

Databases and ontologies

Cas-Database: web-based genome-wide guide RNA library design for gene knockout screens using CRISPR-Cas9

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Abstract

Motivation: CRISPR-derived RNA guided endonucleases (RGENs) have been widely used for both gene knockout and knock-in at the level of single or multiple genes. RGENs are now available for forward genetic screens at genome scale, but single guide RNA (sgRNA) selection at this scale is difficult.

Results: We develop an online tool, Cas-Database, a genome-wide gRNA library design tool for Cas9 nucleases from *Streptococcus pyogenes* (SpCas9). With an easy-to-use web interface, Cas-Database allows users to select optimal target sequences simply by changing the filtering conditions. Furthermore, it provides a powerful way to select multiple optimal target sequences from thousands of genes at once for the creation of a genome-wide library. Cas-Database also provides a web application programming interface (web API) for advanced bioinformatics users.

Availability and implementation: Free access at <http://www.rgenome.net/cas-database/>.

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Supplementary information: [Supplementary data](#) are available at *Bioinformatics* online.

1 Introduction

RNA guided endonucleases (RGENs) derived from the Type II clustered regularly interspaced short palindromic repeats (CRISPR)-Cas (CRISPR associated) system, an adaptive immune response in bacteria and archaea, have been usefully harnessed in many genome engineering applications such as gene knockout and knock-in in various organisms (Doudna and Charpentier, 2014; Kim and Kim, 2014; Sander and Joung, 2014; Shalem *et al.*, 2015). Recently, a few groups have undertaken genome-wide Cas9-mediated genetic screens (Chen *et al.*, 2015; Gilbert *et al.*, 2014; Hart *et al.*, 2015; Koike-Yusa *et al.*, 2014; Konermann *et al.*, 2014; Shalem *et al.*, 2014; Wang *et al.*, 2014; Zhou *et al.*, 2014) in human and other mammalian cells. The selection of target sequences is an initial, rate-limiting step in RGEN applications. We and other groups have developed a number of web-based online tools or command-line programs for single-guide RNA (sgRNA) design

or off-target site identification (Aach *et al.*, 2014; Bae *et al.*, 2014a; Cradick *et al.*, 2014; Doench *et al.*, 2014; Heigwer *et al.*, 2014; Hsu *et al.*, 2013; Lei *et al.*, 2014; Montague *et al.*, 2014; Naito *et al.*, 2015; O'Brien and Bailey, 2014; Park *et al.*, 2015; Sander *et al.*, 2010; Upadhyay and Sharma, 2014; Xiao *et al.*, 2014; Xie *et al.*, 2014). Although they are very useful for choosing sgRNAs that target one gene, selection of sgRNAs at the genome scale is still challenging.

Previously, two genome-wide databases for SpCas9 design (Hodgkins *et al.*, 2015; MacPherson and Scherf, 2015) have been built conceptually. However, although they are useful for selecting sgRNAs that target a small number of genes, neither offers an easy way to select optimal target sequences at once from thousands of genes for genome-wide library construction in a variety of organisms. Here we describe Cas-Database, a web-based genome-wide gRNA design tool for SpCas9 nucleases for genome-scale screening

experiments. Cas-Database contains all available targets of SpCas9 nucleases that recognizes 5'-NGG-3' protospacer-adjacent motif (PAM) sequences in all coding sequence (CDS) regions throughout the whole genome of a selected organism, based on the Ensembl database (Cunningham *et al.*, 2014). Each target site has the following information: GC content, relative cleavage position in the CDS, constitutive exon coverage, a microhomology-associated out-of-frame score (Bae *et al.*, 2014b), and potential off-target sequences with up to 2-nt mismatches. In addition, JBrowse (Skinner *et al.*, 2009) is used to display all available target sites graphically in a zoomable interface with genomic location information for the user's convenience.

Cas-Database basically provides a fast and easy way to select optimal target sequences in genes of interest from a variety of organisms. Additionally, it offers a powerful way to select optimal target sequences in many genes simultaneously. Selecting sgRNA sequences that target each gene is as easy as online shopping because of the use of a 'cart' system, similar to what is used in online shopping malls, which was implemented using cutting-edge web development techniques such as AJAX (asynchronous JavaScript and XML).

Currently, Cas-Database supports sgRNA design in 12 different organisms, including five vertebrates (human, rat, mouse, pig and zebrafish), one insect (*Drosophila melanogaster*), one nematode (*Caenorhabditis elegans*) and five plants (*Arabidopsis thaliana*, tomato, banana, grapes and soybean). All processes required to generate the genome-wide database have now been automated using a scripts pipeline so that we can easily update information about the existing organisms or add new organisms to the database. We are planning to continue to add and support more organisms in the future.

2 Methods

2.1 Target selection for SpCas9 nucleases

The latest whole genome sequence and associated annotation data from each organism in the Ensembl database were automatically retrieved and saved to our server using a homemade program written in Python language. To allow easy access to the annotation database, we used the Biomart protocol (Kasprzyk, 2011; Kasprzyk *et al.*, 2004) for communicating with the Ensembl server. After retrieving the genome sequence and annotation data associated with each organism, we first searched for all possible targets that contain 5'-NGG-3' PAM sequences in CDS regions and then calculated several characteristics of each target, e.g. sgRNA GC content, relative cleavage position in the CDS, common exon coverage throughout the various transcripts of the gene, and a microhomology-associated out-of-frame score.

2.2 Searching for potential off-target sites

After identifying all possible targets for SpCas9 nucleases in CDS regions and calculating the characteristics of each target, we next used the Cas-OFFinder program (Bae *et al.*, 2014a) to search for potential off-target sites that differ from each on-target site by up to 2nt and that contain 5'-NGG-3' or 5'-NAG-3' PAM sequences. Because this step is very time-consuming, we ran Cas-OFFinder on an OpenCL-enabled cluster computer such as Chundoong (<http://chundoong.snu.ac.kr/>) for all sites in parallel. During this process, we periodically validated that each searching node had finished correctly. After validation, all Cas-OFFinder output files were retrieved from the Chundoong server and moved to our local storage site. Then, for each selected target the total number of potential off-targets was counted; information about the genomic location of these potential off-targets,

such as whether they reside in CDS, UTR, intron or intergenic regions, was also evaluated.

2.3 Inserting information into the database

We rearranged all of the resulting data and constructed a SQL database that contains all possible SpCas9 nuclease targets in CDS regions and related characteristics of each target. We chose PostgreSQL (<http://www.postgresql.org/>) as the database server, which showed the best performance in our case.

2.4 Web interface

We built a web interface for Cas-Database using the Python web framework Django (<https://www.djangoproject.com/>). Because of its modern and easily-implemented database application program interface (database API), the algorithm for storing and retrieving data from the PostgreSQL database is simplified and web site maintenance is easy. For creating the HTML part of the interface, we used the web development framework Bootstrap (<http://getbootstrap.com/>) and the JavaScript framework JQuery (<https://jquery.com/>). We also implemented asynchronous data upload and download to achieve a fast response time for each user using the asynchronous JavaScript and XML (AJAX) technique. Because the results are retrieved from the server asynchronously, user requests will be run in parallel rather than in sequence. In conclusion, searching and filtering functions will be operated and displayed immediately after a user changes the input data or filtering condition.

3 Results and discussion

3.1 Cas-Database overview

Cas-Database provides a simple and easy way to design optimal sgRNAs. All available genes from a selected organism are listed on the main page of the Cas-Database website, as shown in Figure 1A. Users can easily search for desired genes by querying with the gene symbol, Ensembl ID or gene description. The search results are instantly displayed on the screen after every keystroke through the use of the AJAX web technique (Fig. 1B).

For a desired gene, users can use the 'Quick Info' function or the 'Add to collection' function. Basically, the 'Quick Info' function is useful for selecting sgRNAs manually for each gene; one can easily preview all available targets in a specific gene by clicking on the 'Quick info' button. Moreover, users can add additional desired genes—just as items are added to a shopping cart online—and select sgRNAs for many genes simultaneously using the 'Add to collection' function. In addition, hundreds or thousands of genes can be imported from text files that contain gene symbols or Ensembl ID lists, with individual entries separated by line breaks or spaces (Fig. 1C).

3.2 Use of the 'Quick Info' function

Cas-Database provides an easy way to preview sgRNAs that target a specific gene. Clicking on the 'Quick info' button immediately provides detailed information about the gene, transcripts and targets in a new dialog box (Fig. 2A–C). Furthermore, the genomic locus of each target is displayed on the graphical genome browser, JBrowse (Skinner *et al.*, 2009) (Fig. 2B). If one clicks on a target in the browser, the corresponding target (Fig. 2C) will be shown on the Cas-Database web page for added convenience.

Cas-Database also offers a powerful filtering function. Using the filtering feature at the top of the web page, as shown in Figure 2A, users can change the filtering conditions and rapidly preview the new results.

A Cas-Database

Optimal selection of guide RNAs at genome-scale.

Cas-Database is a genome-wide gRNA library design tool for Cas9 nucleases from *Streptococcus pyogenes* (SpCas9). It contains **all available targets** of SpCas9 nucleases that recognizes 5'-NGG-3' PAM sequences **in all coding sequence (CDS) regions**. Users can select optimal target sequences from thousands of genes at once simply by changing the filtering conditions (see [tutorial here](#)). In addition, Cas-Database also provides a [web application programming interface \(web API\)](#) for advanced use.

Organism Type: Vertebrates Organism: Homo sapiens (GRCh38/hg38) - Human

Category: All Search:

Total 20247 genes are listed.

Symbol	Gene ID	Description	Biotype	Action
NDRG2	ENSG00000165795	NDRG family member 2 [Source:HGNC Symbol;Acc:HGNC:14460]	Protein coding	Add to Collection Quick Info

B

Organism Type: Vertebrates Organism: Homo sapiens (GRCh38/hg38) - Human

Category: Symbol Search: C4BP

Total 2 genes are listed.

Symbol	Gene ID	Description	Biotype	Action
C4BPA	ENSG00000123838	complement component 4 binding protein, alpha [Source:HGNC Symbol;Acc:HGNC:1325]	Protein coding	Add to Collection Quick Info
C4BPB	ENSG00000123843	complement component 4 binding protein, beta [Source:HGNC Symbol;Acc:HGNC:1328]	Protein coding	Add to Collection Quick Info

[Import genes from file...](#)

[Select optimal sgRNAs](#)

C

Import genes from file...

You can upload a text file that contains gene symbols or Ensembl IDs separated by line breaks or spaces.

File contains: Gene symbols

Organism: Homo sapiens (GRCh38/hg3)

Text File: [Browse](#)

[Next](#)

Fig. 1. Cas-Database overview. (A) All available genes from selected organisms are listed with their Ensembl ID, description and biotype information on the main page of the Cas-Database web site. Users can preview gene information and manually select optimal sgRNAs for each gene using the 'Quick Info' function, or select sgRNAs for many genes automatically using the 'Add to Collection' function. (B) Cas-Database's top search panel provides a powerful searching function. Users can easily search for a desired gene by querying the gene symbol, Ensembl ID or gene description. The results are updated instantly on the screen via the AJAX web technique. (C) When working with hundreds or thousands of genes, users can upload text files that contain Ensembl IDs or gene symbols separated by line breaks or spaces

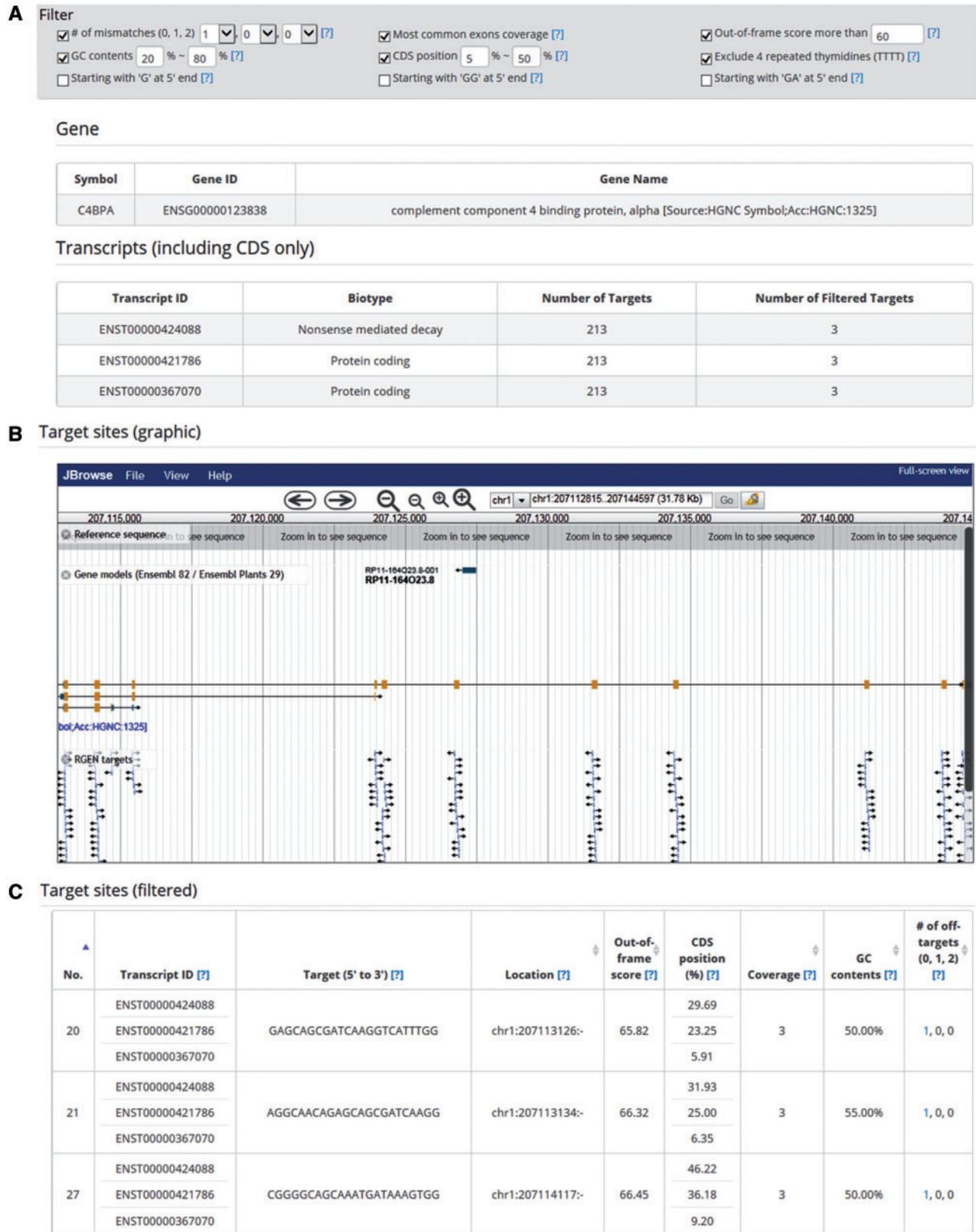


Fig. 2. The 'Quick Info' function of Cas-Database. Clicking on the 'Quick Info' button as described for Figure 1A results in the rapid display of useful information about the selected gene: (A) gene description, transcript variant information that includes the CDS region, (B) location of sgRNAs in the gene visualized by JBrowse and (C) all filtered sgRNAs with useful descriptions. Additionally, all target sites are listed below, with transcript ID, GC content, genomic location, relative position in the CDS, exon coverage throughout all transcript variants of the gene, a microhomology-associated out-of-frame score and the number of mismatched nucleotides. Users can alter filter conditions at the top of the dialog

A Total 20247 genes are listed.

Symbol	Gene ID	Description	Biotype	Action
NDRG2	ENSG00000165795	NDRG family member 2 [Source:HGNC Symbol;Acc:HGNC:14460]	Protein coding	Remove from Collection Quick Info
SMIM22	ENSG00000267795	small integral membrane protein 22 [Source:HGNC Symbol;Acc:HGNC:48329]	Protein coding	Add to Collection Quick Info
RSAD2	ENSG00000134321	radical S-adenosyl methionine domain containing 2 [Source:HGNC Symbol;Acc:HGNC:30908]	Protein coding	Remove from Collection Quick Info
HSPA12B	ENSG00000132622	heat shock 70kD protein 12B [Source:HGNC Symbol;Acc:HGNC:16193]	Protein coding	Add to Collection Quick Info

Collection (8 genes) [Import genes from file...](#) [Select optimal sgRNAs](#)

- ABCC1 (Homo sapiens (GRCh38/hg38) - Human) x
- RSAD2 (Homo sapiens (GRCh38/hg38) - Human) x
- NDRG2 (Homo sapiens (GRCh38/hg38) - Human) x

B Filter

Total count of sgRNAs for each gene:

of mismatches (0, 1, 2) [\[?\]](#)
 Most common exons coverage [\[?\]](#)
 Out-of-frame score more than [\[?\]](#)

GC contents % ~ % [\[?\]](#)
 CDS range % ~ % [\[?\]](#)
 Exclude 4 repeated thymidines (TTTT) [\[?\]](#)

Starting with 'G' at 5' end [\[?\]](#)
 Starting with 'GG' at 5' end [\[?\]](#)
 Starting with 'GA' at 5' end [\[?\]](#)

Card View Detail View

MARCH11 of Homo sapiens (GRCh38/hg38) - Human

Selected guide-RNAs (5' to 3')

TTTACAGAGTGTTAAGCGCTGG

N/A

N/A

CCDC125 of Homo sapiens (GRCh38/hg38) - Human

Selected guide-RNAs (5' to 3')

CCGAAGTCTTCTTACTCCTTGG

TGATAAAGTGGCTTTCAGAGG

N/A

DGKH of Homo sapiens (GRCh38/hg38) - Human

Selected guide-RNAs (5' to 3')

CTCTTCGACAGTTACAGAAGG

TACCTTCGACAGGACAGGCATGG

AGGTGCTACAGTACCTTCGCAGG

ESPN of Homo sapiens (GRCh38/hg38) - Human

Selected guide-RNAs (5' to 3')

N/A

N/A

N/A

[Remove genes with green indicators](#) [Download filtered sequences...](#)

Fig. 3. The 'Quick Info' function of Cas-Database. The 'Add to Collection' function of Cas-Database. (A) Cas-Database provides a unique and easy way to select many genes at once by the implementation of a 'cart' system. Users can collect desired genes by clicking on the 'Add to Collection' button or by uploading a text file as described in the Figure 1C legend. Note that one can also select genes from different organisms. By clicking on the 'Select optimal gRNAs' button, users will proceed to the next step. (B) The results page will list all available sgRNAs filtered by the default conditions on the top panel. A colored indicator represents the selection status for each gene, e.g. green (selected completely), yellow (selected partially) or red (not selected at all). A user can download only the genes for which sgRNAs were successfully designed, and repeat the process for the remaining genes by changing the filter criteria and clicking on the 'Remove "green" genes from list' button

Filtering criteria include GC content, relative position in the CDS, exon coverage throughout all transcript variants of the gene, a microhomology-associated out-of-frame score (Bae *et al.*, 2014b), excluding four thymidine nucleotides in tandem (Braglia *et al.*, 2005) and the number of mismatched nucleotides.

3.3 Use of the 'Add to Collection' function

Cas-Database also provides a novel function that allows the selection of sgRNAs from hundreds or thousands of genes at once

through the use of a 'cart' system. Users can either collect desired genes on the main web page by clicking on the 'Add to Collection' button (Fig. 3A) or by uploading a text file as discussed above and shown in Figure 1C. After all desired genes are collected, clicking on the 'Select optimal sgRNAs' button will open the results page, which will list all available sgRNAs filtered by the default conditions as shown in Figure 3B. Users can easily change the filtering conditions, including the total count of sgRNAs for each gene, in the filter section that appears at the top of the resulting page (Fig. 3B). Because

the AJAX technique is used, the retrieving processes run independently of each other, resulting in fast loading speeds; e.g. the loading time for 1000 genes is about 2 min in the default conditions.

Whether enough sgRNAs have been selected for each gene after filtering is indicated by colored indicators—green (selected completely), yellow (selected partially) or red (not selected at all), as shown in Figure 3B. In this step, users can download either sequences for all selected sgRNAs targeting each gene or for some sgRNAs selected completely from each gene with a green indicator. In other words, users can download sequences only for genes for which sgRNAs were selected completely under the initial filtering conditions. After that, users can eliminate genes with green indicators from the list by clicking the ‘Remove green genes from list’ button, and then can alter the filtering conditions again. After resetting the filtering conditions, users can download the genes with green indicators again and repeat this process until sgRNAs have been selected completely in the most remained genes. Finally, if a few genes are still left, users can manually select targets for those genes using the ‘Quick info’ function on the main page. As a result, users can select optimal sgRNAs from hundreds or thousands of genes and download a list of sgRNAs for every gene with the target specific information.

3.4 Web API

For users familiar with bioinformatics, Cas-Database also provides a web application programming interface (web API). If users send queries through hypertext transfer protocol (HTTP) requests to our database server, the results will be returned in the JavaScript oriented notation (JSON) format. Thus, researchers can easily create their own simple homemade scripts for automated data retrieval. Details about the web interface are described in the [Supplementary data](#).

3.5 Update of Cas-Database

The entire process of database creation, from retrieving an organism’s genome sequence and its associated annotation data from the ENSEMBL server to creating a new database, is totally automated by our homemade scripts. The time to build a new database depends on the organism’s genome size, e.g. the creation of a new database for zebrafish (1.32 GB) took about three days.

Currently, Cas-Database supports 12 different model organisms, including five vertebrates (human, rat, mouse, pig and zebrafish), one insect (*D.melanogaster*), one nematode (*C.elegans*) and five plants (*A.thaliana*, tomato, banana, grapes and soybean), and we are planning to update Cas-Database continuously to obtain the most recent versions of genomic sequences for the existing organisms. We also plan to add data from other organisms in the ENSEMBL database. In addition, we will update the database to allow for alternative CRISPR/Cas nucleases such as Cprf1 (Zetsche et al., 2015).

4 Conclusion

Cas-Database is an easy-to-use web-based tool for designing sgRNAs for SpCas9 nucleases on a genome scale. It can be applied to construct optimal sgRNA libraries that target thousands of coding sequences in 12 different organisms for genome-wide knockout screening experiments. Because Cas-Database contains all available targets in CDS regions as well as target-related information, including data about potential off-target sites, users can easily access the data through the interactive web interface or web API. The web interface was made using cutting-edge web development techniques

such as AJAX, so the website is highly responsive to user input and the output results load quickly.

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