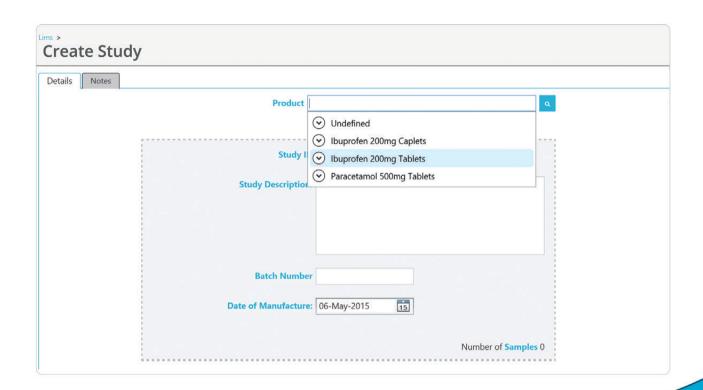


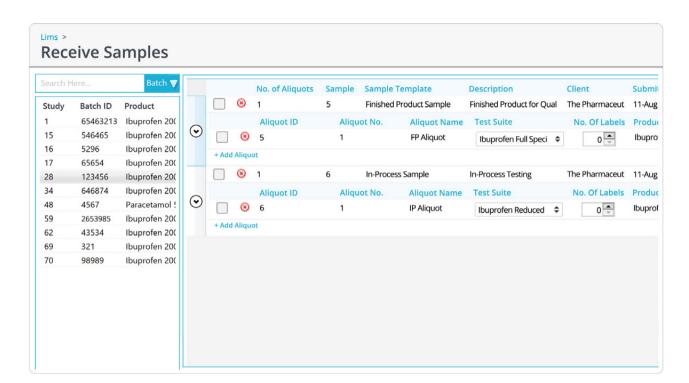
LOGIN!

Users can securely and confidently login to LabHQ LIMS through a user-friendly interface. With username and password protection, access to your data will be controlled. Give your clients read-only access to their own data through the web-client. With their own username and password they can too enjoy using LabHQ just like you!

STEP 1. SAMPLE SUBMISSION

Whether you're testing a batch of product as a support function to manufacturing or you're testing multiple batches as a contract laboratory, LabHQ allows you to submit samples easily. Fields are pre-populated based on the permissions for the user logged in. LabHQ can store your test methods, test schedule, and product specifications in a version-controlled system enabling you and your clients to submit samples for testing with ease.







Customise LabHQ with your own language using our easy to configure localisation file.

STEP 2. RECEIVE SAMPLES

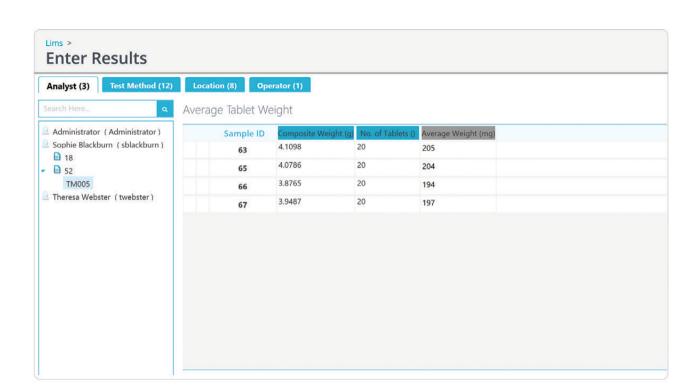
When samples physically arrive to the lab, this stage allows you to reconcile with the submission details ensuring the product, batch numbers, and test schedule were correctly entered. You can also print labels to ensure that each sample displays the appropriate information while it's being tested or stored waiting for testing.

STEP 3. CREATE WORK

So you've suited up with a lab coat and goggles and you're ready to go to work. What's next on the list?

You can choose how to organise your work. Do you prefer to do one test on multiple samples, or take one sample through all the testing it requires? Either way, LabHQ provides you with the flexibility to work however you please. You can group which samples you wish to analyse and prioritise based on turnaround or on a first in first out basis.

Lims > **Create Work** Sample Test Method Name ▼ Study Aliquot Analyst name Username Administrator Administrator Test Method ID **Test Method Description** Aliquot ID Product Name **Product Description** Sophie Blackburn sblackburn V Ibuprofen 200mg Tal TM004 Disintegration of Tablets and Capsules 54 Ibuprofen 200mg Tablets Theresa Webster twebster V TM004 Disintegration of Tablets and Capsules 55 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal V TM004 Disintegration of Tablets and Capsules 63 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal V TM004 Disintegration of Tablets and Capsules 64 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal TM004 Disintegration of Tablets and Capsules 66 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal TM004 Disintegration of Tablets and Capsules 67 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal TM004 125 Disintegration of Tablets and Capsules Ibuprofen 200mg Tablets Ibuprofen 200mg Tal TM004 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal Disintegration of Tablets and Capsules 126 TM004 Disintegration of Tablets and Capsules 129 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal TM004 Disintegration of Tablets and Capsules 104 Ibuprofen 200mg Tablets Ibuprofen 200mg Tal Ibuprofen 200mg Tal TM004 Disintegration of Tablets and Capsules 105 Ibuprofen 200mg Tablets



STEP 4. ENTER RESULTS

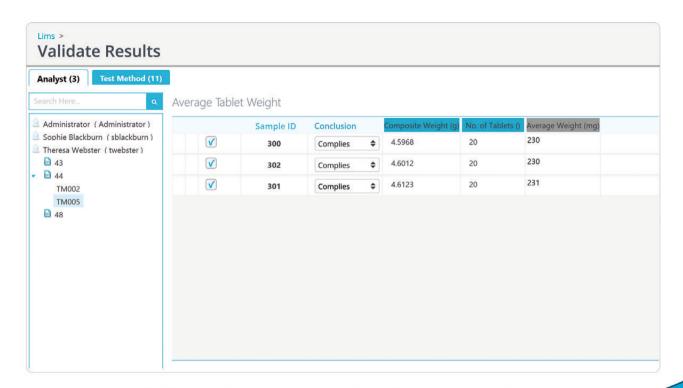
After completing tests on a sample, entering the results on LabHQ gives you the opportunity to check the result against the specification. Any out of trend, or out of specification results can be investigated immediately to help achieve turnaround times.

Want to reduce human error? Why not try our results import module with equipment integration? Want traceability of the stock or equipment used during testing? Then go for our stock and equipment modules, which seamlessly integrate with LIMS and store all of your data in one place.

STEP 5. VALIDATE RESULTS

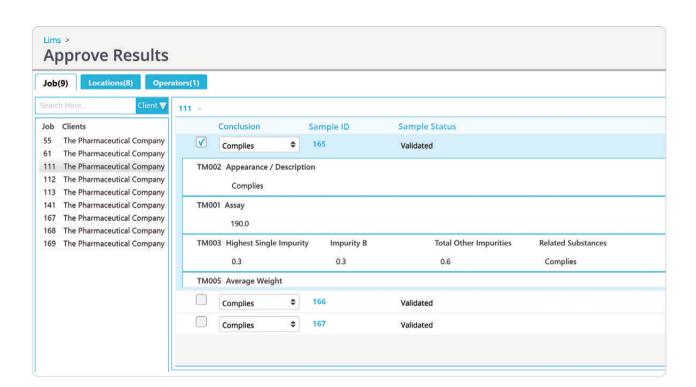
Labs operating a quality management system may implement the "four-eyes" rule to checking results.

LabHQ drives an independent review of results before formal approval and reporting. Having another person review your work minimises the number of transcription errors.





Is your sample lifecycle shorter? or longer? Workflows are conveniently customisable in LabHQ.

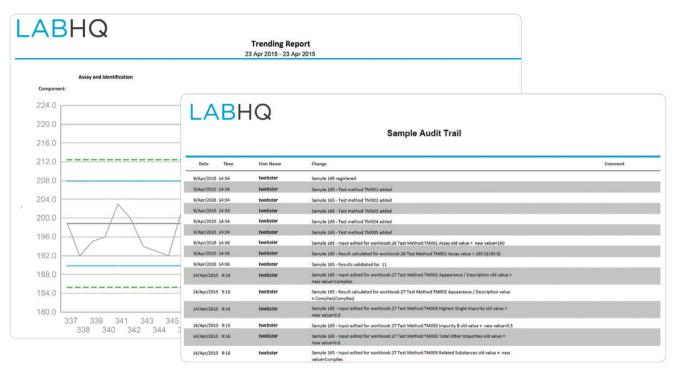


STEP 6. APPROVE RESULTS

Once all results are entered for a sample and validated, the sample is then ready for approval. As the final step in the sample lifecycle, approve results is the last check before the results are formally reported. The sample conclusion is then applied as "Complies" or "Does not comply" providing an unambiguous summary of the sample's testing outcome.

STEP 7. REPORT RESULTS

Generate certificates of analysis or trend reports with one simple click. LabHQ has made it easier for lab users to generate meaningful reports accurately and quickly providing a professional finish to the sample analysis process. For quality reviews, LabHQ also provides a suite of system audit trails ensuring that each action is appropriately recorded. Where changes are made the reason is logged against the user, date and time.





Want your data presented differently? Simply customise your reports in LabHQ.

WANT MORE FUNCTIONALITY TO HELP YOU MANAGE THE LAB?

Automate the supporting processes to your quality control analysis through seamless integration with our complementary LabHQ modules.



Environmental Monitoring



Equipment Maintenance



Stock Management



Stability Study Management

CONTACT US

We'd Love to Hear From You!

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