

Survey: BI4/25
month: October 2025
Participant-No: 0004965
valid from: 11.10.25



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diaglobal GmbH
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Bonn, 29. October 2025

Certificate

We confirm that you have participated in the survey for neonatal bilirubin in serum.
This certificate is according to the current version of the RiliBÄK valid until the end of April 2026.
You have met the requirements of the survey for the following analytes:

Total bilirubin (2)

Three handwritten signatures in blue ink. From left to right: "K. Kohse", "A. Kessler", and "M. Enders".

Prof. Dr. Dr. K. P. Kohse
EQAS scheme director

Dr. Anja Kessler
Head of Reference Institute

Dr. Marika Enders
EQAS-Board

The number in parentheses characterizes the analytical method used.
The assignment of the number to the respective method and/or the respective instrument
is to be taken from the total evaluation.

This certificate is valid in conjunction with the final report dated 29.10.25.
This is available for download in the Rfb-Online system.



Listing and Evaluation of all your results

Explanations

Certificate

A certificate is issued (given) for an analyte only if the basis for an evaluation of the accuracy is given by the guidelines of the German Medical Association and/or if an evaluation is possible in analogy to these guidelines (see comments on the evaluation) and, if both results for an analyte are within the given acceptance limits. (marked as '+' below C)

Certificate of participation

In the participation certificate all analytes which are included in the list on this page are listed.

If all analytes are listed on the certificate no participation certificate is printed.

Legend:	C=Certification, M=No of method, R=your result, D=difference (R-T) Dmax= maximum allowable amount of difference in measurement, partly given by annex 1 of the guideline of the BÄK. T = target value, either reference method value or assigned value, LL UL = lower resp. upper limit	Certification:	+ = fulfilled (quotient D/Dmax <= 1.0) - = not fulfilled (quotient D/Dmax > 1.0) ± = certification cancelled because of technical and/or analytical reasons
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	C	M	R	D/Dmax	T	LL	UL						
Total bilirubin [mg/dl]	+	2	A B	12.4 4.94	0.05 -0.02	12.3 4.96	9.55 3.86	15.0 6.06					