Step by step instructions

TRI 742 (TRI / TRI conc.)

Single measurement



1. Insert capillary with 1 μ L sample into cuvette



2. Eject sample several times with micropipetter into cuvette



3. Screw cap on Turn cuvette upside down several times



4. Switch photometer on with ON/ENTER key Wait for device check, confirm with ON/ENTER Select the required test, confirm with ON/ENTER



5. Zero point adjustment: Insert cuvette with sample (Fig. 3) into photometer, zero point is stored in memory Remove cuvette after signal

tone



6. Replace screw cap with starter cap



7. Turn cuvette upside down several times



8. First press ON/ENTER Then insert cuvette into photometer



9. Time is displayed, wait for measured value

Step by step instructions

TRI 742 (TRI / TRI conc.)



Serial measurement

Number of samples per series: Up to 20 samples at the same time



1. Eject all samples one after the other several times with micropipetter into cuvette



2. Screw all caps on again Turn cuvettes upside down several times

N4 1	
M1 Sample 01 insert	IRI
	$ \rightarrow$

3. Switch photometer on with ON/ENTER key Wait for device check, confirm with ON/ENTER Select the required test, confirm with ON/ENTER



4. Zero point adjustment: Insert cuvettes with samples (Fig. 2) one after the other into photometer, all zero points are stored in memory Note: Ensure the correct order of the samples!



5. After the zero point adjustment of the last cuvette replace all screw caps with starter caps



6. Turn all cuvettes simultaneously upside down, repeat several times



7. First press ON/ENTER key Then insert 1st cuvette into photometer Time is displayed, wait for measured value



8. Read the measured value of the 1st cuvette, remove cuvette Insert 2nd cuvette, read the measured value, remove cuvette, and so on

M2	TRI
Sample 19	
5.28 g/dL	
	\rightarrow

9. Insert the last cuvette, read the measured value, remove cuvette Note: Ensure the correct order of the samples!