

Reads to Discovery

ChIP-Seq Analysis



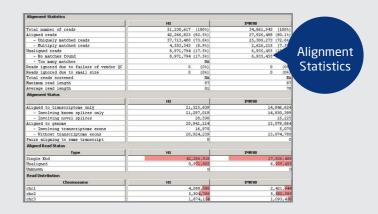
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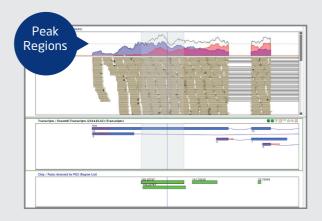
Alignment with Strand NGS aligner

- Handles variable length reads and paired reads as well.
- Allows arbitrary number of gaps and mismatches.
- Options for trimming adaptors, low quality bases, and screening reads against standard screening databases.



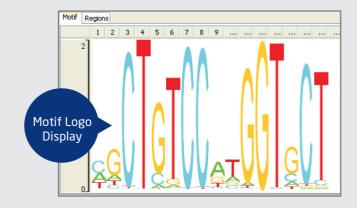
Peak Detection

- Detect enriched regions in ChIP sample as compared to the control.
- Detect peaks of transcription factor regulatory sites using PICS/MACS.
- Identify genes regulated by TF binding sites.
- Detect histone modification sites using the enriched region detection method.



Motif Detection

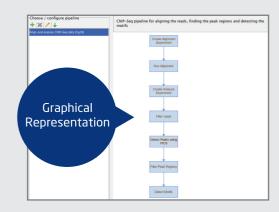
- Identify motifs in detected peak regions using GADEM.
- Import peak regions to detect motifs.
- Import motifs in JASPAR format.
- Scan for motifs in the entire genome or in regions of interest.





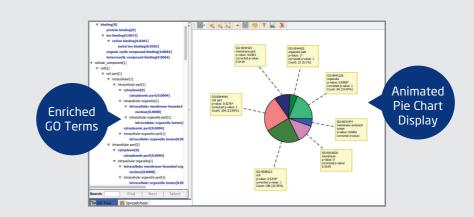
Pipeline Execution

- Pipelines that run in the background.
- Analysis pipeline that includes filters and peak detection.
- Additional pipelines that start from alignment of raw reads or direct import of aligned samples.
- Customization of pipelines for individual runs.



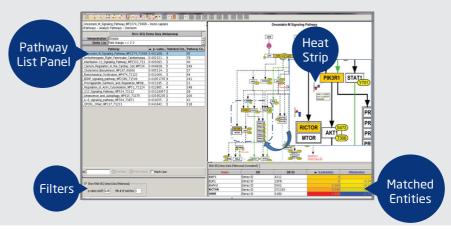
GO Enrichment

- Annotate regions with genic information to discover those affected by peak regions.
- GO Enrichment analysis to detect enriched Gene Ontology terms.



Pathway Analysis

- Single Experiment Analysis (SEA) identifies enriched pathways for the genes from a single experiment type.
- Multi-Omic Pathway Analysis (MOA) from multiple genomics and transcriptomics experiments.
- Overlay differentially expressed entities on curated pathways.
- Choose from curated pathways like WikiPathways, BioCyC pathways, BioPAX pathways or literature-derived networks like NLP and MeSH.
- Find significant pathways for differentially expressed genes.



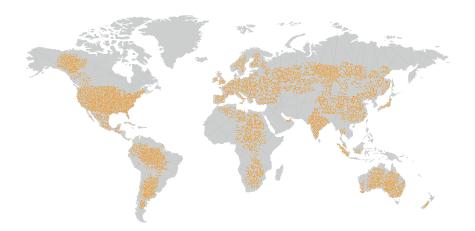


About Strand

A History of Innovative Genomic Research

Strand Life Sciences is a global genomic profiling company and leader in precision medicine diagnostics, aimed at empowering cancer care and genetic testing for inherited diseases. Strand works with physicians and hospitals to enable faster clinical decision support for accurate molecular diagnosis, prognosis, therapy recommendations, and clinical trials. The Strand Center for Genomics & Personalized Medicine is India's 1st and only CAP & NABL accredited NGS laboratory.

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A Trusted Partner to Companies Worldwide

For 15 years, our genomics products and solutions have facilitated the work of leading researchers and medical geneticists in over 2,000 laboratories and 100 hospitals around the world.

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