

**National Advisory Council for Human Genome Research
Concept Clearance for Genomic Technology Development PAR
September 16 - 17, 2019**

Title: Genomic Technology Development PAR (R01, R21 and R43/R44)

Purpose: The purpose of this set of Novel Genomic Technology Development Program Announcements with special Review criteria (PARs) is to catalyze and support major advances in genomic technologies. Progress in basic research and clinical applications of genomics, and more broadly in biomedical research, has been greatly facilitated by significant and sustained genomics technology throughput increases, cost decreases, and improvements in ease of use. The proposed work solicited should allow comprehensive genomic analysis of features not assayable today, completely new ways to assay genomic features, or an increase of no less than an order of magnitude in an existing technology in terms of data quality, throughput, efficiency or comprehensiveness (individually or in combination). Work solicited under the accompanying Technology Development RFAs Concept Clearance will be explicitly excluded from this set of PARs.

New or renewal/modified initiative: Renewal

Some changes reflecting input from town halls, workshops and other strategic planning forums are anticipated for the revised PARs. Switching to every other Council review rather than the current annual cycle is also anticipated to accelerate progress.

Current PARs:

R01: <https://grants.nih.gov/grants/guide/pa-files/PAR-18-777.html>
R21: <https://grants.nih.gov/grants/guide/pa-files/PAR-18-778.html>
R43/R44: <https://grants.nih.gov/grants/guide/pa-files/PAR-18-779.html>

Mechanism of Support: R01, R21 and R43/R44 (see Appendix 1).

Appendix 1: Mechanism Overview

Appendix 2: Novel Genomic Technology Development PAR grants awarded (FY16-FY19).

Genomic Technology Development PAR

Appendix 1. Mechanism Overview

Program Announcement with special Review criteria (PAR)

| FOA set | Receipt dates | Activity Code(s) | Max award duration |
|--------------------|----------------------|-------------------------|---------------------------|
| Genomic Technology | FY21-FY23 | R01, R21 & R43/R44 | 4 years |

Genomic Technology Development PAR

Appendix 2. Novel Genomic Technology Development PAR grants (FY16-FY19).

| Grant | PI | Title |
|-----------------------|----------------------------|---|
| <u>R01</u> | | |
| HG009269 | Bernstein, Bradley | Single-molecule systems for decoding combinatorial chromatin modifications |
| HG009276 | Regev, Aviv | DNA microscopy for spatially resolved genomic analyses in intact tissue |
| HG009283 | Blainey, Paul | Arrayed single-cell readout of pooled genetic perturbation libraries |
| HG009285 | Mali, Prashant | Next-generation Functional Genetic Screening of Un-screenable Traits |
| HG009761 | Zhang, Feng | Programmable RNA-targeting tools |
| HG010318 | Kim, Sanggu | On-site, high-fidelity target sequencing and absolute quantitation for HIV-1 surveillance |
| HG010211 | Zeitlinger, Julia | A transposase system for integrative ChIP-exo and ATAC-seq analysis at single-cell resolution |
| HG010647 | Chen, Fei | A single cell spatial genomics platform for multi-modal characterization of tissue organization |
| HG010634 | Ecker, Joseph | Development of methods for multi-omic analysis of DNA methylation and chromatin architecture in single cells |
| HG010646 | Kohli, Rahul | Non-destructive epigenetic sequencing with DNA deaminase enzymes |
| HG010632 | Shendure, Jay | Versatile, exponentially scalable methods for single cell molecular profiling |
| <u>R21</u> | | |
| HG009268 | Bulyk, Martha | Surveying transcription factor pioneer interactions with nucleosomal DNA |
| HG009264 | Churchman, Lee | Global measurement of splicing kinetics |
| HG009274 | Ecker, Joseph | Multidimensional Epigenomic Single Cell Analyses |
| HG009256 | Lu, Chang | Ultrasensitive microfluidic ChIP-MethylC-seq for integrative analysis of histone modification and DNA methylation |
| HG009255 | Powers, Scott | High-throughput genetic interaction sequencing in mammalian cells |
| HG009749 | Buenrostro, Jason | In situ ATAC-seq, a novel technology for structural epigenomics |
| HG009758 | Davis, Ronald | Ultra high-throughput DNA synthesis via nano-optical conveyer belts |
| HG009742 | Farnham, Peggy | Development of a novel promoter tagging technology to identify enhancer targets |
| HG009750 | Mitra, Robi | SINGLE-CELL ANALYSIS OF PIONEER BINDING AND FUNCTION DURING LINEAGE REPROGRAMMING |
| HG009748 | Smibert, Peter | Massively Parallel Multi-Modal Single-Cell Phenotyping Using a Portable Device |
| HG010200 | Bulyk, Martha | AVATAR: highly parallel analysis of variation in transcription factors and their DNA binding sites |
| HG010195 | Lu, Chang | Drop-BS: high-throughput single-cell bisulfite sequencing on a microfluidic droplet platform |
| <u>R43/R44</u> | | |
| HG009768 | Keogh, Michael-Christopher | Next-Generation Detection Reagents for Chromatin Immunoprecipitation |
| HG010237 | Jelinek, Mary | Epi-seq: Multiplexed ChIP-seq for personalized medicine and drug discovery |
| HG009763 | Seligmann, Bruce | TempO-Seq Profiling of RNA Epitranscriptomic Modifications |