



Valab[®] Manufacturer's Information

Information for medical laboratory accreditation

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Subject

This document is for medical laboratories that use the Valab® expert system for computer aided biological validation. It provides the medical laboratory with information from the manufacturer, the VALAB company, concerning the use of Valab® in an accredited medical laboratory.

This document serves as a complement to the document "[Qualification of Valab® by the Medical Laboratory](#)" (RD2) which comprises a guide providing an approach and a set of example procedures for qualifying the use of Valab® in the laboratory and performing the control of the system in terms of monitoring, maintenance and traceability of modifications.

For more details on the use of Valab® and on the description of its interface with the LIS, please refer respectively to the Valab® User Manual and Valab® Developer Manual provided with Valab®.

References

Update changes

Update of the document for the 16th version of Valab®, impacting mainly the chapters "[LDAP directory connexion](#)", "[BI module ValView](#)" and "[VALAB Services Provider \(VSP\)](#)".

Valab® web site www.valab.com

Click the following link to [find the current version of this document on the Downloads page of the Valab® web site \[www.valab.com\]\(http://www.valab.com\)](#).

Reference documents

RD1	Medical laboratories — Requirements for quality and competence • ISO 15189
RD2	Qualification of Valab® by the Medical Laboratory (available at www.valab.com)
RD3	Valab® User Manual (provided in Valab® "DOC" subdirectory)
RD4	Valab® Developer Manual (provided in Valab® "DOC" subdirectory)
RD5	Valab® - Backup and Restore (available at www.valab.com)
RD6	Manuel Qualité VALAB (Valab® Quality Manual - only available in French) (available at www.valab.com)
RD7	Privacy Policy of the VALAB Company (available at www.valab.com) • VALAB
RD8	Accreditation requirements according to standard NF EN ISO 15189 • Cofrac / SH REF 02
RD9	Accreditation technical guide for medical laboratories • Cofrac / SH GTA 01
RD10	Accreditation technical guide to assess the IT systems in medical biology • Cofrac / SH GTA 02
RD11	Articles L.6211-1 and following of the French Code of Public Health • CSP
RD12	Articles L.5221-1 and following of the French Code of Public Health • CSP
RD13	General data protection regulation (GDPR) • Regulation (UE) 2016/679

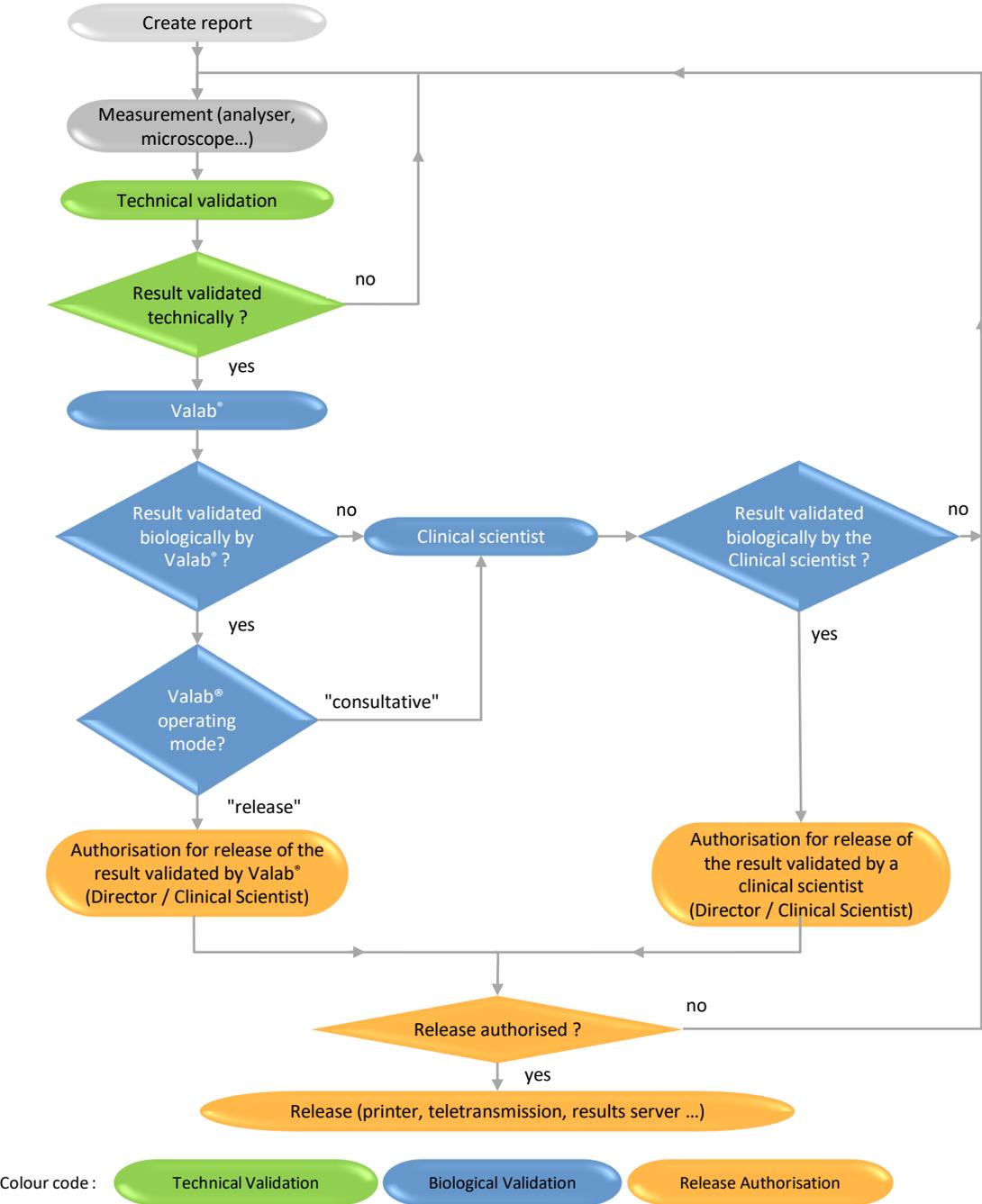
Acronyms

Acronyms	Meaning
LIS	Laboratory Information System
ML	Medical Laboratory
MMI	Man-Machine Interface
QMS	Quality Management System
RCV	Reference Change Value
RD	Reference Document
See	Refers to

Reminder of the Valab® integration into the Medical laboratory

Functional integration overview

The following overview shows the functional integration of Valab® in the laboratory process for the validation of patient test results.



General introduction

Functionality

Valab® is a computer-aided biological validation tool. It assists the clinical scientist with the systematic review of results, their assessment in conformity with available clinical information about the patient, and authorisation for their release. Valab® acts as a filter which identifies any incoherent reports and/or reports containing alerts or critical values for which the intervention of the clinical scientist represents a true biological and medical added value before release, if applicable.

The main objectives of the tool are to improve the safety, reproducibility, quality and turnaround times for results. Moreover, in the context of a medical laboratory with several clinical scientists, the Valab® software tool allows to standardise the act of validation and to assess the practice of biological validation by consensualising the different clinical scientists around the use of a single tool.

In all cases, it is important to indicate that all the medical test results released by the medical laboratory are done so under the responsibility of the clinical scientist, including those verified with the help of the Valab® computer aided validation software. In this respect, all the results reports issued by the medical laboratory are assumed to have been validated by the clinical scientist and must bear his/her signature (first name, surname and signature configured in the LIS). Inscriptions of the form "validated by expert system" are not acceptable.

Means of control

As a complement to the present document, the VALAB company provides its users with a guide for the "Qualification of Valab® by the Medical Laboratory" ([RD2](#)).

This guide provides an approach and a set of example procedures allowing to set up the control of the use of the tool. The recommendations concern in particular the customisation of the parameter settings, the initial qualification, re-qualification after modification, continuous monitoring, the keeping of records, the impact on the QMS documentation of the medical laboratory, and user training.

Advice and expertise

The VALAB company accompanies the user with its expertise during the period of customisation and initial qualification, and afterwards through a maintenance agreement:

- analysis of the statistical indicators with regard to the autoverification of each test
- proposals for parameter setting modifications to the Valab® key contact clinical scientists
- assistance through remote maintenance

Integrated operation

The connection with the LIS and integration of the expertise results of each test on the biological validation screens of the LIS exists for many laboratory information systems, without changing the work habits of the laboratory.

Conclusion

Valab® provides a solution to perform, under the responsibility of the clinical scientist and in conformity with regulatory, legal and normative requirements ([RD1](#) and [RD8](#)), the systematic review of all the results before their

release. This approach also offers the possibility to approach biological validation using consistent and uniform criteria in the context of technical platforms and multi-site laboratories.

Regulatory, legal and normative context

The biological validation activity must consider the current and upcoming regulatory, legal and standard requirements. They blend in with the broader context of quality, the good practices of the profession and the laboratories accreditation. The corresponding reference documents are:

- ISO standard 15189 ([RD1](#)): “Medical laboratories — Requirements for quality and competence • ISO 15189”
- Cofrac document SH REF 02 ([RD8](#)): “Accreditation requirements according to standard NF EN ISO 15189 • Cofrac / SH REF 02”

Analysis of these texts converges in the idea that the validation and authorisation for release of results is an intellectual, systematic and efficient act under the responsibility of the clinical scientist:

- validation must consider the coherence of all the data known for the patient
- validation must be exhaustive and allow the review of all the results, normal and abnormal
- validation must make use of innovation to improve the efficiency of the medical laboratory

History and method of initial qualification of Valab[®]'s expert kernel

History

- In 1986 the Valab[®] project was initiated in the Clinical Chemistry ML of Rangueil University Hospital, Toulouse, France, as a prototype for Biochemistry. The knowledge representation model came from research. Information technology allows to reproduce the logic, complexity and combinatory power of human reasoning by weighting each piece of information considered during the verification process.
- From 1989 the EREMS company developed and marketed the Valab[®] product in the context of a technology transfer initiative supported by the ANVAR (French Agency for Innovation). The evolution of Valab[®] continued to benefit from a partnership with the Toulouse Hospitals for the development of new applications in Haematology, Blood Gases, and Coagulation. Valab[®] became a registered trademark in 1990.
- In 1992, the CNEH (French National Centre for Hospital Expertise, Paris, France) performed a multi-centre assessment confirming the validity of the autoverification results provided by Valab[®]'s expert kernel (TEP / ANVAR procedure, see section "[Initial qualification of the expert kernel by the manufacturer](#)").
- In 1999 the CNEH performed a multi-centre assessment confirming the organisational and financial benefits for the medical laboratory to be obtained by using Valab[®] (procedure ACE-GBM / ANVAR).
- In 2004 the VALAB department of the EREMS company became independent as the VALAB company which now develops, markets and maintains the Valab[®] product. The VALAB company was certified ISO 9001 in 2004 for the design, development, marketing, installation, training and after-sales service of expert systems and software in the field of computer-aided validation of test results in ML.

Initial qualification of the expert kernel by the manufacturer

A methodology for validation of the expert kernel was defined right from the first versions of Valab[®], the validation phase representing a mandatory stage to guarantee the reliability and safety of any operational system. This methodology was used in particular during the multi-centre study carried out in 1992 by the CNEH (TEP / ANVAR procedure). This investigation allowed to confront the expertise results provided by Valab[®] with those provided by the clinical scientists of five user sites on a pool of around 300 patient reports per site (i.e. a total of 1500 reports), and confirmed the validity of the expertise results provided by the expert kernel of Valab[®].

Methodology

The guiding principle was to evaluate the performance of the expert system with quantitative criteria by using a prospective protocol based on the comparison of the Valab[®] expertise results versus those of the clinical scientists.

The methodology takes its inspiration from the epidemiological assessment methods of screening tests, widely used in medicine. Valab[®] is therefore considered to be a system capable of performing a test of the coherency of results.

Biological validation determines the acceptability of a result or a set of results which are either validated or blocked. This Boolean nature allows to assimilate Valab[®] autoverification to a screening test of the reports requiring human intervention for their validation. The test is positive when the report is blocked, negative when it is validated. In this way the following equivalences are considered:

- | | | |
|---------|----------------|--------------------------|
| ■ V(+): | True Positive | Report rightly blocked |
| ■ V(-): | True Negative | Report rightly validated |
| ■ F(+): | False Positive | Report wrongly blocked |
| ■ F(-): | False Negative | Report wrongly validated |

The interpretation of the performance of the test is obtained after adding up the numbers present in each response class [V(+), V(-), F(+), F(-)] by calculating the assessment criteria of the screening methods: sensitivity, specificity, positive and negative predictive values. These different epidemiological criteria correspond to the following definitions:

- Sensitivity: % of reports to be blocked for which the test is positive = $(V+) / [(V+) + (F-)]$
- Specificity: % of reports to be validated for which the test is negative = $(V-) / [(V-) + (F+)]$
- Positive predictive value: % of reports to be blocked among the positive tests = $(V+) / [(V+) + (F+)]$
- Negative predictive value: % of reports to be validated among the negative tests = $(V-) / [(V-) + (F-)]$

Protocol

The assessment is based on the verification of a pool of reports submitted in parallel for validation by the clinical scientists and by Valab[®]. The theoretical assessment phase begins with the creation of a significant and representative sample (e.g. 500 patients' reports).

The procedure then consists in carrying out a blind validation of the reports of the sample, by Valab[®] and by the clinical scientists participating in the study. In order to recreate the natural work conditions of the clinical scientist, the reports are verified sequentially and individually: in series of 100 at a time, without knowing the response either of the other experts or of Valab[®]. The data is collected from individual response notebooks. The exploitation of all this data consists in making a synthesis for each report verified, by bringing together the

information contained in each individual response notebook. The experiment shows that around 20% of the reports do not receive a unanimous assessment. The establishment of a reference response for the verification of each report is obtained when the response is unanimous. For the other reports, a consensus meeting is organised to determine the reference response.

The interpretation of this assessment protocol is based on the comparison of the performance obtained by Valab® and by the clinical scientists. For this, the confrontation of all the responses of each individual notebook with that of the reference notebook allows to add up the number present in each response class [V(+), V(-), F(+), F(-)] for each of the participants, the clinical scientists and Valab®.

The final stage consists in calculating the assessment criteria of the screening methods: sensitivity, specificity, positive and negative predictive values. The global performance of Valab® must be evaluated by the scores obtained for the different numerical criteria in comparison with the clinical scientists.

As the consequences of a validation error can be serious in the case of an inappropriate validation (false negative), the obtention by Valab® of a better score for sensitivity testifies to a low rate of false negatives, therefore putting safety first.

Interpretation

Without being exhaustive, the epidemiological approach offers the possibility of assessing the performance of a test through quantitative criteria. The pertinence of this protocol resides on the one hand in the assimilation of the biological validation to a test situation, on the other hand in the implementation of a prospective and comparative procedure between the Valab® system and the clinical scientists.

Several protocols of this nature have now been conducted and published by the VALAB company, by the laboratories using Valab®, and also by the CNEH. The results obtained have always met the requirements of reliability and safety by demonstrating excellent performance in terms of sensitivity. This demonstrates the capability of the Valab® computer-aided validation tool to not validate reports wrongly.

Conclusion

The initial validation of Valab® in collaboration with users gave rise, in France, to two multi-centre assessments by the CNEH (TEP / ANVAR procedure in 1992 and ACE-GBM / ANVAR procedure in 1999).

Using Valab® has given rise to many publications in congresses and scientific journals with reviewing committees (see section "[Bibliography and press articles on Valab®](#)").

Using the Valab® computer-aided validation system in 2025 represents the state of the art in the light of the following data:

- Nearly 40 years developing, maintaining and commercialising
- Over 350 medical laboratories / multi-site structures using Valab® in France, Europe, French-speaking Africa and South America
- All the medical laboratories / legal entities users in France are accredited (ISO 15189)
- Over 800,000 patient reports autoverified every day
- International distributors

The industrial qualification of successive versions of the Valab® software (see section "[Development cycle and qualification of each version of Valab®](#)") is subjected to a full test plan in conformity with the requirements of our QMS. It integrates in particular, for each version and incrementally:

- The use of a methodology inspired from the Agile Scrum method
- Unitary and integration tests
- System tests

- Non-regression testing
- Verification matrix

Development cycle and qualification of each version of Valab®

In conformity with the description of the VALAB company's Research and Development process, each new version of Valab® is developed according to a method inspired from the Agile Scrum method.

This method guarantees the control of the development process for each new version of the product. The average development cycle duration for a major version of Valab® is 24 months.

The following paragraphs describe the main steps in this development cycle.

Specification

The specification of each new version of Valab® is defined in a specification document specific to each version. This document is organised by topics / specification categories (user interface, functionality, protocol, verification expertise, bug corrections, architecture...). It takes into account the following factors:

- The problems detected by users and recorded in problem reports
- The results of satisfaction surveys conducted with our customers
- The result of the monitoring of regulatory, legal, normative changes and technical, technological and business advances
- The internal and external ideas for product improvements

Each specification is identified by a unique identifier which is then included in the traceability matrix for system tests (see section "[System tests](#)").

Planning

The developing of the Valab® next versions planning is formalised into a RoadMap document along with the definition of the chosen delivery dates.

Once the content of a version is specified, the estimated delivery date is reexamined each time a sprint ends according to the speed of the development team and what remains to be done.

According to the calculation result, the version content may be increased or decreased (along with an update of the specifications). Decision can be made of modifying the chosen date of the version release.

Initialisation

The initialisation (or design) stage allows to study the architecture to be implemented for the new version of Valab® in order to meet the specifications and the schedule of the new version.

The design takes the form of several version-specific analysis documents.

The general architecture of the software is created in UML (Unified Modelling Language).

Realisation

The realisation (or coding) stage is carried out using the documents from the initialisation phase, in conformity with the procedures of the Research and Development process of the VALAB company (coding rules, software configuration management procedure, development procedure, software design and coding).

Verification and Validation

This paragraph explains how the verification and validation stages are qualifying each new version of Valab®.

Unit tests

The unit tests allow to validate singly / independently each elementary object of the Valab® software architecture. These unit tests are implemented as source code. New unit tests are added with each new version of the product.

Tools allow to "play" the unit tests automatically and at will throughout the coding phase in order to validate the new code as it is developed.

All the unit tests are then re-run at the end of the development cycle, they must be run successfully before the system tests can be executed. The results of the unit tests are then reported in the test reports of the version (see section "[System tests](#)").

Integration tests

Integration tests verify how different code units work together. While individual units may work well on their own, integration testing ensures that they work together coherently. These integration unit tests are implemented as source code. They are enhanced with each new version of the product.

Tools allow to "play" the integration tests automatically and at will throughout the coding phase in order to validate the new code as it is developed.

All the integration tests are then re-run at the end of the development cycle, they must be run successfully before the system tests can be executed. The results of the integration tests are then reported in the test reports of the version (see section "[System tests](#)").

Coding review

A coding review is performed at the end of the development cycle, it is recorded in a coding review report.

This review relies in particular on the use of metrics generated by tools for the automatic analysis of code complexity, respect of coding rules, comment inclusion rates, and good practice in relation to the development language. These metrics are checked throughout the entire coding phase in order to detect any deviations at this level.

System tests

The system tests stage allows to qualify the finished product as it will be delivered to the customer. It is formalised in a test and validation document (test and validation plan) and in the form of a source code for automated systems tests (typically, API tests).

These system tests are enhanced with each new version with new tests allowing to qualify the new product functionality. As a complement, the existing tests used for the qualification of previous versions of the product are verified and adapted to any modifications and / or corrections applied to existing product functionality.

The approach is to test each function of the finished product by using test scenarios to validate whether the product meets the specifications. These tests also include tests of backward compatibility with previous versions and non-regression tests allowing to ensure that the product has not regressed with regard to previous versions both at the level of verification expertise and on the level of the other functionality of the software.

The validation test document contains reports that indicate the result of each system test (manual and automated) and a report that indicates the result of the unit tests and integration tests.

As a complement, in order to check that all the specifications of the new version have been tested, a traceability matrix is used which contains the unique identifier of each specification (see section "[Specification](#)") and the tests which are associated with it. This ensures that all the specifications have been tested.

When each of the unit, integration and system tests has been successfully executed or exceptionally accepted by waiver (with an error report for minor problems, if applicable), the version is considered as qualified.

Release of the version

At the end of the development cycle, when all the steps of the cycle have been passed with success, the product is released under the responsibility of the Director of the VALAB company. The Research and Development process of VALAB then makes the qualified version available to the VALAB Order Delivery and Customer Support processes with a view to its release. The version is then considered to be released.

This new version is first of all released to a limited panel of our customers under a maintenance agreement, then progressively to all of our existing customers as well as to new customers.

Registering the new version source components

At the end of the development cycle, the source components of the new Valab® version (software source code, related documentation, transfer document allowing to regenerate the product deliverables) are registered with a specialised third-party company.

Valab® autoverification

Principle

Valab® is an expert system born out of research in artificial intelligence and knowledge modelling. It does not approach biological autoverification like a classic algorithmic system, but globally and contextually just like a clinical scientist does.

The "intelligent" model at the origin of Valab® consists in implementing all the processes related to the interpretation of a result according to a strategy approaching the logic, complexity and combinatorial nature of human reasoning. The Valab® software allows to weigh up each piece of information taken into account in the verification.

The acceptability of a result depends on a true combinatorial analysis and not on a classification in a pathological model. This last point makes it difficult to provide an exhaustive description of the verification logic because in no way does the result depend on the application of an exclusive rule, on the contrary, it depends on the application of all the applicable rules.

A simple example illustrates this aspect:

Given a patient with the clinical information "hepatitis" and an extremely disturbed ASAT (GOT) result at 800 UI 37: this information is compatible, and a simple rule allows to model it.

However, adding an ALAT (GPT) with a normal result of 20 UI 37° in the same report makes the overall report incoherent and it is no longer validated.

Only a "combinatorial" approach allows to describe complex models in this way.

The following information categories are used by the Valab® autoverification in the context of a global reasoning strategy for validation:

- Demographics (age, sex)
- Context (prescriber and/or department, emergency, hospitalised)
- Therapeutic and clinical information (medical data)
- Technical information (observation data)
- Inter-parametric correlation (coherence of the results)
- Anteriority and inter-parametric kinetics (variation and kinetics of the results)

The strategy implemented in Valab®, with a global and systematic approach for the validation of each patient report, corresponds to the Quality requirements of the profession. This approach allows to both validate a "normally abnormal" result and block an "abnormally normal" result.

As a computer-aided biological validation system, Valab® represents an intelligent filter at the service of the clinical scientist and brings additional safety and efficiency to the organisation of a medical laboratory. To this end, the implementation and use of such a system participates both in the improvement of the quality of the results returned to the patient and in the accreditation strategy of the laboratory.

Knowledge modelling methodology

The cognitive engineering methodology for knowledge modelling, used to model the Valab® verification expertise, comprises three main steps:

- 1) Knowledge acquisition:

- Theoretical part: bibliographical research
 - Experimental part: observation and analysis of experts (originally, the teams of Pr. P. Valdiguié, Pr. J.X. Corberand and Pr. B. Boneu – Toulouse University Hospital, France) in situ in order to decompose their reasoning into basic rules
- 2) Modelling:
- This consists in building a software model allowing to exploit the basic rules, previously identified in step 1, and to reproduce the complex reasoning of the expert
- 3) Qualification:
- This step consists in validating the model obtained in step 2. For more details on the qualification methodology, see section "Initial qualification of the expert kernel by the manufacturer"

Traceability and determinism

An indispensable factor for the ethical acceptability of an expert system concerns its operating reliability and its traceability in the context of accreditation for obvious reasons of responsibility.

The technical characteristics of the tools at the base of the Valab[®] application (compilation process and consistency analysis) have allowed to verify the logical coherence of a knowledge base model. The exclusive use of this model allows to guarantee the lack of contradictions between the rules and to automatically generate the algorithmic and deterministic equivalent of an expertise set initially described in the form of rules.

As Valab[®] is a deterministic system, the decision process is always the same for identical operating conditions (version, parameter settings, correspondence tables, parameter values,...).

Upon a user's request for a problem related to the laboratory's responsibility concerning an autoverification result for a specific report, we apply, in a confidentiality framework, an exceptional procedure.

This procedure allows to reproduce all the elementary reasoning implemented in the decision process, and to provide the user with a comprehensible list of all the rules applied in this particular case.

The Benelux countries were precursors with regard to laboratory accreditation. In 2009, half of the laboratories using Valab[®] in Benelux were already accredited according to the ISO 15189 (RD1) standard: 30 laboratories out of the 60 equipped with Valab[®]. Here is an example of a procedure set up in October 1996 with the CCKL (Dutch body for the improvement and maintaining of Quality in the health care system):

Certification / accreditation procedure (NL) :

- ⇒ Valab[®] is a deterministic system,
 - ⇒ Decision process is always the same if environmental conditions are similar (value of parameters, dictionaries, parameter settings...),
 - ⇒ In case of customer's question about the expertise on a specific report, we can establish an exceptional procedure as following :
 - * get the number of the Valab[®] version,
 - * get the complete parameter settings (backup),
 - * get the precise data of the report inside Valab[®] (file *.ptd),
 - * identify the parameter(s) with the understanding problem to be justified,
 - * With this information we are able to produce the whole elementary reasoning used in order to explain the global decision,
 - * So our response could be to furnish the list, clearly readable, of all the rules triggered in this case,
 - ⇒ Of course, such a heavy procedure would have to be exceptional, only when there is a responsibility problem for the laboratory.
-

Implementation and control of Valab[®]

The operating environment of Valab[®] benefits from a set of functionality and documentation allowing to meet the requirements for Quality and accreditation in the laboratory.

Additionally, all of the members of staff that use the Valab[®] software must be trained on how it works and how to use it.

The means of control to be implemented can be based on a risk analysis (see section "Example of risk analysis" of the [RD2](#) "Qualification of Valab® by the medical laboratory").

Customisation of the setup

During the Valab® deployment process, the key contact clinical scientists of the medical laboratory are accompanied by the "biological expertise" Customer Support group of the VALAB company and/or by the support of the distributor when analysing production data in order to finalise the customisation of the Valab® parameter settings. This phase takes place in several successive work sessions over a sufficient and significant period of activity of the medical laboratory in order to obtain a Valab® operation that is suited to the population, the specifics and the requirements of the medical laboratory.

The following paragraphs describe the main parameters of Valab® that can be customised by the medical laboratory.

"Global" consultative mode

The "Global" consultative mode, when activated, allows to present all the reports verified by Valab® during a LIS validation session and systematically requires a manual validation of each report by the clinical scientist .

This option is activated in particular at the startup of a new site and during the period necessary for the customisation of the parameter settings and the initial qualification of the Valab® autoverification by the medical laboratory (see [RD2](#) "Qualification of Valab® by the Medical Laboratory").

Consultative mode specific to one particular test

It is possible to put an individual test in consultative mode ("consultative mode" for a specific test). Reports containing such a test will be verified but they will never be validated by Valab®, and they will be systematically proposed to the clinical scientist in a manual validation session by the LIS.

This option can be activated for example during the connection of a new test to Valab® to allow the medical laboratory to qualify the corresponding autoverification over a given period of time.

Threshold test

Each test can be declared as a threshold test. In this case, a change of state of the test (from "normal" to "abnormal", and inversely) causes the refusal of this test by Valab®. A sub-option enables to cause the rejection only in case of a change in the state of a test from "normal" to "abnormal".

Unit

The units in which each test is sent by the LIS and/or displayed in Valab® are programmable.

Intervals referred to as "Biological (or clinical) reference interval limits for a man of 25 years of age" and "demographic rules"

In [RD1](#) – “Medical laboratories — Requirements for quality and competence • ISO 15189”, the *biological reference intervals or reference intervals* are defined as *the specified intervals of the distribution of values taken from a biological reference population*. These are the intervals which are shown on the report and allow to situate the result of a test. *Clinical decision limits* are defined as *test results indicating an increased risk of negative clinical outcomes or diagnosing the presence of a specific pathology*.

In Valab®, the intervals called "Biological (or Clinical) reference interval limits for a man of 25 years of age" and the "demographic rules" correspond in part to *the limits and conditions of implementation of modalities for biological validation with a computer-aided validation software* mentioned in [RD8](#) – “Accreditation requirements according to standard NF EN ISO 15189 • Cofrac / SH REF 02” and can be associated with *the clinical decision limits* as defined in [RD1](#) – “Medical laboratories — Requirements for quality and competence • ISO 15189”. They are established by the laboratory following customisation of the setup conducted by each medical laboratory during initial qualification and continuous monitoring, in particular by exploiting the statistical data for the laboratory generated by Valab®. In the end, the setup parameters retained correspond to the population, specifics and requirements of the medical laboratory regarding biological validation. The use of these intervals is strictly internal to the medical laboratory.

The ranges defined in Valab® fix the limits inside which a result cannot be interpreted as being pathological in the clinical sense, and outside of which it cannot be validated by Valab® unless it is justified by another item in the report (values of other tests, pathology, treatment, known previous result, etc.). It is therefore clear that these intervals may be different from the biological reference values indicated on the results report (e.g. cholesterol).

The "Biological (or clinical) reference interval limits for a man of 25 years of age" and the "demographic rules" retained must be precisely recorded in the laboratory's QMS and all of the clinical scientists must be informed.

Note that the initial settings for the tests provided with Valab® were established based on bibliographic research (see the bibliographical references in section "[Valab® biological bibliography - Extract](#)").

"Expertise" limits for the results without anteriority

"Expertise" limits for the results without anteriority forbid the validation of a test having no anteriority (1 ° realization of this test for the patient or too former anteriority) and a result which is under and beyond a certain value whatever is the coherence of the data.

"Alert" or "critical" limits

The "alert" or "critical" limits prohibit the validation of a test below or above a certain value. Whatever the coherence of the data, they constitute the absolute limits of the field of competence of the system, requiring the mandatory submission of reports with such results to the clinical scientist for validation.

Delta check / RCV

The RCV ("Reference Change Value") is an acceptable variation percentage between the test result and the previous known value (anteriority) for each specific test. The RCV represents an essential component of the interpretation of a variation (analytical and intra-individual variation). It is calculated using the following formula (see the bibliographical references in section "[Valab® biological bibliography - Extract](#)"):

$$RCV = \sqrt{2} * 1.96 * \sqrt{(CV\% \text{ reproducibility analytics}^2 + CV\% \text{ intra individual}^2)}$$

The RCV's provided with Valab® can be individually consulted and modified for each test. They have been established on the basis of bibliographic research (see the bibliographical references in section "[Valab® biological bibliography - Extract](#)").

The actual acceptability of a variation is modulated by the expert analysis of the data contained in the report which generates a dynamic RCV adapted to the clinical situation of the patient.

Valab® proposes to program a maximum validity period for the previous result for each test. If this limit is exceeded, the previous result is ignored by the autoverification.

"Critical" Delta check / RCV

In order to control the effects of the dynamic RCV, the setting of a "critical" maximum delta is also proposed to the user. It introduces the concept of the highest variation above which a result cannot be validated by Valab® and the report will be submitted to the clinical scientist. Different "critical" maximum amplitude values can be defined both for an increase and a decrease. This criterion can be activated or deactivated for each test.

Sensitivity of the autoverification

Setting the sensitivity parameter allows the user to fine-tune the autoverification of each test by acting on the weight of the rules without modifying the internal logic of the system.

There are sensitivity settings for the rules applicable in correlation (for tests without a previous result) and for the rules applicable in anteriority (for tests with a previous result).

Each sensitivity setting can be adjusted (the neutral value is 1) in order to make the system more permissive (value greater than 1) or more restrictive (value less than 1) with regard to the acceptability of a result.

Correspondence tables

Valab®'s correspondence tables allow to match the "symbolic/qualitative" data known by the LIS with the corresponding data in Valab®:

- Table for Prescribers – specialties – departments
- Table for Sexes
- Table for Clinical Information about the patient
- Table for Complementary Information - technical
- Tables for Quantitative tests / alarms or analyser measurement limits
- Tables for Qualitative tests (symbolic tests)

Users enter information into Valab®'s correspondence tables using the data extracted from their LIS.

The "Reject unknown labels" option can be activated independently for each Valab® correspondence table and allows users to set up Valab® so that it "blocks" all of the reports that are sent to it containing "symbolic/qualitative" data that is not identified in its correspondence tables. This allows to ensure that Valab® will not validate reports containing "symbolic/qualitative" data that it does not know sent to it by the LIS.

Management of user access rights

Access to Valab® is protected by a unique login and password combination for each user account. A user account must be created for each person authorised to have access to Valab®.

LDAP directory connection

It is possible to connect Valab® to your LDAP directory.

Sessions and user accounts

At Valab® startup, no user session is opened in order to avoid unwanted actions. Access to any Valab® functionalities is only possible after the opening of a session. Each access is associated to a user account with a password and customizable rights.

If no action is performed after 5 minutes (default setting), the session is closed automatically.

The sessions and the user accounts also allow to manage the traceability of all the actions made in the configuration of the system.

Different user types

An administrator of the accounts allocates to the various users of Valab® the access profiles to the functionalities (consultation and / or setting rights) adapted to their level of intervention.

Protected functions

All the functionalities are protected in consultation and setting. When a user opens a session he only has access to the functionalities in consultation and /or setting which are adapted to his level of intervention defined by the administrator in his profile access.

Confidentiality and archiving

An option coupled with a right level allows to mask the patients' names from the headers of the reports received from the LIS. On the other hand, only users with a sufficient level access can consult patients' reports expertised by Valab® and the backlog associated to connection exchanges between Valab® and the LIS.

For reasons of traceability and/or data processing in simulation activities, local archiving functions are available for a temporary period that can be set (maximum of 24 months) and deactivated by the user. These files containing sensitive data are encrypted in 128-bit AES by Valab®. The traceability and archiving of the patient reports is ensured elsewhere by the LIS of the medical laboratory and by the implementation of regular backups.

Traceability of user access and modifications

A logbook allows to keep a history and traceability of all the modifications of parameter settings performed by the user.

The logbook records:

- all the modifications made to the parameter settings
- all of the user logons / logouts
- all of the patient report consultations and their associated trace histories if applicable
- the date, the time of day and the user account from which actions have been performed
- previous values before modification

Backup and restore

The document [RD5](#) "Valab® - Backup and Restore (available at www.valab.com)", provides all necessary details to implement the backup and restoration of Valab® computer data.

We recommend that you keep a copy of the backups on another site of the medical laboratory (remote backups).

Activity statistics

The use of Valab® generates a set of statistical indicators for the autoverification results:

- number and percentage of reports validated
- number, percentage and cause of refusal for each test (correlation, anteriority, biological reference intervals or clinical decision values, "alert" or "critical" limits)
- number and percentage of validation for each test

This data is archived to create a history and a database which can be exploited a posteriori and during the entire period of activity of the system.

It allows in particular the implementation of continuous monitoring procedures by watching out for a drift between successive observation periods, for example month by month (see [RD2](#)).

There are also alarms, with a threshold that can be defined by validation rate and the reason for refusal for each test, to highlight those with a validation or refusal rate that is too high. This helps to customise the fine-tuning of the system by the user.

BI module ValView

This optional module offers 8 key indicators grouped in 4 dashboards to help you further the efficiency of Valab®, facilitate the continuous monitoring and improve your control over Valab®:

- Dashboard – *Laboratory activity*: contains two indicators to visualize Valab®'s global activity:
 - a graph showing the evolution of global statistical indicators over a defined period (number of reports received, examined and validated per day, month, quarter or year)
 - a graph showing the hourly flow of runs submitted to Valab® by the LIS on a given day
- Dashboard – *Interpretation and reference*: contains three statistical indicators that enable standardized interpretation and continuous monitoring over a defined period with daily, monthly, quarterly, or yearly granularity:

- a table listing global statistical indicators and their drifts (autoverification rate, validation rate and reject rate (by type))
 - two graphs showing the evolution of autoverification rate and validation rate relative to the respective reference rate defined by the ML
- Dashboard – *Top blocked tests*: contains two indicators identifying tests with the highest rejection rates and those causing the most laboratory report blockage over a defined period with daily, monthly, quarterly, or yearly granularity:
 - a table listing the most frequently rejected tests, by type of rejection (correlation, anteriority or critical values)
 - a table listing the most frequently rejected tests among the most commonly performed tests. The definition of “most common” is user-configurable
- Dashboard – *Drift per test*: contains an indicator that enables identification and standard interpretation of most commonly performed tests with the highest validation rate drift over a defined period with daily, monthly, quarterly, or yearly granularity. The definition of “most common” is user-configurable:
 - a table lists the upward drift and the other table the downward drift

VALAB Services Provider (VSP)

License and services server of the VALAB company.

If you own a Valab® user license of VSP type, your license is hosted by the VSP.

This optional hosting of your license by the VALAB company out of your information system offers the following advantages:

- Allows 12 days of overruns per year of the number of reports/day for the level of your license. Each overrun day allowed permits to process up to the double of the reports/day limit of your license (e.g. useful the day after a public holiday)
- Allows for the monitoring of the basis activity of your Valab® by the customer Support of the VALAB company and/or by the support of your distributor for a better reactivity when applicable
- Is unfazed by material or OS updates and compatible with the servers farms or high availability clusters
- Eases the activation of complementary options for your user license (license amendment, short or long term rental amendments...)

Documentation of Valab® and its interfaces

Among the documentation provided with Valab®, there are reference documents to ensure that the user maintains control of the tool and its interfaces and describing the means for monitoring to be implemented:

- [RD3 – “Valab® User Manual \(provided in Valab® "DOC" subdirectory\)”](#): Describes all the functionality of the software that the user needs to know to understand the operation of Valab® and be proficient in its use;
- [RD4 – “Valab® Developer Manual \(provided in Valab® "DOC" subdirectory\)”](#): Presents all the technical characteristics and organisational recommendations to be taken into account by LIS companies to help them develop a connection and integration of Valab® in conformity with our requirements;
- [RD2 – “Qualification of Valab® by the Medical Laboratory \(available at \[www.valab.com\]\(http://www.valab.com\)\)”](#): Provides a guide describing an approach and a set of example procedures to set up the use of Valab® in the medical laboratory and to ensure its control in terms of continuous monitoring, maintenance and tracing of modifications :

- Qualification of the connection and integration with the LIS
- Initial qualification of the autoverification and of the setup customisation
- Requalification after modification
- Continuous monitoring: activity statistics, sampling of patient reports, pool of typical reports, maintenance
- Impact on the medical laboratory's QMS documentation
- Awareness and training of users

Each new major version of Valab® has its own updated User Manual and Developer Manual.

Training of users

All laboratory personnel who use the Valab® software must be trained on how it works and how to use it. For this purpose, when Valab® is installed, the future key-contact users of the system are trained by an agent of the VALAB company.

The term "user" must be understood in a broad sense, to cover not only the direct users of Valab® (key contact staff authorised to set up, check or otherwise interact with the software), but all the laboratory staff validating reports containing tests connected to Valab®.

Training is recommended both for new users and for existing users when they update to a new major release of Valab®.

Bibliography

Valab® biological bibliography - Extract

Click the following link to [consult the Valab® biological bibliography - Extract on www.valab.com](http://www.valab.com).

Bibliography and press articles on Valab®

Click the following link to [consult the press articles on Valab® on www.valab.com](http://www.valab.com).