

H8 THALASSAEMIA ANALYSIS MODE

Specifications

Methodology	High Performance Liquid Chromatography(HPLC)
Test Item	HbF, HbA1c, HbA2, Hb Variants(HbE, HbD, HbS, HbC, Hb Ottawa, Hb G-Taipei, Hb Tamano, Hb Maputo, Hb G-Honolulu, Hb G-Coushatta, Hb Q-Thailand, Hb New York, HbJ-Bangkok, Hb J-Wenchang-Wuming, Hb J-Baltimore, Hb H.)
Precision	HbA1c (CV ≤ 1.5%), HbA2 (CV ≤ 4.0%)
Test Speed	6.0 Mins/sample
Sample Type	Whole Blood / Diluted Blood
Sample Volume	Whole blood: 5μL; Diluted Blood: 750μL (Dilution Ratio 15:1500μL)
Auto Sample Loader	10 Positions
Piercing Function	Yes
Sample Pretreatment	Automatic
Photometer	415nm+500nm
Reagent Pack	Eluent A, Eluent B, Eluent C, Hemolysin L, Calibrator, QC Material
Display	10.1 " TFT True Color LCD Touch Screen
Storage	4000 Sample Results
Lis Connection	USB, LAN
Operation Condition	Temperature 10 ~ 30 °C (50 ~ 86 °F); Humidity ≤ 80%
Power	AC 100-240V 50/60HZ 150VA
Dimensions	600mm x 360mm x 540mm (23.6"H x14.2 "W x 21.3" D)
Weight	49KG (108 lbs)



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Lifotronic

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THALASSAEMIA ANALYSIS MODE

A Perfect solution for Thalassaemia Screening

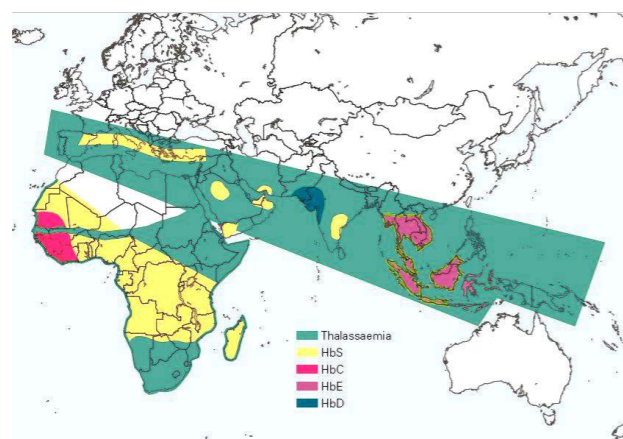


What is Thalassemia?

Thalassaemia is a blood disorder passed down through families (inherited) in which the body makes an abnormal form or inadequate amount of hemoglobin. Hemoglobin is the protein in red blood cells that carries oxygen. The disorder results in large numbers of red blood cells being destroyed, which leads to anemia.

β -thalassaemia trait may develop either thalassaemia major or thalassaemia intermedia. Individuals with thalassaemia major are usually diagnosed within the first 2 years of life and require regular blood transfusions to survive.

Thalassaemia affects approximately 4.4 of every 10,000 live births throughout the world. It causes males and females to inherit the relevant gene mutations equally because it follows an autosomal pattern of inheritance with no preference for gender.



Thalassaemia Distribution in the world

What is the meaning of thalassaemia screening?

Thalassaemia is a blood condition that can cause health problems for your baby. It's passed from parents to children through changes in genes. You and your partner can have carrier screening before or during pregnancy to find out if you have the gene change that causes thalassaemia.

H8 Features

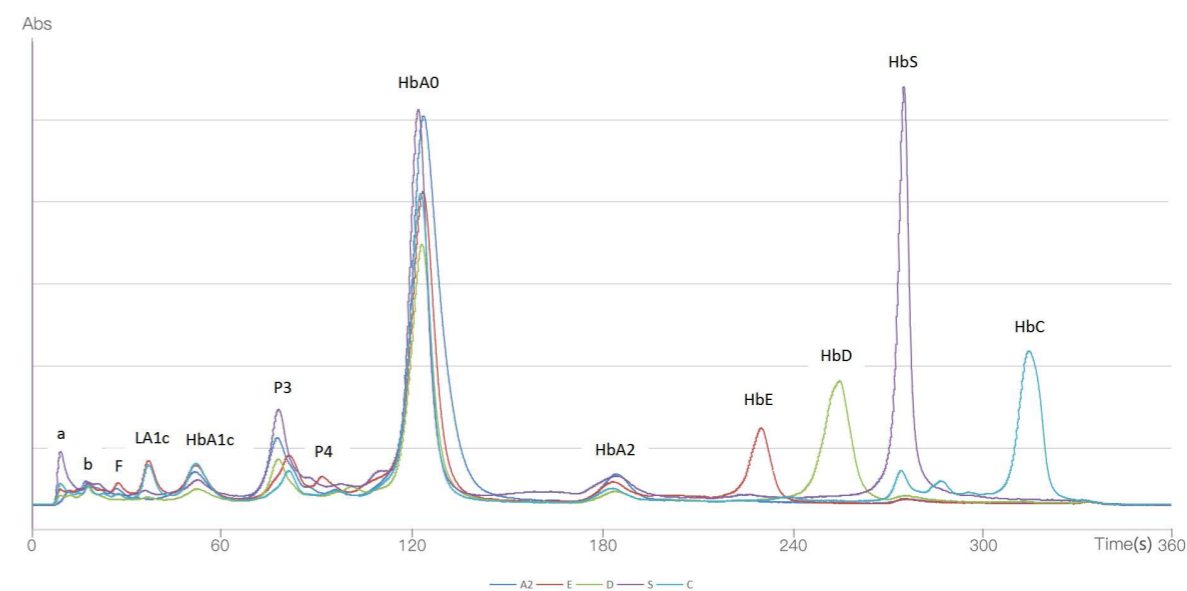
- High Performance Liquid Chromatography(HPLC)
- Report HbF, HbA1c, HbA2 and Hb Variants(HbE, HbD, HbS, HbC)
- Quantitative measure HbA2 and HbF in 6.0 mins
- High resolution chromatography and high reliable system
- Easy to operate and maintenance
- No interference from common hemoglobin variants



About HPLC

High-Performance Liquid Chromatography (HPLC) is the gold standard for hemoglobin testing as it separates mixtures directly and measures the absorbance points continually with superior resolving power. Lifotronic H8, a HPLC thalassaemia analyzer able to report HbF, HbA1c, HbA2 while eliminate interference of such Hb variants as HbE, HbD, HbS and HbC, creates significant and unique values for patients, physicians and laboratories.

H8 Thalassemia Mode Chromatogram



H8 Thalassemia Graph