

GemCode Library Services

ACCESS LONG-RANGE DATA FROM SHORT-READ ILLUMINA SEQUENCING!!!

- We start with very long gDNA, make GemCode Illumina Library and use the Illumina short read NGS sequencing (HiSeq2500) to get LONG RANGE DATA!
- Linked Reads enable haplotyping, structural variation detection and >12 Mbp haplotype blocks!
- Low gDNA inputs allow accurate human chromosome phasing from ~1ng of gDNA.
- Powerful Software Tools Analysis Pipeline and Visualization Package from 10X Genomics.



Amplicon Express
makes very long gDNA for
maximum results!

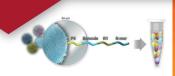




GEMCODE PLATFORM WORKFLOW

HMW gDNA Extraction*

*Amplicon Express performing the gDNA extraction maximizes the length and usefulness of the Linked Reads data.



The GemCode Instrument uses a network of microfluidic channels and pressures to create >100,000 GEMs gel bead-containing emulsions in minutes!

GemCode Library Prep

Library Quantification

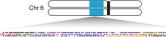
The DNA is delivered "ready to run" on any Illumina HiSeg2500.

Illumina Sequencing

Linked Reads Data Analysis

The GemCode Technology partions gDNA into more than 100,000 fractions, each containing about 0.3 percent of the genome, and has 750,000 different barcodes available.

Linked Reads find the IMPORTANT mutation(s) in your sample!



A 50 Mb phase block from NA12878 within the ${\bf q}$ arm of Chr 6 is shown.

Hap 1

ap 2

Phased Linked-Reads from within the phase block are shown. Haplotype 2 contains a 70 kb deletion as seen by the haplotype-specific loss of coverage.

GemCodes technology is now compatible with Polyploid Genomes!!!!





For orders and enquiries, please contact us:

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