

## FANCA/CCP16 FISH Probe Kit

### Introduction

The FANCA/CCP16 FISH Probe Kit is designed to detect the human FANCA gene located on chromosome band 16q24.3, along with the number of chromosome 16 copies per cell. Mutations in FANCA – also known as FA, FA1, FAA, FAH, FA-H, FACA or FANCH – are the most common cause of Fanconi anemia, have also been observed in cancers such as lung, colon, breast, prostate, bladder urothelial and other malignancies.

### Intended Use

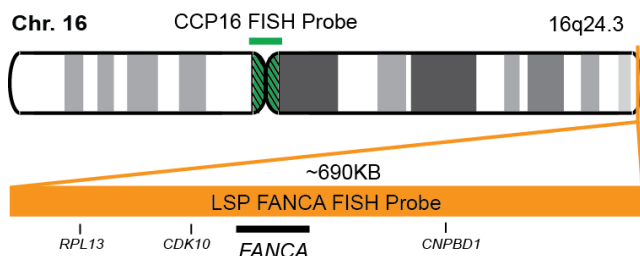
To measure the copy number of the human *FANCA* gene located on chromosome band 16q24.3.

### Cont.

### Color

LSP FANCA FISH Probe      CytoOrange  
CCP16 (Pericentromeric) FISH Probe      CytoGreen

### Probe Design



Not to Scale

LSP FANCA FISH Probe covers a chromosomal region, which includes the entire *FANCA* gene. CCP16 FISH Probe is designed to serve as a control to determine the number of chromosome 16 copies per cell.

### Cat. No.

### Volume

CT-PAC282-10-OG

10 Tests (100 µL)

### Signal Pattern Interpretation

#### Normal Patterns

2O2G

#### Abnormal Patterns

Other Patterns

- 1) Bravo-Navas S, et al. *FASEB J.* 33(9):10477-10489 (2019).
- 2) Hoskins EE, et al. *Oncogene.* 28(5):674-85 (2009).
- 3) Savino M, et al. *Hum Mutat.* 22(4):338-9 (2003).
- 4) Lensch MW, et al. *Blood.* 102(1):7-16 (2003).
- 5) Wijker M, et al. *Eur J Hum Genet.* 7(1):52-9 (1999).

\* CE IVD only available in certain countries. All other countries are either ASR or RUO. Please contact your local dealer or our headquarters for more information.

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