

Supplementary Information

**Double nicking by RNA-guided CRISPR Cas9 for Enhanced
Genome Editing Specificity**

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PRIMER SEQUENCES

SURVEYOR assay

primer name	genomic target	primer sequence (5' to 3')
SUV901	<i>EMX1</i>	CCATCCCCTTCTGTGAATGT
SUV902	<i>EMX1</i>	GGAGATTGGAGACACGGAGA
DYRK1A-F	<i>DYRK1A</i>	GGAGCTGGTCTGTTGGAGAA
DYRK1A-R	<i>DYRK1A</i>	TCCAATCCATAATCCCACGTT
GRIN2B-F	<i>GRIN2B</i>	CAGGAGGGCCAGGAGATTTG
GRIN2B-R	<i>GRIN2B</i>	TGAAATCGAGGATCTGGGCG
F1	<i>VEGFA</i>	CAAAGGACCCAGTCACTCC
R1	<i>VEGFA</i>	GAGGAGGGAGCAGGAAAGTG
F2	<i>VEGFA</i>	GACACTTCCCAAAGGACCCC
R2	<i>VEGFA</i>	TGAGAGCCGTTCCCTCTTTG
F3	<i>VEGFA</i>	GACAGGGGCAAAGTGAGTGA
R3	<i>VEGFA</i>	TTCATGGTTTCGGAGGCC
F4	<i>VEGFA</i>	TGAGTGACCTGCTTTTGGGG
R4	<i>VEGFA</i>	GTTTCATGGTTTCGGAGGCC

Next-generation deep sequencing and HR

primer name	primer sequence (5' to 3')
EMX1-F	GGAGGACAAAGTACAAACGGC
EMX1-R	ATCGATGTCCTCCCCATTGG
EMX1-HR-F	CCATCCCCTTCTGTGAATGT
EMX1-HR-R	GGAGATTGGAGACACGGAGA
EMX1-OT1-F	TGGGAGAGAGACCCCTTCTT
EMX1-OT1-R	TCCTGCTCTCACTTAGACTTTCTC
EMX1-OT2-F	GACATTCCTCCTGAGGAAAA
EMX1-OT2-R	GATAAAATGTATTCCTTCTCACCATTC

EMX1-OT3-F	CCAGACTCAGTAAAGCCTGGA
EMX1-OT3-R	TGGCCCCAGTCTCTCTTCTA
EMX1-OT4-F	CACGGCCTTTGCAAATAGAG
EMX1-OT4-R	CATGACTTGGCCTTTGTAGGA
EMX1-OT5-F	TGGGGTTACAGAAAGAATAGGG
EMX1-OT5-R	TTCTGAGGGCTGCTACCTGT
VEGFA-F	TGAAGCAACTCCAGTCCCAA
VEGFA-R	CCCGGCTCTGGCTAAAGAG
VEGFA-OT1.1-F	TGGGTGTGCACATCTAAGGA
VEGFA-OT1.1-R	CCACTGAGTCAACTGTAAGCA
VEGFA-OT2.1-F	GCAGAGGAATATGTGACATGAGG
VEGFA-OT2.1-R	TACTCCCTGCTGTCTCTCC
VEGFA-OT3.1-F	TTCTGGCCAAGTCGATTCC
VEGFA-OT3.1-R	GGATACCAGCATGGGCTACC
VEGFA-OT4.1-F	ACTCTTGAGATTGGAACGGGA
VEGFA-OT4.1-R	CCACACTTATCTACGCCCA
VEGFA-OT5.1-F	AGCCAGAAGAGAACATCCACG
VEGFA-OT5.1-R	AGTTGCTCTTTGTTGAGAGGGA
MECP2-F	TGACCGGGGACCTATGTATGA
MECP2-R	ACAAGACTTGCTCTTACTTACTTGA

qRT-PCR for Cas9 and sgRNA expression

primer name	primer sequence (5' to 3')
sgRNA reverse-strand synthesis	AAGCACCGACTCGGTGCCAC
EMX1.3 sgRNA qPCR F	AGTCCGAGCAGAAGAAGAAGTTT
EMX1.3 sgRNA qPCR R	TTTCAAGTTGATAACGGACTAGCCT
Cas9 qPCR F	AAACAGCAGATTCGCCTGGA

Cas9 qPCR R	TCATCCGCTCGATGAAGCTC
GAPDH qPCR F	TCCAAAATCAAGTGGGGCGA
GAPDH qPCR R	TGATGACCCTTTTGGCTCCC

Multiplex nicking screening primers

Deletion size (kb)	Forward primer (5' to 3')	Reverse primer (5' to 3')
0.5	AGGTTGTTGCTGTTGCTTTACA	AGCACGGTTAATTTGCATACATCT
1	AGGTTGTTGCTGTTGCTTTACA	TCCAGGCAGTTTTCTTCTGGT
2	AGGTTGTTGCTGTTGCTTTACA	CCAGAGAGCCAGCATTCCAA
6	AGGTTTCACCTGGTTTGGGG	AGGGCTCCCACTAGAAGAGG

VEGFA off-target sites

sgRNA	Off-target site	Off-target sequence (5' to 3')
96	OT-1.1	GGATGGAGGGAGTTTGCTCCTGG
98	OT-2.1	CGCCCTCCCCACCCCGCCTCCGG
98	OT-3.1	TACCCCCACACCCCGCCTCTGG
98	OT-4.1	GGGCCCTCCACCCCGCCTCTGG
98	OT-5.1	CTACCCCTCCACCCCGCCTCCGG

AACAGAACTTCATGCAGCTGATCCACGACGACAGCCTGACCTTTAAAGAGGACATCCAGAAAGCCCAGGTGTCCGGC
CAGGGCGATAGCCTGCACGAGCACATTGCCAATCTGGCCGGCAGCCCCGCCATTAAGAAGGGCATCCTGCAGACAGTG
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CAGACCACCCAGAAGGGACAGAAGAACAGCCGCGAGAGAATGAAGCGGATCGAAGAGGGCATCAAAGAGCTGGGCAGC
CAGATCCTGAAAGAACACCCCGTGGAAAACACCCAGCTGCAGAACGAGAAGCTGTACCTGTACTACCTGCAGAATGGG
CGGGATATGTACGTGGACCAGGAAGTGGACATCAACCGGCTGTCCGACTACGATGTGGACCATATCGTGCCTCAGAGC
TTTCTGAAGGACGACTCCATCGACAACAAGGTGCTGACCAGAAGCGACAAGAACCGGGGAAGAGCGACAACGTGCC
TCCGAAGAGGTGCTGAAGAAGATGAAGAAGTACTGGCCGGCAGCTGCTGAACGCCAAGCTGATTACCCAGAGAAAGTTC
GACAATCTGACCAAGGCCGAGAGAGGCGGCCTGAGCGAAGTGGATAAGGCCGGCTTCATCAAGAGACAGCTGGTGGAA
ACCCGGCAGATCACAAAGCAGTGGCACAGATCCTGGACTCCCGGATGAACACTAAGTACGACGAGAATGACAAGCTG
ATCCGGGAAGTGAAGTGATCACCTGAAGTCCAAGCTGGTGTCCGATTTCCGGAAGGATTTCCAGTTTTACAAAGTG
CGCGAGATCAACAAGTACCACCACGCCACGACGCCTACCTGAACGCCGTGCTGGGAACCGCCCTGATCAAAAAGTAC
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AGCAAAGAGTCTATCCTGCCAAGAGGAACAGCGATAAGCTGATCGCCAGAAAGAAGGACTGGGACCCTAAGAAGTAC
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AAAGAGGTGCTGGACGCCACCCTGATCCACCAGAGCATCACCGGCTGTACGAGACACGGATCGACCTGTCTCAGCTG
GGAGGCGACAAGCGTCTGCTGCTACTAAGAAAGCTGGTCAAGCTAAGAAAAGAAA

> 3xFLAG-NLS-SpCas9n-NLS (the D10A nickase mutation is labeled in red)

ATGGACTATAAGGACCACGACGGAGACTACAAGGATCATGATATTGATTACAAAGACGATGACGATAAGATGGCCCCA
AAGAAGAAGCGGAAGGTGGTATCCACGGAGTCCCAGCAGCCGACAAGAAGTACAGCATCGGCCTGGCCATCGGCACC
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ACCCTGACCAATCTGGGAGCCCCTGCCGCCTTCAAGTACTTTGACACCACCATCGACCGGAAGAGGTACACCAGCACC
AAAGAGGTGCTGGACGCCACCCTGATCCACCAGAGCATCACCGGCCTGTACGAGACACGGATCGACCTGTCTCAGCTG
GGAGGCGACAAGCGTCCTGCTGCTACTAAGAAAGCTGGTCAAGCTAAGAAAAAGAAA

> dsODN ligation insert fwd

AGCAGGCCAATGGGGAGGACATCGATGTCACCTCCAATGACTAcTACCCATACGATGTTCCAGATTACGCTATGGACT
ATAAGGACCACGACGGAGACTACAAGGATCATGATATTGATTACAAAGACGATGACGATAAGaagcttgaaATGGCAT
CAATGCAGAAGCTGATCTCAGAGGAGGACCTGtaa

> dsODN ligation insert rev

TAGTCATTGGAGGTGACATCGATGTCCTCCCCATTGGCCTGCTTTACAGGTCCTCCTCTGAGATCAGCTTCTGCATTG
ATGCCATTTCAAGCTTCTTATCGTCATCGTCTTTGTAATCAATATCATGATCCTTGTAGTCTCCGTCGTGGTCCTTAT
AGTCCATAGCGTAATCTGGAACATCGTATGGGTAG

The RNA-guided Cas9 nuclease can tolerate certain mismatches to the DNA target and thereby promote undesired off-target mutagenesis. Cas9 nickase mutants can be combined with guide RNA pairs to introduce targeted double-strand breaks. Paired nicking reduces off-target activity by 50-1,000 fold in cell lines and facilitates gene knockout in mouse zygotes without sacrificing on-target cleavage efficiency. This versatile strategy enables a wide variety of genome editing applications with higher levels of specificity.