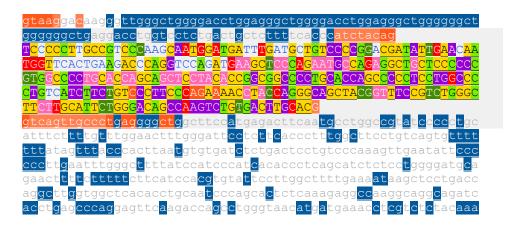
### Ejemplo comparación de resultados predictores in sillico

Cambio de estudio TP53 c.375+1G>A (chr17:7675993 G/A, rs15675554450 NM\_000546.6: c.375+1G>A)

## Exón 4 e intrones adyacentes:



El cambio se encuentra en primera posición justo después del exón 4 (la g en color rojo).

Se va a obtener los resultados que produce analizar esta variable con los diferente predictores y ver cuál de ellos es más preciso.

### NetGene2

Donor splice s	sites, direct				Donor splice sites, direct strand				
	pos 5'->3' 389	phase stra 0 +	nd confidence 0.70	5' exon intron 3' GACTTGCACG^GTCAGTTGCC				hold.	
Donor splice s					Donor splice				
			nd confidence 0.71	5' exon intron 3' AGCTGCCCTG^GTAGGTTTTC			phase strand 2 -	confidence 0.71	5' exon intron 3' AGCTGCCCTG^GTAGGTTTTC
Acceptor spli	e sites, di	rect strand			Acceptor spli	ce sites, di	rect strand		
	pos 5'->3'	phase stra	nd confidence	5' intron exon 3'		pos 5'->3'	phase strand	confidence	5' intron exon 3'
	109	0 +	0.96	CCATCTACAG^TCCCCCTTGC		109	0 +	0.96	CCATCTACAG^TCCCCCTTGC
	128	1 +	0.25	CCGTCCCAAG^CAATGGATGA		128	1 +	0.25	CCGTCCCAAG^CAATGGATGA
	313	0 +	0.93	CCCTTCCCAG^AAAACCTACC		313	0 +	0.93	CCCTTCCCAG^AAAACCTACC
	325	0 +	0.07	AACCTACCAG^GGCAGCTACG		325	0 +	0.07	AACCTACCAG^GGCAGCTACG
	609	2 +	0.15	ACACCCTCAG^CATCTCTCCT		609	2 +	0.15	ACACCCTCAG^CATCTCTCCT
Acceptor splic	ce sites, cor	mplement str	and		Acceptor spli	ce sites, con	mplement strand	ı	
	•		nd confidence			•	phase strand		
360	449	0 -	0.64	GCTGTCCCAG^AATGCAAGAA	360	449	0 -	0.65	GCTGTCCCAG^AATGCAAGAA
43	766	0 -	0.00	AGCCCTCCAG^GTCCCCAGCC	43	766	0 -	0.00	AGCCCTCCAG^GTCCCCAGCC

Se está perdiendo un sitio *donor* en la secuencia mutante. Este coincide con el sitio *donor* del exón 4, por lo que estará afectando al *splicing*, probablemente perdiéndose el exón completo ya que no se detecta otro sitio *donor* en la hebra directa.

## **Splice Site Prediction by Neural Network (NNSplice)**

## Donor site predictions for 10.42.0.139.315469.0:

## Donor site predictions for 10.42.3.123.315859.0:

Start	End	Score	Exon	Intron
382	396	0.59	ttgcad	ggtcagttg

Start End Score Exon Intron

# Acceptor site predictions for 10.42.0.139.315469.0:

# Acceptor site predictions for 10.42.3.123.315859.0:

Start	End	Score	Intron	Exon	Start	End	Score	Intron	Exon
89	129	0.64	gctcttttcaccca	atctac <b>ag</b> tcccccttgccgtcccaagc	89	129	0.64	gctcttttcaccca	tctac <b>ag</b> tcccccttgccgtcccaagc
293	333	0.89	tcatcttctgtcc	cttccc <b>ag</b> aaaacctaccagggcagcta	293	333	0.89	tcatcttctgtccc	ttccc <b>ag</b> aaaacctaccagggcagcta
495	535	0.99	cctgtcagtgttt	ttttat <b>ag</b> tttacccacttaatgtgtga	495	535	0.99	cctgtcagtgtttt	tttat <b>ag</b> tttacccacttaatgtgtga

Se está perdiendo un sitio *donor* en la secuencia mutante. Este coincide con el sitio *donor* del exón 4, por lo que estará afectando al *splicing*, probablemente perdiéndose el exón completo ya que no se detecta otro sitio *donor* en la hebra directa.

## **Spliceman**

gcacg(g/a)tcagt ggtcag gatcag 31909	Kanking (L1)
geacg(g/a)icagt	52%

# **Human Splicing Finder**

Broken WT Donor	Alteration of the WT Donor site, most probably affecting splicing
Site	

Algorithm/Matix	position	sequences	variation
HSF Donor site (matrix GT)	chr17:7675996	- REF : ACGGTCAGT - ALT : ACGATCAGT	87.47 > 60.33 => -31.03%
MaxEnt Donor site	chr17:7675996	- REF : ACGGTCAGT - ALT : ACGATCAGT	8.02 > -0.16 => -102%

## **SVM-BPfinder**

seq_id	agez	ss_dist	bp_seq bp_scr	y_cont ppt_off	ppt_len ppt_scr	svm_scr			
wt	37	78	ctgtgactt	1.28093191374	0.602739726027	58	16	26	-2.3311001
wt	37	66	cggtcagtt	-1.00825174026	0.606557377049	46	16	26	-2.4666109
wt	37	56	ccctgaggg	0.492741774974	0.607843137255	36	16	26	-1.2455025
wt	37	39	ccatgagac	-0.948831341482	0.676470588235	19	16	26	-0.71171041
wt	37	32	acttcaatg	-2.14672886083	0.703703703704	12	16	26	-0.72886012
mut	37	78	ctgtgactt	1.28093191374	0.602739726027	58	16	26	-2.3311001
mut	37	66	cgatcagtt	-2.03508075528	0.606557377049	46	16	26	-2.868663
mut	37	56	ccctgaggg	0.492741774974	0.607843137255	36	16	26	-1.2455025
mut	37	39	ccatgagac	-0.948831341482	0.676470588235	19	16	26	-0.71171041
mut	37	32	acttcaatg	-2.14672886083	0.703703703704	12	16	26	-0.72886012

El cambio que se da entre ambas secuencias está en el BP que contiene la posición de interés. Sin embargo, como las puntuaciones son negativas, no se va a tener en cuenta.

## **Variant Effect Predictor tool**

ENST00000445888 6 c 375-10-A 17.767.993- T	ENST00000445888 &c.375-1G-A 177675932 T											
ENST0000044588 6 c. 375+1G-A 17.7675992 T Ispice_donor_variant TP53 ENSG0000141510 Transcript ENST000044588 6 protein_coding - 1159755454	ENST00000445888 & c. 375+1G-A 17.7675893 T			splice_donor_variant	TP53	ENSG00000141510	Franscript	ENST00000269305.9	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,
ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG0000141510   Transcript   ENST00000420246.6   protein_coding   1158755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG0000141510   Transcript   ENST0000045888.6   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG0000141510   Transcript   ENST0000045888.6   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG0000141510   Transcript   ENST00000505283.6   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG0000141510   Transcript   ENST00000505293.1   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   upstream_gene_variant   TP53   ENSG00000141510   Transcript   ENST00000504290.5   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   upstream_gene_variant   TP53   ENSG00000141510   Transcript   ENST00000504290.5   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   upstream_gene_variant   TP53   ENSG00000141510   Transcript   ENST00000504997.5   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   upstream_gene_variant   TP53   ENSG00000141510   Transcript   ENST00000504997.5   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG00000141510   Transcript   ENST0000050993.5   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675992. T   spice_donor_variant   TP53   ENSG00000141510   Transcript   ENST0000050993.5   protein_coding   1159755445.    ENST00000445888.6 c: 375+1G-A 177675993. T   intron_variant   TP53   ENSG00000141510   Transcript   ENST0000050993.5   protein_coding   1159755445.    ENST000000445888.6 c: 375+1G-A 177675993. T   intron_variant   TP53   ENSG00000141510   Transcript   ENST0000050993.5   protein_coding   1159755445.    ENST000000445888.	ENST00000445886 6c.375+1G-A 17.7675993- T   spice_donor_variant TP53			splice_donor_variant	TP53	ENSG00000141510	ranscript	ENST00000359597.8	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,
ENST00000445888.6c.375+1G>A 17.7675993- T   splice_donor_variant   TP53   ENSC00000141510   Transcript   ENST00000445888.6 protein_coding   Institute	ENST0000445888 6 c 375+1G-A 177675993- T   spice_donor_variant   TP53   ENSC00000141510 Transcript   ENST0000445888 6 protein_coding   5156755445   536138			splice donor variant	TP53	ENSG00000141510	ranscript	ENST00000413465.6	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,
ENST00000445888.6 c 375+1G>A 17.7675993 T   splice_donor_variant   TP53   ENSG0000141510   Transcript   ENST00000455263.6   protein_coding	ENST0000044588 6.c.375+1G>A 17.7675993- T   splice_donor_variant   TP53   ENSG0000141510   Transcript   ENST0000045583.6   protein_coding   -			splice donor variant	TP53	ENSG00000141510	Franscript	ENST00000420246.6	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,
ENST0000044588.6 c. 375+1G>A 17.7675993 T	ENST00000445888.6c.375+1G>A 17.7675993- T   splice_donor_variant   TP53   ENSG00000141510   Transcript   ENST0000503591.1   protein_coding   -   enstruction   enstruction	ENST00000445888.6:c.375+1G>A 17: 761	:7675993- T 75993	splice_donor_variant	TP53	ENSG00000141510	Franscript	ENST00000445888.6	protein_coding	-	-	CS951538, COSV52663569, COSV52668477
C98/563569	ENST00000445888.6:c.375+1G>A   17.7675993   T			splice donor variant	TP53	ENSG00000141510	Franscript	ENST00000455263.6	protein_coding	-	-	COSV52663569, COSV52668477,
CS951538   CS951538   CS952634568   COSV526634569   COSV526634569   COSV52663469   COSV5266346	ENST00000445888.6:c.375+1G>A   17.7675993-   T			splice donor variant	TP53	ENSG00000141510	Franscript	ENST00000503591.1	protein_coding	-	-	CS951538, COSV52663569, COSV52668477.
Telegraph   Tele	ENST00000445888.6:c.375+1G>A   17:7675993   T	ENST00000445888.6:c.375+1G>A 17: 767		upstream_gene_variant	TP53	ENSG00000141510	Franscript	ENST00000504290.5	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,
Total   Transcript   Transcri	CS951538			upstream_gene_variant	TP53	ENSG00000141510	ranscript	ENST00000504937.5	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,
COSV52663569, COSV52663569, COSV5270628  ENST00000445888.6:c.375+1G>A 17:7675993- T intron_variant TP53 ENSG00000141510 Transcript ENST00000509690.5 protein_coding rs1567555445,	ENST00000445888.6:c.375+1G>A 17:7675993- T Intron_variant TP53 ENSG00000141510 Transcript ENST00000509690.5 protein_coding Is1567555445, C3951538, COSV52663569, COSV52663569, COSV52668477,			splice_donor_variant, non_coding_transcript_variant	TP53	ENSG00000141510	Franscript	ENST00000505014.5	retained_intron	-	-	CS951538, COSV52663569, COSV52668477,
	7675993 CS951538 COSV52663569, COSV52668477,			splice_donor_variant	TP53	ENSG00000141510	Franscript	ENST00000508793.5	protein_coding	-	-	COSV52663569, COSV52668477,
COSV52663569, COSV52668477,				intron_variant	TP53	ENSG00000141510	Franscript	ENST00000509690.5	protein_coding	-	-	CS951538, COSV52663569, COSV52668477,

### **ESEfinder**

Para los resultados dónde se encuentra la posición de interés, observamos que se obtienen dos resultados con puntuación positivapara las matrices 5' (362 y 374) y un único resultado para las matrices 3' con puntuación positiva (379):

362 (-447)	GGGACAGCCAAGTCTGTGACTTGCACGgtc	2.17840	362 (-447)	GGGACAGCCAAGTCTGTGACTTGCACGgtc	-30.94350	362 -447) GGGACAGCCAAGTCTGTGACTTGCACGgtc 2.34700 (-44	362 47) GGGACAGCCAAGTCTGTGACTTGCACGgtc -29.97490
374 (-435)	TCTGTGACTTGCACGgtcagttgccctgag	5.64780	374 (-435)	TCTGTGACTTGCACGgtcagttgccctgag -	-13.25930	374 TCTGTGACTTGCACGgtcagttgccctgag 5.72280 (-43	374 TCTGTGACTTGCACGgtcagttgccctgag -11.59380
379 (-430)	GACTTGCACGgtcagttgccctgaggggct	-12.30430	379 (-430)	GACTTGCACGgtcagttgccctgaggggct	1.28560	379 GACTIGCACGgtcagttgccctgaggggct -12.54740 (-43	379 GACTTGCACGgtcagttgccctgaggggct 0.87710

Si observamos las predicciones equivalentes para la secuencia mutante, se observa que la predicción positiva para las matrices 3' (379) sigue siendo positiva igual que una de las predicciones para la matriz 5' (362). Por el contrario, la predicción 374 tiene puntuaciones negativas donde antes eran positivas, por lo tanto se está perdiendo un sitio *donor* en la secuencia mutante, lo que llevara cambios en el *splicing*:

362 (-447)	GGGACAGCCAAGTCTGTGACTTGCACGatc	1.73910	362 (-447)	GGGACAGCCAAGTCTGTGACTTGCACGate	-31.30600	362 GGGACAGCCAAGTCTGTGACTTGCACGAtc 1.81350 362 GGGACAGCCAAGTCTGTGACTTGCACG&	tc-30.30250
374 (-435)	TCTGTGACTTGCACGatcagttgccctgag	-5.33470	374 (-435)	TCTGTGACTTGCACGatcagttgccctgag	-14.58930	374 TCTGTGACTTGCACGatcagttgccctgag -5.40550 374 TCTGTGACTTGCACGatcagttgccctg	ag -12.84420
379 (-430)	GACTTGCACGatcagttgccctgaggggct	-12.07620	379 (-430)	GACTTGCACGatcagttgccctgaggggct	1.43070	379 GACTIGCACGatcagttgccctgaggggct -12.28100 GACTIGCACGatcagttgccctgagggg	ot 0.96920