



Savante is a total solution for nonclinical data query, consolidation, and reporting. Integrated with the industry-standard Pristima[®] XD Core preclinical data suite, it allows organizations to capture and aggregate study data based on CDISC-SEND standards. Flexible data analysis and visualization using TIBCO's Spotfire platform enables self-service of the data by multiple stakeholders.

Savante™

Savante is a seamless solution for nonclinical data query consolidation analytics and reporting. Assemble data sets from multiple sources and format data for CDISC-SEND output and reports. And move beyond basic reporting.

Integration with TIBCO Spotfire enables flexible just-in-time data analysis and visualization.

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Savante is designed to support the workflows of study directors, toxicologists, pathologists, and data scientists, enabling consistent views of study data, regardless of the source. Whether coming from internal sources or external partners, you can be sure the data will be the right format, readily accessible in a variety of formats.

Key Benefits

Savante automates the preclinical data management workflow by allowing you to:

- **Format and consolidate study data based on CDISC-SEND standards.** Take data in from a variety of formats and consolidate it, assembling it into a SEND-compliant format.
- **Aggregate data from multiple sources.** Savante takes in data from a variety of formats and consolidates it, assembling it into a SEND compliant format. The Define.xml file that is required as part of the submission is created automatically.
- **Automate data collection.** Synchronization from Pristima[®] XD Core makes data collection from instruments seamless. Savante can accept XPT format from CROs or other systems, and this data can be augmented with data in CSV or Excel data files.

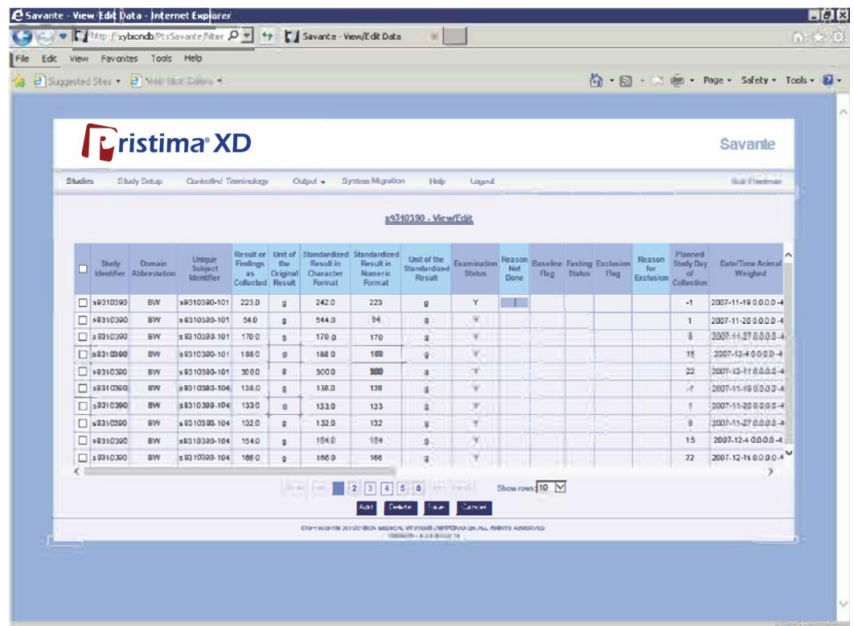


Figure 1. Import multiple data sets into a study and then view/edit each domain as needed, ensuring GLP data integrity.



- Perform flexible data analysis and visualization. Savante offers an integrated solution with TIBCO's Spotfire for making non clinical study data from in-house studies or partners accessible for cross study analysis

Xybion-Your Expert Partner

- Xybion has over 20 years of experience managing laboratory workflows and helping clients streamline compliance and mitigate risk.
- Xybion has extensive global experience and works with companies of all sizes from multi-ple industries.
- Xybion offers deep domain knowledge of FDA regulations and has a long track record of helping clients build effective risk mitigation strategies.

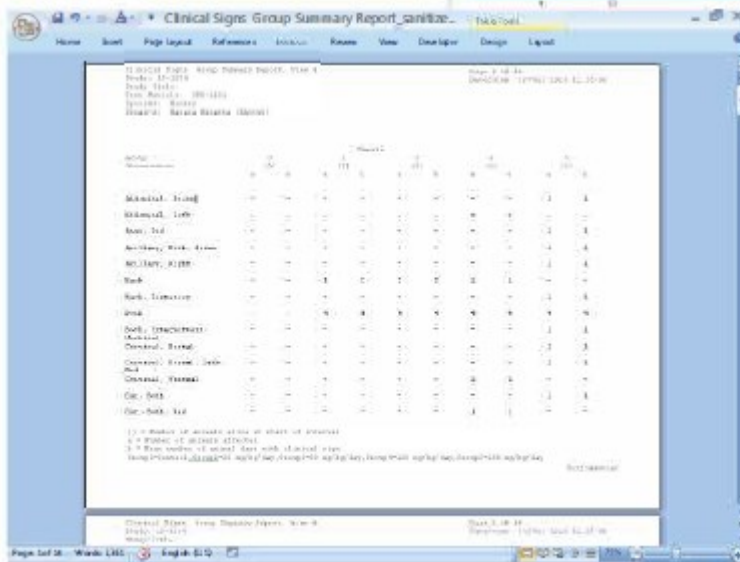
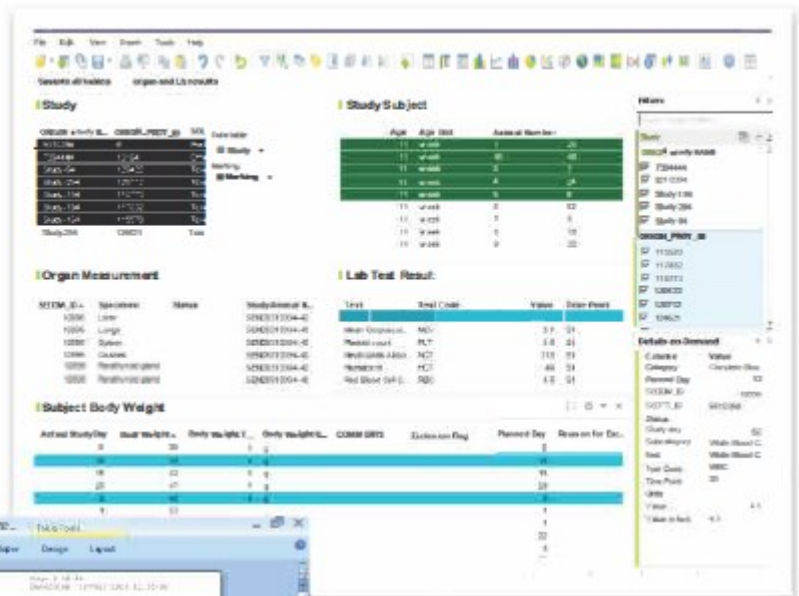


Figure 3. Dashboards can be created to summarize the data . View selections from studies through study groups, study animals and drill down into specific lab

Figure 4. Create reports in Word and PDF formats. These reports, taken from the same data as the SEND submission ,can be part of your submission package or used for internal verification.

