

GUIDE-Seq Results

CCR5 knockout experiment

Generated 04/11/2017

Summary

| Condition | Organism | Cell Type | Genome | Nuclease | Sequencer | Number of Samples |
|---------------|----------|-----------|--------|---------------------|-----------|-------------------|
| CD34-MYC-10nm | Human | U2OS | hg19 | Cas9 (VRER variant) | MiSeq | 1 |
| CD34-MYC-20nm | Human | U2OS | hg19 | Cas9 (VRER variant) | MiSeq | 1 |

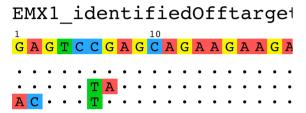
Read Mapping Statistics

| Sample | ample Condition Total Reads | | High Quality Mapped Reads Number of On-target Re | | Number of Off-target Reads | |
|-----------|-----------------------------|------|--|-----|----------------------------|--|
| EMX1-10nm | CD34-MYC-10nm | 4868 | 768 | 489 | 193 | |
| EMX1-20nm | CD34-MYC-20nm | 4868 | 768 | 489 | 193 | |

Sample 1: EMX1-10nm

| C | Cleavage ID | Location | Targeting Status | GUIDE-Seq Reads | Closest Gene | Distance to Closest Gene | Relationship with Closest Gene | Regulatory Region? |
|---|-------------|----------|------------------|------------------------|--------------|--------------------------|--------------------------------|--------------------|
| | | | | | | | | |

Figure

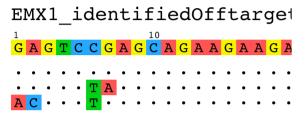


Sequences of off-targets detected by GUIDE-Seq. The full target sequence is displayed in the first row. Each subsequent row indicates a cleaved site detected by GUIDE-Seq. Consensus is represented by a black dot, while nucleotide mismatches are represented by colored squared with the mismatched base. GUIDE-Seq read counts for each cleavage event are listed on the right of each row.

Sample 2: EMX1-20nm

| C | Cleavage ID | Location | Targeting Status | GUIDE-Seq Reads | Closest Gene | Distance to Closest Gene | Relationship with Closest Gene | Regulatory Region? |
|---|-------------|----------|------------------|------------------------|--------------|--------------------------|--------------------------------|--------------------|
| | | | | | | | | |

Figure



Sequences of off-targets detected by GUIDE-Seq. The full target sequence is displayed in the first row. Each subsequent row indicates a cleaved site detected by GUIDE-Seq. Consensus is represented by a black dot, while nucleotide mismatches are represented by colored squared with the mismatched base. GUIDE-Seq read counts for each cleavage event are listed on the right of each row.

Footnotes

Footnotes about where reference data came from goes here