

### TARTARIC ACID

Colorimetric – Method RX ALTONA MANUAL FOOD AND WINE

#### **INTENDED USE**

For the quantitative determination of Tartaric Acid in food and wine. This product is suitable for manual use and on the Rx altona analyser. Applications for a variety of additional analysers are available from www.randoxfooddiagnostics.com.

# FOR THE ANALYSIS OF FOOD AND WINE. Not for diagnostic procedures.

### Cat. No.

 TK4060
 R1.
 Buffer
 I x 60 ml

 R2.
 Chromogen
 I x I5 ml

 CAL
 Standard
 I x 5 ml

 Decolourant
 I x 5 ml

#### **SIGNIFICANCE**

Tartaric Acid occurs naturally in grapes and is one of the most prevalent organic acids, along with L-Malic acid, present in wine. Its concentration is decisive in the definition of the acidity of wines and therefore affecting wine characteristics such as odour, colour and flavor. The determination of tartaric acid is also of interest in stability studies, since it forms complexes with potassium and calcium and the resulting precipitate causes undesirable deposits in the bottle.

### **PRINCIPLE**

This end-point colorimetric test is based on the principle that at acidic pH, tartaric acid (tartrate) reacts with vanadate to produce a coloured complex (metapervanadyl tartrate) which is measured spectrophotometrically at 480-520 nm.

### **SAMPLE**

Wine. Turbid samples should be filtered prior to assay. Decolourant supplied should be added to all coloured samples as outlined in procedure section. Sparkling wine samples should be degassed before analysis.

### SAFETY PRECAUTIONS AND WARNINGS

For the analysis of food and wine. Not for diagnostic procedures. Do not pipette by mouth. Exercise the normal precautions required for handling laboratory reagents.

Solutions R2 and CAL contain Sodium Azide. Avoid ingestion or contact with skin or mucous membranes. In case of skin contact, flush affected area with copious amounts of water. In case of contact with eyes or if ingested, seek immediate medical attention.

Sodium Azide reacts with lead and copper plumbing, to form potentially explosive azides. When disposing of such reagents, flush with large volumes of water to prevent azide build up. Exposed metal surfaces should be cleaned with 10% sodium hydroxide.

Health and Safety data sheets are available on request.

Please dispose of all Biological and Chemical materials according to local guidelines.

The reagents must be used only for the purpose intended by suitably qualified laboratory personnel, under appropriate laboratory conditions.

### STABILITY AND PREPARATION OF REAGENTS

#### RI. Buffer

Contents ready for use. Stable up to the expiry date specified when stored at +2 to  $+8^{\circ}$ C.

### R2. Chromogen

Contents ready for use. Stable up to the expiry date specified when stored at +2 to +8°C.

### CAL. Standard

Contents ready for use. Stable up to the expiry date when stored at +2 to  $+8^{\circ}$ C.

### Decolourant

Contents ready for use. Stable up to the expiry date when stored at +2 to +8°C.

### **MATERIALS PROVIDED**

Buffer Chromogen CAL Decolourant



### **RX ALTONA (FOOD AND WINE) TK4060**

### **RX ALTONA PROCEDURE**

Select Tartaric Acid in the Test Screen. Then select Run Calibration or Run Sample and carry out a water blank as instructed.

### Calibration and White Wine Samples

Pipette into cuvette:			
	S0	SI	White Wine Sample
Sample			40 µl
CAL dil I + 4 (S0)	40 µl		
CAL (SI)		40 µl	

Mix, and incubate for 3 minutes at  $+25^{\circ}$ C. Insert the cuvette into the RX altona flowcell holder when prompted for Sample Blank and press Read. Then add

600 µl

600 µl

600 µl

Chromogen (R2)	150 μΙ	150 µl	150 µl
<b>O</b> ( )	•	•	•

Mix, and incubate for 5 minutes at +25°C. Insert the cuvette into the RX **altona** flowcell holder when prompted for Sample and press Read.

### **Coloured Samples**

Buffer (RI)

Pipette into cuvette	:	
	Red Wine	
	Sample	
Sample	40 µl	
Decolourant	50 μl	
Mix, and incubate for	or I minute at +25°C. Then add	
Buffer (RI)	550 μΙ	

Mix, and incubate for 3 minutes at  $+25^{\circ}$ C. Insert the cuvette into the RX altona flowcell holder when prompted for Sample Blank and press Read. Then add

Chromogen (R2)	I 50 μΙ	

Mix, and incubate for 5 minutes at  $\pm 25^{\circ}$ C. Insert the cuvette into the RX **altona** flowcell holder when prompted for Sample and press Read.

**NOTE:** Bubbles may form in cuvette when completing the coloured samples procedure. User should flick the side of cuvette a few times to disperse these before reading in the analyser.

### **CALIBRATION FOR RX ALTONA**

The use of CAL used neat (as S1) and diluted I+4 (as S0) with double distilled water is recommended for calibration. A 2 point linear calibration is recommended with change in reagent lot or as indicated by quality control procedures.

## FOR MANUAL USE PROCEDURE SEMI MICRO

Wavelength:	520 nm (480-520 nm)
Cuvette:	I cm path length
Temperature:	+20 to +25°C

### **Calibration and White Wine Samples**

Pipette into I ml cuvette

	S0	SI	White Wine Sample
Sample			80 µl
CAL dil I + 4 (S0)	80 µl		
CAL (SI)		80 µl	
Buffer (RI)	1200 µl	Ι200 μΙ	Ι 200 μΙ
Mix, and read absort	ance AI after	3 minutes at +	25°C. Then add
Chromogen (R2)	300 µl	300 µl	300 µl

Mix, and read absorbance A2 after 5 minutes at +25°C.

### **Coloured Samples**

Pipette into I ml cuvette

-	
	Red Wine
	Sample
Sample	80 µl
Decolourant	Ι00 μΙ
Mix, and incubate for	or I minute at +25°C. Then add
Buffer (RI)	Ι Ι 00 μΙ
Mix, and read absor	bance AI after 3 minutes at +25°C. Then add
Chromogen (R2)	300 µl

Mix, and read absorbance A2 after 5 minutes at +25°C

### **MANUAL CALCULATION**

Calculate A2 – A1 for all standards/samples ( $\triangle A$ ).

Plot a linear calibration graph using  $\Delta A$  standards (x axis) versus Standard Concs. g/L (y axis).

Use the linear regression equation achieved (y = mx + c) to calculate the tartaric acid concentration for samples, where y = concentration (g/L) and  $x = \Delta A$  sample.

### SPECIFIC PERFORMANCE CHARACTERISTICS

The following performance characteristics were obtained using a RX **altona** analyser in cuvette mode at +25°C.

### **LINEARITY**

The tartaric acid assay is linear to 10 g/l.

### **SENSITIVITY**

The minimal detectable concentration of tartaric acid with an acceptable level of precision was determined as  $0.51\ g/l$ .



### **PRECISION**

Intra assay prec	ision		
	Level I	Level 2	Level 3
Mean (g/l)	2.82	4.68	8.24
S.D	0.045	0.044	0.151
C.V(%)	1.59	0.94	1.83
n `´	20	20	20
Inter assay prec	ision		
Inter assay prec	ision Level I	Level 2	Level 3
Inter assay prec Mean (g/l)		Level 2 4.73	Level 3 8.27
, ,	Level I		
Mean (g/l)	Level I 2.83	4.73	8.27



THIS PAGE IS INTENTIONALLY BLANK