

Professional Edition

Cloning, graphic maps, primer design, sequence alignment -- Version 9 for Windows

Clone Manager Professional provides a comprehensive, fully integrated set of tools for enzyme operations, cloning simulation, graphic map drawing, primer design and analysis, global and local sequence alignments, similarity searches, and laboratory-sized sequence assembly projects.

You can also use Clone Manager as a quick and easy way to view or edit sequence files, find open reading frames, translate genes, find genes or text in files, or organize primer collections.

Work with Enzymes

- Find enzyme sites, view enzyme information
- Do restriction enzyme analysis
- Create user enzyme lists, transient lists

Clone Manager scans the sequence and identifies enzyme recognition sites, drawing an accurate on-screen working map. Click tabs at the bottom of the molecule viewer window to see the map, restriction map data, sequence, features, or molecule information. Zoom to transiently view just a portion of a larger molecule.



Scrolling map sites list shows enzyme sites, primer sites or features. Enzyme site information includes name, position, type of ends produced and if a single cutter (sc).

A fast and powerful restriction enzyme analysis module has five display styles, including a list of No-Cutters. Filter to see only the results you need. View enzyme properties, suppliers, isoschizomers, or compatible ends information. Update master enzyme lists monthly using REBASE files.

Simulate Cloning

- Cut, ligate DNA, modify molecule ends
- Use Ligate window or Plan Cloning Wizard
- New Gateway® and Topo Cloning Wizards
- New Assembly Cloning Wizard

Clone Manager can cut DNA at enzyme cut sites or at user-specified basepair positions. Molecule features and enzyme sites are automatically recalculated after a cut operation. The program can ligate DNA, checking for compatible ends. You can modify molecule ends, if needed for ligation, and select the cloning orientation.

For fast simulated cloning, you can cut, ligate, and modify ends -- all from the same Ligate display window showing sticky ends and molecule icon maps.

The Plan Cloning Wizard lets you describe your cloning objectives, review the list of possible solutions, and instruct the Wizard to construct the recombinant of choice.

The Gateway or Topoisomerase Cloning Wizards help you to simulate cloning using Gateway® Recombination Cloning or Topo® Cloning techniques. The wizards will help you select the appropriate components, show you the proposed result, and create the final recombinants.

The new Assembly Cloning wizard simulates cloning using Gibson Assembly[™] or related cloning techniques. You can assemble a set of overlapping fragments, previously prepared, or use PCR to create amplified fragments with overlaps, then assemble the fragments.

Draw Graphic Maps

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- Prepare high-resolution graphic maps
- Print, copy to clipboard, export to file
- Enhanced Views of graphic map

Prepare high-resolution circular or linear graphic maps. Control styles and colors for feature types and enzyme sites; control map size and position. For your lab notebook, select to print the map with associated text (description and notes; map features and enzyme sites).

For a publication or presentation, use an Enhanced View of the graphic map to customize a special molecule map. Set individual feature styles, colors, patterns, positions. Drag feature names to reposition, if needed. Remove enzyme sites or features from a map without modifying the original molecule data. Save enhanced views in the molecule file (maximum 6 views/file). Print, copy, export in vector or raster (Tif, Png, Gif, bitmap) formats.

Design Primers

• Search for primers and list results

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- Direct enter (type in) primers
- Position one primer in molecule sequence
- New Sequencing Primer Wizard

Search for PCR primers or pairs, sequencing primers, or probes. Sort results lists by rank (quality) or position, or view summary of search progress showing which criteria eliminated primers.

Use the Direct Entry wizard to enter sequence and other information for primers already in use or purchased commercially. Use the new Sequencing Primer Wizard to design a set of primers for sequencing a molecule region.

Position a primer in the molecule sequence at a target position or just select (highlight) sequence in the molecule viewer to initiate primer design at this position. In the design view, you can see the primer evaluation, the sequence flanking the primer, genes that are nearby, and the location of neighboring primers that will meet the evaluation criteria you have set.



Analyze Primers

- Evaluate using primer criteria
- Do in-depth analyses
- Use the Analyze Mix Wizard

To find or evaluate primers, the program analyzes GC content, melting temperature (Tm), dimers, stability, runs of bases and repeats, hairpins and false priming. Detailed analysis options display primer dimers, summarize cautions, and report homologies between the primer and template (false priming or other homologies).

The Analyze Mix Wizard will analyze primer binding and products formed in a mix of primers and molecules. Just identify the primers and molecules, set the parameters for the analysis, and view results. The Primer Viewer Window puts all primer information in one convenient location. Click tabs at the bottom of the window to see primer information, edit the primer, or analyze the primer. You can save the primer, including description and notes information, to a disk file or to a primer collection (see below).

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Primer Operations

- Find primers that bind to your molecule
- Find molecules with sites for your primer
- Primer Values, Mutagenesis Profiles
- Internet BLAST primer searches

Scan the primers you have created to find which could bind effectively to a molecule of interest. Primer sites can be added to molecule files and displayed on the molecule map or formatted sequence. Or scan molecule files on your computer or local network to look for molecules that have binding sites for a primer of interest.

Use the Primer Values profile to analyze regions of the molecule to assist in identifying areas for primers in difficult situations. Calculate Tm, GC and stability for primers of a length specified and plot these profile values. Use the Mutagenesis Profile to help construct mutant primers, identify restriction enzyme sites that can be introduced, or to check for changes in translation.

Use the Internet BLAST Primer Search to set up a remote search to check primer specificity using the NCBI BLAST server (equivalent to using the BLAST program Search for Short, Nearly Exact Matches to find primer binding sites).

Organize, Use Primers

- Store primers in Primer Collections
- Easy-to-use Create Product module
- Use primer sites on molecule map

Store essential information about many primers in one or more collection files for streamlined storage and access. Create new collection files, as needed, and create folders to organize your collection files. Search in collection files to find a specific primer. Export collection file data in tabdelimited format for use in other applications.

Use the Create Product module to identify two primers and one molecule to produce an amplified product with a PCRtype reaction. Create a new DNA molecule or export the product sequence. Review product size, GC content, melting temperature, annealing temperature.

Select a primer site on your molecule map and export the primer sequence, rename the primer, create the primer from its sequence, or (with a second primer on this map) quickly create an amplified product.

Align Multiple Sequences

- Align many sequences against a reference
- Align many sequences against each other
- Use file names or molecule names
- Save, export, edit alignment data

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Align a group of DNA or protein sequences against a reference sequence. View a summary table of results, aligned sequences in a variety of formats, or a graphic display of areas of significant similarity in each sequence. Enter name, description. Save alignment results.

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A multi-way alignment function lets you do exhaustive pairwise alignments of all sequences and progressive assembly of alignment using Neighbor-Joining phylogeny. Alignment results include a dendrogram which suggests the pattern of relatedness of all of the sequences aligned.

Edit alignment data — replace a base if you believe it is in error, or shift unaligned bases left or right. If the sequence files contain trace data, view the trace for a base in question before editing. Export the consensus sequence derived from an alignment or export the aligned sequences in CLUSTAL W multiple sequence alignment format.

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Score sort 🗸 🔅 🖏 🔛 60% 🖌 🗐 = 💷 🖉 🕄 🧭 🕅					
Multi-way DNA alignment. Sequences: 5. Scoring matrix: Linear (Mismatch 2, OpenGap 4, ExtGap 1)					
Sequence View: Similarity Format, Color areas of high matches at same base position					
CVGEM5ZFM	88	gagageteecaacgegttggatgeatagettgagtattetatagt			
CVGEM7ZFM	85	gagageteecaacgegttggatgeatagettgagtattetatagt			
CVGEM3ZFM	32	tctagagtcgacctgcaggcatgcaagcttgagtattctatagt			
CVGEM11ZM	54	g-gccctctagatgcggcc-gcatgcataagcttgagtattctatagt			
CVGEM9ZFM	47	-cgagctccctatagt			
CVGEM5ZFM	133	gtcacctaaatagcttggcgtaatcatggtcatagc			
CVGEM7ZFM	130	gtcacctaaatagcttggcgtaatcatggtcatagc			
CVGEM3ZFM	76	gtcacctaaatagcttggcgtaatcatggtcatagc			
CVGEM11ZM	100	gtcacctaaatagcttggcgtaatcatggtcatagc			
CVGEM9ZFM	62	gagtcgta-ttagaggccgacttggccaaattcgtaatcatggtcatagc			
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Info Picture Seque	ence LMap				

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Compare Two Sequences

- Find regions of local similarity
- Progressively assemble an alignment

Search two DNA or protein molecules for regions of local similarity using the FastScan method, maximized for score or quality, or the Needleman-Wunsch method.

Results of local alignments include a sortable list or graphic view of similarities found, and aligned sequences or a detailed map for each homology block listed. Move easily to Next or Previous result in sequence/map views.

Assembled alignments begin with a FastScan local similarity search, lock in significant regions of homology found, and then complete the alignment of the remaining sequence. Export merged/spliced consensus sequence.

Scan for Similarities

- Scan sequence files in folders on your computer
- Scan databank library files on your computer

Search for similarities to a sequence by scanning folders of sequence files or databank library files on your personal computer or local network. The program will open each file, scan the sequences found, and record significant regions of similarity encountered. Set the speed and cutoff value to alter the sensitivity of the search. Save and recall lists of frequently-used search locations.

Results include a sortable list or graphic view of regions of similarity found, and aligned sequences or a detailed map for each homology block listed. From the results list, use the Scan Results Wizard to identify molecules to load for use now, save to a disk file for later use, or set up as an assembled multiple sequence alignment.

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Sequence Assembly

- Simple or complex assembly projects
- Locate subclones on reference sequence

Handles laboratory-sized assembly projects. Just identify sequence files to assemble and set assembly parameters, if applicable. Sequence assembly results are shown in a data window with multiple views -- review a table of results information, a picture view of alignment locations, or the aligned sequences with optional accessory data in a split pane. Accessory data includes an assembly overview with a position locator, a histogram of coverage and errors, and trace data, if available in the sequence files.

Save results and later re-open to continue to review or export results data. Export consensus sequences.

Public Domain Database Searches

- Internet BLAST, protein pattern database searches
- Built-in web browser

Initiate remote searches of public domain databases, including NCBI BLAST databases, and display results in a built-in web browser.

The program helps you set up a BLAST search by handling query sequence formatting, in-frame translation, or excision of a subsequence. Pick the BLAST program and database to use and your request is sent to the NCBI server over the internet. View BLAST results, access molecules of interest and save molecule files to disk or load a molecule for immediate use.

You need an internet connection to use these functions.

Other Features

- Integrated sequence editor and file handler
- Extensive sequence formatting options

• New sequence phrase collections

Clone Manager reads files in GenBank, EMBL, FASTA or Clone Manager file formats, or sequence data in simple ASCII text files. You can also import (paste) files from the clipboard or retrieve files from Entrez (NCBI).

The sequence editor can display full or partial sequences in a variety of formats. Display any combination of sequence, enzyme sites, sequence labels, translation, annotations and memorize your favorite style settings.

You can also edit sequences, join sequences, rotate circular molecules, invert or translate sequences, find Open Reading Frames, or find regions in the sequence that match or nearly match a search string. A new module stores sequence phrases in collection files and lets you find matching sites in your molecules.

DataBook is a simple, built-in database to store and organize information about your molecules. Automatic data entry makes it really easy to use.

Analysis Functions

- Open Reading Frame analysis
- DNA Repeats, Dyad Symmetries analyses

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- Hydrophilicity, Protein structure analyses
- Mutagenesis Profile module

Open Reading Frame analysis maps start and stop codons in 6 reading frames and calculates Fickett's TESTCODE scores to help identify coding regions.

Analyze direct or inverted repeats and dyad symmetries to identify regions that might be associated with biologically active sites such as control regions, transcription terminator regions, or RNA processing sites.

Analyze protein or translated DNA for hydrophilicity and surface exposure. Analyze protein structure for alphahelix, beta-sheet (extended), beta-turn, random coil states.

Mutagenesis Profile shows a portion of molecule sequence with one frame of translation and enzyme sites. View possible silent changes, find 'almost' enzyme sites, or view base changes required to change from one amino acid to another. Introduce a mutation and create a new molecule.

System Requirements: Recommended configuration: Windows 10, Windows 8 or Windows 7. Minimum configuration: Windows Vista. 20 Mb hard disk space required for program installation. Concurrent user license installation supports most standard Windows network systems or TCP/IP networks. Upgrades from version 8 available. Call or email for more information.

License Type	Price	Catalog No
Clone Manager Professional 9, Individual User License	\$1150	IL 8901
Clone Manager Professional 9, 4-Seat WorkGroup License	\$1950	WG 8908-4
Clone Manager Professional 9, 8-Seat WorkGroup License	\$3300	WG 8908-8
Clone Manager Professional 9, Concurrent 5-User License	\$3800	CL 8905-5

License Information

Individual User License

For use by one individual or installation on a single computer. Receive program CD, 1 Getting Started manual and 1 Quick Reference guide.

WorkGroup (Seat) License

Install on the licensed number of individual computers (seats) for use by members of the workgroup. Receive program CD, 1 or 2 Getting Started manuals, 4 or 8 Quick Reference guides.

Concurrent 5-User License

Install on a single server for a multi-user computer network. Permits 5 users to concurrently use the program. Receive program CD, 2 Getting Started manuals, 10 Quick Reference guides. (Contact us for larger license information.)

For Research Use Only - Not for use in diagnostic procedures.

Ordering Information

Program disks and manuals will be shipped within 5 days of receipt of order. For standard shipping the following shipping and handling charges will be added to your total order. Expedited shipping can be arranged.

US (continental) - Fed Ex economy service	\$15
International - Individual User licenses	\$35
International - Multi-user licenses	\$65
Light Pack - call for details	

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November 2015. Prices subject to change without notice.