



How well do you know your IWR/IVRS partner?

Six questions to ask your IWR/IVRS provider.

- 1 Does your IWR/IVRS vendor make you write a lengthy programming specification?
- 2 Can your IWR/IVRS vendor set-up a new protocol within weeks instead of months?
- 3 Does your IWR/IVRS support complex visit schedules and data driven dispensing?
- 4 Can your IWR/IVRS provider handle late breaking protocol changes without significant extra cost and time?
- 5 Is your IWR/IVRS fully integrated with your supply forecasting software?
- 6 Does your IWR/IVRS empower you with direct access to a protocol administration system?



BioClinica has the answers. Make your next trial a success.

1 Does your IWR/IVRS vendor make you write a lengthy programming specification?

BioClinica® Trident reduces the costly and slow routine of writing development specifications, programming and validating a new system each time a new study is built. BioClinica has a simple but thorough specification process that captures the needed information for our team to configure your system. BioClinica completes the first draft of your specification from information in the protocol and/or information collected in a kickoff meeting.

🔗 **BioClinica Advantage: BioClinica does the work that other vendors force you to do.**

2 Can your IWR/IVRS vendor set-up a new protocol within weeks instead of months?

By changing the build process from programming to configuring, BioClinica can reduce your set-up time by 75% or more. Configuration simplifies everything — the specification, the build and validation.

🔗 **BioClinica Advantage: No one has a more flexible system.**

3 Does your IWR/IVRS support complex visit schedules and data driven dispensing?

The Trident visit schedule supports screening, data collection, various enrollment options, fixed/variable-based/titration dispensing, visit skipping, optional and unscheduled visits, re-randomization and configurable window enforcement. Also, Trident allows you to replace patient dispensations without affecting the visit schedule.

🔗 **BioClinica Advantage: No one has a more powerful system.**

4 Can your IWR/IVRS provider handle late-breaking protocol changes without significant extra cost and time?

After an impact analysis is conducted, a study can be re-configured in Trident to accommodate most mid-study changes in little time and with minimal cost.

🔗 **BioClinica Advantage: Save time rewriting specs and eliminate surprises in your budget.**

5 Is your IWR/IVRS fully integrated with your supply forecasting software?

Trident interoperates with BioClinica® Optimizer to determine the optimal re-supply parameters that will yield the maximum business value. Since you've configured all the key IWR/IVRS parameters in the simulator, why set it up again in your IWR/IVRS when you can just push a button and upload your modeled, tested and generally accepted study into Trident.

🔗 **BioClinica Advantage: Save an average of a week of IWR/IVRS set-up effort.**

6 Does your IWR/IVRS empower you with direct access to a protocol administration system?

Using the powerful Trident Admin user interface, business users have the ability to view or control aspects of live studies such as shipping logistics, user management, site management, enhanced reports, resupply etc.

🔗 **BioClinica Advantage: Business users can easily monitor and maintain studies without the need for programmers.**

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BIOCLINICA®
Global clinical trial solutions. Real-world results.

Have more questions?

Learn more about how BioClinica can help you manage your next trial with greater efficiency and less risk. Visit us on the web: www.bioclinica.com or call **888.392.7456**.