SCC's Genetics Information Systems Suite™

SoftMolecular®

SoftMolecular—SCC's intuitive workflow solution for the diagnostic molecular genetics laboratory—enables geneticists and pathologists to create customized protocols for managing testing workflows, derive data and images from automated instruments by interfaces, auto-populate report text based on result-driven reporting rules, and easily access patient and family medical and testing histories.

Spotlight Feature:
This innovative software supports both clinical and research applications of Next Generation Sequencing (NGS) whether it involves Targeted Panels, Whole Exome Sequencing (WES), or Whole Genome Sequencing (WGS).

SoftMolecular empowers pathologists, geneticists, and forensic scientists to automate workflow and manage data and analysis stemming from current and future molecular testing methods in order to generate accurate interpretations and meaningful reports with minimized effort. This advanced genetics LIS platform supports both clinical and research applications of Next Generation Sequencing (NGS) including Targeted Panels, Whole Exome Sequencing (WES), or Whole Genome Sequencing (WGS) as well as other common molecular technologies. Robust and user-friendly, SoftMolecular enables users to design and streamline workflows, interface with automated instrumentation, manage patient and family cases, track sample storage, and control inventory. As part of SCC's Genetics Information Systems Suite™, SoftMolecular supports interfacing and communication with other laboratories and health agencies, and represents the most comprehensive suite of genetics LIS applications in the world.

FEATURES AND BENEFITS

Feature: Variant/Mutation Table and Gene Master Table
Benefit: The functionality supports generation of the laboratory's variant data base, categorization of variants, versioning, and linkage to relevant databases (OMIM, NCBI, etc.) for ease of variant entry, management, and interpretation. The Gene Master table allows users to capture information from other Internet databases for variant interpretation on a single gene level.

Feature: NGS reporting provides easy identification of the reportable variants into distinct category sections (reflecting states of pathogenicity, whether the associated disorder is medically actionable, and whether the finding is related to the patient's phenotype). Hyperlinks can be included for pertinent websites associated for each of the variants. Web-based reporting is available as well as options for various levels of reports.
Benefit: Users will be able to provide their clients clear and concise reports representing only the results opted for and with easy access to pertinent Internet information.

Feature: Seamless interfacing with sequencing and fragment analysis instruments
Benefit: Replacing manual data result entry with instrument interfaces eliminates the resource investment and inherent error rate associated with manual entry, thus providing greater efficiency and accuracy.

Feature: Images on reports
Benefit: Images and graphics are available on patient reports providing professional and sophisticated reports for users' clients and patients.

Feature: Ability to choose a re-run of an entire batch or auto-assign a specific sample to the next pending batch with a single mouse-click
Benefit: This feature enables users to manage repeat testing quickly and easily to avoid any delays or mismanagement that can occur with manual processes. This ensures the most optimal turnaround time thus enhancing client satisfaction.

Feature: Ability to link controls to a sample or batch
Benefit: Control documentation will be accurate, can be accessible per patient or batch, and will remain electronically linked for historical reference or investigation. Users will no longer need to sort through manual logs or files to identify or trend control data for quality control troubleshooting.

FEATURES AND BENEFITS CON'T

Feature: Auto-calculation of master mixes with auto-deduction of volume from inventory

Benefit: Reduction of manual calculations and inventory control leading to error-free calculations and effective inventory management

---

Feature: Rules-driven alerts or notifications based on client, patient demographic, test, or result parameters

Benefit: Reduce paper or telephone communications ensuring timely and accurate delivery of special instructions or critical information. Reduce risk of errors and inefficiencies that can be associated with frequent personnel interruptions without interrupting personnel at inopportune times.

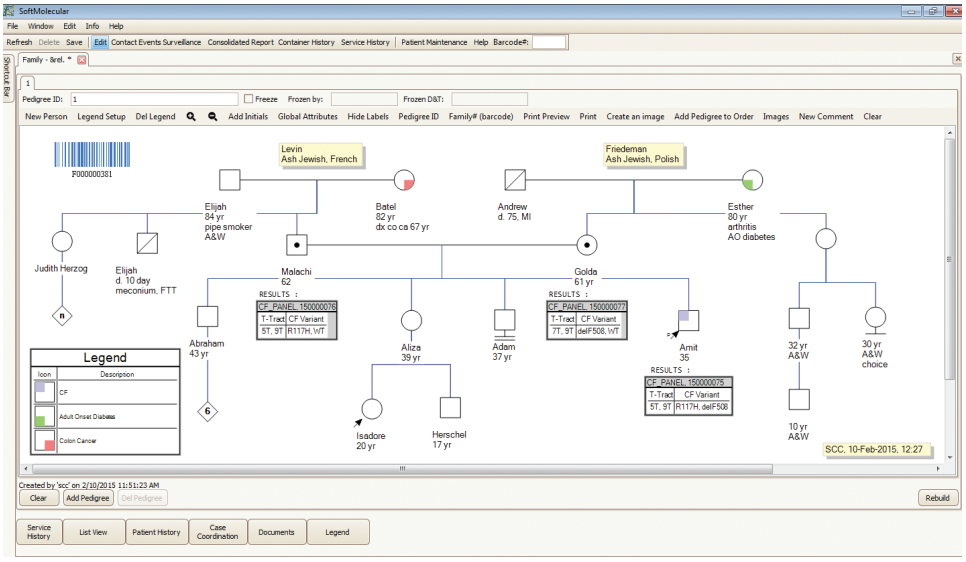
Feature: Auto reflex and Auto rerun testing across technologies based on client-defined rules

Benefit: Eliminates manual error inherent in manual reflex/rerun ordering and reduces turnaround time of tests.

SAMPLE SCREENSHOTS

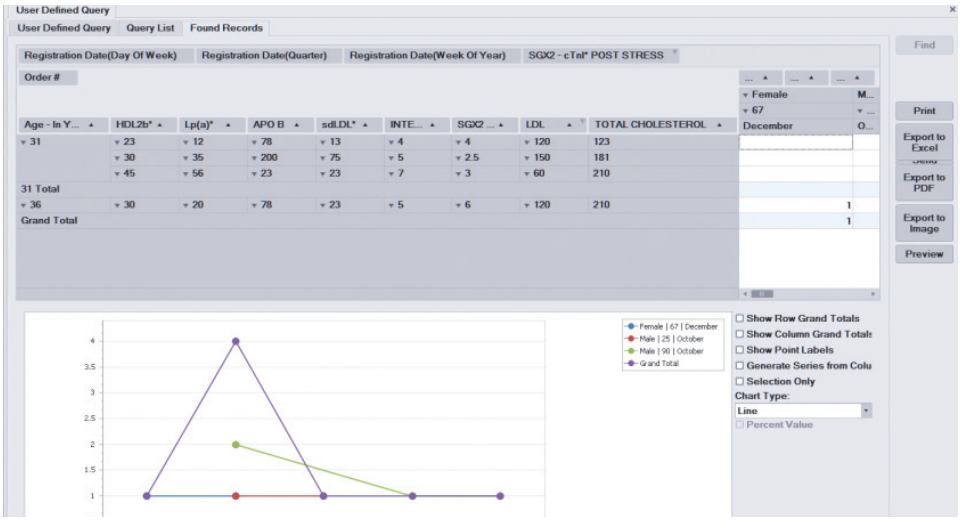
Family Pedigrees

SCC's Genetics Information System Suite™ Pedigree functionality allows linking of family members based on their relationships. User-defined attributes such as carrier or affected status, mutations, and phenotypic traits can be captured and displayed within the pedigree.



Trending Graphs

SoftMolecular offers an extensive library of trending chart templates that can be used to create customized graphs. Users can select the attributes/fields to be included, the number of recent values, and the scale of the graph. SoftMolecular's flexibility enables users to combine different result values on one graph for comparison purposes or have multiple graphs for a single test.



SCC's Genetics Information Systems Suite™
SoftMolecular® • SoftCytogenetics® • SoftPathDx®
SoftFlowCytometry® • SoftBiochemistry®
SoftHLA® • SoftGenePortal™ • SoftIWS®

SCC Soft Computer
5400 Tech Data Drive
Clearwater, Florida 33760

www.softcomputer.com
sales@softcomputer.com