



# DIACHEM Ltd

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# COMPLEMENT C3

## LIQUID REAGENT



Cat. No.: 31430 31432  
4x25 ml 10x25 ml  
(4x20 ml+ 4x5ml) (10x20 ml+ 10x5ml)

### Reagent kit for immuno-turbidimetric determination of Complement C3 in human serum.

#### Principle

Measurement of antigen/antibody reaction by the endpoint method.

#### Reference values

**Serum: 75- 135 mg/dl**

It is recommended that each laboratory should assign its own normal range.

#### Reagents

##### 1. Reagent 1 (R1)

buffer  
sodium-azide (0,50 g/l)

##### 2. Reagent 2 (R2)

antisera  
sodium-azide (0,50 g/l)

#### Precaution

Discard cloudy reagent. Avoid contamination by using clean laboratory material (pipettes, plastic vials for analyzers,... ).  
The reagents contain sodium azide. To avoid the possible build-up of azide compounds, flush waste-pipes with water after the disposal of undiluted reagent.

#### Sample

Use fresh serum. The serum can be stored at 2-8 °C for 48 hours. If stored for a longer period, the sample should be frozen.

### PROCEDURE

The reagents are ready for use.

#### Assay conditions

Wavelength: 340 nm  
Temperature: 37 °C  
Cuvette: 1 cm light path  
Read against: sample blank  
Method: endpoint (increasing)

#### Pipette into cuvette

	Blank	Standard	Sample
1. reagent (R1)	360 µl	360 µl	360 µl
Standard	3 µl or	3 µl	
Sample	3 µl		3 µl

Mix, wait 1 minute then add:

2. reagent (R2)		90 µl	90 µl
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Mix and after 7 minutes incubation read the absorbance against the sample blank.

#### Calibration:

5 level (at the least 3) standard set is necessary for calibration.

S1: Distilled water or physiological salt solution  
S2-S6: 5 level standard set

New calibration is recommended:

- after reagent lot change,
- as required following quality control procedures.

#### Calculation

Based on the calibration curve.

#### Quality control

A quality control program is recommended for all clinical laboratories. The analysis of control material in both the normal and abnormal ranges with each assay is recommended for monitoring the performance of the procedure. Each laboratory should establish corrective measures to be taken if values fall outside the limits.

### PERFORMANCES DATA

The following data were obtained using the Olympus AU600 and Advia 1650 analysers (37°C).

#### Linearity

The test is linear up to 400 mg/dl (4 g/l).

#### Sensitivity

It is recommended that each laboratory establishes its own range of sensitivity as this is limited by the sensitivity of the spectrophotometer used.  
The estimated value of limit of detection (LOD) is 0,3 mg/dl.

#### Precision

Sample	Reproducibility		
	Average concentration (g/l)	SD	CV%
sample I	1,34	0,12	9,30
sample II	1,20	0,08	6,96
sample III	1,07	0,07	6,22

Sample	Repeatability		
	Average concentration (g/l)	SD	CV%
sample I.	1,64	0,01	0,50
sample II.	1,34	0,01	0,72
sample III.	0,96	0,01	0,62

#### Correlation

Comparative studies were done to compare our reagent with another commercial Complement C3 reagent. The results from these studies are detailed below.

Correlation coefficient:  $r=0,9924$   
Linear regression:  $y (g/l)= 1,034x + 0,055$   
(x= other commercial reagent, y= own reagent).

#### Specificity

Bilirubin 20 mg/dl, triglicerydes 2500 mg/dl, haemoglobin 1000 mg/dl don't interfere with the assay up to the given levels.

#### NOTE

Do not use reagents after the expiry date stated on each reagent container label. Do not use products, test solutions and reagents described above for any purpose other than described herein.

**For in vitro diagnostic use only.**

**The following symbols are used on labels**

For in vitro diagnostic use

Use by (last day of the month)

Temperature limitation

Batch Code

Code

#### Bibliography

Dati, F. et al., Lab. Med. 13, 87 (1989)