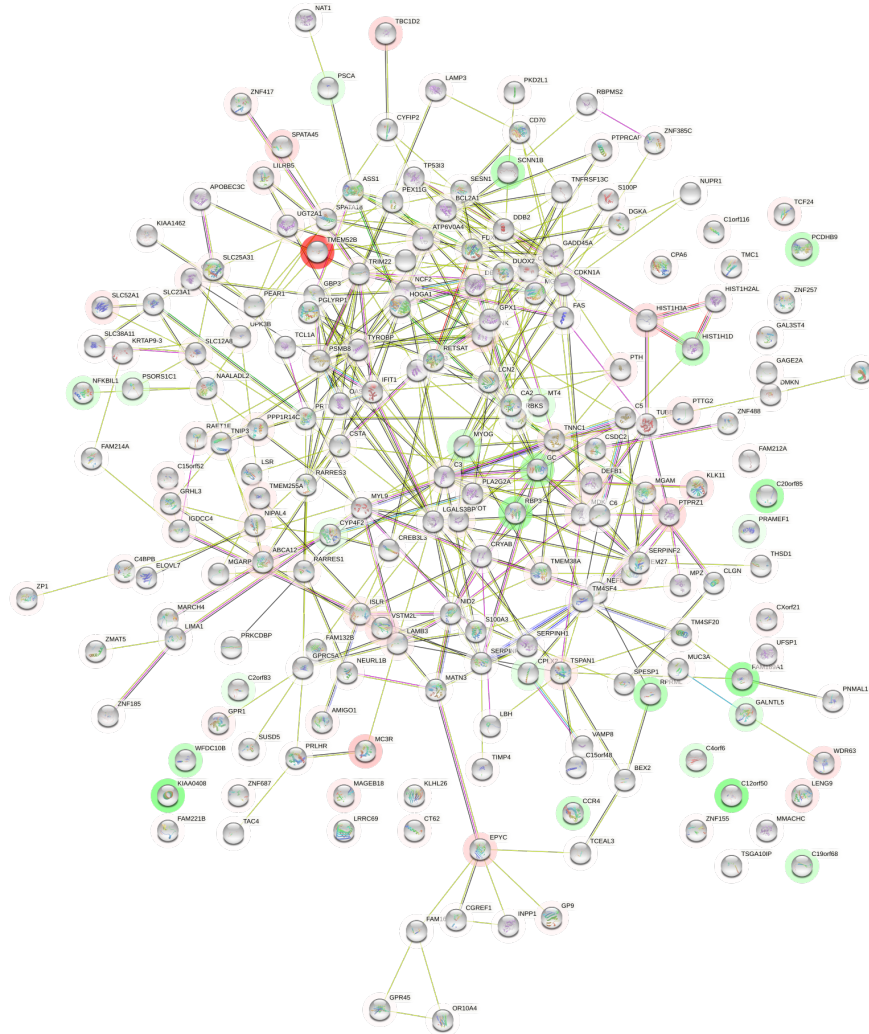
For example, this allows to color in green the genes that are down-regulated and in red the genes that are up-regulated. For this mechanism to work, we provide a function that posts the information on our web server.

```
> # filter by p-value and add a color column
> # (i.e. green down-regulated genes and red for up-regulated genes)
> example1_mapped_pval05 <- string_db$add_diff_exp_color( subset(example1_mapped, pvalue<0.05),
+                                                         logFcColStr="logFC" )

> # post payload information to the STRING server
> payload_id <- string_db$post_payload( example1_mapped_pval05$STRING_id,
+                                     colors=example1_mapped_pval05$color )

> # display a STRING network png with the "halo"
> string_db$plot_network( hits, payload_id=payload_id )
```

proteins: 200  
interactions: 382  
expected interactions: 229 (p-value: 0)



### 3 ENRICHMENT

We provide a method to compute the enrichment in Gene Ontology (Process, Function and Component), KEGG and Reactome pathways, PubMed publications, UniProt Keywords, and PFAM/INTERPRO/SMART domains for your set of proteins all in one simple call. The enrichment itself is computed using an hypergeometric test and the FDR is calculated using Benjamini-Hochberg procedure.

```
> enrichment <- string_db$get_enrichment( hits )
> head(enrichment, n=20)
```

	category	term	number_of_genes	number_of_genes_in_background
1	Process	G0:0006952	34	1296
2	Process	G0:0010951	12	248
3	Process	G0:0051707	31	1256
4	Component	G0:0005576	66	4166
5	Component	G0:0005615	55	3195
6	Component	G0:0070062	41	2099
7	Component	G0:1903561	42	2121
8	TISSUES	BT0:0004850	10	170
9	Keyword	KW-0732	57	3233
10	Keyword	KW-0391	17	522
11	Keyword	KW-0964	37	1818
12	KEGG	hsa04115	6	72
13	WikiPathways	WP4963	8	67

```
ncbiTaxonId
```

1	9606
2	9606
3	9606
4	9606
5	9606
6	9606
7	9606
8	9606
9	9606
10	9606
11	9606
12	9606
13	9606

1	
2	
3	
4	9606. ENSP00000008938, 9606. ENSP00000014914, 9606. ENSP00000187762, 9606. ENSP00000216286, 9606. ENSP00000
5	
6	
7	
8	
9	
10	
11	
12	
13	

```

1
2
3
4 PGLYRP1,GPRC5A,TMEM38A,NID2,C5,RARRES1,C4BPB,CD70,C3,ISLR,SERPINF1,THSD1,EPYC,LGALS3BP,C6,VAMP8,C5
5 PGLYRP1,GPRC5A,TMEM38A,NID2,C5,RARRES1,C4BPB,CD70,C3,ISLR,SERPINF1,THSD1,EPYC,LGALS3BP,C6,VAMP8,C5
6
7
8
9 PGLYRP1,NID2,C5,C4BPB,C3,ISLR,SERPINF1,THSD1,EPYC,LGALS3BP,C6,VAMP8,C5
10
11
12
13

```

	p_value	fdr	description
1	5.07e-07	0.00650	Defense response
2	1.43e-05	0.03780	Negative regulation of endopeptidase activity
3	5.89e-06	0.03780	Response to other organism
4	9.03e-05	0.04080	Extracellular region
5	5.17e-05	0.04080	Extracellular space
6	4.18e-05	0.04080	Extracellular exosome
7	2.40e-05	0.04080	Extracellular vesicle
8	1.54e-05	0.02940	Bone marrow cell
9	1.78e-05	0.01200	Signal
10	3.64e-05	0.01230	Immunity
11	4.61e-05	0.01230	Secreted
12	1.40e-04	0.04690	p53 signaling pathway
13	9.02e-07	0.00061	p53 transcriptional gene network

If you have performed your experiment on a predefined set of proteins, it is important to run the enrichment statistics using that set as a background (otherwise you would get a wrong p-value!). Hence, before to launch the method above, you may want to set the background:

```

> backgroundV <- example1_mapped$STRING_id[1:2000] # as an example, we use the first 2000 genes
> string_db$set_background(backgroundV)

```

You can also set the background when you instantiate the STRINGdb object:

```

> string_db <- STRINGdb$new( score_threshold=200, backgroundV = backgroundV )

```

If you just want to know terms are assigned to your set of proteins (and not necessary enriched) you can use "get\_annotations" method. This method will output all the terms from most of the categories (the exceptions are KEGG terms due to licensing issues and PubMed due to the size of the output) that are associated with your set of proteins.

```

> annotations <- string_db$get_annotations( hits )
> head(annotations, n=20)

```

	category	term_id	number_of_genes	ratio_in_set	species
1	COMPARTMENTS	GOCC:0000109	1	0.005	9606
2	COMPARTMENTS	GOCC:0000139	2	0.010	9606
3	COMPARTMENTS	GOCC:0000151	1	0.005	9606
4	COMPARTMENTS	GOCC:0000228	1	0.005	9606
5	COMPARTMENTS	GOCC:0000307	1	0.005	9606
6	COMPARTMENTS	GOCC:0000323	6	0.030	9606
7	COMPARTMENTS	GOCC:0000502	1	0.005	9606
8	COMPARTMENTS	GOCC:0000785	2	0.010	9606
9	COMPARTMENTS	GOCC:0000786	1	0.005	9606
10	COMPARTMENTS	GOCC:0000791	1	0.005	9606
11	COMPARTMENTS	GOCC:0001533	1	0.005	9606
12	COMPARTMENTS	GOCC:0001650	1	0.005	9606
13	COMPARTMENTS	GOCC:0001669	1	0.005	9606
14	COMPARTMENTS	GOCC:0001725	1	0.005	9606
15	COMPARTMENTS	GOCC:0001726	2	0.010	9606
16	COMPARTMENTS	GOCC:0002133	1	0.005	9606
17	COMPARTMENTS	GOCC:0005576	40	0.200	9606
18	COMPARTMENTS	GOCC:0005577	1	0.005	9606
19	COMPARTMENTS	GOCC:0005579	3	0.015	9606
20	COMPARTMENTS	GOCC:0005604	2	0.010	9606

1  
2  
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16  
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1  
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3  
4  
5  
6  
7  
8

9606. ENSP00000008938, 9606. ENSP00000216286, 9606. ENSP00000223642, 9606. ENSP00000237696, 9606. ENSP00000

```

9
10
11
12
13
14
15
16
17 PGLYRP1,NID2,C5,RARRES1,C4BPB,CD70,C3,SERPINF1,THSD1,LGALS3BP,C6,CSTA,ANXA3,PTH,CA2,TIMP4,DEFB1,PS
18
19
20

```

	description
1	Nucleotide-excision repair complex
2	Golgi membrane
3	Ubiquitin ligase complex
4	Nuclear chromosome
5	Cyclin-dependent protein kinase holoenzyme complex
6	Lytic vacuole
7	Proteasome complex
8	Chromatin
9	Nucleosome
10	Euchromatin
11	Cornified envelope
12	Fibrillar center
13	Acrosomal vesicle
14	Stress fiber
15	Ruffle
16	Polycystin complex
17	Extracellular region
18	Fibrinogen complex
19	Membrane attack complex
20	Basement membrane

## 4 CLUSTERING

The iGraph package provides several clustering/community algorithms: "fastgreedy", "walktrap", "spinglass", "edge.betweenness". We encapsulate this in an easy-to-use function that returns the clusters in a list.

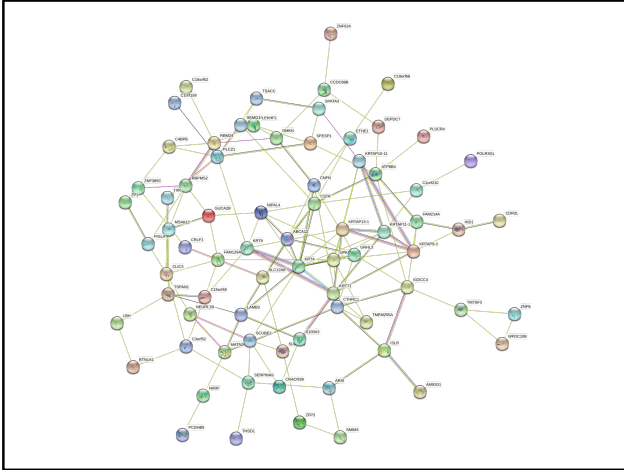
```

> # get clusters
> clustersList <- string_db$get_clusters(example1_mapped$STRING_id[1:600])

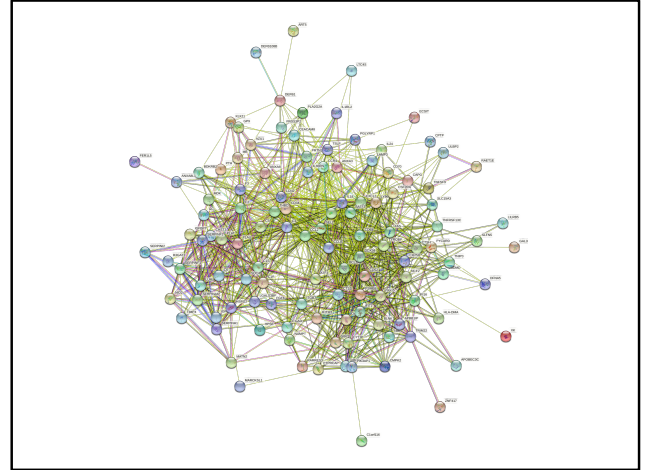
> # plot first 4 clusters
> par(mfrow=c(2,2))
> for(i in seq(1:4)){
+   string_db$plot_network(clustersList[[i]])
+ }

```

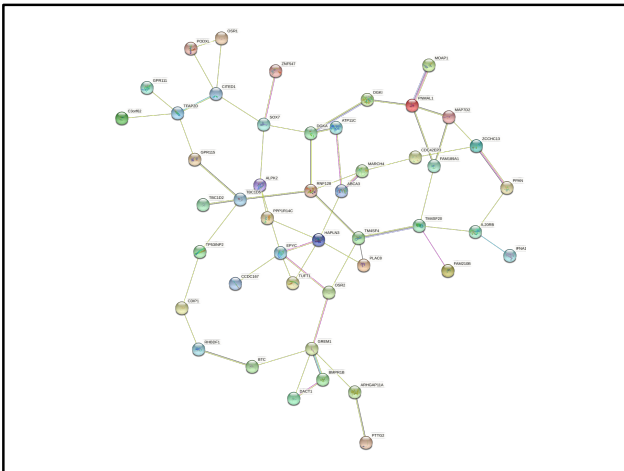
proteins: 74  
interactions: 137  
expected interactions: 13 (p-value: 0)



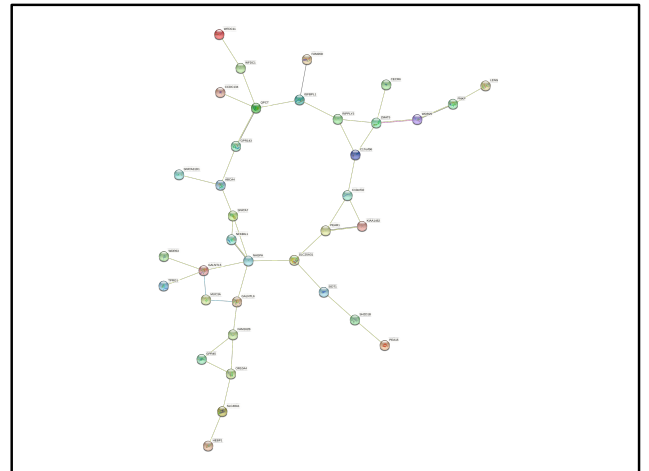
proteins: 119  
interactions: 934  
expected interactions: 175 (p-value: 0)



proteins: 46  
interactions: 59  
expected interactions: 8 (p-value: 0)



proteins: 36  
interactions: 41  
expected interactions: 3 (p-value: 0)



## 5 ADDITIONAL PROTEIN INFORMATION

You can get a table that contains all the proteins that are present in our database of the species of interest. The protein table also include the preferred name, the size and a short description of each protein.

```
> string_proteins <- string_db$get_proteins()
```

In the following section we will show how to query STRING with R on some specific proteins. In the examples, we will use the famous tumor proteins TP53 and ATM .

First we need to get the STRING identifier of those proteins, using our mp method:

```
> tp53 = string_db$mp( "tp53" )
> atm = string_db$mp( "atm" )
```

The mp method (i.e. map proteins) is an alternative to our map method, to be used when you need to map only one or few proteins.

It takes in input a vector of protein aliases and returns a vector with the STRING identifiers of those proteins.

Using the following method, you can see the proteins that interact with one or more of your proteins:

```
> string_db$get_neighbors( c(tp53, atm) )
```

It is also possible to retrieve the interactions that connect certain input proteins between each other. Using the "get\_interactions" method we can clearly see that TP53 and ATM interact with each other with a good evidence/score.

```
> string_db$get_interactions( c(tp53, atm) )
```

	from	to	combined_score
1	9606.ENSEP00000269305	9606.ENSEP00000278616	999
2	9606.ENSEP00000269305	9606.ENSEP00000278616	999

STRING provides a way to get homologous proteins: in our database we store ALL-AGAINST-ALL alignments within all 5090 organisms. You can retrieve all of the paralogs of the protein using "get\_paralogs" method.

```
> # Get all homologs of TP53 in human.
> string_db$get_paralogs(tp53)
```

STRING also stores best hits (as measured by bitscore) between the proteins from different species. "get\_homologs\_bests" lets you retrieve these homologs.

```
> # get the best hits of the following protein in all the STRING species
> string_db$get_homologs_besthits(tp53)
```

... or you can specify the species of interest (i.e. all the blast hits):

```
> # get the homologs of the following two proteins in the mouse (i.e. species_id=10090)
> string_db$get_homologs_besthits(c(tp53, atm), target_species_id=10090, bitscore_threshold=60)
```

## 6 CITATION

Please cite:

Szklarczyk D, Gable AL, Nastou KC, Lyon D, Kirsch R, Pyysalo S, Doncheva NT, Legeay M, Fang T, Bork P, Jensen LJ, von Mering C. 'The STRING database in 2021: customizable protein-protein networks, and functional characterization of user-uploaded gene/measurement sets.' Nucleic Acids Res. 2021 Jan 8;49(D1):D605-12