



PRE-MEETING WORKSHOP

GENOME EDITING
WORKSHOP—
THE EVER-EXPANDING
EDITING TOOLBOX

MAY 15, 2022



EDITAS IS TRANSFORMING MEDICINE



Tristan
Living with sickle cell disease

We are harnessing the power and potential of gene editing to develop medicines for people living with serious diseases around the world.
Visit us at: www.editasmedicine.com.

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GENOME EDITING WORKSHOP— THE EVER-EXPANDING EDITING TOOLBOX

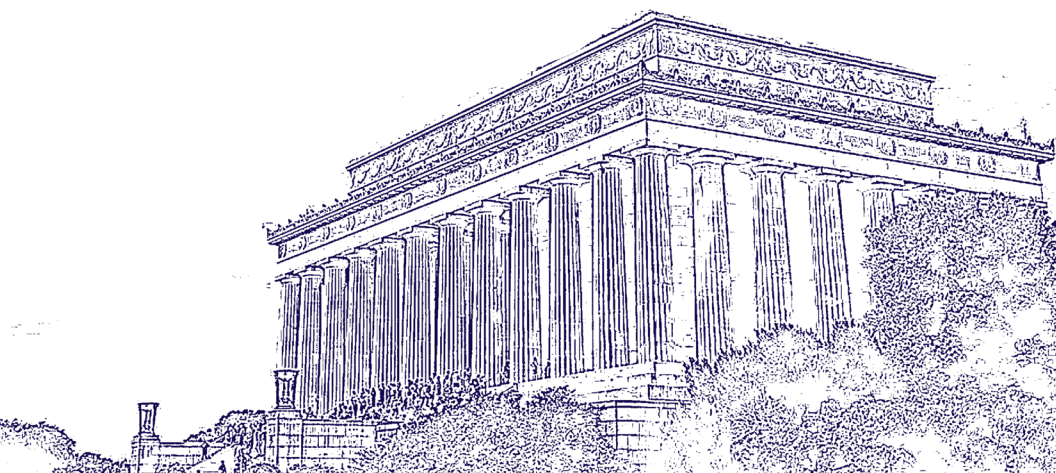
SUNDAY, MAY 15, 2022

All times listed below ET

Co-chairs: Paula Rio, PhD
Thomas Cradick, PhD

Description

Gene editing has become an essential technology in our day-to-day research and it is a promising approach for the application in the clinics. In this workshop we will focus on the most novel gene editing strategies designed and the development of new editing tools by protein engineering. A special attention will be given to HDR improvement by optimized donor templates. Altogether, this workshop will contribute to learning about the ever-expanding gene editing toolbox.



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About Our
Approach



GENETIC MEDICINES FOR GENETIC DISEASES





GENOME EDITING WORKSHOP— THE EVER-EXPANDING EDITING TOOLBOX

Schedule

All times listed below ET

8:00–8:25 AM

Enhancing HDR Efficiency in Human Pluripotent Stem Cells

Sridhar Selvaraj, PhD, Stanford University

8:25–8:50 AM

Cas9-HRs: Manipulating DSB Repair Pathway Choice to Increase HDR

Chris Hackley, PhD, CRISP-HR Therapeutics

8:50–9:15 AM

Improved Methods for CRISPR HDR Research Using Alt-R™ Modified Donors and Alt-R HDR Enhancer V2

Mollie Schubert, MS, Integrated DNA Technologies

9:15–9:40 AM

***In Vivo* Epigenome Editing**

Lucie Guo, PhD, Stanford University

9:40–10:05 AM

Growing the CRISPR Toolbox for Genetic Medicines

David Scott, PhD, Arbor Biotechnologies



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10:05–10:30 AM

Identifying and Engineering Orthogonal and Synthetic RNA-Guided Nucleases (sRGN)

Andre Cohnen, PhD, Bayer AG

10:30–10:45 AM

Break

10:45–11:10 AM

Prime Editing

Gerald Schwank, PhD, University of Zurich

11:10–11:35 AM

Targeting Double-strand Break Indel Byproducts with Secondary Guide RNAs Improves Cas9 HDR-mediated Genome Editing Efficiencies

Zsolt Bodai, PhD, UC San Diego

11:35 AM–12:00 PM

Programmable Transposon-encoded RNA-guided Nucleases for Genome Editing

Soumya Kannan, Massachusetts Institute of Technology

change life stories with genome editing therapies



Our mission

Transforming lives of people
with severe diseases by
developing potentially curative
genome editing treatments.

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