G6PD Deficiency

Quantase™ Neonatal G6PD* Assay

Newborn Screeners Are Talking
“Simpler is better.”

Bio-Rad Is Listening
Quantase™ newborn screening assays feature streamlined protocols that are readily adapted to any laboratory environment. Whether you are looking for manual or automated solutions, Quantase™ assays provide the efficiency you demand. When you are ready to simplify your workflow, choose Bio-Rad.

Quantase™ Neonatal G6PD Assay
Bio-Rad introduces a convenient and precise assay for the screening of G6PD deficiency. The Quantase™ G6PD Assay utilizes a rapid colorimetric procedure designed specifically for dried-blood spot screening, free of a red blood cell lysis step.

* Not available in the US

The Bio-Rad Assay
• Simplified protocol for use on most microplate systems
• Designed for use with dried-blood spots
• Established ranges for newborn screening
• Room temperature procedure incubator not required
• Numerical results eliminate subjective interpretations
• Rapid results in about 45 minutes
G6PD Deficiency

Clinical Background
Glucose-6-phosphate dehydrogenase, an enzyme found in most body cells, is part of the pentose phosphate pathway that metabolizes glucose. G6PD deficiency is a hereditary, sex-linked condition carried on the X chromosome, with clinical disease found mostly in males. G6PD deficiency impairs the stability of red cell membranes and makes red cells susceptible to destruction by strong oxidizing agents. Affected patients are at risk for hemolytic anemia, which can be induced by exposure to certain drugs and foods.

G6PD is one of the most common hereditary enzyme disorders, affecting over 200 million people worldwide. About 10% of all African American males inherit mild forms of G6PD deficiency. Some people of Mediterranean origin inherit severe forms of the disease. The ingestion of fava beans may also elicit hemolytic episodes in some affected Caucasians.

Ordering Information

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