

**Table S14:**

Table lists all sequencing libraries created in this study along with relevant details and mapping parameters.

RNA sample	Library purpose	Primers	Prep kit	Run type	Mapping parameters
N2-CGC B L4-20C (x2)	Gene abundance quantification	Poly(A)	Illumina true seq.	100-single	Tophat2 --b2-very-sensitive -i 30 -l 5000 --read-edit-dist 3 -N 3 --read-realign-edit-dist 0 --segment-length 25 --segment-mismatches 2 --no-coverage-search --min-coverage-intron 30 --max-coverage-intron 5000 --min-segment-intron 30 --max-segment-intron 5000
<i>tdp-1(ok803)</i> L4 20C (x2)	Gene abundance quantification	Poly(A)	Illumina true seq.	100-single	Tophat2 --b2-very-sensitive -i 30 -l 5000 --read-edit-dist 3 -N 3 --read-realign-edit-dist 0 --segment-length 25 --segment-mismatches 2 --no-coverage-search --min-coverage-intron 30 --max-coverage-intron 5000 --min-segment-intron 30 --max-segment-intron 5000
N2-CGC B L4-20C	Alternative splicing analysis	Poly(A)	Illumina true seq.	76-Paired-high depth	Tophat2 --b2-very-sensitive -r 198 --mate-std-dev 18 -i 30 -l 5000 --read-edit-dist 3 -N 3 --read-realign-edit-dist 0 --segment-length 25 --segment-mismatches 2 --no-coverage-search --min-coverage-intron 30 --max-coverage-intron 5000 --min-segment-intron 30 --max-segment-intron 5000
<i>tdp-1(ok803)</i> L4-20C	Alternative Splicing Analysis	Poly(A)	Illumina true seq.	76-Paired-high depth	Tophat2 --b2-very-sensitive -r 195 --mate-std-dev 18 -i 30 -l 5000 --read-edit-dist 3 -N 3 --read-realign-edit-dist 0 --segment-length 25 --segment-mismatches 2

					--no-coverage-search --min-coverage-intron 30 --max-coverage-intron 5000 --min-segment-intron 30 --max-segment-intron 5000
<b>N2-CGC B L4-20C (x3)</b>	A-to-I RNA editing analysis	random primed	Illumina true seq.	100-Paired	Bfast – See <b>Supplemental Table 16</b>
<b><i>tdp-1(ok803)</i> L4 20C (x3)</b>	A-to-I RNA editing analysis	random primed	Illumina true seq.	100-Paired	Bfast – See <b>Supplemental Table 16</b>
<b><i>adr-2(gv42)</i> L4 20C</b>	A-to-I RNA editing analysis	random primed	Illumina true seq.	100-Paired	Bfast – See <b>Supplemental Table 16</b>
<b>N2-CGC Young Adult-15C input RNA and J2-IP'ed RNA x3)</b>	J2-IP enrichment identification	random primed	scriptseq v2 strand specific	100-single	<i>Tophat --b2-very-sensitive -i 30 -l 5000 --read-edit-dist 3 -N 3 --read-realign-edit-dist 0 --segment-length 25 --segment-mismatches 2 --no-coverage-search --min-coverage-intron 30 --max-coverage-intron 5000 --min-segment-intron 30 --max-segment-intron 5000.</i>
<b><i>tdp-1(ok803)</i> Young adult (Input RNA and J2-IP'ed RNA x3)</b>	J2-IP enrichment identification	random primed	scriptseq v2 strand specific	100-single	<i>Tophat --b2-very-sensitive -i 30 -l 5000 --read-edit-dist 3 -N 3 --read-realign-edit-dist 0 --segment-length 25 --segment-mismatches 2 --no-coverage-search --min-coverage-intron 30 --max-coverage-intron 5000 --min-segment-intron 30 --max-segment-intron 5000.</i>
<b>N2- CGC B Young Adult-20C (x2 and X1 with RNase)</b>	Identification of TDP-1 sites of co-transcriptional association	-	ChIP-Seq DNA Sample Prep Kit (IP-102-1001)	50-single	bowtie -l 47 -n 1 -y --best

<p><b>Human M17 cells mock treated or treated with Tardbp siRNAs</b></p>	<p>J2-IP enrichment identification</p>	<p>random primed</p>	<p>Scriptseq v2 strand specific (RT 55-59C)</p>	<p>100- Paired</p>	<p>Trim 1 off 5', 10 off 3', filter 20  tophat2 --b2-very-sensitive -N 2 --read-realign-edit-dist 0 --library-type frsecondstrand -p 10 -r 157 --mate-std-dev 50 --segmentlength 25 --segmentmismatches 2 --nocoverage-search</p>
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