

Sodium Single Reagent

REF: 303 001 (2 x 25 ml) 50 test
 REF: 303 002 (4 x 25 ml) 100 test
 REF: 303 003 (2 x 100 ml) 200 test
 REF: 303 004 (2 x 500 ml) 1000 test

Intended Use

Spectrum-Diagnostics Sodium reagent is intended for the in-vitro quantitative diagnostic estimation of sodium in human serum on manual systems.

Background

Sodium and Potassium are the major cations of extracellular and intracellular fluids respectively. Sodium maintains the normal distribution of water and the osmotic pressure in the various fluid compartments. Potassium influences the acid base balance and osmotic pressure including water retention. Increased sodium levels are found in severe dehydration and excessive treatment with sodium salts. Decreased levels are found in severe polyurea, metabolic acidosis, diarrhoea and renal insufficiency. Increased potassium levels are found in renal failure, dehydration, shock and adrenal insufficiency. Decreased levels are found in malnutrition, gastrointestinal fluid loss, and hyperactivity of the adrenal cortex.

Method

Colorimetric method.

Assay Principle

The Present method is based on reaction of sodium with a selective chromogen producing a chromophore whose absorbance varies directly as the concentration of sodium in the test specimen.

Reagents

Reagent (R) Color Reagent

Chromogen	0.03 gm/L
EDTA	25 mmol/L
Dimethyl sulfoxide (DMSO)	75 mmol/L
preservatives	0.05%
Antifoam	0.01 %

Standard (S) Sodium 150 mEq/l

Precautions and Warnings

Do not ingest or inhale. In case of contact with eyes or skin; rinse immediately with plenty of soap and water. In case of severe injuries; seek medical advice immediately.

Reagent Storage and Stability

Reagents and standard are ready-to-use. When stored at 15 - 25°C; they are stable up to the expiry date stated on the label. The remaining stability after opening the bottles is 1 month at 15 - 25 °C.

Sample

Freshly drawn non-hemolysed serum is the specimen of choice. Heparinised plasma can also be used. Serum Sodium is stable for at least 24 hours at room temperature and two weeks at 2-8°C. Urine diluted 1+1 with distilled water can be used for Sodium estimation.

SYMBOLS IN PRODUCT LABELLING

	Authorised Representative		Temperature Limitation
	For in-vitro diagnostic use		Use by/Expiration Date
	Batch Code/Lot number		CAUTION. Consult instructions for use
	Catalogue Number		Manufactured by
	Consult instructions for use		

System Parameters

Wavelength	630 nm
Optical path	1 cm
Assay type	colorimetric end-point
Direction	Increase
Sample: Reagent Ratio	1:100
e.g.: Reagent volume	1 ml
Sample volume	10 µl
Temperature	Room temperature
Zero adjustment	Against reagent blank
Linearity	180 mEq/l
Incubation	5 min.
Blank absorbance limit	1.2

Procedure

Pipette into clean test tubes:

	Blank	Standard	Sample
Reagent (R)	1 ml	1 ml	1 ml
Standard	10 µl
Sample	10 µl

Mix well, let stand for 5 minutes at R.T.

Read absorbances ,A standard and A sample against Reagent Blank at 630 nm.

Calculation

$$\text{Serum Sodium Conc. (mEq/l)} = \frac{A_{\text{Sample}}}{A_{\text{Standard}}} \times 150$$

Expected Values

Serum 135 – 150 mEq/l.

Note:

It is recommended for each laboratory to establish and maintain its own reference values. The given data are only an indication.

Spectrum Diagnostics does not interpret the results of a clinical laboratory procedure; interpretation of the results is considered the responsibility of qualified medical personnel. All indications of clinical significance are supported by literature references.

Sensitivity

When run as recommended, the minimum detection limit of the assay 55 mEq/l.

Linearity

The assay is linear up to 180 mEq/l.

Waste Disposal

This product is made to be used in professional laboratories. Please consult local regulations for a correct waste disposal. **S56:** dispose of this material and its container at hazardous or special waste collection point.

S57: use appropriate container to avoid environmental contamination.
S61: avoid release in environment. refer to special instructions/safety data sheets.

References

1. Tietz, N.W., Fundamentals of clinical Chemistry, W.b. Saunders Co. Phila, P.A. p. 874.
2. Henry R.F., et, al, Clinical Chemistry Principles and Technics. 2nd Ed, Harper and Row, Harper and Row, Hargersein, M.D. (1974)
3. Maruna RFL., Clin Chem. Acta. 2:581, (1958)
4. Trinder, P:Analyst, 76:596, (1951)

ORDERING INFORMATION	
CATALOG NO.	QUANTITY
303 001	2 x 25 ml 50 Test
303 002	4 x 25 ml 100 Test
303 003	2 x100 ml 200 Test
303 004	2 x500 ml 1000 Test



Egyptian Company for Biotechnology (S.A.E)

Obour city industrial area. block 20008 piece 19 A. Cairo. Egypt.

Tel: +202 4489 2248 - Fax: +202 4489 2247

www.spectrum-diagnostics.com

E-mail:info@spectrum-diagnostics.com



MDSS GmbH
Schiffgraben 41
30175 Hannover, Germany



IFUFCC59

Rev.(7), 23/2/2019