



ChIP-Seq & MDB-Seq Service

Never Ending Service



BIO-Experiment & Analysis Service

Microarray

- ❖ Gene Expression
- ❖ miRNA
- ❖ CGH / CNV
- ❖ ChIP on chip
- ❖ Methylation
- ❖ Agilent/Affymetrix
- ❖ Exiqon

NGS

- ❖ RNA-Seq
- ❖ small RNA-Seq
- ❖ ChIP-Seq
- ❖ MBD-Seq
- ❖ Exome-Seq
- ❖ Target DNA-Seq
- ❖ Illumina/IonProton

qRT-PCR

- ❖ mRNA
- ❖ microRNA
- ❖ PCR Array
- ❖ Qiagen
- ❖ Genosensor
- ❖ Others

Antibody Array

- ❖ Protein Profiling
- ❖ Phosphorylation
- ❖ FullmoonBio
- ❖ RayBiotech
- ❖ Others

nature COMMUNICATIONS

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NATURE COMMUNICATIONS | ARTICLE

Metabolic engineering of *Corynebacterium glutamicum* for L-arginine production

Seok Hyun Park, Hyun Uk Kim, Tae Yong Kim, Jun Seok Park, Suok-Su Kim


[Affiliations](#) | [Contributions](#) | [Corresponding author](#)

Microarray



❖ Meaningful Data from Any Genome

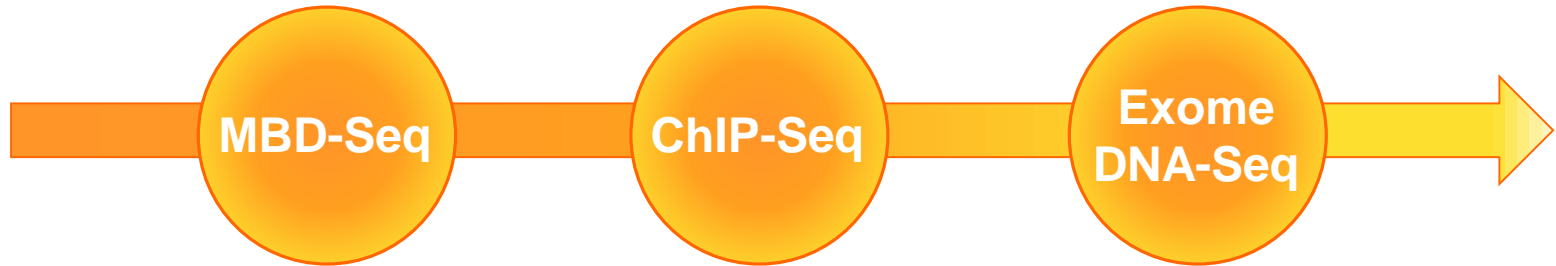
NGS



❖ Next Generation Sequencing



DNA

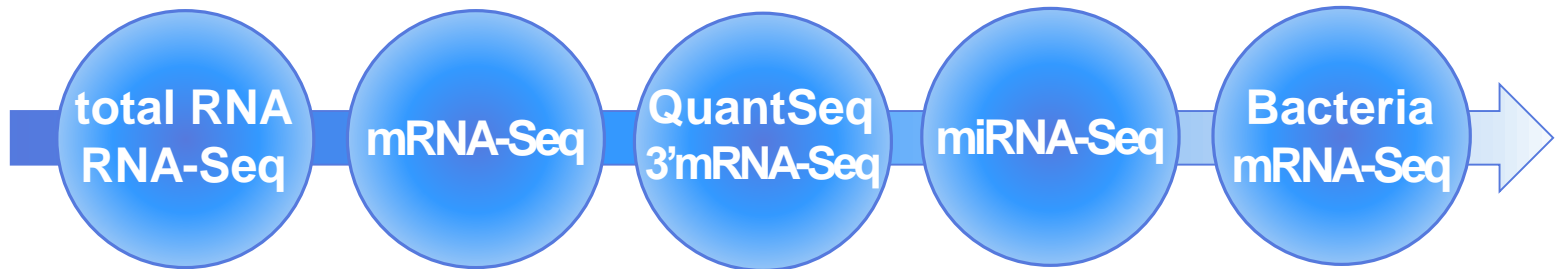


- DNA Methylation Profiling
- Epigenome 분석
- ~ 2 μ g gDNA

- DNA-Protein Interactions
- Histone modifications
- ~20 μ l IP-DNA

- Whole Exome Sequencing
- SNP, CNV, Mutation
- ~ 2 μ g gDNA

RNA



- mRNA-Seq 분석외 lncRNA 분석
- Whole Transcriptome
- ~ 2 μ g total RNA

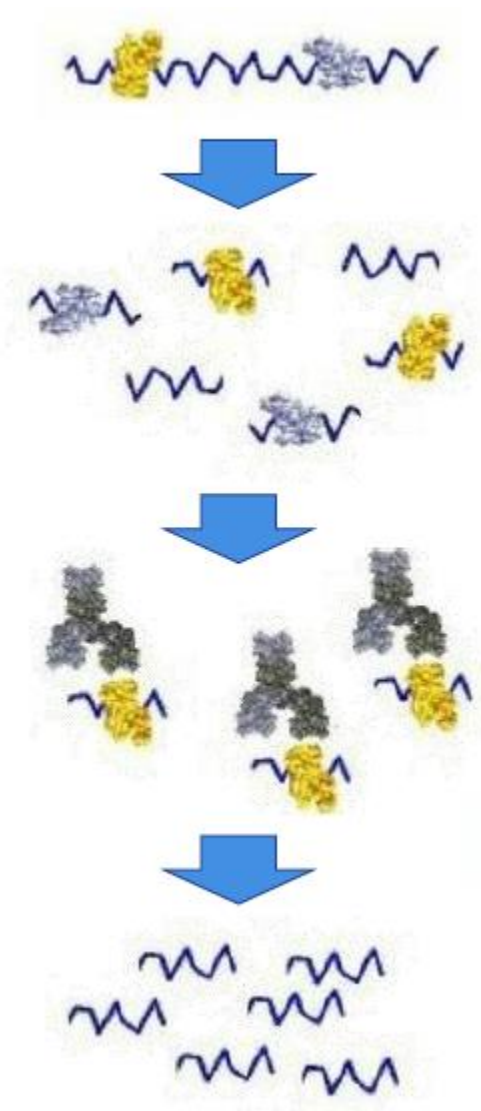
- Gene Expression, Isoform, GO/Pathway
- Splicing, Gene Fusion
- ~ 2 μ g total RNA

- mRNA Expression Profiling
- Low Quantity & Quality RNA 샘플 실험가능
- ~ 2 μ g total RNA

- microRNA Expression Profiling
- Target Gene 분석외
- ~ 2 μ g total RNA

- Gene Expression Profiling
- GO/Pathway/Network
- ~ 2 μ g total RNA

ChIP-Seq Sample Preparation



Cross-link protein on DNA

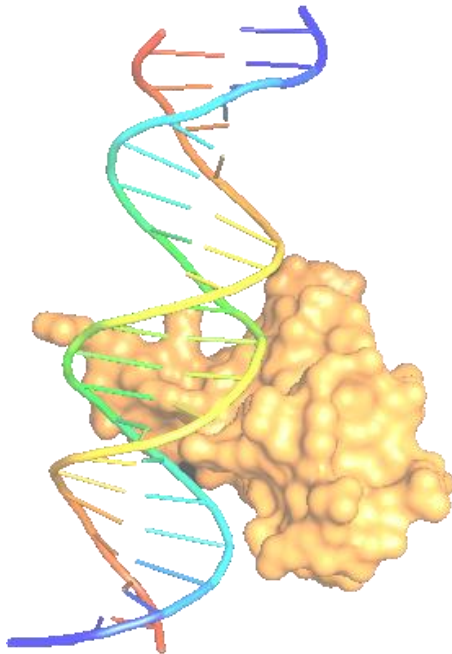
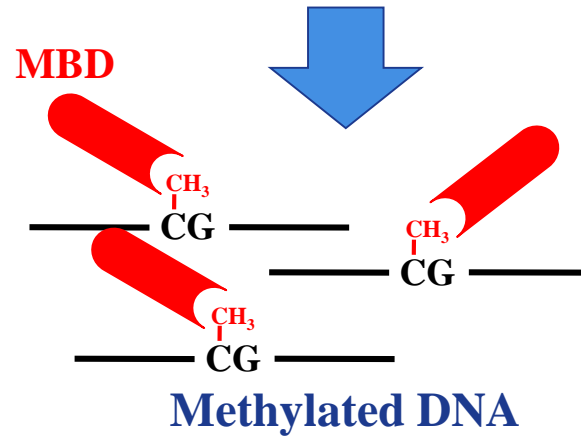
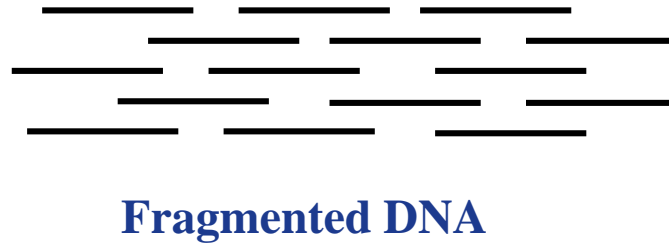
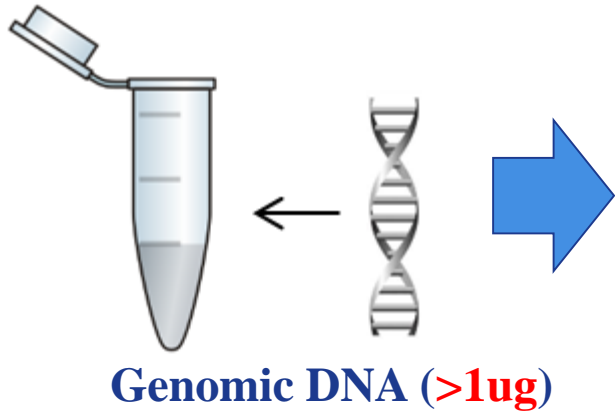
Cut DNA

Immunoprecipitate

Isolate the fragment DNA from complex



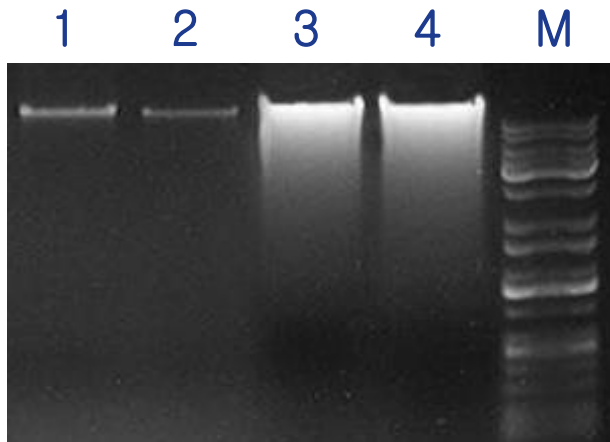
MBD-Seq Sample Preparation



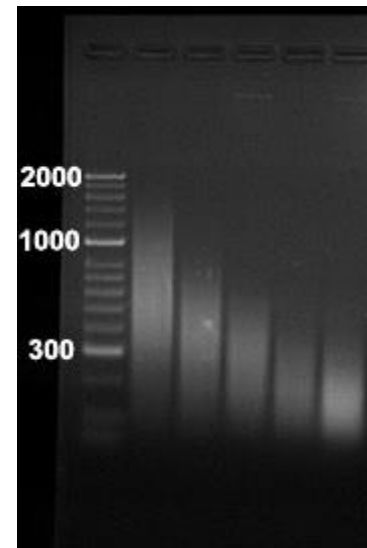
DNA Sample Prep / Shearing



Genomic DNA의 전기영동



Genomic DNA의 Shearing



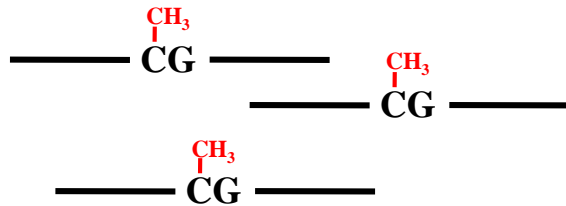
1, 2 : 실험에 좋은 샘플의 예
3, 4 : 사용할 수 없는 샘플의 예

다양한 조건에서 shearing을 진행하여 알맞은 size의 DNA 단편 획득 (300~500nt)

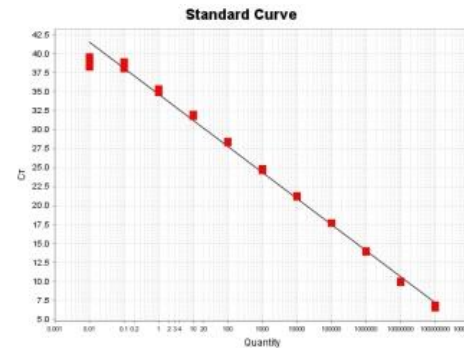
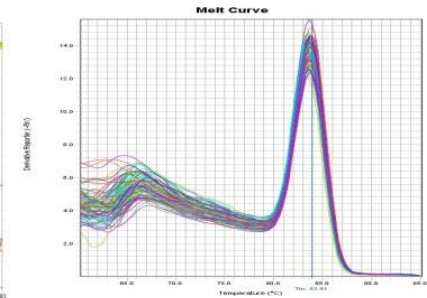
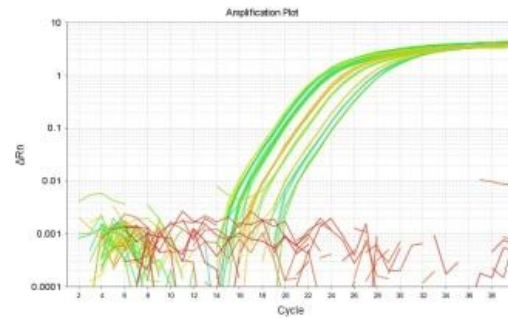
Sample Quality Control (QC)



Methylated DNA



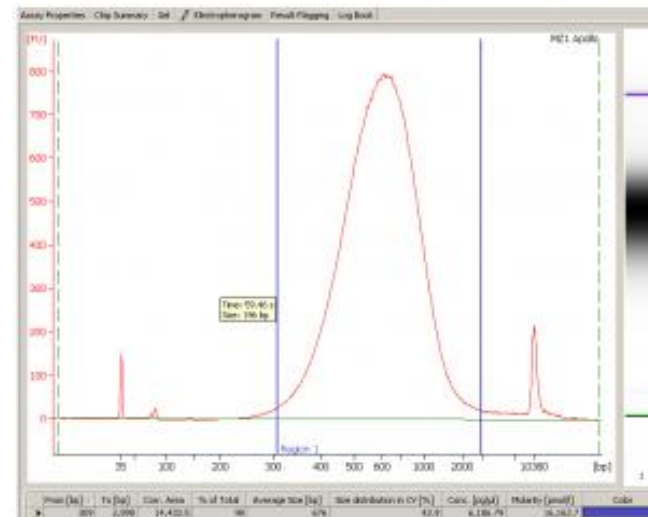
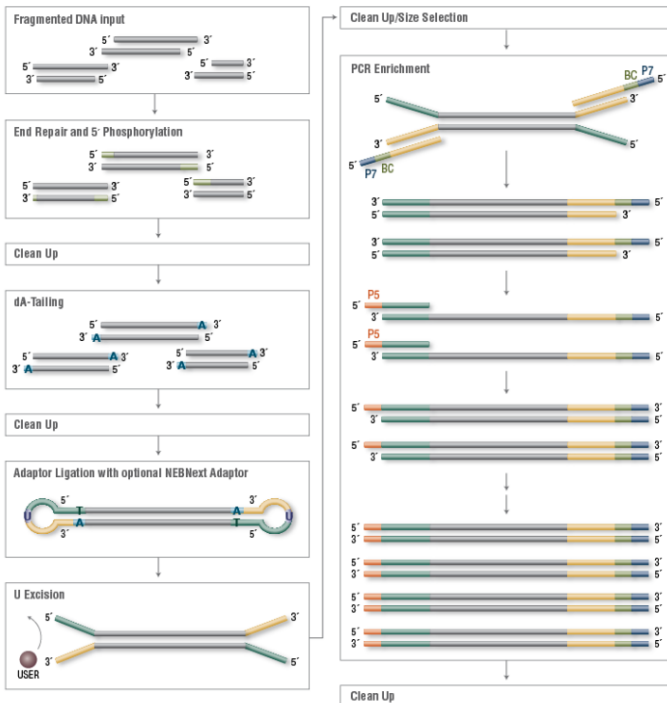
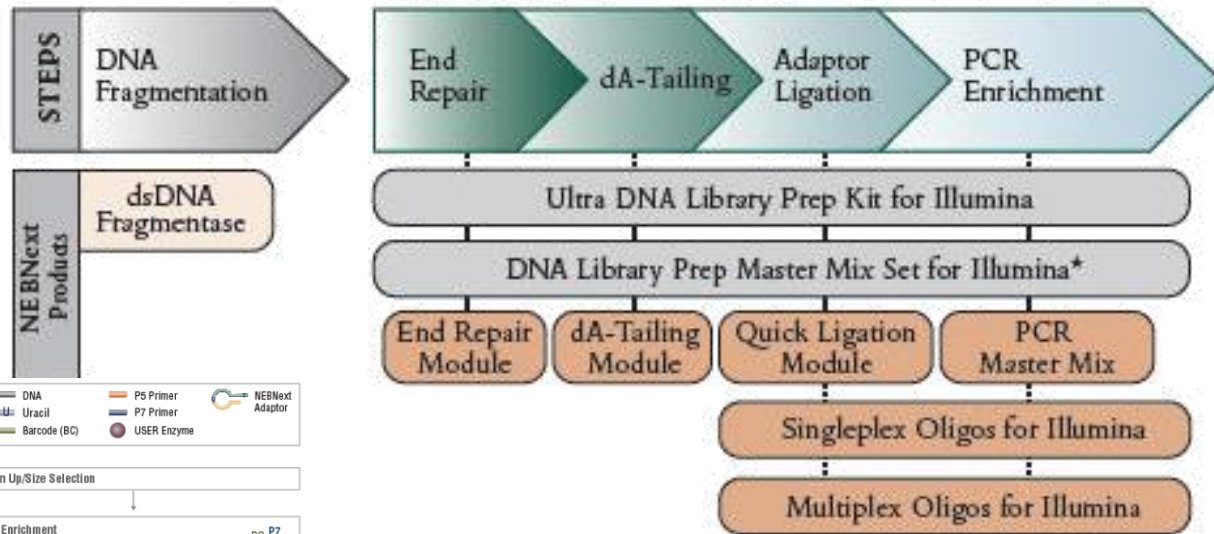
qRT-PCR



Known methylated region/
known unmethylated region 에 대한 qRT-PCR

Quantitative Analysis
Comparative Analysis ($\Delta\Delta C_T$)

Library Generation

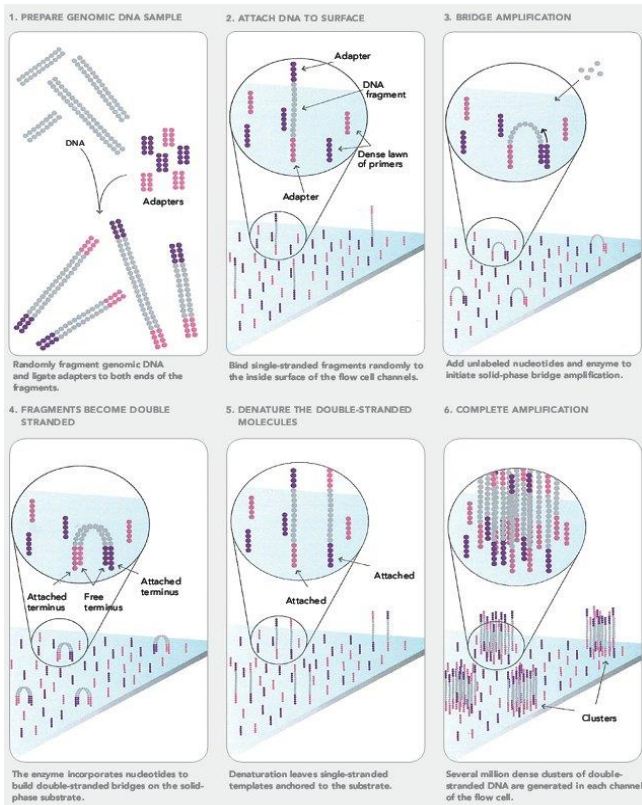


Electropherogram of DNA library

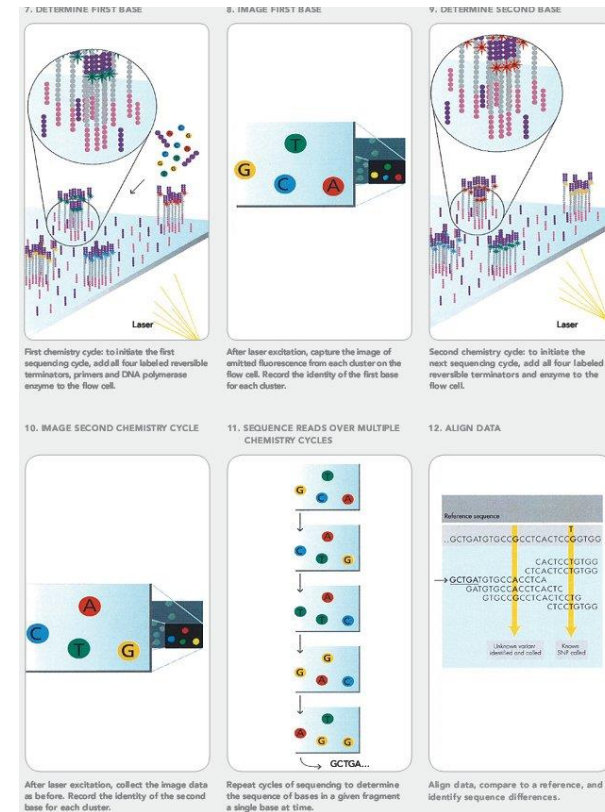
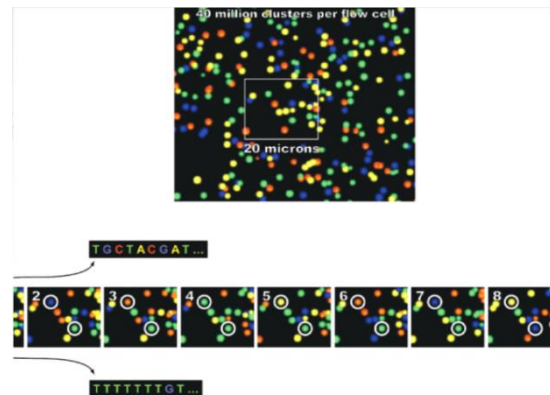
Sequencing



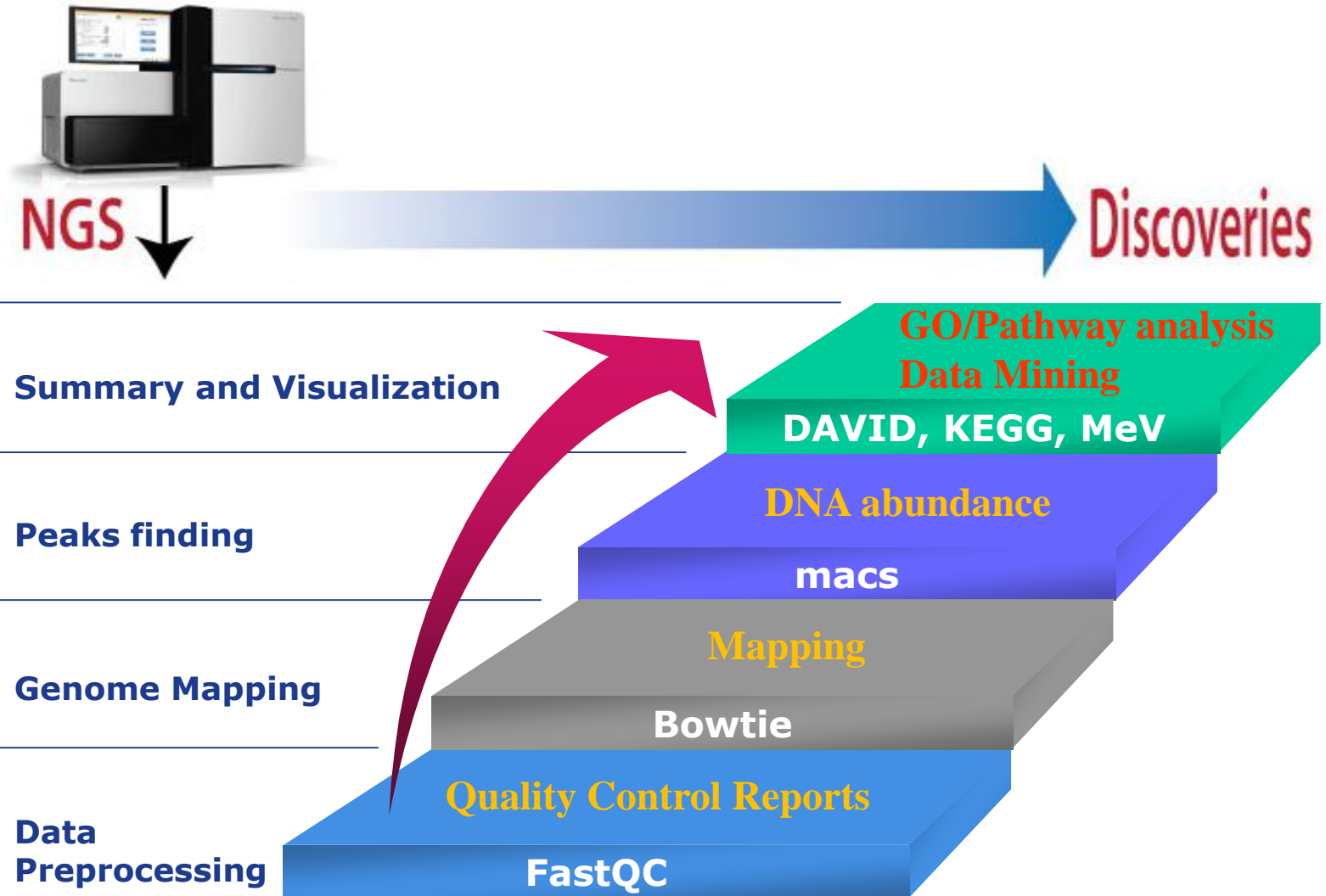
Run Format	Read Length	# of Reads	High-Quality Output (Gb)	
			/lane	/FlowCell
Single End	1 x 50bp	~150 million/lane	7.5 Gb	135-150 Gb
	1 x 100bp		15 Gb	270-300 Gb
Paired End	2 x 50bp	~300 million/lane	15 Gb	270-300 Gb
	2 x 100bp		30 Gb	540-600 Gb



Illumina HiSeq-2500



Work Flow of Data Analysis



Other Data Analysis Tools for Data Processing, Mining, Visualization : MISO, Quick GO, ClueGO, CytoScape, GSEA, UCSC Genome, IGV, IPA, Others

Data Analysis – Data preprocessing

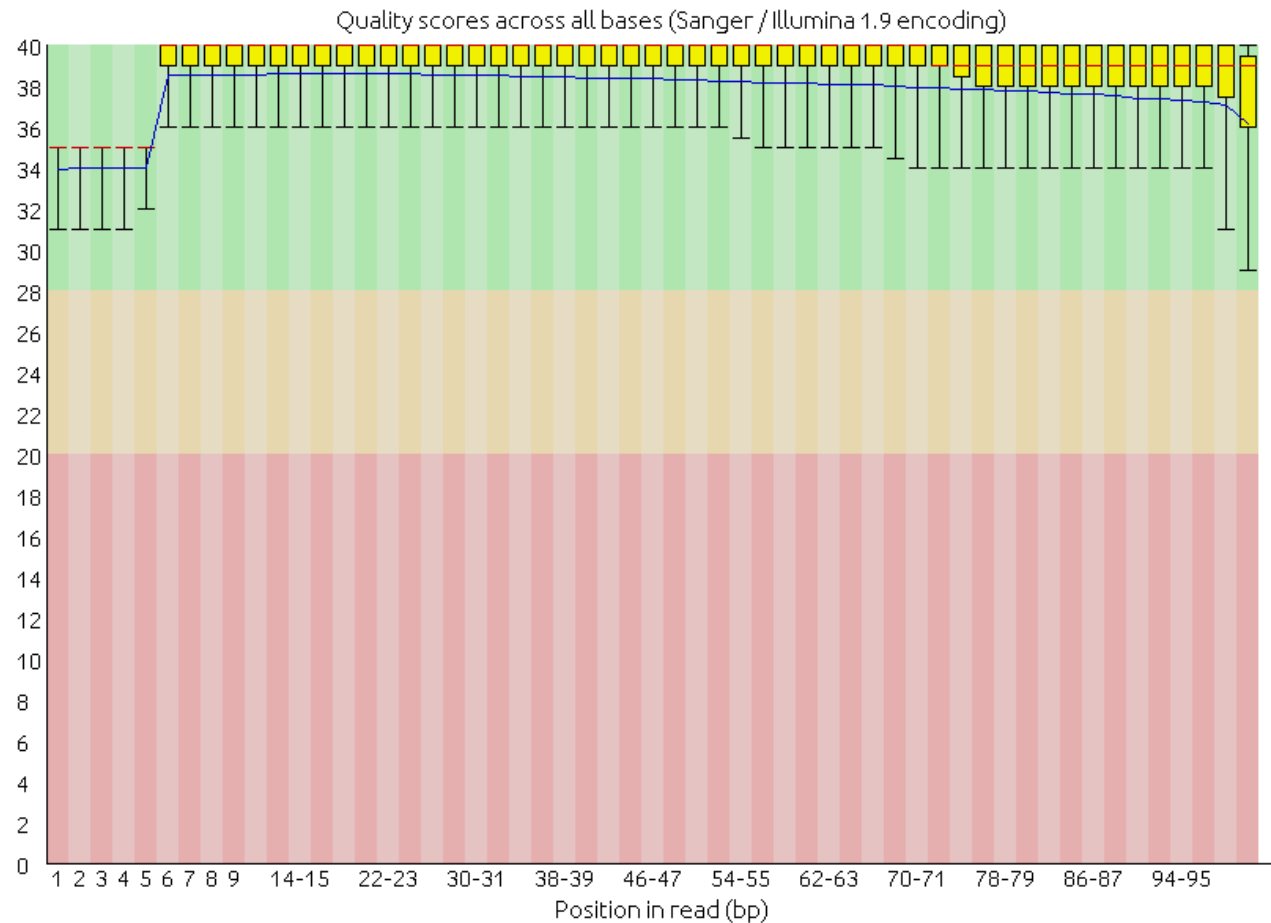


FastQ 파일 안에서 각 position의 base에 대한 quality 값의 영역을 보여 줌

Summary

- ✓ [Basic Statistics](#)
- ✓ [Per base sequence quality](#)
- ✓ [Per tile sequence quality](#)
- ✓ [Per sequence quality scores](#)
- ✗ [Per base sequence content](#)
- ! [Per sequence GC content](#)
- ✓ [Per base N content](#)
- ✓ [Sequence Length Distribution](#)
- ✓ [Sequence Duplication Levels](#)
- ✓ [Overrepresented sequences](#)
- ✓ [Adapter Content](#)
- ✗ [Kmer Content](#)

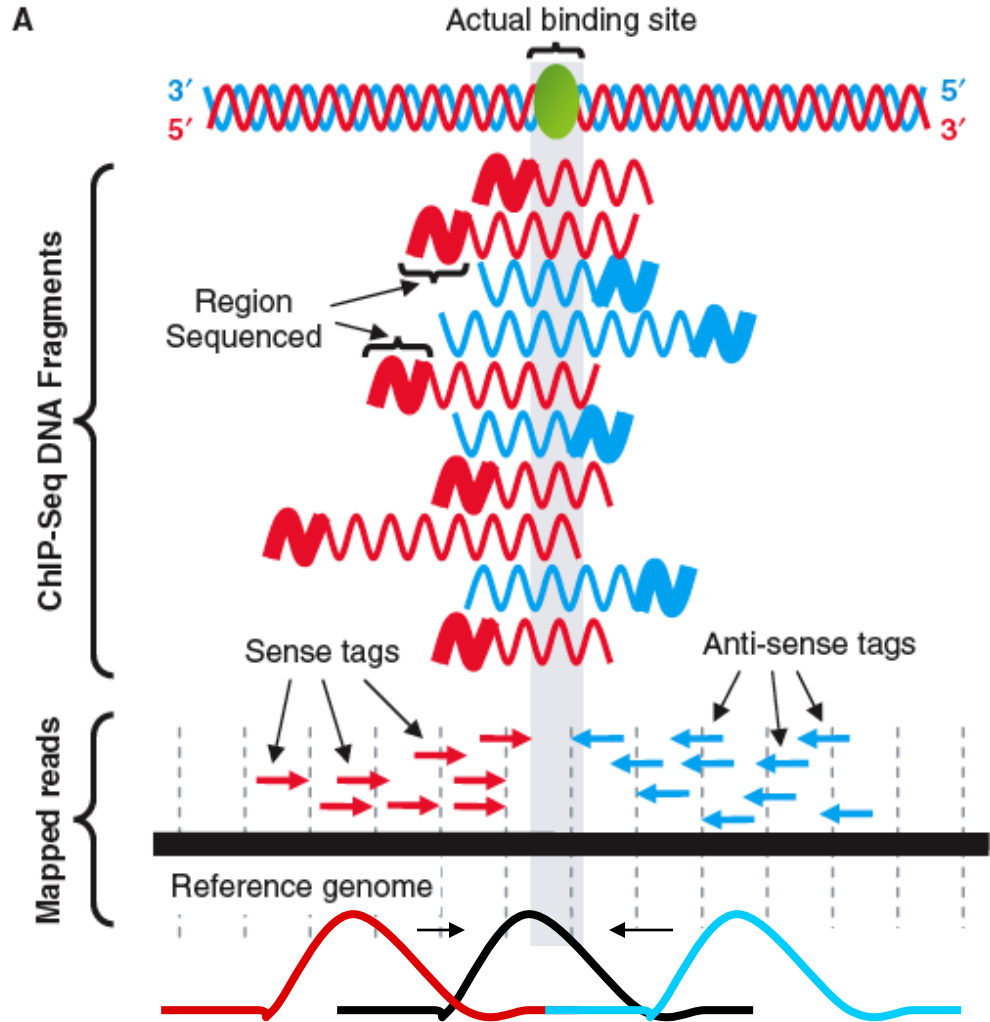
✓ Per base sequence quality



How to find peak

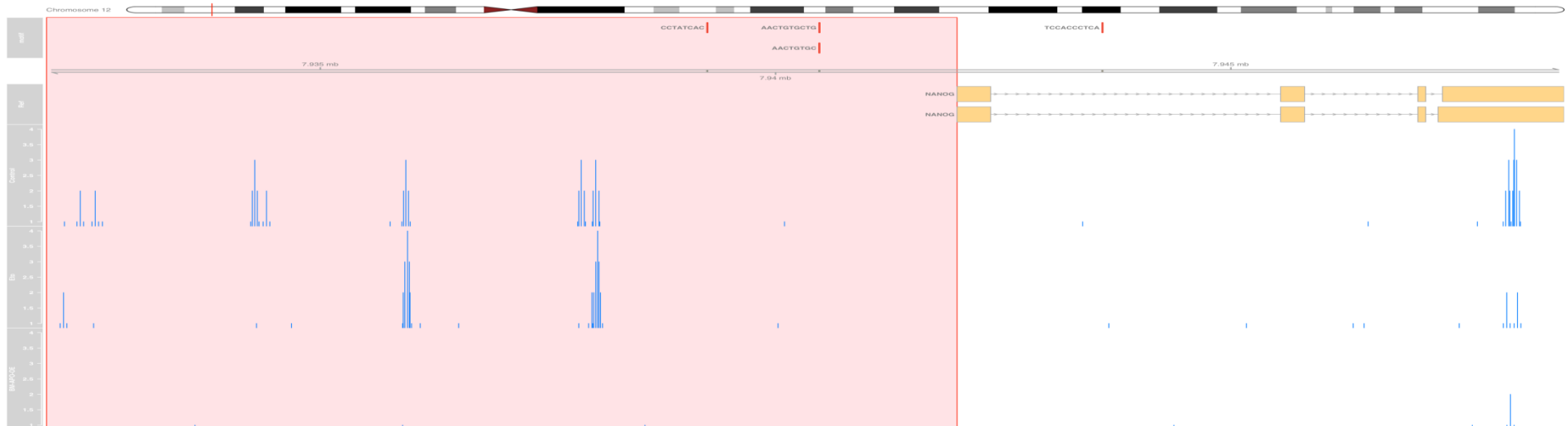
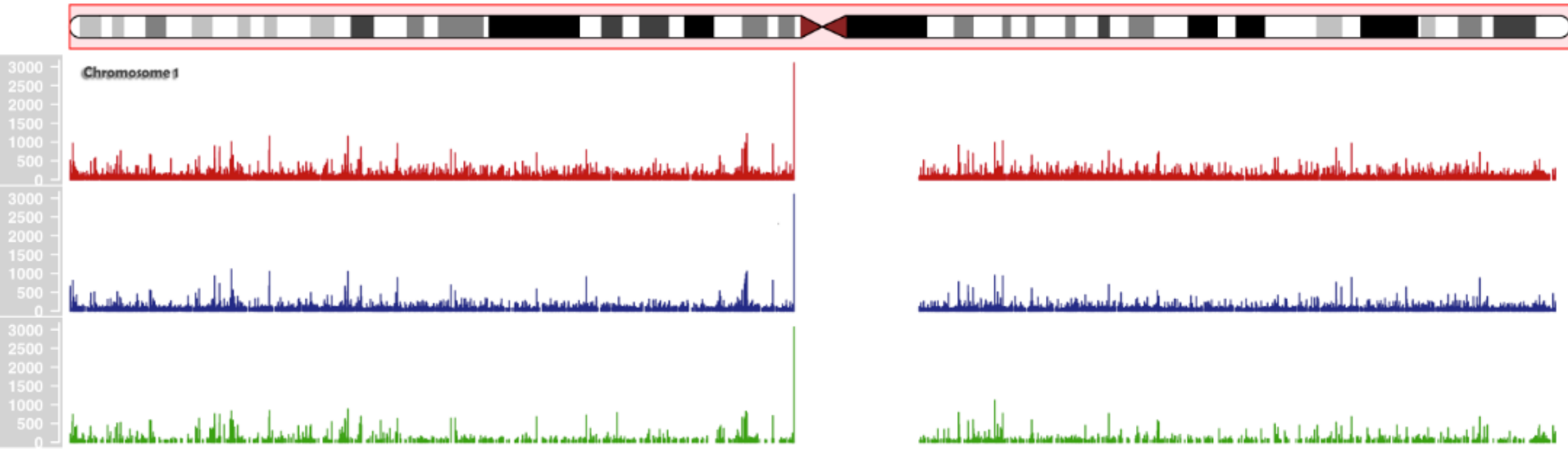


- Fragments contain the TF binding site at a (mostly) random position within them.
- Reads are (randomly) from left or right edges (sense or antisense) of fragments.
- Thus peak for sense tags will be 1/2 the fragment length upstream...
- Binding site position = mid-way between sense tag peak & antisense tag peak.
- To get binding site peak, shift sense downstream by 1/2 fragsize & antisense upstream by 1/2 fragsize.

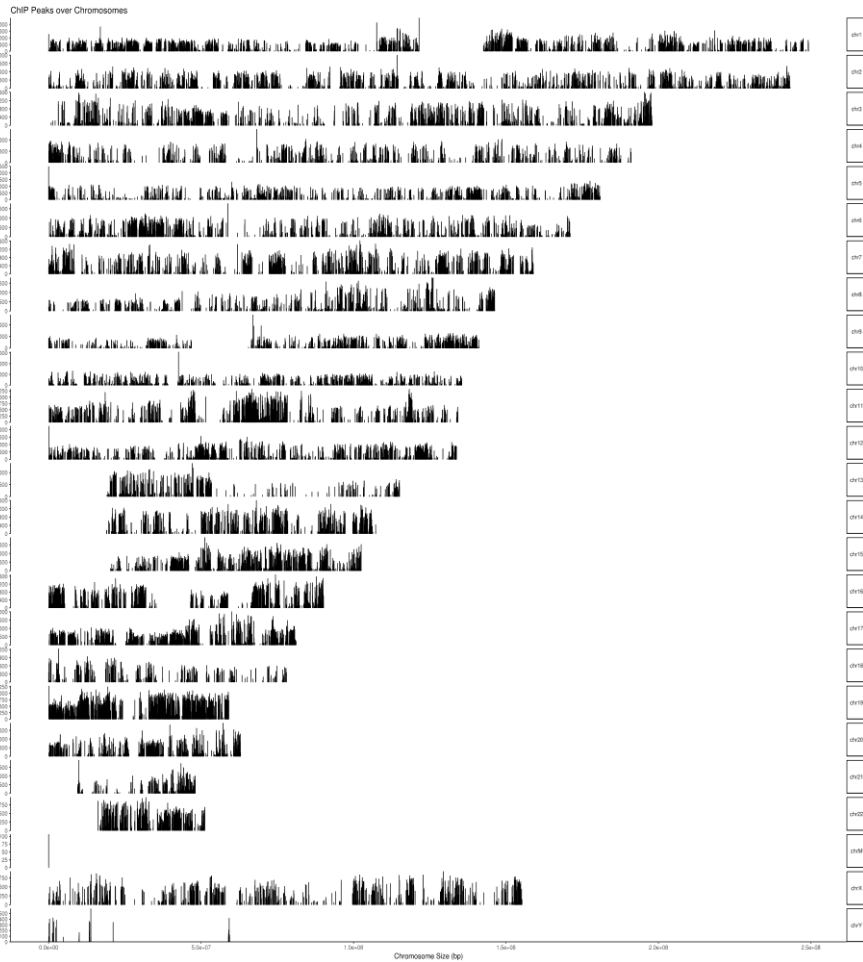


Adapted from slide set by: Stuart M. Brown, Ph.D., Center for Health Informatics & Bioinformatics, NYU School of Medicine & from Jothi, et al. Genome-wide identification of in vivo protein-DNA binding sites from ChIP-Seq data. NAR (2008), 36: 5221-31

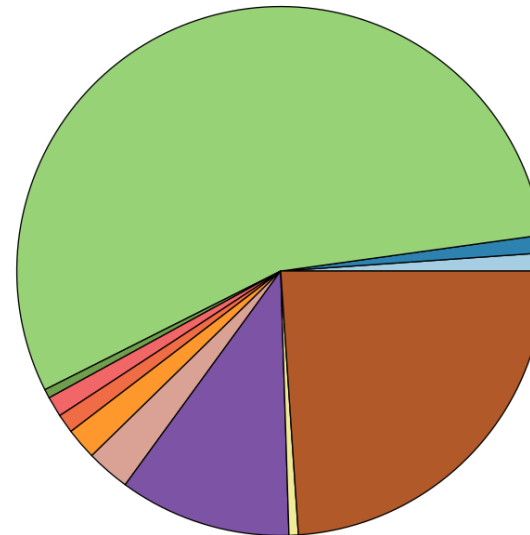
Peak Image



Visualization Data

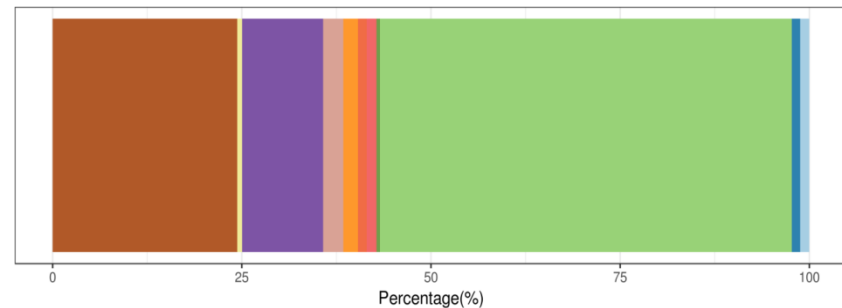


chr1
chr2
chr3
chr4
chr5
chr6
chr7
chr8
chr9
chr10
chr11
chr12
chr13
chr14
chr15
chr16
chr17
chr18
chr19
chr20
chr21
chr22



- Promoter (1-2kb) (1.1%)
- Promoter (2-3kb) (1.08%)
- Promoter (<=1kb) (55.23%)
- 5' UTR (0.55%)
- 3' UTR (1.25%)
- 1st Exon (1.17%)
- Other Exon (1.93%)
- 1st Intron (2.67%)
- Other Intron (10.53%)
- Downstream (<=3kb) (0.57%)
- Distal Intergenic (23.93%)

Feature Distribution



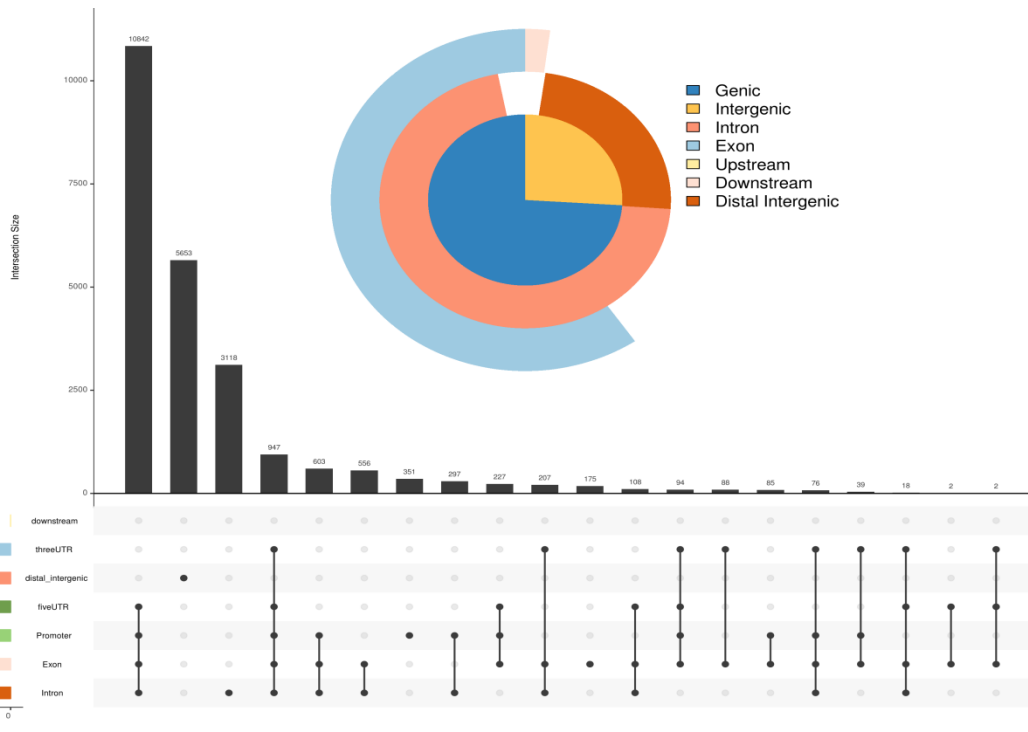
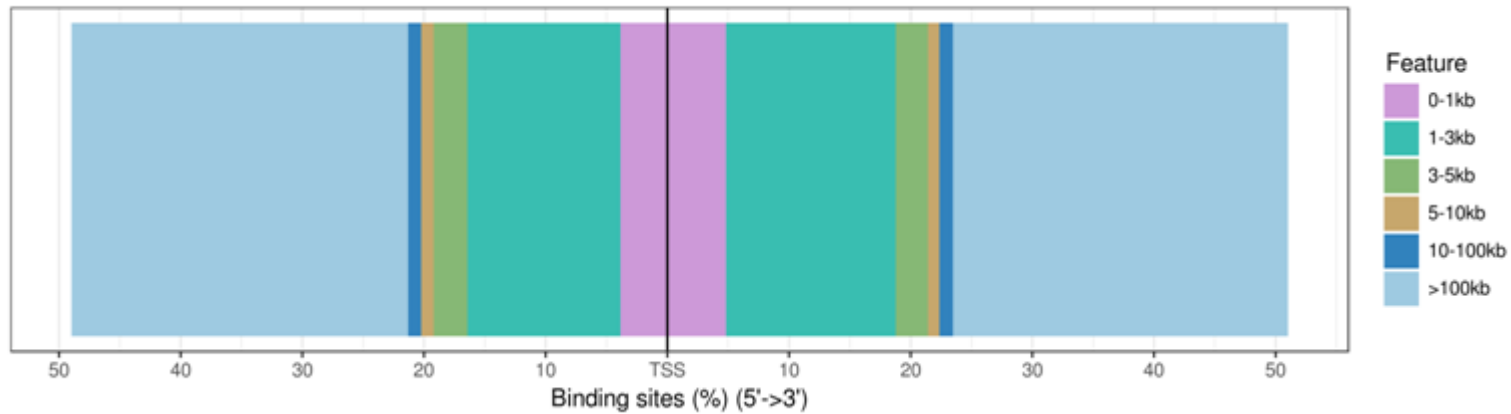
Feature

- Promoter (1-2kb)
- Promoter (2-3kb)
- Promoter (<=1kb)
- 5' UTR
- 3' UTR
- 1st Exon
- Other Exon
- 1st Intron
- Other Intron
- Downstream (<=3kb)
- Distal Intergenic

Visualization Data



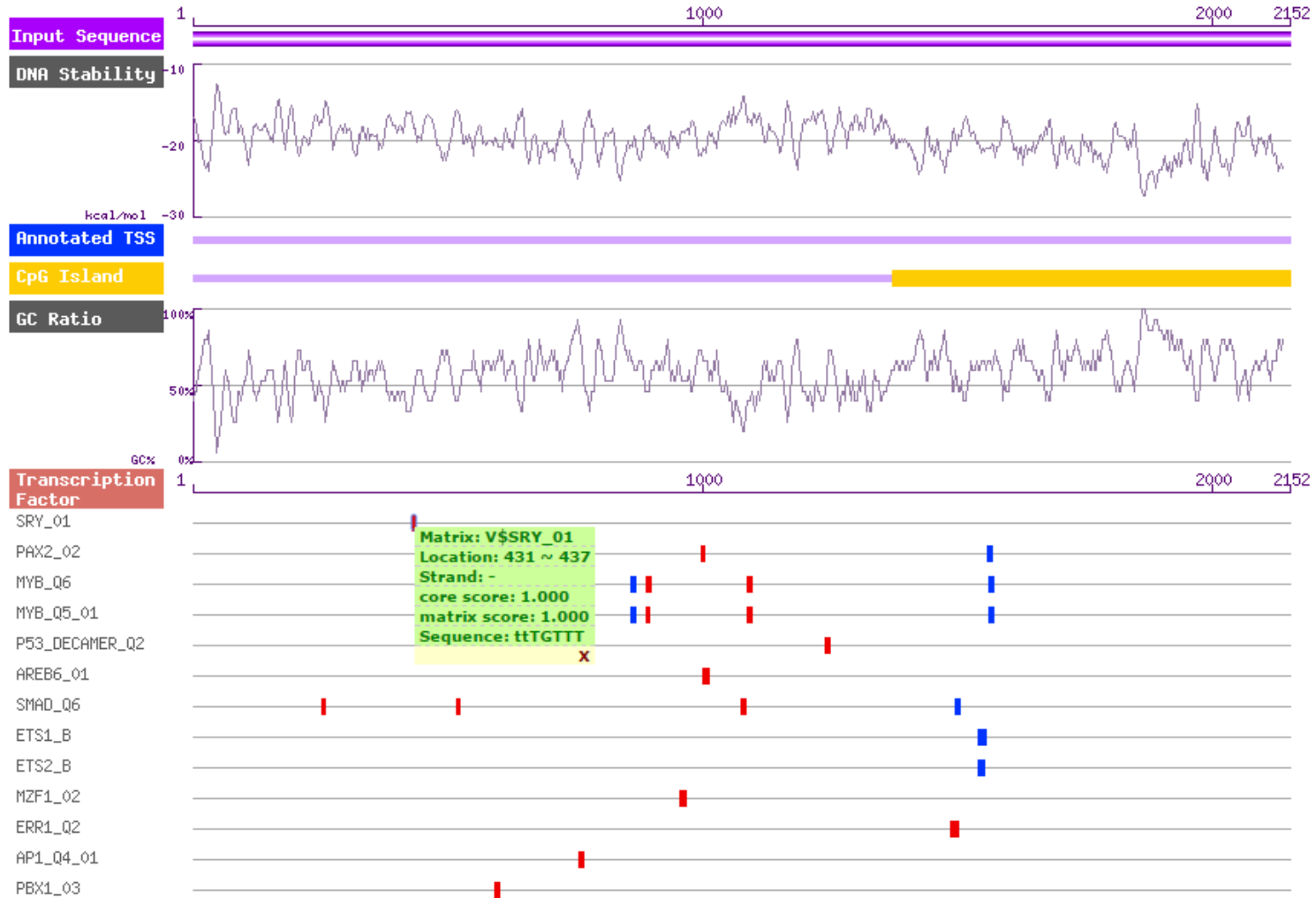
Distribution of transcription factor-binding loci relative to TSS



Transcription Factor Binding



[\[Sequence Homologs\]](#)



DATA ANALYSIS TOOLS



고객을 최우선으로 하고
직원이 즐겁게 일할 수 있는
국내 최고의 바이오 기업으로
성장해 나가겠습니다.

감사합니다.



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