

# In situ hybridization

ZytoPure® FISH Probes for lung cancer



## ZytoPure® FISH Probes

ZytoMed Systems is proud to offer the new product line **ZytoPure® FISH**, a reliable and accurate tool for the detection of genetic aberrations in FFPE tissue sections, cell samples and metaphase spreads by fluorescence *in situ* hybridization. **ZytoPure® FISH Probes** display brilliant signals, low background, and an excellent signal to noise

ratio. They are intended to be used in combination with the **ZytoPure® FISH Accessory Kit** comprising all reagents necessary for performing a FISH procedure on FFPE sections. The kit contains just 4 reagents and includes a robust and easy to follow protocol. All **ZytoPure® FISH Probes** as well as the **ZytoPure® FISH Accessory Kit** are CE/IVD-labelled.

### ► Background information: ZytoPure® FISH Probes for lung cancer

Rearrangements of receptor tyrosine kinases (RTK) ALK, ROS1, and RET determine clinically important molecular subgroups of non-small cell lung cancer (NSCLC). About 3% - 5% of all NSCLCs show an ALK rearrangement, and 1% - 2% show ROS1 or RET rearrangements [1]. In all fusion variants, the 3' TK domain is fused to the promotor regions and 5' coding regions of the fusion partner. Hence, rearranged RTKs are permanently active, sending proliferation signals into the cell, and thereby promoting

non-controlled cell growth. Several very efficient TK inhibitors (TKIs) were launched in recent years [2], and as a consequence the median survival of an ALK positive NSCLC patient treated with two ALK inhibitors is now more than 5 years [3]. Amplification of the MET gene is the second most common resistance mechanism of lung adenocarcinomas against EGFR TKIs [4]. In addition, primary MET amplification is also found in a small percentage of treatment-naïve lung adenocarcinomas [5].

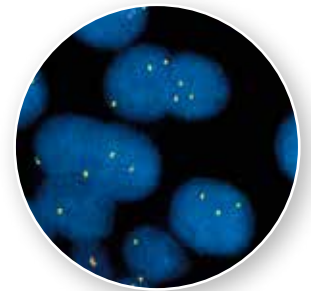
### ► Product information

ZytoPure® FISH Probes for lung cancer

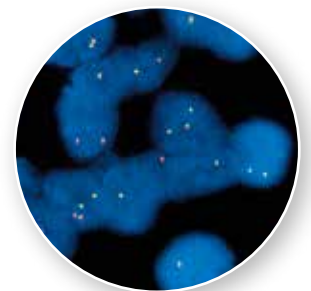
Description	Labeling	CE/IVD	Volume	Cat. No.
<b>ALK Break Apart ZytoPure® FISH Probe</b>	green/orange	✓	20 µl	F2C002-002
			100 µl	F2C002-010
<b>ROS1 Break Apart ZytoPure® FISH Probe</b>	orange/green	✓	20 µl	F2C003-002
			100 µl	F2C003-010
<b>RET Break Apart ZytoPure® FISH Probe</b>	orange/green	✓	20 µl	F2C004-002
			100 µl	F2C004-010
<b>MET/CEP7 ZytoPure® FISH Probe</b>	green/orange	✓	20 µl	F2C005-002
			100 µl	F2C005-010

### Ancillary Reagents

Description	CE/IVD	Volume	Cat. No.
<b>ZytoPure® FISH Accessory Kit</b> (Pepsin Solution, 4 ml; FISH Pretreatment Buffer, 500 ml; 20x FISH Wash Buffer, 2 x 50 ml; DAPI/Antifade Solution, 800 µl)	✓	1 Kit (20 tests)	FA-Kit1-20
<b>Pepsin Solution</b>	✓	4 ml	FA-001-004
<b>FISH Pretreatment Buffer (rtu)</b>	✓	500 ml	FA-002-500
<b>20x FISH Wash Buffer</b>	✓	50 ml	FA-003-050
<b>DAPI/Antifade Solution</b>	✓	800 µl	FA-004-008
<b>Fixogum (Rubber Cement)</b>	-	50 g	ZY-FX50
		125 g	ZY-FX125



ALK Break Apart ZytoPure® FISH Probe, lung cancer specimen showing solely ALK fusion signals



ALK Break Apart ZytoPure® FISH Probe, lung cancer specimen with ALK translocation, as indicated by single red signals in addition to fusion signals



Human metaphases hybridized with the ROS1 Break Apart ZytoPure® FISH Probe

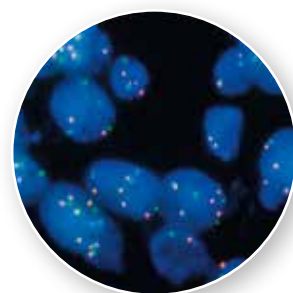
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## Other ZytoPure® FISH Probes

Description	Labeling	CE/IVD	Volume	Cat. No.
<b>ERBB2/CEP17 ZytoPure® FISH Probe</b>	orange/green	✓	20 µl	F2C001-002
			100 µl	F2C001-010
<b>MDM2/CEP12 ZytoPure® FISH Probe</b>	green/orange	✓	20 µl	F2C006-002
			100 µl	F2C006-010
<b>USP6 Break Apart ZytoPure® FISH Probe</b>	green/orange	✓	20 µl	F2C007-002
			100 µl	F2C007-010
<b>EWSR1 Break Apart ZytoPure® FISH Probe</b>	orange/green	✓	20 µl	F2C008-002
			100 µl	F2C008-010
<b>SS18 Break Apart ZytoPure® FISH Probe</b>	orange/green	✓	20 µl	F2C009-002
			100 µl	F2C009-010
<b>FUS Break Apart ZytoPure® FISH Probe</b>	green/orange	✓	20 µl	F2C010-002
			100 µl	F2C010-010



MET/CEP7 ZytoPure FISH Probe®, lung carcinoma showing amplification of both MET and CEP7

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## ZytoPure® Fluorochromes

Fluorochrome	Excitation	Emission
<b>PureGreen</b>	498 nm	532 nm
<b>PureOrange</b>	550 nm	576 nm

## ► Literature

- [1] Thunnissen E *et al.* Virchows Arch 464:347-358, 2014
- [2] Ou SH *et al.* Oncologist 17:1351-1375, 2017
- [3] Duruisseaux M *et al.* Oncotarget 8:21903-21917, 2017
- [4] Savic S, Bubendorf L Arch Pathol Lab Med 140:1323-1330, 2016
- [5] Schildhaus HU *et al.* Clin Cancer Res 21:907-915, 2015