

Gesellschaft für Biochemica und Diagnostica mbH

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Urgent Field Safety Notice

New Calibrator values of AUTOCAL LOT 0019 for CHOLESTEROL liquicolor and revised Kit Standard values for CHOLESTEROL liquicolor Recall

October 16th, 2025

Attention:

Distributors of HUMAN and end users of:

Details on affected devices:

Product Name	REF	LOTs including all sublots
AUTOCAL	13160	0019

when used to calibrate the following CHOLESTEROL reagent kits.

Product Name	REF	Size
CHOLESTEROL liquicolor	10028300	2 x 100 tests
	10028600	3 x 150 tests
	10028	4 x 100 mL
	10017	4 x 30 mL
	10019	3 x 250 mL

And the kit standards included in the following reagent kits:

Product Name	REF	Size
	10028	4 x 100 mL
CHOLESTEROL liquicolor	10017	4 x 30 mL
	10019	3 x 250 mL

Description of the problem:

In the course of internal quality assurance activities, we noted that the concentration for CHOLESTEROL liquicolor in AUTOCAL REF 13160 LOT 0019 has increased over time. Based on this observation the usage of AUTOCAL LOT 0019 in combination with the CHOLESTEROL liquicolor reagent leads to falsely low Cholesterol values in patient samples up to 12% These discrepancies occur when the above-mentioned Cholesterol reagents are calibrated either with AUTOCAL LOT 0019 or with the kit standard included in the CHOLESTEROL liquicolor kits.

As a measure the Cholesterol value in AUTOCAL LOT 0019 was revised as indicated below:

Product	REF	LOT	Old concentration	New concentration	Deviation
AUTOCAL	13160	0019	311 mg/dl	348 mg/dl	12 %

FORM 7.2-02.04-02

Vers. 06

Gültig ab: 29.09.2022



As a consequence, the Cholesterol values for corresponding kit standards were revised as well, as indicated below:

Kit standard	Kit REF	Kit LOT	Old	New	Deviation
LOT	TXIC IXE	Tall 201	concentration	concentration	in %
			in mg/dL	in mg/dL	/0
	10017	24006	3 -	3 -	
0.400	10017	24007	405	214	10
0120	40000	24010	195		
	10028	24011			
	10017	24008			10
	10017	24009			
0121	10019	24002	195	214	
	10028	24012			
	10020	24013			
	10017	24010			
		24011			
0122	40000	24014	196	213	9
	10028	24015			
	10017	24016 24012			
	10017	24012	-		
0123	10019	24003	195	216	11
	10028	24017			
	10017	25001			
	10017	25001	-	214	10
0124	10019	24019	194		
		25001			
		25002			
	10017	25002		215	11
0125		25003	193		
	10028	25003			
	10017	25004		216	11
		25005			
0126	10019	25002	193		
0126	10028	25004	193		
		25005			
		25006			
0127	10017	25006	193	212	10
	10028	25007			
		25008			
	40047	25009			
	10017	25007	-	215	12
0128	10019	25003	- 192		
		25010			
		25011			



Additionally, the target values and ranges for the HUMAN Quality Controls Serodos, Serodos plus, HumaTrol N and HumaTrol P were revised and are accessible on https://www.human.de/

Advice on action to be taken by:

Distributor:

Please inform your customers about the issue of the affected Lots and the revised cholesterol values. Please fill in the attached Reply Form confirming receipt of this Urgent Field Safety Notice and send it to support@human.de

User:

Please use the revised cholesterol values from now on and ensure that the instructions resulting from this Urgent Field Safety Notice are implemented in the laboratory accordingly. End users should confirm receipt of this Urgent Field Safety Notice to the local distributor

Transmission of this Urgent Field Safety Notice:

This notice needs to be passed on all those who need to be aware within your organization or to any organization where the potentially affected devices have been transferred.

Please maintain awareness on this notice and resulting action for an appropriate period to ensure effectiveness of this corrective action.

The Federal Institute for Drugs and Medical Devices (BfArM) and the National Competent Authorities of European countries which are affected by the recall have received a copy of this Urgent Field Safety Notice.

Contact reference person:

(For distributors only. Distributors should provide their own detailed contact information to their end users):

Petzold, Tony

e-mail: support@human.de Telephone: +49-6122-9988-383

We regret the inconvenience.

With kind regards,

Petzold, Tony

Customer Support & Applications

Dr. Märker-Stilz, Tina

Product Management Clinical Chemistry

Attachment Reply Form

Gültig ab: 29.09.2022



Reply Form

Urgent Field Safety Notice

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Please return by e-mail this filled in and signed Reply Form latest until October, 26, 2025 to:

support@human.de

Date:

I confirm receipt of this Urgent Field Safety Notice and have informed all end users, who have obtained the affected products, in writing about the problem and the HUMAN recommendations.

If requested by national regulations, I have informed the respective authorities about the problem. (Note: to comply with European regulatory requirements HUMAN will inform European competent authorities directly.)

For distributors in the European Economic Area (EEA) and Turkey:

Please also provide the Urgent Field Safety Notice in your national language, which you have sent out to your end customers, as HUMAN will be approached by your national competent authority to provide this.

Company:	
Name:	Signature: