### London Institute of Medical Sciences

# MRC London Institute of Medical Sciences, Imperial College, ICTEM Building 2<sup>nd</sup> floor, Hammersmith Campus, London - <u>Web</u>: http://genomics.lms.mrc.ac.uk

# NGS – Illumina seg

### **HiSeq2500**





### High Output v4 mode or Rapid

- 50-125bp reads, single or paired-end
- 8 lanes per flow cell
- 2 lanes for Rapid flow cell
- up to 400 Gigabases per run
- up to 250 millions reads per lane

### **Applications**

RNA-seq, ChIP-seq, Whole genome, exome seq, Whole methylome seq, Targeted resequencing



Quality score distribution for a 2x100bp High Output Hiseg run of TruSeg Illumina RNA-seq libraries.

### NextSea



### High or Mid Output

- 50-150bp reads, s
- up to 130 or 400 r mid or high output

### Applications

RNA-seq, ChIP-seq, exome seq, Whole m Targeted re-sequenci



Qscore heatmap of a Ne run of a RNA-seq pool c

### NGS – Library



- **DNA-seq** library prep using NEB kits
- **ChIP-seq** library prep using NEB kit
- Amplicon-seq using PCR
- Other applications on demand

### Nanopore seque

### Read Length Histog Summary read length distributio

- Native DNA and RNA sequencing
- Long reads 10-100 kilobases
- Bacterial strain sequencing
- Native methylation marks







# LMS Genomics Laboratory

uencing	platforms		S
<b>1500</b>	MiSeq		Ch • F
Image: Weight of the second	MSeq"	MiSeq flow cell	• S • S • F • F • V
t single or paired-end millions reads per	<ul><li>50-300bp reads, single</li><li>up to 7 Gigabases pe</li><li>up to 25 million reads</li></ul>	or paired end r run s per run	FA
Whole genome, ethylome seq, ing	<section-header><list-item><list-item><list-item><list-item><list-item></list-item></list-item></list-item></list-item></list-item></section-header>	quencing g ng Run quality metrics and Q30 quality plot for a Miseq 2x150bp run of a pool of 96 amplicons prepared by Fluidigm multiplexed PCR method.	Flu 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
prep ser	vice	Restricted to LMS users	
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# ingle-cell analysis

### romium 10X Genomics droplet system

- For targeting 10,000 cells
- Single-cell RNA-seq
- scATAC-seq
- Feature Barcoding technology for cell surface protein expression and CRISPR perturbations
- /isium Spatial transcriptomics





**CS sorted single cells analysis** in 96- or 384-well plates and library preparation using nanoliter volumes on Mosquito HV robot

uidigm microfluidics C1 System and Biomark

### pen access equipment

- **Bioanalyser & Tapestation**
- Nanodrop and Qubit
- Covaris
- **qPCR** machines
- Mosquito HV nanolitre robot
- **Biomek Fx robot**
- Fluidigm Access Array system

### Short-hairpin RNA library clone picking service

We host the **Dharmacon GIPZ Lentiviral short-hairpin RNA (shRNA) libraries for Human and Mouse**. The libraries provide a RNA tool capable of producing RNA interference with powerful viral delivery for targeting highly characterized genes in the human and mouse genomes. Selected clones can be picked and cultured on demand.

# ioinformatics and Computing Support

work closely with the LMS Computing and Bioinformatics group for data management and analysis.

- NGS data processing and analysis QC, demultiplexing, alignment and downstream analysis.
- Dedicated Isilon File store for NGS data storage and data archiving at Imperial HPC
- LMS Bioinformatics contacts: Mahdi Karimi, Marian Dore, Gopu Dharmalingam, Sanjay Khadayate, Yi-Fang Wang

# ontact details

ad: Laurence Game - Lab staff: Ivan Andrew, JP Haywood





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# **Recent Publications**

 Single-cell imaging and RNA sequencing reveal patterns of gene expression heterogeneity during fission yeast growth and adaptation.
Saint M, Bertaux F, Tang W, Sun XM, Game L, Köferle A, Bähler J, Shahrezaei V, Marguerat S. Nat Microbiol. 2019 Mar;4(3):480-491

• Subclonal mutation selection in mouse lymphomagenesis identifies known cancer loci and suggests novel candidates. Webster P, Dawes JC, Dewchand H, Takacs K, Iadarola B, Bolt BJ, Caceres JJ, Kaczor J, Dharmalingam G, Dore M, Game L, Adejumo T, Elliott J, Naresh K, Karimi M, Rekopoulou K, Tan G, Paccanaro A, Uren AG. Nat Commun. 2018 9(1):2649.

 Complex multi-enhancer contacts captured by genome architecture mapping Beagrie RA, Scialdone A, Schueler M, Kraemer DC, Chotalia M, Xie SQ, Barbieri M, de Santiago I, Lavitas LM, Branco MR, Fraser J, Dostie J, Game L, Dillon N, Edwards PA, Nicodemi M, Pombo A. **Nature 2017** 534; 519-524.

## **Imperial College** London