

# Package: annotaR (via r-universe)

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**Title** Tidy, Integrated Gene Annotation

**Version** 0.1.1

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**Description** A framework for intuitive, multi-source gene and protein annotation, with a focus on integrating functional genomics with disease and drug data for translational insights. Methods used include g:Profiler (Raudvere et al. (2019) <doi:10.1093/nar/gkz369>), biomaRt (Durinck et al. (2009) <doi:10.1038/nprot.2009.97>), and the Open Targets Platform (Koscielny et al. (2017) <doi:10.1093/nar/gkw1055>).

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add_disease_links	<i>Add disease association data</i>
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### Description

Augments an annotaR object with disease association data from the OpenTargets platform.

### Usage

```
add_disease_links(annotaR_object, score_threshold = 0.5)
```

### Arguments

`annotaR_object` A tibble, typically from `annotaR()`, containing a 'gene' column with HGNC symbols.

`score_threshold` Minimum association score (from 0 to 1) to include. Defaults to 0.5.

### Value

A new tibble with the original data joined with disease association columns (`disease_name`, `association_score`).

### Examples

```
annotaR(c("TP53", "EGFR")) %>%
  add_disease_links(score_threshold = 0.8)
```

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add_drug_links	<i>Add known drug association data</i>
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### Description

Augments an annotaR object with known drug/compound data from the OpenTargets platform. This includes the drug name, type, mechanism of action, and clinical trial phase.

### Usage

```
add_drug_links(annotaR_object)
```

**Arguments**

`annotaR_object` A tibble, typically from `annotaR()`, containing a 'gene' column with HGNC symbols.

**Value**

A new tibble with the original data joined with drug association columns (e.g., `drug_name`, `drug_type`, `mechanism_of_action`, `phase`).

**Examples**

```
annotaR(c("EGFR", "BRAF")) %>%
  add_drug_links()
```

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`add_go_terms`

*Add GO functional enrichment data*

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**Description**

Augments an `annotaR` object with functional enrichment data from `g:Profiler`. It performs a Gene Ontology (GO) analysis on the gene list and joins the results.

**Usage**

```
add_go_terms(annotaR_object, organism = "hsapiens", sources = c("GO:BP"), ...)
```

**Arguments**

`annotaR_object` A tibble, typically the output of `annotaR()`. Must contain a 'gene' column.

`organism` The organism name to use for the query (e.g., "hsapiens"). Passed to `gprofiler2::gost`.

`sources` A vector of data sources to query. Defaults to GO Biological Process. See `gprofiler2::gost` for options.

... Additional parameters passed on to `gprofiler2::gost`.

**Value**

A new tibble with the original 'gene' column joined with functional annotation columns (e.g., `term_id`, `term_name`, `p_value`, `source`).

**Examples**

```
annotaR(c("TP53", "EGFR")) %>%
  add_go_terms()
```

annotaR

*Create an annotaR object*

---

**Description**

Initializes the annotation pipeline by creating a tibble from a character vector of gene symbols. This is the entry point for a typical annotaR workflow.

**Usage**

```
annotaR(genes)
```

**Arguments**

genes            A character vector of HGNC gene symbols (e.g., c("TP53", "BRCA1")).

**Value**

A tibble with a single column 'gene', ready to be used in downstream annotation functions.

**Examples**

```
my_genes <- c("TP53", "EGFR", "BRCA1")
annotaR(my_genes)
```

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plot\_enrichment\_dotplot

*Plot GO Enrichment Results as a Dot Plot*

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**Description**

Creates a publication-ready dot plot from the results of an `add_go_terms()` call. The plot shows the top enriched terms, with dot size representing the number of genes and color representing the p-value.

**Usage**

```
plot_enrichment_dotplot(
  annotaR_object,
  n_terms = 20,
  title = "Top GO Enrichment Results"
)
```

**Arguments**

- `annotaR_object` An object processed by `add_go_terms()`. Must contain `term_name`, `p_value`, and `gene` columns.
- `n_terms` The maximum number of top terms to display, ordered by p-value. Defaults to 20.
- `title` The title of the plot.

**Value**

A ggplot object.

**Examples**

```
# Create a dummy annotaR object with enrichment data
annotated_data <- tibble::tibble(
  gene = c("TP53", "TP53", "EGFR"),
  term_name = c("Cell cycle", "Apoptosis", "Cell cycle"),
  p_value = c(0.001, 0.005, 0.001),
  source = "GO:BP",
  intersection = "TP53,EGFR"
)

plot_enrichment_dotplot(annotated_data)
```

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