

TFAZZIN/CCPX FISH Probe Kit

Introduction

The TFAZZIN/CCPX FISH Probe Kit is designed to detect the human *TFAZZIN* gene located on chromosome band Xq28, along with the number of chromosome X copies per cell. Mutations in this gene – also known as *EFE*, *TAZ*, *BTHS*, *EFE2*, *G4.5*, *Taz1*, *CMD3A* or *LVNCX* – have been associated with cervical cancer and a number of clinical disorders including Barth syndrome, dilated cardiomyopathy (DCM), hypertrophic DCM, endocardial fibroelastosis, and left ventricular noncompaction (LVNC).

Intended Use

To measure the copy number of the human *TFAZZIN* gene located on chromosome band Xq28.

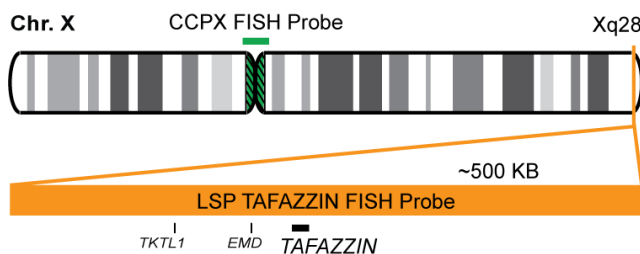
Cont.

Color

LSP TFAZZIN FISH Probe
CCPX FISH Probe

CytoOrange
CytoGreen

Probe Design



Not to Scale

LSP TFAZZIN FISH Probe covers a chromosomal region which includes the entire *TFAZZIN* gene. CCPX FISH Probe, derived from chromosome X-specific alpha satellite DNA, is designed to serve as a control to determine the number of chromosome X copies per cell.

Cat. No.

Volume

CT-PAC436-10-OG

10 Tests (100 µL)

Signal Pattern Interpretation

Normal Patterns

101G (Male)
202G (Female)

Abnormal Patterns

Other Patterns

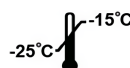
- 1) Fixler DE, et al. *Am J Cardiol.* 26(2):208-13 (1970).
- 2) Barth PG, et al. *Am J Med Genet A.* 126A(4):349-54 (2004).
- 3) Avdjieva-Tzavella DM, et al. *Genet Couns.* 27(4):495-501 (2016).
- 4) Deel MD, et al. *Clin Cancer Res.* 24(11):2616-2630 (2018).
- 5) Petit PX, et al. *Cells.* 9(10):2333 (2020).

* 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.

DCN032

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V2024.09.01

T-07-10-PAC436-OG-EN