

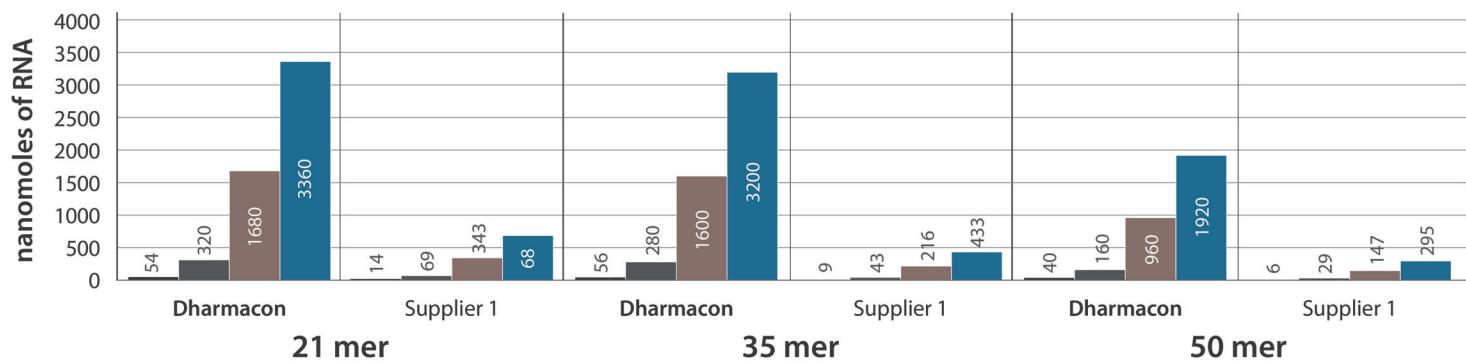
## Meet your custom RNA requirements with our nearly limitless synthetic options.

If your NMR, crystallography, or RNA binding study requires a non-standard chemical modification or a commercially unavailable modified nucleobase, we can help. For more than 20 years our chemists have utilized Dharmacon™ proprietary 2' ACE chemistry to synthesize RNA with superior yield and quality giving you the flexibility you need.

**Better RNA yields give you more for your money.**

RNA yield (HPLC-purified)

Synthesis scale ■ 0.25  $\mu$ mol ■ 1  $\mu$ mol ■ 5  $\mu$ mol ■ 10  $\mu$ mol



The yield for RNA oligos of three different lengths was compared for all available synthesis scales between Dharmacon and a competitor.

# New to single-cell sequencing?



Takara Bio has the solutions you need.

- **SMART-Seq® HT kit**—streamlined full-length cDNA library prep
- **SMART-Seq Stranded kit**—complete RNA library prep that captures coding and noncoding RNAs
- **SMARTer® PicoPLEX® Gold kit**—accurate detection of SNVs and CNVs from single cells
- **SMARTer™ ICELL8® system**—automated high-throughput single-cell isolation, selection, and processing

SMARTer  
NGS



To learn more:  
[takarabio.com/SMARTerNGS](http://takarabio.com/SMARTerNGS)

Takara Bio USA, Inc.  
United States/Canada: +1.800.662.2566 • Asia Pacific: +1.650.919.7300 • Europe: +33.(0)1.3904.6880 • Japan: +81.(0)77.565.6999

For Research Use Only. Not for use in diagnostic procedures.

© 2019 Takara Bio Inc. All rights reserved. All trademarks are the property of Takara Bio Inc. or its affiliate(s) in the U.S. and/or other countries or their respective owners. Certain trademarks may not be registered in all jurisdictions. Additional product, intellectual property, and restricted use information is available at [takarabio.com](http://takarabio.com).

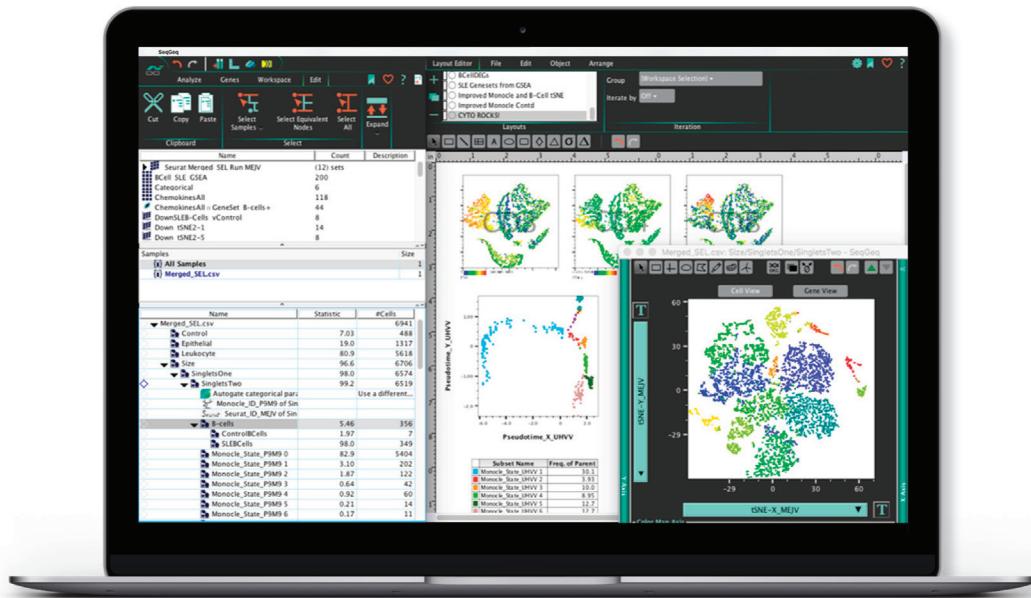


Clontech **Takara** cellartis



The world is waiting for your results.  
Unlock your scRNA seq data in minutes.

SeqGeq (seek-geek) is a desktop bioinformatics platform that makes complex scRNA seq analysis accessible with an intuitive interface. With SeqGeq, you control your analysis—no more writing R scripts to visualize your data.



Easily share your results for publication and collaboration. With our wide library of available plugins on the FlowJo Exchange, you can superpower your analysis with the most cutting-edge informatics tools.

**Discover the difference**

[www.flowjo.com/solutions/seqgeq](http://www.flowjo.com/solutions/seqgeq)



BD and FlowJo are one!  
Supporting you and your research from design to discovery.





**arbor**  
**biosciences**

*Advancing Agricultural Genomics*



**my Baits®**

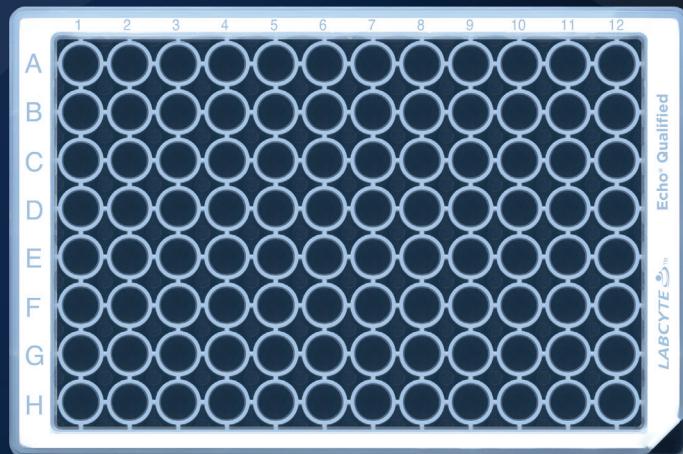
**NEW ➤ myBaits® Expert Wheat Exome Panel**

*Developed in collaboration with the IWGSC*

A comprehensive wheat exome panel targeting the complete high-confidence exon-annotated genome in hexaploid wheat based on the Chinese Spring genome assembly from the International Wheat Genome Sequencing Consortium (IWGSC).

- Probes designed using the IWGSC genome assembly and annotations
- Targets over 200 Mbp, including all high-confidence annotated exons
- Compatible with any NGS library preparation system

Coming soon...



## 96 New Reasons to Buy an Echo® Liquid Handler

Introducing the Echo® Qualified 96-well Microplate

A favorite for genomics applications, the Echo® 525 Liquid Handler saves reagents, sample, and time. Transfer of nanoliter volumes with high accuracy and precision enables assay miniaturization while maintaining data quality. Coming soon, Echo Qualified 96-well Microplates will link the unsurpassed performance of the Echo System with upstream sample preparation steps performed in a 96-well format.

- Qualified by Labcyte for reproducible acoustic performance
- High transparency with high contrast well locators for quick sample identification
- Compatible with the Echo 525 Liquid Handler

For more information, visit [www.labcyte.com/echo-525](http://www.labcyte.com/echo-525).

© 2018 LABCYTE INC. All rights reserved. Labcyte®, Echo®, and the Labcyte logo are registered trademarks or trademarks of Labcyte Inc., in the U.S. and/or other countries.

FOR RESEARCH USE ONLY. Not for use in diagnostic procedures.

**LABCYTE**   
The Future of Science is Sound

[info-us@labcyte.com](mailto:info-us@labcyte.com)

# Design Engineer Innovate

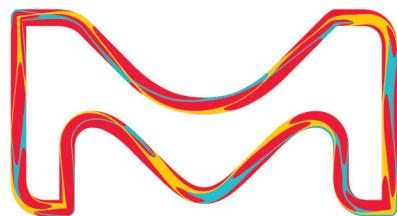
with MISSION™ CRISPR

Be confident in your  
genome editing.

The SygRNA® synthetic, crRNA:tracrRNA and one-part sgRNA systems accelerate genome editing. Pair with Cas9 protein, mRNA, or established Cas9 expressing cell lines. Deliver by a variety of methods including microinjection, electroporation, and lipofection.

Guaranteed performance with  
MISSION™ CRISPR predesigned  
knockout gRNAs

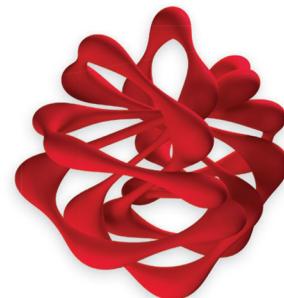
To find out more, visit  
[SigmaAldrich.com/SygRNA](https://SigmaAldrich.com/SygRNA)



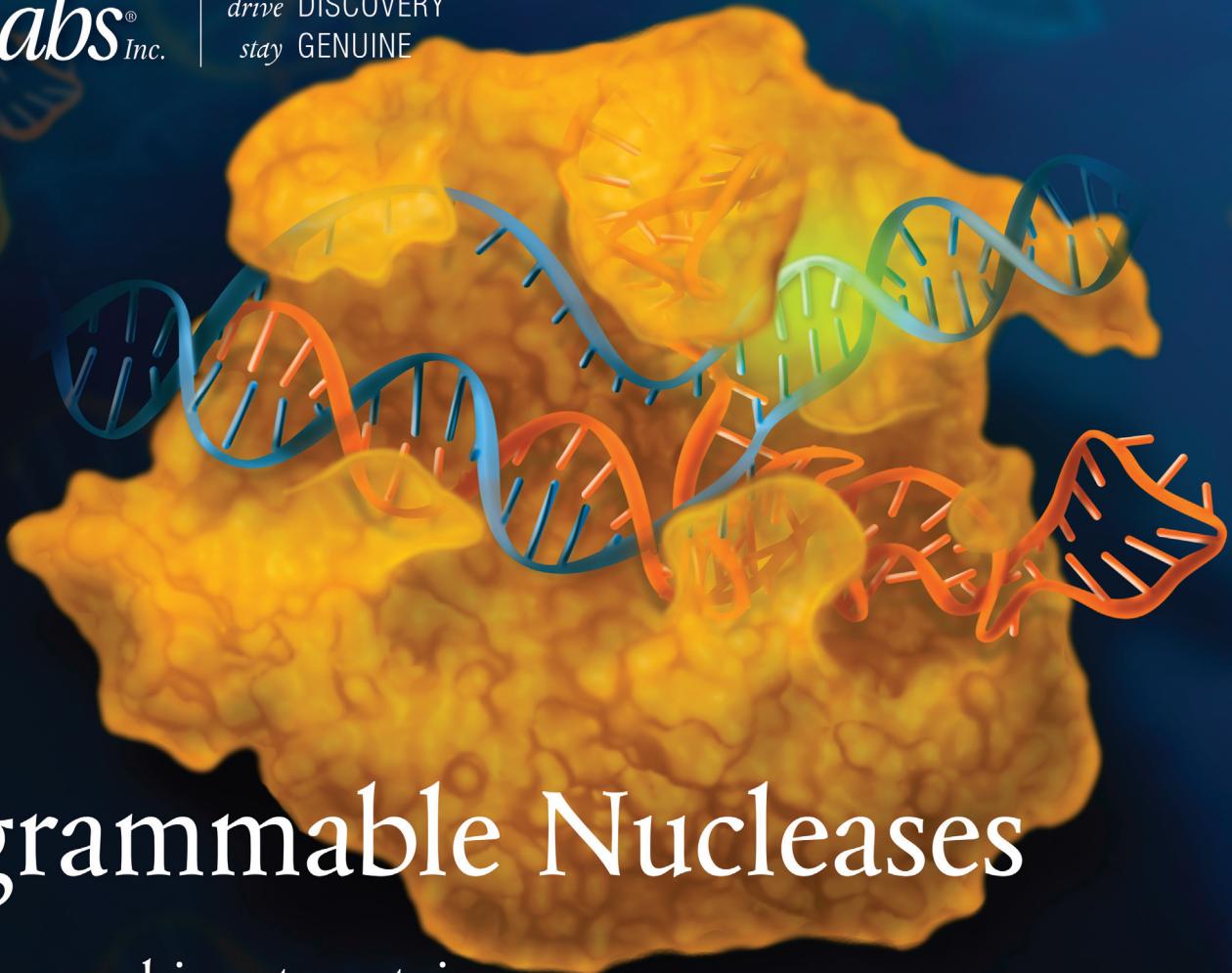
© 2019 Merck KGaA, Darmstadt, Germany and/or its affiliates. All Rights Reserved. MilliporeSigma, the vibrant M, Sigma-Aldrich, MISSION, and SygRNA are trademarks of Merck KGaA, Darmstadt, Germany or its affiliates. All other trademarks are the property of their respective owners. Detailed information on trademarks is available via publicly accessible resources.

2019 - 19308 1/2019

The life science  
business of Merck  
KGaA, Darmstadt,  
Germany operates as  
MilliporeSigma in the  
U.S. and Canada.



**Sigma-Aldrich®**  
Lab & Production Materials



# Programmable Nucleases

Purified, recombinant proteins  
for genome editing applications

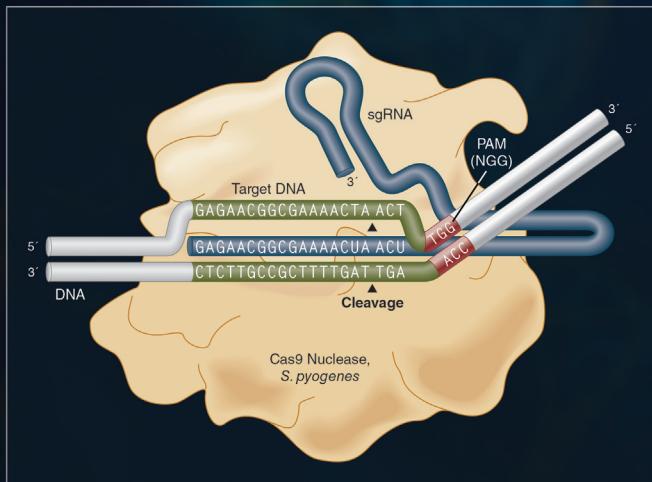
Cas nucleases, key proteins involved in CRISPR/Cas genome editing, are now available from New England Biolabs<sup>®</sup>. These Cas nucleases expand the portfolio of products from NEB<sup>®</sup> that support CRISPR workflows, including sgRNA cloning, transcription and HDR template construction.

Download our Genome Editing Brochure  
at [www.neb.com/GenomeEditing](http://www.neb.com/GenomeEditing)

**NOW AVAILABLE:**

EnGen<sup>®</sup> Spy Cas9 Nickase, EnGen Spy dCas9 (SNAP-tag<sup>®</sup>), EnGen Lba Cas12a (Cpf1) and EnGen Sau Cas9

Schematic representation of Cas9 Nuclease, *S. pyogenes* sequence recognition and DNA cleavage

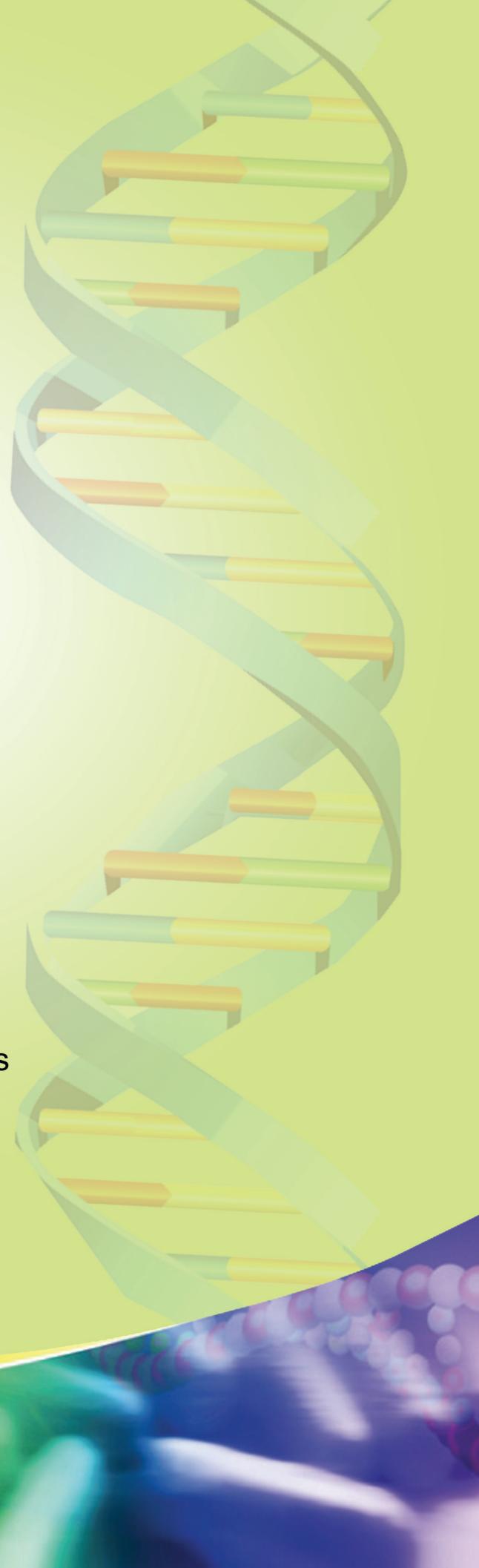


## Custom Oligonucleotides

- ◆ Regular oligos
- ◆ Long oligos
- ◆ Phosphorothioated oligos (S-Oligos)
- ◆ Modified oligos
- ◆ Fluorescent oligos
- ◆ Taqman probes
- ◆ Molecular beacon
- ◆ Oligo pool & microarray

## Custom Peptide Synthesis

- ◆ Purities from desalt to 98%
- ◆ Acetylation/Amidation
- ◆ Phosphorylated peptides
- ◆ Fluorescein/Biotin labeled peptides
- ◆ Specialty peptides with unnatural amino acids
- ◆ Cyclic peptides
- ◆ KLH/BSA/OVA Conjugation
- ◆ Multiple Antigenic Peptides
- ◆ Peptide nucleic acid (PNA)



**AACR** American Association  
for Cancer Research®  
**ANNUAL  
MEETING**  
**2020 · SAN DIEGO**

APRIL 24-29

**TURNING SCIENCE  
INTO LIFESAVING CARE**

Join us in San Diego for the latest innovative and inspiring cancer research from around the world...the **AACR ANNUAL MEETING 2020!**

**Abstract Submission Deadline:** Thursday, December 5, 2019

**Early Registration Deadline:** Friday, December 20, 2019

**Become a Member!**

Join the AACR and receive a discount on registration.



Continuing Medical Education Activity -  
AMA PRA Category 1 Credits™ available

The AACR Annual Meeting highlights the work of the greatest minds in cancer science and medicine from institutions all over the world. This meeting presents the many scientific discoveries across the breadth of cancer research—from prevention, early detection, and interception; to cancer biology, translational, and clinical studies; to survivorship, population science, and advocacy. This year's program, with the theme of "Turning Science into Lifesaving Care," will be a comprehensive, cutting-edge scientific event that you will not want to miss!

**We look forward to seeing you!**

**AACR.ORG • #AACR20**



190103IN



## Postdoctoral Positions

### About SCISSOR

Single-Cell In Situ Spatial Omics at subcellular Resolution (SCISSOR) is a well-supported multidisciplinary program that aims to introduce new paradigms for cancer biology and diagnostics, using spatial and non-spatial omics technologies. Our team comprises of computational biologists (lead: Shyam Prabhakar), oncologists (lead: Iain Tan), biotechnologists (lead: Kok Hao Chen), and pathologists (lead: Tony Lim) with a track record of combining cutting-edge computational and experimental approaches to infer disease mechanisms and develop clinical applications (Chen et al., *Science* 2015; Li et al., *Nat Genet* 2017; Sun et al., *Cell* 2016; Fukawa et al., *Nat Med* 2016; del Rosario et al., *Nat Methods* 2015; Kumar et al., *Nat Biotechnol* 2013; Ku et al., *Lancet Oncol* 2012).

We are looking for bright, motivated individuals who are interested in working on cutting-edge research projects that leverage single cell and spatial omics. Our interdisciplinary team combines experimental biology, technology development and computational biology to address major questions in cancer biology.

### Position 1

#### Postdoctoral fellow: Machine Learning and Mathematical Analysis of Spatial Transcriptomics Data

Successful candidates will develop and apply algorithms for the analysis of large-scale cancer data. This will be a unique opportunity to lead computational analysis of new types of data in the nascent field of spatial transcriptomics.

##### **Requirements:**

- Strong programming skills
- Expertise in mathematics, computer science, statistics, engineering, machine learning, signal processing, computational genomics, or a related field
- General quantitative intuition
- Strong publication record
- Strong communication skills
- The ability to work closely with clinicians and experimental biologists

### Position 2

#### Postdoctoral fellow: Assay Development, Cancer Markers and Mechanisms

Successful candidates will have the opportunity to lead experimental design and execution for a spatial transcriptomics study looking at DNA and RNA changes in a variety of human cancers at subcellular resolution.

##### **Requirements:**

- Expertise in cancer biology, immunology, genomics or related fields
- Skilled in molecular and cellular assays
- Strong publication record
- Team player and strong communication skills (oral and written)
- The ability to work closely with clinicians and computational biologists

##### **Benefits:**

The Genome Institute of Singapore offers a competitive salary and a complete benefits package that ensures a very high living standard in one of the most modern cities in the world.

### **About the Organisation**

The Genome Institute of Singapore (GIS), A\*STAR Research Entities is the national flagship program for genomic science in Singapore. GIS is located within the Biopolis, the biomedical research hub of Singapore, which houses in close proximity research institutes under the Agency of Science, Technology and Research (A\*STAR), biotech startups and international pharmaceutical corporations. The applicant would have the opportunity to interact with scientists, bioinformaticians, clinicians, engineers and other professionals from all over the world in a vibrant, intellectually stimulating and scientifically curious setting. You will be part of a vibrant scientific community where you will have the opportunity to share your ideas and demonstrate your skills and passion for scientific research. You can find out more about the Genome Institute of Singapore online: <https://www.a-star.edu.sg/gis/>.

### **Why Singapore?**

Singapore, a city-state with one of the highest standards of living in the world, is an international hub for the biomedical sciences. Singapore is a tropical city with a rich Asian heritage and modern style of living, and is an ideal gateway to explore Asia providing a unique experience and an excellent quality of life.

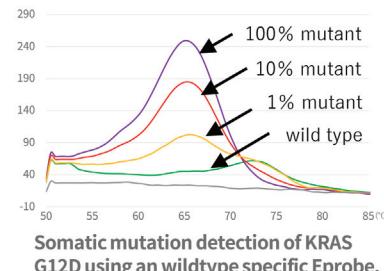
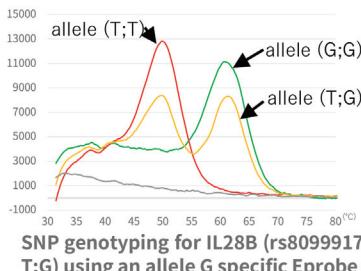
### **How to Apply**

To apply, please email your CV and names of references to: [prabhakars@gis.a-star.edu.sg](mailto:prabhakars@gis.a-star.edu.sg), [arulrayan@gis.a-star.edu.sg](mailto:arulrayan@gis.a-star.edu.sg)

# A novel solution for SNP/somatic mutation detection

Eprobe is a **DNA-based fluorescent probe** which emits fluorescence when specifically binding to a complementary strand. Melting curve analysis after PCR can detect **SNP genotype** and **somatic mutations**. Two fluorescent dyes (thiazole orange and thiazole pink) are available.

- High resolution SNP detection**—Increased  $T_m$  (approx.10°C) by the thiazole orange enables a shorter probe design and a clearer distinction of SNPs
- Simple and highly sensitive somatic mutation detection**—sensitive detection of somatic mutations (down to 0.1%) can be achieved by suppression of PCR amplification of wild-type alleles by Eprobe (PCR clamping)
- Compatible with most real time PCR instruments**—fluorescence emitted by Eprobe can be detected using a filter for SYBR® Green I\* \*SYBR® is a registered trademark of Molecular Probes, Inc.
- Easy to use online design tools**—a design tool for a primer/Eprobe (E-design, [www.dnaform.com/edesign2/](http://www.dnaform.com/edesign2/)) and a thermodynamic calculation tool (ECHO, [www.dnaform.com/devel/echo/thermodynamics/](http://www.dnaform.com/devel/echo/thermodynamics/)) are available



Fluorophore (excitation/emission)	1.5 nmol	3.0 nmol	5.0 nmol	10.0 nmol
Thiazole orange (510 nm / 530 nm)	19,000 JPY 38,000 JPY	30,000 JPY 60,000 JPY	45,000 JPY 90,000 JPY	70,000 JPY 140,000 JPY
Thiazole pink (570 nm / 590 nm)	45,000 JPY	70,000 JPY	110,000 JPY	170,000 JPY

Special offer for new customers  
50% OFF the list price!  
All Thiazole orange-labeled products



Learn more at

[www.dnaform.jp/en/products/fluorescent\\_oligonucleotide/eprobe\\_eprimer/](http://www.dnaform.jp/en/products/fluorescent_oligonucleotide/eprobe_eprimer/)

# The CellRaft<sup>®</sup> AIR

## Image, Sort, Isolate.

.....



Expand your single cell analysis research with imaging-based sorting.

The CellRaft AIR System integrates single cell imaging and viable cell isolation, with no minimum input requirement – and works with virtually any cell type.



- Phenotyping by imaging provides more information than RFUs alone
- Perform 'on-array' functional assays like co-culture or drug challenges
- Verify live single cells and eliminate empties, doublets, or dying cells
- Flexible for any downstream process – NGS, multi-omics, expansion culture
- Inform, QC, and cross-reference your genomic data analysis with cell images

Learn more about the CellRaft AIR system and how it can accelerate your single cell analysis research at [cellmicrosystems.com](http://cellmicrosystems.com)



**CELL**  
Microsystems<sup>®</sup>