

We have provided the raw counts as supplementary files in GEO under accession ID [GSE112037](#)

Supplementary file	Size	Download	File type/resource
GSE112037_RASL_count.txt.gz	66.2 Kb	<a href="#">(ftp)</a> <a href="#">(http)</a>	TXT
GSE112037_darts_read_counts.txt.gz	2.9 Mb	<a href="#">(ftp)</a> <a href="#">(http)</a>	TXT
GSE112037_kallisto.tar.gz	16.7 Mb	<a href="#">(ftp)</a> <a href="#">(http)</a>	TAR

The first two files are junction counts for RASL-seq and RNA-seq, respectively.

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In the RASL-seq count file, each column is a sample (e.g. PC3E\_rep1), and each row is an event. Each entry represents the inclusion counts and skipping counts separated by a comma (e.g. 20,19 means inclusion counts I=20, skipping counts S=19). See the header snippet of RASL count file below.

ID	PC3E_rep1_RASL	PC3E_rep2_RASL	PC3E_rep3_RASL	GS689_rep1_RASL	GS689_rep2_RASL	GS689_rep3_RASL
chr6:+:130370538:130370900:130370975:130372393	20,19	11,18	23,18	30,24	28,27	15,13
chr13:+:96377506:96409897:96410050:96412293	1504,0	1215,2	1795,3	1960,2	2235,5	1193,2
chr3:+:66396832:66413283:66413353:66419901	70,1	86,4	77,1	80,2	72,0	46,1
chr6:+:43470087:43470325:43470355:43471138	0,475	0,629	2,502	0,439	0,416	5,314

RASL-PSI is calculated as  $I/(I+S)$ . This is because only one inclusion junction is profiled in RASL-seq. Refer to the Method part in our paper (PubMed: 30923373) for the details in PSI calculation.

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The RNA-seq count file follows the DARTS-BHT format. I1,S1 are inclusion and skipping junction counts from PC3E triplicates; I2,S2 are inclusion and skipping junction counts from GS689 triplicates. Last two columns are the effective lengths for inclusion and skipping counts.

ID	I1	S1	I2	S2	inc_len	skp_len
chr16:+:19125446:19125962:19126257:19131428	2,8,9	0,0,0	0,5,3	0,1,0	202	101
chr12:+:118454697:118455494:118455620:118455799	23,28,24	1,3,0	20,13,10	4,2,0	202	101
chr12:+:118454697:118455494:118455620:118456876	31,36,30	618,392,679	35,21,10	408,330,374	202	101
chr12:+:118454697:118455494:118455657:118455799	25,23,24	1,3,0	20,14,7	4,2,0	202	101
chr12:+:118454697:118455494:118455657:118456876	23,23,24	618,392,679	18,13,7	408,330,374	202	101

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In both files, the ID column is the unique identifiers for exon-skipping events in hg19 genome. The coordinates in the ID are joint by colons in the order of:

chromosome, strand, upstreamExonEnd, cassetteExonStart, cassetteExonEnd, downstreamExonStart.

The two count files can be inner-joint using the ID columns.