

# gb GENETIC HFE

## Clinical implications

Hereditary hemochromatosis (HHC) is a hereditary metabolic disorder caused by HFE gene mutation with an autosomal recessive type of inheritance and incomplete penetration (which means it does not have to always be shown in a phenotype). It is one of the most common hereditary disorders with an indicated prevalence of the majority C282Y mutation in a homozygous condition in 1:300–400 in the Caucasian population. HFE protein is bound to beta-2-microglobulin and this complex blocks the transferrin receptor in the cells of the duodenum. There is no receptor blockade in the complex consisting of the mutated form of the protein which results in permanent increased iron absorption from the gastroin-

testinal tract into the blood stream and its accumulation in parenchymatous tissues and organs, especially in liver, pancreas, heart, gonads and in the skin. These organs can then be irreversibly damaged.

## Principle of detection

The kit is intended for detection of mutations C282Y, H63D and S65C in the gene HFE in human genomic DNA. Detection is based on **real-time polymerase chain reaction (qPCR) using fluorescently labelled probes (allelic discrimination)**.

## Available products




| Cat. No. | Product        | rxn |
|----------|----------------|-----|
| 3208-025 | gb GENETIC HFE | 25  |
| 3208-050 | gb GENETIC HFE | 50  |

1 kit contains reagents to provide 25 or 50 PCR reactions (20 µl volume of each reaction).

## Parameters of the diagnostic kit

- *in vitro* diagnostics
- CE IVD marked
- ready-to-use assay
- sample concentration 1-100 ng/µl
- positive and negative controls included
- FAM and HEX channels detection
- identical amplification profile as gb HEMO, gb GENETIC, gb PHARM kits

## Content of the diagnostic kit

| * Component  | Conc.                  | Purpose          |
|--|------------------------|------------------|
|  Assay qPCR HFE C282Y   | 1.25×                  | Detection assay  |
|  Standard WT HFE C282Y  | 10 <sup>4</sup> cop/µl | Positive Control |
|  Standard MUT HFE C282Y | 10 <sup>4</sup> cop/µl | Positive Control |
|  Standard HET HFE C282Y | 10 <sup>4</sup> cop/µl | Positive Control |
|  Assay qPCR HFE H63D    | 1.25×                  | Detection assay  |
|  Standard WT HFE H63D   | 10 <sup>4</sup> cop/µl | Positive Control |
|  Standard MUT HFE H63D  | 10 <sup>4</sup> cop/µl | Positive Control |
|  Standard HET HFE H63D  | 10 <sup>4</sup> cop/µl | Positive Control |
|  Assay qPCR HFE S65C    | 1.25×                  | Detection assay  |
|  Standard WT HFE S65C   | 10 <sup>4</sup> cop/µl | Positive Control |
|  Standard MUT HFE S65C  | 10 <sup>4</sup> cop/µl | Positive Control |
|  Standard HET HFE S65C  | 10 <sup>4</sup> cop/µl | Positive Control |
|  Deionized Water        |                        | Negative Control |

\* Lid colour



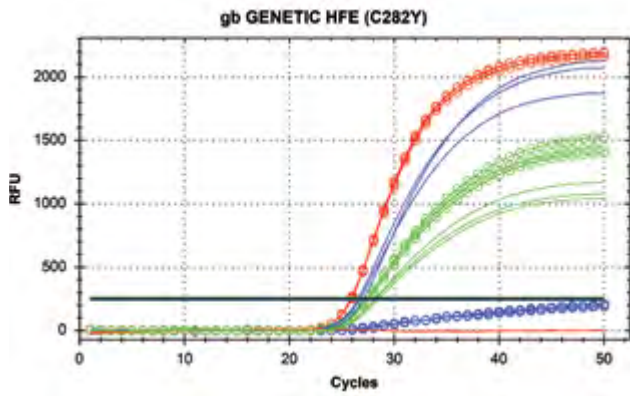


Fig. 1 – Detection of HFE (C282Y) standards on CFX96 device; blue line – wild type; red line – mutant; green line – heterozygote; smooth line – FAM channel; dotted line – HEX channel

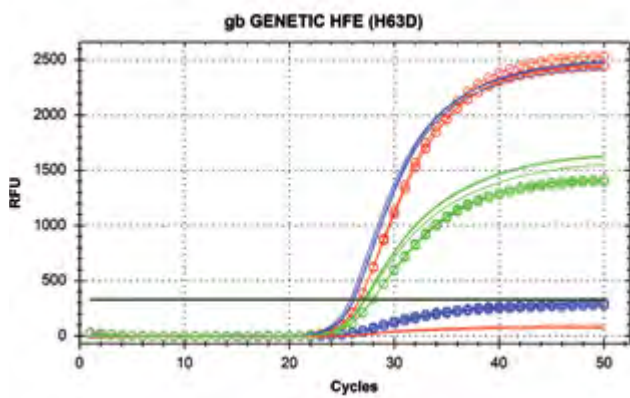


Fig. 2 – Detection of HFE (H63D) standards on CFX96 device; blue line – wild type; red line – mutant; green line – heterozygote; smooth line – FAM channel; dotted line – HEX channel

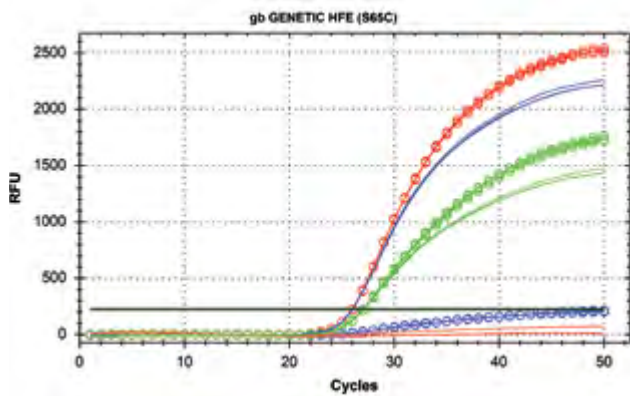


Fig. 3 – Detection of HFE (S65C) standards on CFX96 device; blue line – wild type; red line – mutant; green line – heterozygote; smooth line – FAM channel; dotted line – HEX channel

## Validated for cyclers

- Rotor-Gene 3000/6000/Q (Corbett Research, Qiagen)
- iCycler iQ5/CFX96/CFX96 Touch (Bio-Rad)
- ABI 7500/7500 Fast/7900HT (Applied Biosystems)
- SmartCycler (Cepheid)
- MIC (Bio Molecular Systems)
- AriaMx (Agilent Technologies)
- Light Cycler 480/Cobas z480 (Roche Diagnostics)

