



Abstract
submission
deadline:

November 1,
2019

Early
registration
deadline:

December 13,
2019

This meeting will bring together academic and industry scientists and clinicians working on developing and implementing gene therapy and cell-based therapy for a wide variety of conditions. Topics will include in vivo gene therapy with CRISPR and other platforms, ex vivo genetic engineering of immune cells and stem cells for translational applications, development of methods for delivering genes and editing machinery to desired target cells, and strategies to circumvent barriers such as host immune reactions against introduced vectors, proteins, and cells.

Keynote Speaker

Alta Charo, *USA*

Confirmed Speakers

Daniel Anderson, *USA*

Alessandra Biffi, *USA*

Nathalie Cartier, *France*

Yvonne Chen, *USA*

Beverly Davidson, *USA*

Viviana Gradinaru, *USA*

Katherine High, *USA*

Anna Kajaste-Rudnitski, *Italy*

David Liu, *USA*

Crystal MacKall, *USA*

Eric Olson, *USA*

Malin Parmar, *Sweden*

Matthew Porteus, *USA*

Waseem Qasim, *UK*

Sonja Schrepfer, *USA*

Luk Vandenberghe, *USA*

Organizers

Jennifer Doudna, UC Berkeley, UCSF Gladstone Institutes, USA

Luigi Naldini, San Raffaele Telethon Institute for Gene Therapy, Università Vita-Salute San Raffaele, Italy

April Pawluk, *Cell*

Jonathan Saxe, *Cell Stem Cell*

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