
grasp Documentation

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A Simple GRASP (grasp.nhlbi.nih.gov) API based on SQLAlchemy and Pandas.

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License	MIT License, made at Stanford, use as you wish.
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For an introduction see the [github readme](#)

For table information see the [wiki](#)

Basic Usage

This module contains a Python 3 API to work with the GRASP database. The database must be downloaded and initialized locally. The default is to use an sqlite backend, but postgresql or mysql may be used also; these two are slower to initialize (longer write times), but they may be faster on indexed reads.

The GRASP project is a SNP-level index of over 2000 GWAS datasets. It is very useful, but difficult to batch query as study descriptions are heterogenous and there are more than 9 million rows. By putting this information into a relational database, it is easy to pull out bite-sized chunks of data to analyze with [pandas](#).

Commonly queried columns are indexed within the database for fast retrieval. A typical query for a single phenotype category returns several million SNPs in about 10 seconds, which can then be analyzed with pandas.

To read more about GRASP, visit the [official page](#).

For reference information (e.g. column, population, and phenotype lists) see [the wiki](#).

For complete API documentation, go to the [documentation site](#)

1.1 Installation

Use the standard installation procedure:

```
git clone https://github.com/MikeDacre/grasp.git
cd grasp
python ./setup.py install --user
```

This code requires a grasp database. Currently sqlite/postgresql/mysql are supported. Mysql and postgresql can be remote (but must be set up with this tool), sqlite is local.

Database configuration is stored in a config file that lives by default in `~/grasp`. This path is set in `config.py` and can be changed there is needed.

A script, `grasp`, is provided in `bin` and should automatically be installed to your `PATH`. It contains functions to set up your database config and to initialize the grasp database easily, making the initial steps trivial.

To set up your database configuration, run:

```
grasp config --init
```

This will prompt you for your database config options and create a file at `~/grasp` with those options saved.

You can now initialize the grasp database:

```
grasp init study_file grasp_file
```

The study file is available in this repository ([grasp2_studies.txt.gz](#)) It is just a copy of the official [GRASP List of Studies](#) converted to text and with an additional index that provides a numeric index for the non pubmed indexed studies.

Both files can be gzipped or bziped.

The grasp file is the raw unzipped file from the project page: [GRASP2fullDataset](#)

The database takes about 90 minutes to build on a desktop machine and uses about 3GB of space. The majority of the build time is spent parsing dates, but because the dates are encoded in the SNP table, and the formatting varies, this step is required.

1.2 Usage

The code is based on SQLAlchemy, so you should read their [ORM Query tutorial](#) to know how to use this well.

It is important to note that the point of this software is to make bulk data access from the GRASP DB easy, SQLAlchemy makes this very easy indeed. However, to do complex comparisons, SQLAlchemy is very slow. As such, the best way to use this software is to use SQLAlchemy functions to bulk retrieve study lists, and then to directly get a pandas dataframe of SNPs from those lists.

Tables are defined in *grasp.tables* Database setup functions are in *grasp.db* Query tools for easy data manipulation are in *grasp.query*.

1.2.1 Tables

This module provides 6 tables:

[Study](#), [Phenotype](#), [PhenoCats](#), [Platform](#), [Population](#), and [SNP](#) (as well as several association tables)

1.2.2 Querying

The functions in *grasp.query* are very helpful in automating common queries.

The simplest way to get a dataframe from SQLAlchemy is like this:

```
df = pandas.read_sql(session.query(SNP).statement)
```

Note that if you use this exact query, the dataframe will be too big to be useful. To get a much more useful dataframe:

```
studies = grasp.query.get_studies(pheno_cats='t2d', primary_pop='European')
df = grasp.query.get_snps(studies)
```

It is important to note that there are **three** ways of getting phenotype information: - The Phenotype table, which lists the primary phenotype for every study - The PhenoCats table, which lists the GRASP curated phenotype categories, each Study has several of these.

- The phenotype_desc column in the SNP table, this is a poorly curated column directly from the full dataset, it roughly corresponds to the information in the Phenotype table, but the correspondance is not exact due to an abundance of typos and slightly differently typed information.

1.3 Example Workflow

```

from grasp import db
from grasp import tables as t
from grasp import query as q
s, e = db.get_session()

# Print a list of all phenotypes (also use with populations, but not with SNPs (too_
↳many to display))
s.query(t.PhenoType).all()

# Filter the list
s.query(t.PhenoType).filter(t.PhenoType.phenotype.like('%diabetes%')).all()

# Get a dictionary of studies to review
eur_t2d = get_studies(only_disc_pop='eur', primary_phenotype='Type II Diabetes_
↳Mellitus', dictionary=True)

# Filter those by using eur.pop() to remove unwanted studies, and then get the SNPs_
↳as a dataframe
eur_snps_df = get_snps(eur, pandas=True)

# Do the same thing for the african population
afr_t2d = get_studies(only_disc_pop='afr', primary_phenotype='Type II Diabetes_
↳Mellitus', dictionary=True)
afr.pop('Use of diverse electronic medical record systems to identify genetic risk_
↳for type 2 diabetes within a genome-wide association study.')
afr_snps_df = get_snps(afr, pandas=True)

# Collapse the matrices (take median of pvalue) and filter by resulting pvalue
eur_snps_df = q.collapse_dataframe(eur_snps_df, mechanism='median', pvalue_filter=5e-
↳8)
afr_snps_df = q.collapse_dataframe(afr_snps_df, mechanism='median', pvalue_filter=5e-
↳8)

# The new dataframes are indexed by 'chr:pos'

# Plot the overlapping SNPs
snps = q.intersect_overlapping_series(eur_snps_df.pval_median, afr_snps_df.pval_
↳median)
snps.plot()

```

GRASP Console Script

A Simple GRASP (grasp.nhlbi.nih.gov) API based on SQLAlchemy and Pandas

Author	Michael D Dacre < mike.dacre@gmail.com >
Organization	Stanford University
License	MIT License, use as you wish
Created	2016-10-08
Version	0.3.0b1

Last modified: 2016-10-17 00:18

This is the front-end to a python grasp api, intended to allow easy database creation and simple querying. For most of the functions of this module, you will need to call the module directly.

```
usage: grasp [-h] {search,conf,info,init} ...
```

Sub-commands:

search (s, lookup) Query database for variants by location or id

Query for SNPs in the database. By default returns a tab-delimited list of SNPs with the following columns: 'id', 'snpid', 'study_snpid', 'chrom', 'pos', 'phenotype', 'pval'

The `--extra` flag adds these columns: 'InGene', 'InMiRNA', 'inLincRNA', 'LSSNP'

The `--study-info` flag adds these columns: 'study_id (PMID)', 'title'

The `--db-snp` flag uses the myvariant API to pull additional data from db_snp.

```
usage: grasp search [-h] [--extra] [--study-info] [--db-snp] [--pandas] [-o]
                  [--path]
                  query
```

Positional arguments:

query rsID, chrom:loc or chrom:start-end

Options:

--extra Add some extra columns to output

--study-info Include study title and PMID

--db-snp Add dbSNP info to output

--pandas Write output as a pandas dataframe

-o, --out File to write to, default STDOUT.

--path PATH to write files to

conf (config) Manage local config

```
usage: grasp conf [-h]
                  [--db {sqlite,postgresql,mysql} | --get-path | --set-path_
↪PATH | --init]
```

Options:

--db Set the current database platform.
Possible choices: sqlite, postgresql, mysql

--get-path Change the sqlite file path

--set-path Change the sqlite file path

--init Initialize the config with default settings. Will ERASE your old config!

info Display database info

Write data summaries (also found on the wiki) to a file or the console.

Choices: all: Will write everything to separate rst files, ignores all other flags except ‘-path‘ phenotypes: All primary phenotypes. phenotype_categories: All phenotype categories. populations: All primary populations. population_flags: All population flags. snp_columns: All SNP columns. study_columns: All Study columns.

```
usage: grasp info [-h] [-o] [--path]
                  {phenotype_categories,populations,phenotypes,study_columns,
↪snp_columns,all,population_flags}
```

Positional arguments:

display Choice of item to display, if all, results are written to independant rst files, and optional args are ignored
Possible choices: phenotype_categories, populations, phenotypes, study_columns, snp_columns, all, population_flags

Options:

-o, --out File to write to, default STDOUT.

--path PATH to write files to

init Initialize the database

```
usage: grasp init [-h] [-n] study_file grasp_file
```

Positional arguments:

study_file GRASP study file from: github.com/MikeDacre/grasp/blob/master/grasp2_studies.txt

grasp_file GRASP tab delimited file

Options:

-n, --no-progress Do not display a progress bar

Library (API Documentation)

This code is intended to be primarily used as a library, and works best when used in an interactive python session (e.g. with jupyter) alongside pandas. Many of the query functions in this library returns pandas dataframes.

Below is a complete documentation of the API for this library. The functions in query will be the most interesting for most users.

3.1 grasp.tables

GRASP table descriptions in SQLAlchemy ORM.

These tables do not exist in the GRASP data, which is a single flat file. By separating the data into these tables querying is much more efficient.

This submodule should only be used for querying.

```
class grasp.tables.SNP (**kwargs)
    Bases: sqlalchemy.ext.declarative.api.Base
    An SQLAlchemy Table for GRASP SNPs.
    Study and phenotype information are pushed to other tables to minimize table size and make querying easier.
    Table Name: snps
    Columns: Described in the columns attribute
    int
        The ID number of the SNP, usually the NHLBIkey
    str
        SNP loction expressed as 'chr:pos'
    hvg_ids
        A list of HGVS IDs for this SNP
    columns
        A dictionary of all columns 'column_name'=>('type', 'desc')
    ConservPredTFBS
    CreationDate
    EqtlMethMetabStudy
    HUPfield
```

HumanEnhancer

InGene

InLincRNA

InMiRNA

InMiRNABS

LSSNP

LastCurationDate

NHLBIkey

NearestGene

ORegAnno

PolyPhen2

RNAedit

SIFT

UniProt

chrom

columns = `OrderedDict([('id', ('BigInteger', 'NHLBIkey')), ('snpid', ('String', 'SNPid')), ('chrom', ('String', 'chr')), ('p`

A description of all columns in this table.

dbSNPClinStatus

dbSNPMAF

dbSNPfxn

dbSNPinfo

dbSNPvalidation

display_columns (*display_as='table', write=False*)

Return all columns in the table nicely formatted.

Display choices: table: A formatted grid-like table tab: A tab delimited non-formatted version of table

list: A string list of column names

Parameters

- **display_as** – {table,tab,list}
- **write** – If true, print output to console, otherwise return string.

Returns A formatted string or None

get_columns (*return_as='list'*)

Return all columns in the table nicely formatted.

Display choices: list: A python list of column names dictionary: A python dictionary of name=>desc

long_dict: A python dictionary of name=>(type, desc)

Parameters **return_as** – {table,tab,list,dictionary,long_dict,id_dict}

Returns A list or dictionary

get_variant_info (*fields='db SNP', pandas=True*)

Use the myvariant API to get info about this SNP.

Note that this service can be very slow. It will be faster to query multiple SNPs.

Parameters

- **fields** – Choose fields to display from: docs.myvariant.info/en/latest/doc/data.html#available-fields Good choices are 'db SNP', 'clinvar', or 'gwassnps' Can also use 'grasp' to get a different version of this info.
- **pandas** – Return a dataframe instead of dictionary.

Returns A dictionary or a dataframe.

hvg IDs

The HVGS ID from myvariant.

id

paper_loc

phenotype_cats

phenotype_desc

population

population_id

pos

pval

snp_loc

Return a simple string containing the SNP location.

snpid

study

study_id

study_snpid

class grasp.tables.**Phenotype** (***kwargs*)

Bases: sqlalchemy.ext.declarative.api.Base

An SQLAlchemy table to store the primary phenotype.

Table Name: phenos

Columns: phenotype: The string phenotype from the GRASP DB, unique. alias: A short representation of the phenotype, not unique. studies: A link to the studies table.

int

The ID number.

str

The name of the phenotype.

alias

id

phenotype

studies

```
class grasp.tables.PhenoCats (**kwargs)
    Bases: sqlalchemy.ext.declarative.api.Base
    An SQLAlchemy table to store the lists of phenotype categories.
    Table Name: pheno_cats
    Columns: category: The category from the grasp database, unique. alias: An easy to use alias of the category,
        not unique. snps: A link to all SNPs in this category. studies: A link to all studies in this category.

    int
        The PhenoCat ID
    str
        The category name
    alias
    category
    id
    snps
    studies

class grasp.tables.Platform(platform)
    Bases: sqlalchemy.ext.declarative.api.Base
    An SQLAlchemy table to store the platform information.
    Table Name: platforms
    Columns: platform: The name of the platform from GRASP. studies: A link to all studies using this platform.

    int
        The ID number of this platform
    str
        The name of the platform
    id
    platform
    studies

class grasp.tables.Population(population)
    Bases: sqlalchemy.ext.declarative.api.Base
    An SQLAlchemy table to store the platform information.
    Table Name: populations
    Columns: population: The name of the population. studies: A link to all studies in this population. snps: A
        link to all SNPs in this populations.

    int
        Population ID number
    str
        The name of the population
    id
    population
```


3.2 grasp.db

Functions for managing the GRASP database.

`get_session()` is used everywhere in the module to create a connection to the database. `initialize_database()` is used to build the database from the GRASP file. It takes about an hour 90 minutes to run and will overwrite any existing database.

`grasp.db.get_session(echo=False)`

Return a session and engine, uses config file.

Parameters `echo` – Echo all SQL to the console.

Returns

A SQLAlchemy session and engine object corresponding to the grasp database for use in querying.

Return type session, engine

`grasp.db.initialize_database(study_file, grasp_file, commit_every=250000, progress=False)`

Create the database quickly.

Study_file Tab delimited GRASP study file, available here:
github.com/MikeDacre/grasp/blob/master/grasp_studies.txt

Grasp_file Tab delimited GRASP file.

Commit_every How many rows to go through before committing to disk.

Progress Display a progress bar (db length hard coded).

3.3 grasp.query

A mix of functions to make querying the database and analyzing the results faster.

`grasp.query.get_studies(primary_phenotype=None, pheno_cats=None, pheno_cats_alias=None, primary_pop=None, has_disc_pop=None, has_rep_pop=None, only_disc_pop=None, only_rep_pop=None, query=False, count=False, dictionary=False, pandas=False)`

Return a list of studies filtered by phenotype and population.

There are two ways to query both phenotype and population.

Phenotype: GRASP provides a ‘primary phenotype’ for each study, which are fairly poorly curated. They also provide a list of phenotype categories, which are well curated. The problem with the categories is that there are multiple per study and some are too general to be useful. If using categories be sure to post filter the study list.

Note: I have made a list of aliases for the phenotype categories to make them easier to type. Use `pheno_cats_alias` for that.

Population: Each study has a primary population (list available with ‘get_populations’) but some studies also have other populations in the cohort. GRASP indexes all population counts, so those can be used to query also. To query these use `has_` or `only_` (exclusive) parameters, you can query either discovery populations or replication populations. Note that you cannot provide both `has_` and `only_` parameters for the same population type.

For doing population specific analyses most of the time you will want the `excl_disc_pop` query.

Argument Description: Phenotype Arguments are 'primary_phenotype', 'pheno_cats', and 'pheno_cats_alias'.

Only provide one of pheno_cats or pheno_cats_alias

Population Arguments are *primary_pop*, *has_disc_pop*, *has_rep_pop*, *only_disc_pop*, *only_rep_pop*.

primary_pop is a simple argument, the others use bitwise flags for lookup.

The easiest way to use the following parameters is with the `_ref.PopFlag` object. It uses py-flags. For example:

```
pops = _ref.PopFlag.eur | _ref.PopFlag.afr
```

In addition you can provide a list of strings corresponding to PopFlag attributes.

Note: the *only_* parameters work as ANDs, not ORs. So *only_disc_pop*=`'eurlafr'` will return those studies that have BOTH european and african discovery populations, but no other discovery populations. On the other hand, *has_* works as an OR, and will return any study with any of the specified populations.

Parameters

- **primary_phenotype** – Phenotype of interest, string or list of strings.
- **pheno_cats** – Phenotype category of interest.
- **pheno_cats_alias** – Phenotype category of interest.
- **primary_pop** – Query the primary population, string or list of strings.
- **has_disc_pop** – Return all studies with these discovery populations
- **has_rep_pop** – Return all studies with these replication populations
- **only_disc_pop** – Return all studies with ONLY these discovery populations
- **only_rep_pop** – Return all studies with ONLY these replication populations
- **query** – Return the query instead of the list of study objects.
- **count** – Return a count of the number of studies.
- **dictionary** – Return a dictionary of title->id for filtering.
- **pandas** – Return a dataframe of study information instead of the list.

Returns A list of study objects, a query, or a dataframe.

`grasp.query.get_snps(studies, pandas=True)`

Return a list of SNPs in a single population in a single phenotype.

Studies A list of studies.

Pandas Return a dataframe instead of a list of SNP objects.

Returns Either a DataFrame or list of SNP objects.

`grasp.query.get_variant_info(snp_list, fields='db SNP', pandas=True)`

Get variant info for a list of SNPs.

Parameters

- **snp_list** – A list of SNP objects or SNP rsIDs
- **fields** – Choose fields to display from: docs.myvariant.info/en/latest/doc/data.html#available-fields Good choices are 'db SNP', 'clinvar', or 'gwassnps' Can also use 'grasp' to get a different version of this info.

- **pandas** – Return a dataframe instead of dictionary.

Returns A dictionary or a dataframe.

```
grasp.query.find_intersecting_phenotypes(primary_pops=None, pop_flags=None,
                                         check='cat', pop_type='disc', exclusive=False,
                                         list_only=False)
```

Return a list of phenotypes that are present in all populations.

Can only provide one of primary_pops or pop_flags. pop_flags does a bitwise lookup, primary_pops queries the primary string only.

By default this function returns a list of phenotype categories, if you want to check primary phenotypes instead, provide check='primary'.

Parameters

- **primary_pops** – A string or list of strings corresponding to the *tables.Study.phenotype* column
- **pop_flags** – A *ref.PopFlag* object or list of objects.
- **check** – cat/primary either check categories or primary phenos.
- **pop_type** – disc/rep Use with pop_flags only, check either discovery or replication populations.
- **exclusive** – Use with pop_flags only, do an exclusive rather than inclusion population search
- **list_only** – Return a list of names only, rather than a list of objects

Returns A list of *table.Phenotype* or *table.PhenoCat* objects, or a list of names if *list_only* is specified.

```
grasp.query.collapse_dataframe(df, mechanism='median', pvalue_filter=None,
                               protected_columns=None)
```

Collapse a dataframe by chrom:location from get_snps.

Will use the mechanism defined by 'mechanism' to collapse a dataframe to one indexed by 'chrom:location' with pvalue and count only.

This function is agnostic to all dataframe columns other than:

```
['chrom', 'pos', 'snpid', 'pval']
```

All other columns are collapsed into a comma separated list, a string. 'chrom' and 'pos' are merged to become the new colon-separated index, snpid is maintained, and pval is merged using the function in 'mechanism'.

Parameters

- **df** – A pandas dataframe, must have 'chrom', 'pos', 'snpid', and 'pval' columns.
- **mechanism** – A numpy statistical function to use to collapse the pvalue, median or mean are the common ones.
- **pvalue_filter** – After collapsing the dataframe, filter to only include pvalues less than this cutoff.
- **protected_columns** – A list of column names that will be maintained as is, although all duplicates will be dropped (randomly). Only makes sense for columns that are identical for all studies of the same SNP.

Returns

Indexed by chr:pos, contains flattened pvalue column, and all original columns as a comma-separated list. Additionally contains a count and stddev (of pvalues) column. stddev is nan if count is 1.

Return type DataFrame

```
grasp.query.intersect_overlapping_series(series1, series2, names=None, stats=True,
                                         plot=False)
```

Plot all SNPs that overlap between two pvalue series.

Parameters

- **series** – A pandas series object
- **names** – A list of two names to use for the resultant dataframes
- **stats** – Print some stats on the intersection
- **plot** – Plot the resulting intersection

Returns with the two series as columns

Return type DataFrame

3.4 grasp.config

Manage a persistent configuration for the database.

```
grasp.config.config = <configparser.ConfigParser object>
```

A globally accessible ConfigParser object, initialized with CONFIG_FILE.

```
grasp.config.CONFIG_FILE = '/Users/dacre/.grasp'
```

The PATH to the config file.

```
grasp.config.init_config(db_type, db_file='', db_host='', db_user='', db_pass='')
```

Create an initial config file.

Parameters

- **db_type** – 'sqlite/mysql/postgresql'
- **db_file** – PATH to sqlite database file
- **db_host** – Hostname for mysql or postgresql server
- **db_user** – Username for mysql or postgresql server
- **db_pass** – Password for mysql or postgresql server (not secure)

Returns NoneType

Return type None

```
grasp.config.init_config_interactive()
```

Interact with the user to create a new config.

Uses readline autocompletion to make setup easier.

```
grasp.config.write_config()
```

Write the current config to CONFIG_FILE.

3.5 grasp.info

Little functions to pretty print column lists and category info.

`get_{phenotypes,phenotype_categories,populations}` all display a dump of the whole database.

`get_population_flags` displays available flags from PopFlag.

`display_{study,snp}_columns` displays a list of available columns in those two tables as a formatted string.

`get_{study,snp}_columns` return a list of available columns in those two tables as python objects.

`grasp.info.display_snp_columns` (*display_as='table', write=False*)

Return all columns in the SNP table as a string.

Display choices: `table`: A formatted grid-like table `tab`: A tab delimited non-formatted version of table `list`: A string list of column names

Parameters

- **display_as** – {table,tab,list}
- **write** – If true, print output to console, otherwise return string.

Returns A formatted string or None

`grasp.info.display_study_columns` (*display_as='table', write=False*)

Return all columns in the Study table as a string.

Display choices: `table`: A formatted grid-like table `tab`: A tab delimited non-formatted version of table `list`: A string list of column names

Parameters

- **display_as** – {table,tab,list}
- **write** – If true, print output to console, otherwise return string.

Returns A formatted string or None

`grasp.info.get_phenotype_categories` (*list_only=False, dictionary=False, table=False*)

Return all phenotype categories from the PhenoCats table.

List_only Return a simple text list instead of a list of Phenotype objects.

Dictionary Return a dictionary of phenotype=>ID

Table Return a pretty table for printing.

`grasp.info.get_phenotypes` (*list_only=False, dictionary=False, table=False*)

Return all phenotypes from the Phenotype table.

List_only Return a simple text list instead of a list of Phenotype objects.

Dictionary Return a dictionary of phenotype=>ID

Table Return a pretty table for printing.

`grasp.info.get_population_flags` (*list_only=False, dictionary=False, table=False*)

Return all population flags available in the PopFlags class.

List_only Return a simple text list instead of a list of Phenotype objects.

Dictionary Return a dictionary of population=>ID

Table Return a pretty table for printing.

`grasp.info.get_populations` (*list_only=False, dictionary=False, table=False*)
Return all populatons from the Population table.

List_only Return a simple text list instead of a list of Phenotype objects.

Dictionary Return a dictionary of population=>ID

Table Return a pretty table for printing.

`grasp.info.get_snp_columns` (*return_as='list'*)
Return all columns in the SNP table.

Display choices: list: A python list of column names dictionary: A python dictionary of name=>desc long_dict:
A python dictionary of name=>(type, desc)

Parameters `return_as` – {table,tab,list,dictionary,long_dict,id_dict}

Returns A list or dictionary

`grasp.info.get_study_columns` (*return_as='list'*)
Return all columns in the SNP table.

Display choices: list: A python list of column names dictionary: A python dictionary of name=>desc long_dict:
A python dictionary of name=>(type, desc)

Parameters `return_as` – {table,tab,list,dictionary,long_dict,id_dict}

Returns A list or dictionary

3.6 grasp.ref

ref.py holds some simple lookups and the *PopFlags* classes that don't really go anywhere else. Holds reference objects for use elsewhere in the module.

class `grasp.ref.PopFlag`
Bases: `flags.Flags`

A simplified bitwise flag system for tracking populations.

Table Columns

The two important tables with the majority of the data are Study and SNP. In addition, phenotype data is stored in Phenotype and PhenoCats, population data is in Population, and platforms are in Platform.

Contents

- *Table Columns*
 - *Study*
 - *SNP*
 - *Phenotype*
 - *PhenoCats*
 - *Population*
 - *Platform*

4.1 Study

To query studies, it is recommended to use the `query.get_studies()` function.

Column	Description	Type
id	id	Integer
pmid	PubmedID	String
title	Study	String
journal	Journal	String
author	1st_author	String
grasp_ver	GRASPversion?	Integer
noresults	No results flag	Boolean
results	#results	Integer
qtl	IsEqtl/meQTL/pQTL/gQTL/Metabolmics?	Boolean
snps	Link to all SNPs in this study	relationship
phenotype_id	ID of primary phenotype in Phenotype table	Integer
phenotype	A link to the primary phenotype in the Phenotype table	relationship
phenotype_cats	A link to all phenotype categories assigned in the PhenoCats table	relationship
datepub	DatePub	Date

Continued on next page

Table 4.1 – continued from previous page

Column	Description	Type
in_nhgri	In NHGRI GWAS catalog (8/26/14)?	Boolean
locations	Specific place(s) mentioned for samples	String
mf	Includes male/female only analyses in discovery and/or replication?	Boolean
mf_only	Exclusively male or female study?	Boolean
platforms	Link to platforms in the Platform table. Platform [SNPs passing QC]	relationship
snp_count	From “Platform [SNPs passing QC]”	String
imputed	From “Platform [SNPs passing QC]”	Boolean
population_id	Primary key of population table	Integer
population	GWAS description, link to table	relationship
total	Total Discovery + Replication sample size	Integer
total_disc	Total discovery samples	Integer
disc_pop_flag	A bitwise flag that shows presence/absence of discovery populations	Integer
european	European	Integer
african	African ancestry	Integer
east_asian	East Asian	Integer
south_asian	Indian/South Asian	Integer
hispanic	Hispanic	Integer
native	Native	Integer
micronesian	Micronesian	Integer
arab	Arab/ME	Integer
mixed	Mixed	Integer
unpecified	Unspec	Integer
filipino	Filipino	Integer
indonesian	Indonesian	Integer
total_rep	Total replication samples	Integer
rep_pop_flag	A bitwise flag that shows presence/absence of replication populations	Integer
rep_european	European.1	Integer
rep_african	African ancestry.1	Integer
rep_east_asian	East Asian.1	Integer
rep_south_asian	Indian/South Asian.1	Integer
rep_hispanic	Hispanic.1	Integer
rep_native	Native.1	Integer
rep_micronesian	Micronesian.1	Integer
rep_arab	Arab/ME.1	Integer
rep_mixed	Mixed.1	Integer
rep_unpecified	Unspec.1	Integer
rep_filipino	Filipino.1	Integer
rep_indonesian	Indonesian.1	Integer
sample_size	Initial Sample Size, string description of integer population counts above.	String
replication_size	Replication Sample Size, string description of integer population counts above.	String

4.2 SNP

Column	Description	Type
id	NHLBIkey	BigInteger
snpid	SNPid	String
chrom	chr	String
Continued on next page		

Table 4.2 – continued from previous page

Column	Description	Type
pos	pos	Integer
pval	Pvalue	Float
NHLBIkey	NHLBIkey	String
HUPfield	HUPfield	String
LastCurationDate	LastCurationDate	Date
CreationDate	CreationDate	Date
population_id	Primary	Integer
population	Link	relationship
study_id	Primary	Integer
study	Link	relationship
study_snpid	SNPid	String
paper_loc	LocationWithinPaper	String
phenotype_desc	Phenotype	String
phenotype_cats	Link	relationship
InGene	InGene	String
NearestGene	NearestGene	String
InLincRNA	InLincRNA	String
InMiRNA	InMiRNA	String
InMiRNABS	InMiRNABS	String
dbSNPfxn	dbSNPfxn	String
dbSNPMAF	dbSNPMAF	String
dbSNPinfo	dbSNPalleles	String
dbSNPvalidation	dbSNPvalidation	String
dbSNPClinStatus	dbSNPClinStatus	String
ORegAnno	ORegAnno	String
ConservPredTFBS	ConservPredTFBS	String
HumanEnhancer	HumanEnhancer	String
RNAedit	RNAedit	String
PolyPhen2	PolyPhen2	String
SIFT	SIFT	String
LSSNP	LS	String
UniProt	UniProt	String
EqtlMethMetabStudy	EqtlMethMetabStudy	String

4.3 Phenotype

All available phenotypes are available on the [Phenotypes wiki page](#)

- id
- phenotype
- studies (link to Study table)
- snps (link to SNP table)

4.4 PhenoCats

All phenotype categories are available on the [Phenotype Categories wiki page](#)

- id
- population
- alias
- studies (link to Study table)
- snps (link to SNP table)

4.5 Population

- id
- population
- studies (link to Study table)
- snps (link to SNP table)

All population entries are available on the [Populations wiki page](#)

4.6 Platform

- id
- platform
- studies (link to Study table)
- snps (link to SNP table)

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