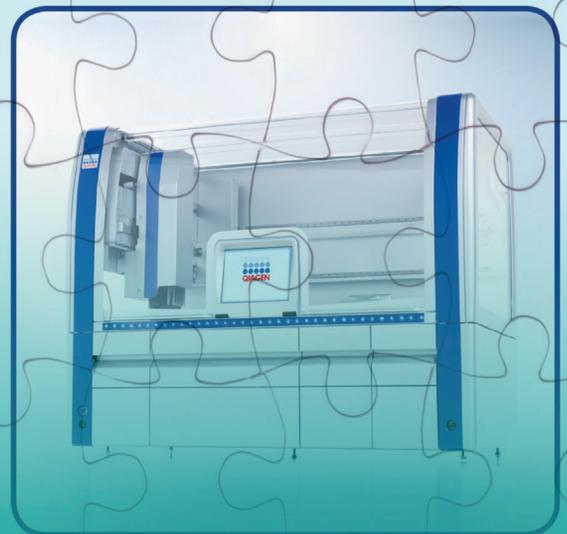


All the pieces are in place

Discover the **PAXgene® Blood ccfDNA System**: a complete, integrated preanalytical workflow solution for circulating cell-free DNA

Sample
Collection &
Stabilization

Circulating
DNA
Isolation



- ⊗ **Non-crosslinking preservation** of ccfDNA levels
- ⊗ **Seamless integration** with proven ccfDNA isolation kits
- ⊗ Streamlined with **primary tube sampling** during automated processing
- ⊗ Designed for sensitive research assays, including analysis of **circulating tumor DNA** and **non-invasive prenatal testing**



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Custom Oligonucleotides

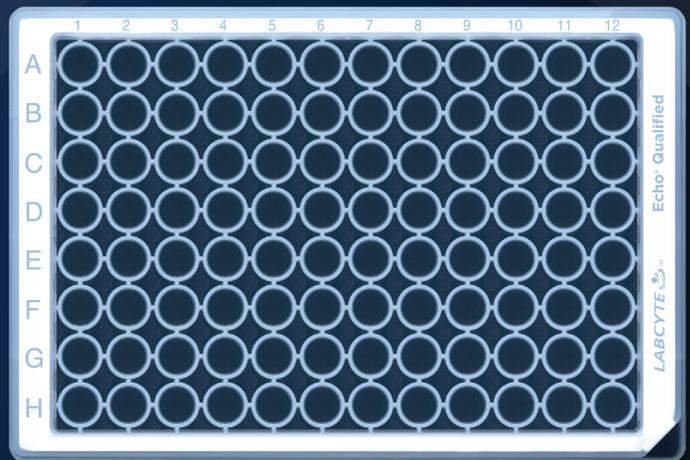
- ◆ Regular oligos
- ◆ Long oligos
- ◆ Phosphorothioated oligos (S-Oligos)
- ◆ Modified oligos
- ◆ Fluorescent oligos
- ◆ Taqman probes
- ◆ Molecular beacon
- ◆ Oligo pool & microarray

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)



Coming soon...



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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

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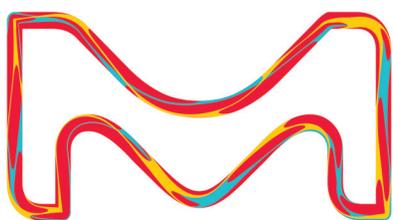
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Be confident in your
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The SygRNA® synthetic, crRNA:tracrRNA and one-part sgRNA systems accelerate genome editing. Pair with Cas9 protein, mRNA, or established Cas9 expressing cell lines. Deliver by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

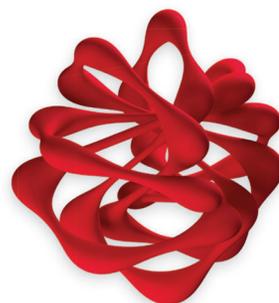
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CSHL Meetings & Courses



aerial view of CSHL sandspit and inner harbor during meeting social

2019 Meetings

- Ubiquitins, Autophagy & Disease** April 23 - 27
- Telomeres & Telomerase** April 30 - May 4
- The Biology of Genomes** May 7 - 11
- Mechanisms of Metabolic Signaling** May 14 - 18
- Retroviruses** May 20 - 25
- 84th Symposium: RNA Control & Regulation** May 29 - June 3
- Microbiome** July 18 - 21
- Cell Death** August 13 - 17
- Eukaryotic mRNA Processing** August 20 - 24
- Mechanisms of Eukaryotic Transcription** August 27 - 31
- Eukaryotic DNA Replication & Genome Maintenance** September 3 - 7

Microbial Pathogenesis and Host Response

September 10 - 14

Stem Cell Biology

September 17 - 21

Biology of Cancer: Microenvironment & Metastasis

September 24 - 28

Neurobiology of *Drosophila*

October 1 - 5

Genome Engineering: Frontiers of CRISPR/Cas

October 10 - 13

Yeast Research: Origins, Insights, Breakthroughs

October 23 - 26

Genome Informatics

November 6 - 9

Single Cell Analyses

November 13 - 16

Zebrafish Neural Circuits & Behavior

November 20 - 23

Plant Genomes, Systems Biology & Engineering

December 4 - 7

Development & 3D Modeling of the Human Brain

December 9 - 12

2019 Fall Courses

- Workshop on Cereal Genomics** October 15 - 21
- Programming for Biology** October 15 - 30
- X-Ray Methods in Structural Biology** October 15 - 30

Advanced Sequencing Technologies & Applications

November 5 - 17

The Genome Access Course

November 11 - 13

Scientific Writing Retreat

November 13 - 17

Computational Genomics

December 4 - 11

meetings.cshl.edu

Preview of our early 2020 Meetings

- Systems Biology: Global Regulation of Gene Expression** March 11 - 14
- Neuronal Circuits** March 18 - 21
- The PARP Family & ADP-ribosylation** April 1 - 4
- Gene Expression & Signaling in the Immune System** April 14 - 18

- Protein Homeostasis in Health & Disease** April 21 - 25
- Genome Organization & Nuclear Function** April 28 - May 2
- The Biology of Genomes** May 5 - 9
- Regulatory & Non-Coding RNAs** May 12 - 16
- Retroviruses** May 18 - 23

SAVE THE DATE!

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AACR ANNUAL MEETING 2020

THE SAN DIEGO CONVENTION CENTER APRIL 25-29



For more information
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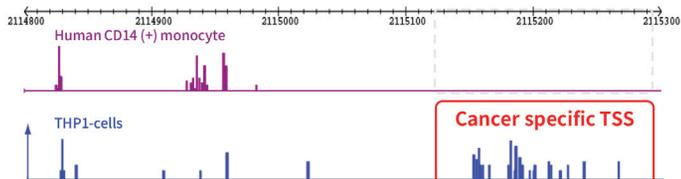
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FINDING CURES TOGETHER®

Promoter / Enhancer Annotation in the NGS era

Cap Analysis of Gene Expression (CAGE) is a new NGS library preparation method using “cap-trapping” technology which enables you to detect and quantify **transcription start site (TSS)** of RNA pol II transcripts including **mRNAs**, **lincRNAs** and **enhancer RNAs**.

- **Accurate promoter annotation**—reliable estimation of promoter positions and their activities based on precise TSS information
- **Estimation of transcription factor binding sites**—genome-wide motif search around TSS which have different expression profiles among samples
- **Detection of active enhancers**—identify active enhancers by detection of bidirectional enhancer RNAs
- **Development of new biomarkers**—TSS variants are valuable candidate of biomarkers even in the case that there are no difference at the gene expression level
- **Accurate quantification of gene expression**—PCR-free library preparation process without fragmentation allow for more reliable quantification of gene expression than RNA-seq



CAGE expression pattern of a histone H3 methyltransferase gene of human CD14(+) monocytes and THP-1 leukemia monocytic cells.

CAGE library preparation & analysis services

Library preparation for Illumina sequencers	500 USD/sample
Sequencing (Illumina HiSeq/ NextSeq)	250 USD/sample
Bioinformatics analysis	250 USD/sample

CAGE library preparation kit

8 samples (Cat. 52003-8)	2,000 USD
48 samples (Cat. 52003-48)	10,000 USD

More than 250 papers using CAGE have been published!
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