MARVIN YOUR STUDY!

xclinical
Countries with medical studies using XClinical’s software

Founded in Munich as an international EDC software vendor

Total studies realised for the medical sector: Phase I–IV pharmaceutical trials, medical device trials, non-interventional studies and registry studies

Patients in therapeutic areas such as Cardiovascular, Oncology, Ophtalmology, Dermatology, Respiratory, Endocrinology, Gastrointestinal, CNS, Reproductive Health, etc.

Different languages in 4 offices worldwide
Imagine a software solution that simplifies and speeds up your entire data management process. A one-stop shop for all your clinical trials.

Our goal is to support the end-to-end process from study setup to regulatory submission, resulting in faster delivery of vital medications and devices to the market. We believe that every day saved can make a difference.
Our Software

Marvin for your trial process: The end-to-end solution.

The eClinical Software Solution Marvin is a CDISC-certified Electronic Data Capture (EDC) system that includes numerous modules such as Randomization (IWRS), Patient Reported Outcomes (WebPRO), Double Data Entry (DDE), Clinical Trial Management (CTM), CDISC Tabulation, Reporting, Coding and more.

XClinical’s product portfolio manages your clinical data from CRF Design to Archiving. With Marvin clinical trials become easy, streamlined and cost-effective.
Take the Test! How does your eClinical Software Solution score?

<table>
<thead>
<tr>
<th>Feature</th>
<th>Other Solutions</th>
<th>marvin</th>
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<tbody>
<tr>
<td>eCRFs in every browser, every language</td>
<td>◯</td>
<td>✓</td>
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<tr>
<td>Set-up without programming</td>
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<td>✓</td>
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<td>IWRS &amp; Drug Management</td>
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<td>WebPRO</td>
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<tr>
<td>Web-based MedDRA &amp; WHODrug Coding</td>
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<td>✓</td>
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<td>Device Data Import</td>
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<tr>
<td>Clinical Trial Management System</td>
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<tr>
<td>Risk-based Monitoring</td>
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<tr>
<td>Double Data Entry</td>
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<tr>
<td>Standard / Customized / Ad Hoc Reports</td>
<td>◯</td>
<td>✓</td>
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<tr>
<td>Auto generated annotated CRFs, DVP, etc.</td>
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<td>✓</td>
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<td>eLearning</td>
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<tr>
<td>SDTM mapping / define.xml</td>
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<td>✓</td>
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<tr>
<td>CDISC end-to-end</td>
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<td>Easy integration with other systems</td>
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... and more!
Marvin Benefits

- For universal use: Multi language capabilities, incl. special character sets. Runs on every browser – in every language
- CDISC throughout the process, provides audit trail and archive files in XML format
- Combination of EDC and DDE in the same study (hybrid)

- Easy to use: True all-in-one platform and the best user experience – no programming needed
- Friendly and professional; offering high-end support services
- Cost-effective solution; individualized for your needs

- Fast setup and automation – quick response times. Implements your requests within weeks, not months. Every day can make a difference!
- Applying highest standards in data security – storage of patient identification in compliance with data privacy laws

Enjoy working with us.

Marvin optimally automates processes.

**COMPLIANT & CERTIFIED:**
- 21 CFR Part 11 compliant
- ICH GCP compliant
- CDISC ODM certified since 2007
Meet our business development team.

Dr. Philippe Verplancke :: Global Head of Business Development and Co-Founder
“We founded XClinical in 2002 with the motivation to support medical research using modern technology. And with us as a medium-sized organization, our customers can get a more personal support and greater flexibility.”

Cathy Hlinka :: VP Business Development USA
“The Marvin software platform is unique in the market largely because we listen to our customers and implement their feedback. The ability to collaborate with the end users who are experts in the industry allows XClinical to build software that people need.”

Jérôme Zakka Bajjani :: Business Development Manager
“Our goal is to produce software that is easy to use and flexible to configure to match many various scenarios for conducting your clinical trial.”

Rupert Sedlmayr :: Business Development Manager
“Our key focus is to enable our customers to address all aspects of this very complex and dynamic market. We are proud to provide them with quality software for managing their clinical trials end-to-end with ease.”

Contact Us – Follow Us.

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