



Streamline monitoring work in clinical trials and clinical research

QLIFICA monitoring is a service that specializes in monitoring in the field of quality control and quality assurance in GCP and Clinical Trials Act.

In addition to improving monitoring efficiency, various functions are installed to support compliance with regulations and monitoring plans.

Monitor

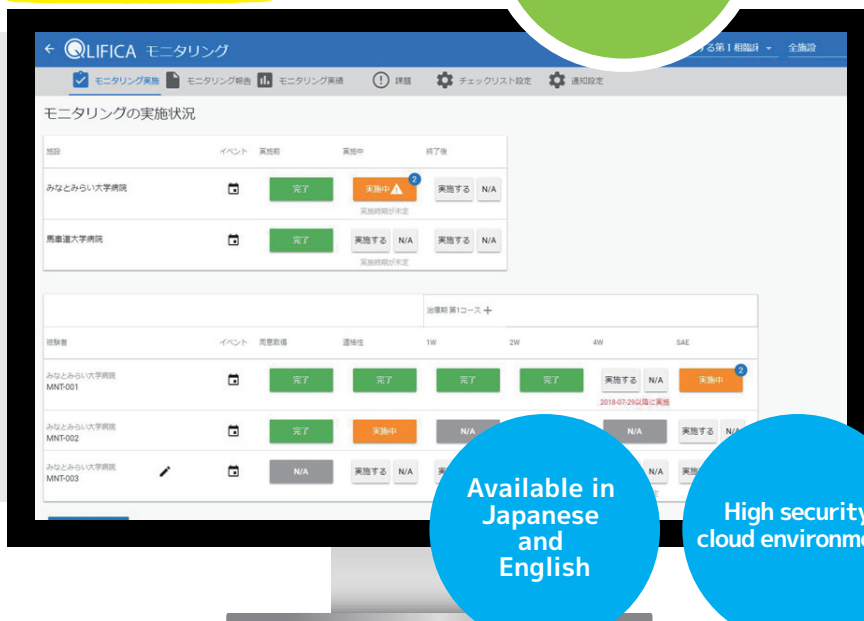
- Implementation of monitoring based on a plan
- Creation and submission of monitoring report
- Check and resolve issues



Project manager

- Creation of monitoring plan
- Review of monitoring report
- Progress management of studies and monitoring



Free trial for **90** days

Apply for a free trial

Please contact us from the contact information below or access our QR code and visit our website.

Also an online demonstration that you can try out is available on our website.

Available in
Japanese
and
English

High security
cloud environment

Validation
that meets
the requirements
of the ER / ES
guidelines



Point 1

Easy to use

Simple operation and procedure assistance ensure reliable implementation even for users with little work experiences.

Point 2

Save money

No limit to the number of studies that can be registered.
The cost will be charged per user license, so you can easily introduce the system.

Share progress with your team

You can grasp how much procedures and case monitoring are progressing, and also check the resolution status of issues in the list.
The monitoring records and history can be downloaded in Excel format.

Work on a PC or iPad wherever you go

As long as you have internet access, you can conduct on-site monitoring using a checklist based on your monitoring plan. You can also use the travel time to create a monitoring report and submit it online.

Get notification of monitoring implementation

Depending on the progress of the study, we will guide you what kind of monitoring should be done next and notify you of the timing of the monitoring by e-mail.

Reduce the time and burden of reporting

The monitoring report can be automatically generated from the monitoring record and the creation time can be shortened. And the request email of monitoring report review is automatically sent, so you can work smoothly.

Functions

Registration of trial related information / Management of study progress / Creating a monitoring plan / Confirmation of monitoring status / Monitoring implementation / Creation and submission of monitoring reports / Review of monitoring report / Download monitoring results / Various email notifications

Contact us Feel free to contact us by phone, email or our website.



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QLIFICA

