

# Devyser Compact v3

Designed for accuracy, speed and ease of use

# Key advantages of the Devyser Compact v3 QF-PCR kit:

- Rapid prenatal aneuploidy analysis of chromosomes 13, 18, 21, X and Y
- No tissue culture needed
- Proven QF-PCR technology
- Twenty six (26) highly informative genetic markers analysed in one single PCR mix
- Two (2) unique X-chromosome counting markers for reliable detection of Turner syndrome
- Requires minute amounts of genomic DNA
- Cost-effective solution with less hands-on time required

# Designed for accuracy, speed and ease of use

For fast, precise and cost-effective prenatal diagnosis, QF-PCR is the technology of choice. Devyser brings you more than the usual QF-PCR benefits in a single, ready-to-use kit. The CE-IVD certified Devyser Compact v3 kit relies on quantitative, multiplex PCR amplification of genetic markers for prenatal diagnosis of the most common chromosome aneuploidies. In addition to detecting Down, Edwards, Patau and Klinefelter syndromes, the Devyser Compact v3 kit includes two dedicated X-counting markers for reliable detection of Turner syndrome.



#### Your key benefits include:

# Results in less than five hours

The complete procedure from sample to results takes less than five hours, allowing your laboratory to provide results in just one day.

### Quick and easy implementation

All reagents needed to perform QF-PCR are provided in a single kit.

### Cost-efficiency

This very simple procedure requires less than 90 minutes hands-on time from sample to results.

#### Fewer re-runs

Devyser's excellent assay performance and large number of highly informative markers significantly reduce the number of re-runs.

#### Easy handling

All reagents are ready to use reducing the time needed for PCR set-up and the risk of contamination.

#### Easy results interpretation

GeneMapper and Genemarker plugins for easy results interpretation and reporting.

#### Reliable detection of Turner syndrome

Two unique X-chromosome counting markers for reliable detection of Turner syndrome are included in Devyser Compact v3



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# **Technical Specification**

| Intended Use                                      | The Devyser Compact v3 kit is an in vitro diagnostic product for detection of whole chromosome aneuploidies of chromosomes 13, 18, 21, X and Y |
|---|--|
| C ← -labelled for IVD use                         | Yes  |
| Detection format                                  | ABI/Life Technologies Capillary electrophoresis instruments: 310, 3100, 3130, 3500, 3730   |
| Complies with Best Practice guidelines for QF-PCR | Yes  |
| Total number of markers                           | 26 genetic markers in one single multiplex PCR reaction.   |
| Reaction volume                                   | 25 µL  |
| DNA input   | 2 – 30 ng / PCR reaction is recommended for optimal results  |
| Ready to use for PCR                              | Yes  |
| Accessories                                       | DNA Size marker:<br>560 SIZER ORANGE, 500 loadings (8-A402)  |
| Article ID  | 25 test kit: 8-A017.3-25<br>50 test kit: 8-A017.3-50<br>100 test kit: 8-A017.3-100   |

# About Devyser

Devyser AB is specialized in the development, manufacturing and sales of diagnostic kits and reagents based on DNA analytical procedures, including PCR.

Devyser AB operates under strict quality control protocols and has an ISO13485:2003 (medical device) certification. Products are CE-labelled according to IVDD 98/79/EC for use in in-vitro diagnostic procedures. The main areas of application for our products are prenatal diagnostics and clinical genetics.

