# Concept Clearance for Genomic Technology Development

Michael W. Smith May 18, 2015 NACHGR

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## Workshop Participants

#### April 10

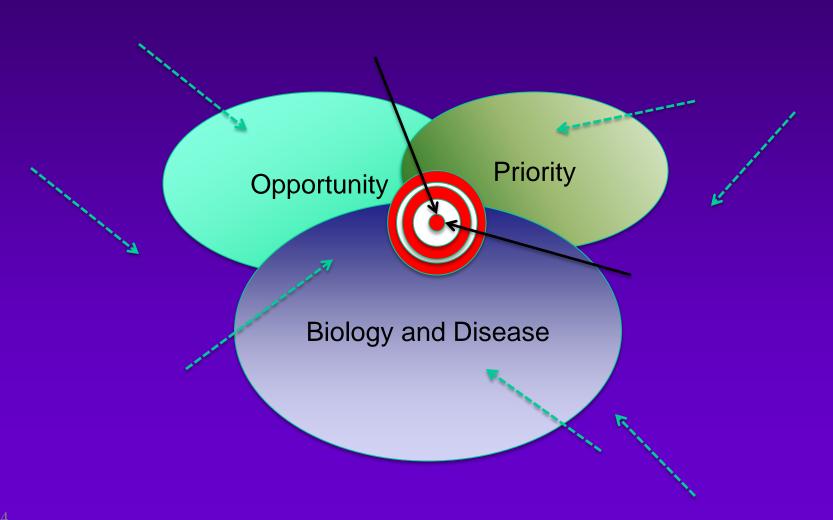
Mark Chee
Joe Ecker
Andy Feinberg
Chuan He
Jonas Korlach
Chris Mason
Steve McCarroll
Chad Nusbaum
Mike Snyder
Marc Vidal



#### April 16

George Church Steve Fodor Hanlee Ji Pui-Yan Kwok Andre Marziali Deirdre Meldrum Donna Muzny John Nelson Steve Quake Aviv Regev Jay Shendure **David Walt Bob Waterston** 

# Workshop charge



### General Workshop Recommendations

- Technology development continues to be sorely needed across a wide swath of genomics
  - A strong need for new DNA and direct RNA sequencing methods
  - Many and diverse areas of opportune and relevant priorities
  - Details provided in the initiatives
- Technology opportunities best identified by investigator interests
- Applications benefit from dedicated review and solicitation

### Concept - Overall Goals

- Facilitate development of novel genomic technologies
- Specifically solicit novel technologies for nucleic acid sequencing technology development
- Provide a new avenue of support and dedicated review for broadly defined genomic technology development

#### Initiatives

1. Novel Nucleic Acid Sequencing Technology Development

2. Genomic Technology Development Awards

# 1- Novel Nucleic Acid Sequencing Technology Development

- <u>Purpose</u>: Develop revolutionary nucleic acid sequencing technologies
- Scope and Objectives: Support new DNA and direct RNA sequencing technologies and chemistries to develop transformative and foundational nucleic acid sequencing technologies with long read lengths, high accuracy and low cost.
- Mechanism of Support and Budget
  - \$2M TC in additional RPG funding/year; four years (FY16-19);
     RFAs
  - 2-4 awards each year. R01s up to 4 yrs; R21s up to 3 yrs
  - SBIR/STTR funds will be in addition to those listed in the previous bullets

Activity	FY16	FY17	FY18	FY19	FY20	FY21	FY22
Novel Nucleic Acid Sequencing Technologies	\$2M	\$4M	\$6M	\$8M	\$6M	\$4M	\$2M

# 1- Novel Nucleic Acid Sequencing Technology Development

#### Examples of potential research topics include:

- Novel chemistries, physics or instrumentation for entirely new ways to perform DNA sequencing. Examples of important sequencing needs include:
  - Exhaustive and quantitative sequencing of every DNA and/or RNA molecule in a sample
  - Very long reads (e.g., ≥150 Kb) with accuracy and error structure sufficient to de novo assemble human genomes
- Direct RNA sequencing of full length transcripts without a cDNA intermediate
- Several orders of magnitude improvements to existing sequencing technologies
- All of the above w/o sacrificing cost

### **Timeline For Initiatives**

Event	Date
Concept Clearance Council Review:	May 2015
FOAs Released:	Summer 2015
Scientific Review:	Winter 2016
Council Review:	May 2016
Funding:	Summer 2016

#### 2- Genomic Technology Development Awards

- <u>Purpose</u>: Catalyze investigator-initiated genomic technology development that impacts the field within five to seven years.
- Scope and Objectives: Enable a wide swath of genomic technology development for, e.g.:
  - High throughput/high information content functional analysis with genomic readout
  - Single cell methods for everything on the list
  - Foundational technologies (e.g., sample prep, novel sensors)
  - Transcriptome analysis
  - Functional analysis
- Mechanisms of Support and Budget
  - \$3M TC in additional RPG funding/year; four years (FY16-19); PARs
  - 3-6 awards each year. R01s up to 4 yrs; R21s up to 3 yrs
  - SBIR/STTR funds will be in addition to those listed in the previous bullets

Activity	FY16	FY17	FY18	FY19	FY20	FY21	FY22
Genomic Technology Development Awards	\$3M	\$6M	\$9M	\$12M	\$9M	\$6M	\$3M

#### 2- Genomic Technology Development Awards

#### Examples of potential research topics include:

- DNA, RNA, epigenome, protein, etc. from the same sample
- High throughput genome modifications (by recombination and transient assays), for replacement, activation and inhibition, with genomic readout
- Scaling DNA sequencing, transcriptome, and functional technologies to operate on 10<sup>4</sup> samples (with ultimate goal of 10<sup>8</sup> cost-effectively) for, e.g., single cell/small samples and for population studies
- In situ methods (tissue context) for DNA, epigenome, RNA, and protein
- Measuring proximal transcription dynamics, and transcriptome dynamics over time, from cells to organs
- Generating sequence tags at orders of magnitude lower costs than currently available

### **Budget Summary\***

Activity**	FY16	FY17	FY18	FY19	FY20	FY21	FY22
Novel Nucleic Acid Sequencing Technologies	\$2M	\$4M	\$6M	\$8M	\$6M	\$4M	\$2M
Genomic Technology Development Awards	\$3M	\$6M	\$9M	\$12M	\$9M	\$6M	\$3M
Total	\$5M	\$10M	\$15M	\$20M	\$15M	\$10M	\$5M

<sup>\*</sup> The current concept funds at the previous \$1000 genome level, but is split between sequencing RFAs (\$2M new grants per year) and the wider genomic PARs (\$3M new grants per year) for technology development.

Note: SBIR/STTR funds will be in addition to the amounts listed above.

<sup>\*\*</sup> R01s of up to four years and R21s of up to three years are anticipated.

### Genomic Tech Dev Budget

\$1,000 Genome RPG spending.

FY06	FY07	FY08	FY09	FY10	FY11	FY12	FY13	FY14
\$22.8M	\$22.2M	\$15.8M	\$20.8M	\$18.1M	\$15.7M	\$15.1M	\$14.2M	\$13.8M

FY15	FY16	FY17
\$9.9M	\$5.5M	\$1.2M

#### Proposed Genomic Technology Development Concept

Activity**	FY16	FY17	FY18	FY19	FY20	FY21	FY22
Novel Nucleic Acid Sequencing Technologies	\$2M	\$4M	\$6M	\$8M	\$6M	\$4M	\$2M
Genomic Technology Development Awards	\$3M	\$6M	\$9M	\$12M	\$9M	\$6M	\$3M
Total	\$5M	\$10M	\$15M	\$20M	\$15M	\$10M	\$5M

### Thank you's to:

Bianca Patel

Katya Vaydylevich

**Chris Wellington** 

Elise Feingold

Mike Pazin

Adam Felsenfeld

Bettie Graham

Jeff Schloss

Hanlee Ji

Joe Ecker

Jay Shendure

& 21 other

workshop/survey

participants