# Cancer gene expression illuminated

## Stellaris® RNA FISH Probes for cancer research

Visualisation and quantification of single RNA transcripts, using RNA fluorescent *in situ* hybridisation (FISH), offers the unique ability to observe altered disease-specific expression in fixed cells or intact tissue. Gene expression profiles are complex and can vary immensely between regions of tissue, cells, regions of a single nucleus, and therefore require advanced methods of analysis.

#### Custom or DesignReady probe sets

LGC, Biosearch Technologies<sup>™</sup> offers a free, online probe designer (www.biosearchtech.com/ <u>stellarisdesigner</u>) which allows you to craft Stellaris RNA FISH probes with optimal binding properties for your target RNA sequence. A custom Stellaris RNA FISH probe set is a blend of up to 48 oligos each labeled with a fluorophore.

We also offer DesignReady probe sets which are professionally designed and go through rigorous bioinformatics analysis to ensure specificity. These made-to-order probe sets include cancer specific targets across a wide range of model organisms.

More information and current protocols are available online (www.biosearchtech.com/stellaris).

| DesignReady cancer sets include: |           |        |        |
|----------------------------------|-----------|--------|--------|
| ADAM17                           | EGFR      | HOTAIR | MYC    |
| AKT1                             | ERBB2 & 3 | HOXA5  | NCOA3  |
| AR                               | ERG       | JAG1   | PCA3   |
| BCL2                             | EZH2      | JUN    | PIK3CA |
| BRAF                             | FOS       | KIT    | MTOR   |
| BRCA1 & 2                        | GREB1     | KLK3   | TERT   |
| CDK4                             | H19       | MALAT1 | TERC   |
| CTNNB1                           | HIF1A     | MKI67  | TP53   |

# Detect, localise, and quantify RNA molecules in cells or intact tissue

Biosearch Technologies' Stellaris RNA FISH probes, can quantify and localise RNA within cells, tissue sections, or whole mount tissue where the morphology is preserved. A Stellaris FISH probe set comprises multiple oligonucleotides targeting a single RNA target. The fluorescently labeled probes bind along the target transcript, and combine to produce a punctate signal for an individual RNA molecule.

Image of simultaneous Stellaris RNA FISH and immunofluorescence assay. ERBB2 mRNA (green), ERBB2 (HER2) protein (magenta) and DNA (red) in human adenocarcinoma (SK-BR-3) cells. (AV Orjalo Biosearch Tech.)

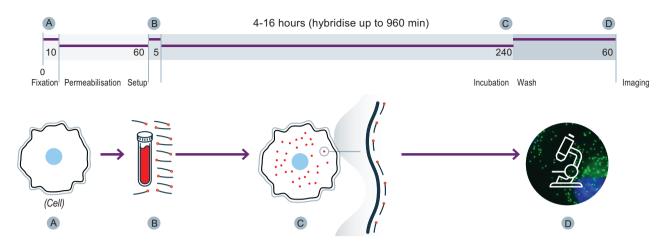






#### The Stellaris RNA FISH method

#### TIME FRAMES (MINUTES)



#### **Publication highlights**

Over 33% of publications citing Stellaris RNA FISH publish in Cell, Nature, or Science journals. Visit our Citation Center for the complete list.

Post-transcriptional modifications contribute to the upregulation of Cyclin D2 in multiple myeloma. Misiewicz-Krzeminska *et al.* Clinical Cancer Research. 2016. Url: <u>https://dx.doi.org/10.1158/1078-0432.CCR-14-2796</u>

G1/S inhibitors and the SWI/SNF complex control cell-cycle exit during muscle differentiation. Ruijtenberg *et al.* Cell. 2015. Url: <u>https://dx.doi.org/10.1016/j.cell.2015.06.013</u>

Genome-wide profiling of p53-regulated enhancer RNAs uncovers a subset of enhancers controlled by a IncRNA. Leveille *et al.* Nature Communications. 2015. Url: <u>https://dx.doi.org/10.1038/ncomms7520</u>

Niche appropriation by Drosophila intestinal stem cell tumors. Patel *et al.* Nature Cell Biology. 2015. Url: <u>https://dx.doi.org/10.1038/ncb3214</u>

### Integrated tools. Accelerated science.

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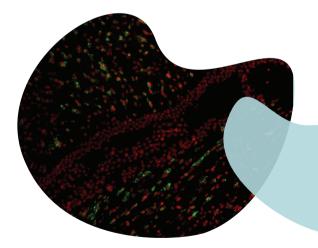
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FuseFISH: robust detection of transcribed gene fusions in single cells. Semrau *et al.* Cell reports. 2014. Url: <u>https://dx.doi.org/10.1016/j.celrep.2013.12.002</u>

Proteostatic control of telomerase function through TRiCmediated folding of TCAB1. Freund *et al.* Cell. 2014. Url: <u>https://dx.doi.org/10.1016/j.cell.2014.10.059</u>

Single-molecule transcript counting of stem-cell markers in the mouse intesting. Itzkovitz et al. Nature Cell Biology. 2011. Url: <u>https://dx.doi.org/10.1038/ncb2384</u>





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