

## LASERGENE VERSION COMPARISON

| Lasergene Molecular Biology  | 15 | 16 | 17 |
|--|----|----|----|
| Sanger Sequence Assembly and Analysis  |    |    |    |
| New 64-bit SeqMan Ultra application with better performance and increased capacity                 |    |    | •  |
| Improved algorithms for increased accuracy of Sanger assemblies                                    |    |    | •  |
| Access to data from dbNSFP, 1000 Genomes, and ESP's Exome Variant Server for variant analysis      |    |    | •  |
| New hybrid reference-guided/de novo assembly workflow  |    |    | •  |
| Assess read alignment, coverage, and SNPs  | •  | •  | •  |
| Assemble reads <i>de novo</i> or against one or more reference sequences                           | •  | •  | •  |
| Cloning, Primer Design, and Gene Detection   |    |    |    |
| Enhanced support for VectorNTI, Geneious, and Clone Manager Suite file types                       |    | •  | •  |
| PCR site-directed mutagenesis with ability to predict impact of mutation on protein structure      |    | •  | •  |
| Access SeqBuilder Pro as a standalone application  |    | •  | •  |
| Agarose gel simulation in SeqBuilder Pro   | •  | •  | •  |
| Automated clone verification workflow  | •  | •  | •  |
| Batch sequence editing built-in to SeqBuilder Pro  | •  | •  | •  |
| New SeqBuilder Pro application for sequence editing and analysis                                   | •  | •  | •  |
| Quick search and selection of enzymes  | •  | •  | •  |
| Collapsible and sortable feature and primer lists  | •  | •  | •  |
| Plasmid auto-annotation (single or batch) using curated database                                   | •  | •  | •  |
| Automated virtual cloning: site-directed, TA, TOPO, Gateway, InFusion, GeneArt and Gibson Assembly | •  | •  | •  |
| Design customized primer pairs   | •  | •  | •  |
| Pairwise and Multiple Sequence Alignment   |    |    |    |
| Perform BLAST and Entrez searching from MegAlign Pro alignments                                    |    | •  | •  |
| Updated views and enhanced user controls in MegAlign Pro   |    | •  | •  |
| Export editable images to PowerPoint   | •  | •  | •  |
| Profile alignment functionality to merge alignments or additional sequences                        | •  | •  | •  |
| MAFFT, MUSCLE, Mauve, and Clustal Omega alignment algorithms                                       | •  | •  | •  |
| Align sequences and create phylogenetic trees  | •  | •  | •  |

Add our Genomics & Protein Applications to Complete Your Lasergene Package! See reverse side for details.

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| Lasergene Genomics  | 15 | 16 | 17 |
|---|----|----|----|
| Next-Gen Sequence Assembly and Alignment  |    |    |    |
| 64-bit SeqMan Ultra application for project analysis with better performance and increased capacity |    |    | •  |
| Redesigned SeqMan NGen with better guidance for project setup                                       |    |    | •  |
| Auto-analysis of hardware and data to determine if assembly should be run locally or on the Cloud   |    |    | •  |
| Ability to polish Canu or Spades assemblies of PacBio and Oxford Nanore long read data              |    |    | •  |
| miRNA quantitation workflow to quantitate and analyze miRNA gene expression levels                  |    |    | •  |
| Integrated DNASTAR Cloud Assemblies for projects that exceed the capacity of your hardware          |    | •  | •  |
| Quick access to NGS and Sanger project setup in the new DNASTAR Navigator                           |    | •  | •  |
| Enhanced RNA-Seq statistics with DESeq and edgeR from Bioconductor                                  | •  | •  | •  |
| Visualization, browsing, isoform analysis and multi-sample comparison in GenVision Pro              | •  | •  | •  |
| RNA-Seq analysis for model and non-model organisms  | •  | •  | •  |
| Automatic mRNA annotation using RefSeq  | •  | •  | •  |
| Gene panel workflows with control validation  | •  | •  | •  |
| Reference-guided assembly for any size genome on a desktop computer                                 | •  | •  | •  |
| De novo genome and transcriptome assembly   | •  | •  | •  |
| Variant Detection and Analysis  |    |    |    |
| Ability to compare and analyze multiple VCF files from other NGS software pipelines                 |    |    | •  |
| Model detected variants on protein structure with updated protein design workflow*                  |    | •  | •  |
| Combine variant and PDB annotations to predict potential effects on protein structure*              | •  | •  | •  |
| Combine SNP and CNV analysis with any workflow  | •  | •  | •  |
| Access to allele and genotype frequencies for SNPs  | •  | •  | •  |
| SNP detection accuracy >99.8%   | •  | •  | •  |
| Direct comparison to dbSNP and GERP and dbNSFP databases  | •  | •  | •  |
| Lasergene Protein   | 15 | 16 | 17 |
| Macromolecular Structure, Motion, and Function  |    |    |    |
| Automated hot spot scanning to locate residues that are important for protein fold stability*       |    | •  | •  |
| Protein design tools to improve fold stability and developability*                                  |    | •  | •  |
| Integrated BLAST and Entrez searching in Protean 3D   |    | •  | •  |
| Mutation modeling and neighbor search to analyze SNP impact on protein structure                    | •  | •  | •  |
| Protein composition analysis  | •  | •  | •  |
| Export editable images to PowerPoint and PDF  | •  | •  | •  |
| Access to NovaDock® protein-protein docking, analysis and visualization*                            | •  | •  | •  |
| Access to NovaFold Antibody structure prediction*   | •  | •  | •  |
| Protein function and ligand binding site prediction*  | •  | •  | •  |
| B-cell epitope prediction   | •  | •  | •  |
| * Additional subscription may be required to acess protein modeling and design programs             |    |    |    |

## DNASTAR