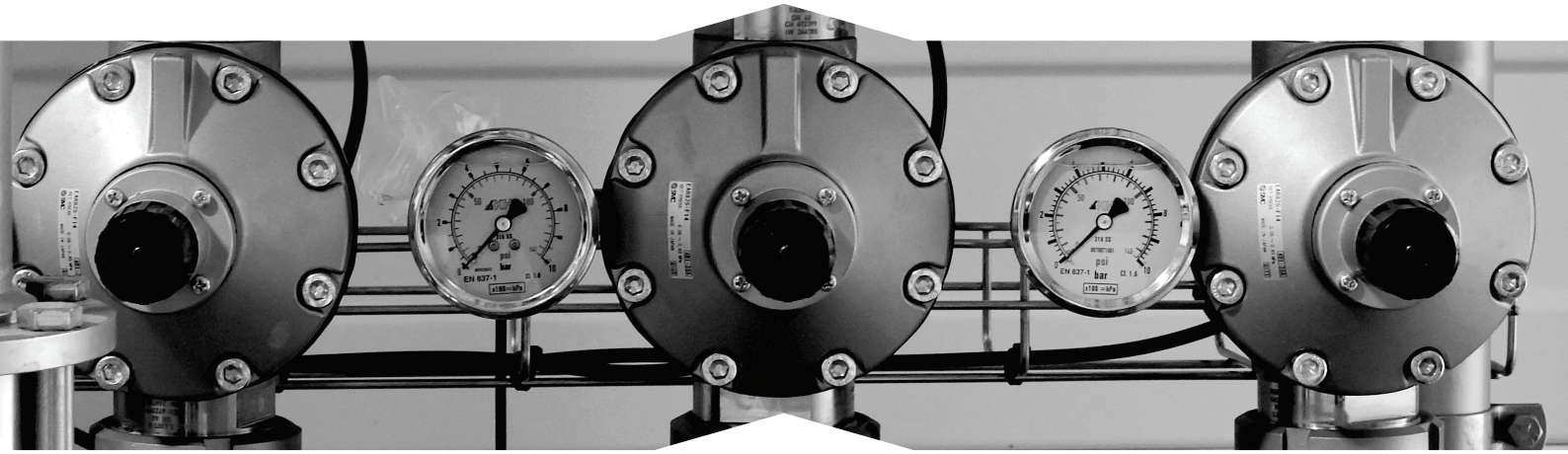
OCS Life Sciences



DATA CONVERSION FOR REGULATORY SUBMISSION

OCS Life Sciences has a specialised team for the conversion of clinical trial data for regulatory submissions. Equipped with innovative software our scientifically educated data managers and clinical programmers relieve you from the burden, allowing you to focus on all other aspects of your submission.

Our services for data conversion include:
– Project planning and management
– SDTM and ADaM datasets
– Define-XML for SDTM and ADaM
– Study and Analysis Data Reviewer's Guides
– Annotated Case Report Forms

DATA INTEGRATION AND POOLING

Even when you are not submitting to the authorities you will greatly benefit from integrating or pooling your data in CDISC-compliant SDTM and ADaM datasets.

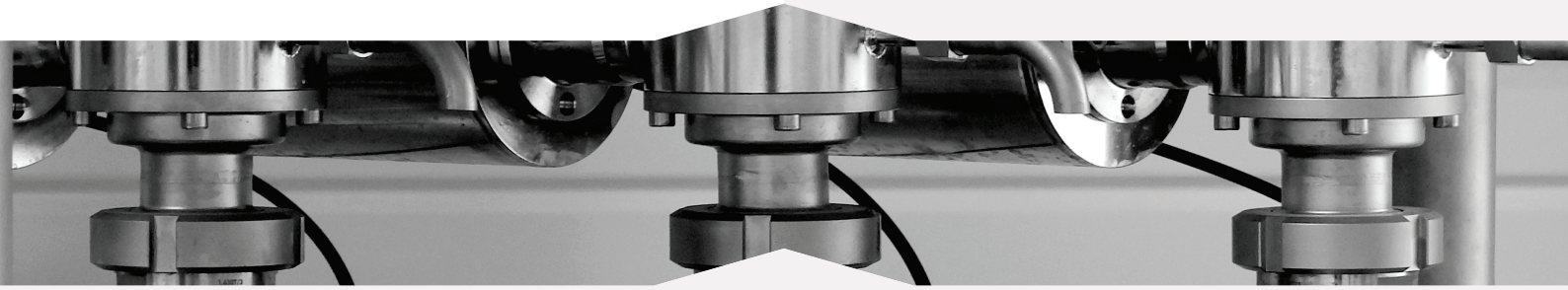
– Publications on consolidated findings
– Enhanced precision in efficacy analysis
– Identification of rare safety signals
– Better exploration of drug interactions
– Starting point for ISS and ISE reports
– Structured basis for data visualisation

THE KEY TO OUR SUCCESS

The key to our success is the way we engage with our clients. We have been most succesful not working *for* our clients, but by working side by side *with* our clients.

You get to be consulted in every decision that you wish to influence, and can leave everything else with us.

It's that simple.



Keen to find out more?
Visit ocs-lifesciences.com/data-conversion or get in touch with Melanie Schopp at melanie.schopp@ocs-consulting.com

www.ocs-lifesciences.com
info.nl@ocs-consulting.com
+31 (0)73 523 6000