

# CERTIFICATE OF ANALYSIS

Product name	XN CHECK
Code No.	1101 (Level L1)
Lot No.	4189
Exp. date (YYYY-MM-DD)	2024-10-06
Date of release* (YYYY-MM-DD)	2024-07-03

Item of analysis and testing	Acceptance criteria	Results (pass/fail)
Appearance	Red liquid	pass
Assay value	Specifications fulfilled (For details refer to assay sheet**)	pass

\* Date of Release is the date upon which the product manufacturing process terminates and the distribution process starts.

\*\* Please choose the assay sheet for your analyser type

This product was manufactured under FDA QSR, ISO 9001/ISO 13485 and CMDR conditions.

This is to certify that the above product has been produced and passed the inspection according to the specifications of Sysmex Europe SE.

Sysmex Europe SE  
C. Dahmen  
Product Management Quality Control

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# CERTIFICATE OF ANALYSIS

Product name	XN CHECK
Code No.	1102 (Level L2)
Lot No.	4189
Exp. date (YYYY-MM-DD)	2024-10-06
Date of release* (YYYY-MM-DD)	2024-07-03

Item of analysis and testing	Acceptance criteria	Results (pass/fail)
Appearance	Red liquid	pass
Assay value	Specifications fulfilled (For details refer to assay sheet**)	pass

\* Date of Release is the date upon which the product manufacturing process terminates and the distribution process starts.

\*\* Please choose the assay sheet for your analyser type

This product was manufactured under FDA QSR, ISO 9001/ISO 13485 and CMDR conditions.

This is to certify that the above product has been produced and passed the inspection according to the specifications of Sysmex Europe SE.

Sysmex Europe SE  
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Product Management Quality Control

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# CERTIFICATE OF ANALYSIS

Product name	XN CHECK
Code No.	1103 (Level L3)
Lot No.	4189
Exp. date (YYYY-MM-DD)	2024-10-06
Date of release* (YYYY-MM-DD)	2024-07-03

Item of analysis and testing	Acceptance criteria	Results (pass/fail)
Appearance	Red liquid	pass
Assay value	Specifications fulfilled (For details refer to assay sheet**)	pass

\* Date of Release is the date upon which the product manufacturing process terminates and the distribution process starts.

\*\* Please choose the assay sheet for your analyser type

This product was manufactured under FDA QSR, ISO 9001/ISO 13485 and CMDR conditions.

This is to certify that the above product has been produced and passed the inspection according to the specifications of Sysmex Europe SE.

Sysmex Europe SE  
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