

# Haemoglobin (Ready-To-Use) **Drabkin's Solution**

REF:611 001 (1 x 250 ml) 100 test REF:611 002 (2 x 250 ml) 200 test REF:611 003 (2 x 500 ml) 400 test

#### **Intended Use**

Spectrum Diagnostics haemoglobin reagent is intended for the invitro quantitative, diagnostic determination of haemoglobin in human

# **Background**

Haemoglobin (Hb) is the red pigmented protein located in the erythrocytes and consists of four subunits. Its main function is the transport of oxygen and carbon dioxide in blood. In normal human adults, at least 96 % of the haemoglobin is HbA. HbA2 is usually about 2.5 – 3 % of total haemoglobin. Fetal hemoglobin (HbF predominates during fetal life and diminishes rapidly during the first year of postnatal life. In normal adults less than 1 % is HbF.Blood haemoglobin concentration may be diminished as a consequence of hemorrhage or hemolysis or as a result of impaired blood formation in the bone marrow.

#### Method

Colorimetric method using Drabkin's solution.

# **Assay Principle**

Haemoglobin is oxidized by potassium ferricyanide which is converted into stable cyanomethaemoglobin by potassium cyanide. The absorbance of the cyanomethaemoglobin is monitored at 540 nm.

# Reagents

Reagent

Potassium ferricyanide Potassium phosphate 0.62 mmol/l 1.04 mmol/l Potassium cyanide 1.54 mmol/l

Harmful (Xn): R20/21/22: Harmful by inhalation, in contact with skin and if swallowed. S7: Keep container tightly closed. S28.1: After contact with skin, wash immediately with plenty of water. S45: In case of accident or if you feel unwell, seek medical advice immediately. The amount of cyanide present in one bottle of reagent

is appreciábly less than thé minimum lethal dose for an adult. However, hydrogen cyanide is liberated by acidification. Never allow reagent to come in contact with acid.

For further information, refer to the Haemoglobin reagent material safety data sheet.

# **Precautions and Warnings**

Do not ingest or inhalate. 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.

# Deterioration

Failure to recover control values within the assigned range may be an indication of reagent deterioration.

### Reagent Preparation, Storage and Stability

Reagent is supplied ready to Use. Reagent is stable until expiration date stated on label when stored at 15 - 25 °C.Once opened, the reagent is stable for 3 months at the specified temperature if contamination is avoided.

#### SYMBOLS IN PRODUCT LABELLING

ECREP Authorised Representative 📮 Use by/Expiration Date LOT Batch Code/Lot number REF Catalogue Number Consult instructions for use X (Xi) - Irritant

Temperature Limitation

For in-vitro diagnostic use A CAUTION. Consult instructions for use Manufactured by

# **Specimen Collection and Preservation**

Anticoagulated venous or capillary blood . Blood may be anticoagulated with EDTA, or fluoride. Blood can be taken directly from a finger or heel puncture without use of anticoagulant.

**Stability**: 7 days at 2 – 8 °C 4 days at 20 – 25 °C

#### **System Parameters**

Wavelength 540 nm (Hg 546 nm) Optical path Assay type **End-point** Direction Increase Reagent Ratio Reagent volume Sample: 1:250 2.5 ml e.g .: 10 μl 20 – 25 °C Sample volume

Temperature Incubation time 5 minutes

Against reagent blank Zero adjustment

#### **Procedure**

### Pipette into test tubes

Reagent	2.5 m
Blood sample	10 μ

Mix well and rinse the blood pipette several times with the reagents and incubate for 5 minutes at 20-25 °C. Measure absorbance of specimen (Aspecimen) against reagent blank.

### Calculation

Haemoglobin concentration (g/dL) = Aspecimen x 36.77

Haemoglobin concentration (mmol/L)= Aspecimen x 22.83

# **Quality Control**

Normal and abnormal commerical control blood of known concentrations should be analyzed with each run.

### **Performance Characteristics**

### Precision

Within run (Repeatability)

	Level 1	Level 2
n	20	20
Mean (g/dL)	10	14
SD	0.23	0.182
CV%	2.3	1.3

# Run to run (Reproducibility)

, ,	Level 1	Level 2
n	20	20
Mean (g/dL)	11.1	14.1
SD	0.322	0.296
CV%	2.9	2.1

# **Methods Comparison**

A comparison between Spectrum Diagnostics Haemoglobin reagent and a commercial reagent of the same methodology was performed on 40 human blood samples. A correlation (r) of 0.983 was obtained.

#### **Expected values**

```
15.2 - 23.5 g/dL
10.3 - 16.6 g/dL
10.0 - 12.9 g/dL
     1 – 6 days
14 – 50 days
2 – 10 months
                                                                              ( 9.4 – 14.6 mmol/L)
( 6.4 – 10.3 mmol/L)
( 6.1 – 8.0 mmol/L)
      1 – 15 years
                                   11.0 - 14.3 g/dL
                                                                              (6.8 - 8.8 \, \text{mmol/L})
                                   12.0 - 16.0 g/dL
14.0 - 18.0 g/dL
Adults: Women
                                                                             (7.5 – 9.9 mmol/L)
(8.7 – 11.2 mmol/L)
               Men
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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.

#### **Waste Disposal**

This product is made to be used in professional laboratories. Please consult local regulations for a correct waste disposal.

\$56: dispose of this material and its container at hazardous or special waste collection point.

\$57: use appropriate container to avoid environmental contamination.

**S61:** avoid release in environment. refer to special instructions/safety

## References

- International committee for standardization in haematology.Brit.
   J. Haemat., 1967:13 (Suppl.) 71.
   Van Kampen, E. J. and Zijlstra, W.G., Clin. Chem. Acta.,
   1961:6:538 544.
   Tietz NW, Ed. Clinical guide to laboratory tests. 2ND ED.
   Philadelphia: WB Saunders; 1990:566.

ORDERING INFORMATION		
CATALOG NO.	QUANTITY	
611 001 611 002 611 003	1 x 250 ml 2 x 250 ml 2 x 500 ml	



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