

Databases for Gene Expression analysis

Perform a THOROUGH in silico analysis before doing benchwork

What is the difference between databases and papers?

For your pet gene:

determine all names using genecards

how many known exonic SNPs are there

find two commercial antisera for the protein

get the complete sequence for your pet gene using the UCSC database and

compare the exon intron-organisation with the one predicted by exonscan

There are multiple databases for expression analysis

<http://www.science.co.il/biomedical/RNA-Databases.asp>

Biomedical directory
Biomedical Databases
Carbohydrate Databases
Enzyme Databases
Genome Databases
Genome, Plant
Genome, Prokaryotic
Glossaries
Lipid Databases
Metabolic Pathways
Mutation Databases
Nomenclature
Nomenclature - Gene
Organelle Databases
Plasmid Databases
Protein Databases
RNA Databases
Sequence Databases
Species Databases
Structure Databases
Virus Databases
Primer Tools
Protein Tools
Sequence Tools

Abbreviation	Database
	5S ribosomal RNA database
ASTRA	Alternative Splicing and Transcription Archives
AAARS	Aminoacyl-tRNA synthetases database
	Codon Usage Database
	Comparative RNA Web Site
Mamit-tRNA	Compilation of mammalian mitochondrial tRNA genes
	Compilation of tRNA sequences and sequences of tRNA genes
PolyA_DB	Database for mammalian mRNA polyadenylation
snoRNABase	Database of Human H/ACA and C/D Box snoRNAs
BPS	Database of RNA Base pair Structures
Recode	Database of translational recoding events
	DNA and RNA Molecular Weights and Conversions
	European Ribosomal RNA Database
FSDB	Frameshift Database
	Free Energy and Enthalpy Tables for RNA Folding
fRNAdb	Functional RNA Database
sRNAmap	Genomic maps for small non-coding RNAs, their regulators and their targets in microbial genomes
GtRDB	Genomic tRNA Database
	Greengenes: 16S rDNA gene database and tools
H-InvDB	H-Invitational database of human genes and transcripts
RNAdb	Mammalian noncoding RNA database
	Methylation Guide snoRNA Database
miRBase	microRNA Sequence Database
	Modomics: a database of RNA modification pathways
ncRNA	Noncoding RNA database
RNaseP	Ribonuclease P Database
RDP	Ribosomal Database Project of 16S rRNA sequences
	Ribosomal RNA Mutation Database
RAG	RNA As Graphs Web resource
Rfam	RNA families database
	RNA Modification Database
STRAND	RNA secondary STRucture and statistical ANALysis Database
	Silva rRNA database
	Small Subunit rRNA Modification Database
SCOR	Structural Classification of RNA
tmRDB	tmRNA Database
	tmRNA Website
	Viral RNA Structure Database
	Yeast Small Nucleolar RNA (snoRNA) Database

Five useful sites

Genecards, <http://www.genecards.org/>
Fast overview over a gene, link to reagents

UCSC genome browser, <http://genome.ucsc.edu/>
Most comprehensive (and best to use) database

Fast DB, <http://www.fast-db.com/fastdb2/frame.html>
Specialized in splicing, very easy to use,
[Log in: Stefan@stamms-lab.net](mailto:Stefan@stamms-lab.net) , PW: OUSMqllerX

Gil Ast lab page, <http://www.tau.ac.il/~gilast/>
Links to splicing analysis tools

Christopher Burge lab, <http://genes.mit.edu/burgelab/software.html>
Links to splicing analysis tools

Genecards

GeneCards® Version 3
The Human Gene Compendium

WEIZMANN INSTITUTE OF SCIENCE with XENNET

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TRA2B Gene
protein-coding **GFIs: 50**
GC03M185632

transformer 2 beta homolog (Drosophila)
(Previous names: **splicing factor, arginine/serine-rich 10 (transformer 2 homolog, Drosophila)**)
Symbol approved by the [HUGO Gene Nomenclature Committee \(HGNC\) database](#)
(Previous symbol: **SFRS10**)

Services
Jump to Section...

Aliases & Descriptions
transformer 2 beta homolog (Drosophila)^{1,2}
Htra2-beta²
splicing factor, arginine/serine-rich 10 (transformer 2 homolog, Drosophila)^{1,2}
transformer-2 protein homolog B^{2,3}
TRA-2 beta^{2,3}
SFRS10^{2,3}
SFRS10^{2,3}
TRAN2B²
DKFZp686F1820²
TRA2-BETA²
transformer-2 protein homolog beta²
transformer-2-beta²
transformer 2 homolog²
Splicing factor, arginine/serine-rich 10³
TRA2-beta³
hTRA2-beta³

External Ids: HGNC: 10781¹ Entrez Gene: 6434² Ensembl: ENSG00000136527⁷ UniProtKB: P62993³

Search outside databases for aliases for TRA2B gene
Previous GC identifier: GC03M187110

Summaries for TRA2B gene
(According to [Entrez Gene](#), [Toxins Bioscience](#), [Wikipedia's Gene Wiki](#), [UniProtKB/Swiss-Prot](#), and/or [UniProtKB/TrEMBL](#))
[About This Section](#)
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UniProtKB/Swiss-Prot: TRA2B_HUMAN_P62995
Function: Sequence-specific RNA-binding protein which participates in the control of pre-mRNA splicing
Gene Wiki entry for TRA2B (SFRS10)

Regulatory elements:
Search SABiosciences Regulatory transcription factor binding sites for TRA2B [improved](#)

Epigenetics:
SABiosciences Methyl-Profiler DNA Methylation qPCR Primer Assays for TRA2B: [MePH05827-2A](#)

Genomic Location:
Genomic View: [UCSC Golden Path with GeneCards custom track](#)

Genomic Views for TRA2B gene
(According to [GeneLoc](#) and/or [HGNC](#), and/or [Entrez Gene \(NCBI build 37\)](#), and/or [miRBase](#),
Genomic Views according to [UCSC](#) and [Ensembl \(release 58\)](#),
Regulatory elements and Epigenetics data according to

5/11 Alternative Splicing Database (ASD) splice patterns (SP) for TRA2B (see all 11)

ExUns: 1A 1B 1C 2 3 4A 4B 5 6A 6B 7A 7B 7C 7D 8A 8B 9 10A 10B 11 12 13A 13B 13C

SP1:

SP2:

SP3:

SP4:

SP5:

[About this scheme](#)

ECgene alternative splicing isoforms for **TRA2B**

18 Ensembl transcripts including schematic representations:

ENST00000453386 ENST00000456380 ENST00000487615 ENST00000492417 ENST00000463328 ENST00000466832
ENST00000382191 ENST00000480461 ENST00000465245 ENST00000485530 ENST00000471134 ENST00000477939
ENST00000259043 ENST00000493864 ENST00000414862 ENST00000342294 ENST00000392502 ENST00000448735

TRA2B expression in normal and diseased human tissues

[Applied Biosystems TaqMan @ Gene Expression Assays for TRA2B](#)

1 GeneNote / 2 GeneAnnot

11 probe-sets matching TRA2B gene

Affymetrix probe-set	Array	GeneAnnot data		GeneNote data		GeneTide data					
		# genes	Sensitivity	Specificity	Correlation	Length	Gb_Accession	Consensus	Uniqueness	Score	Rank
140_s_at ²	U95-A	1	1.00	1.00	0.70	1.32	--	--	--	--	--
70770_at ²	U95-D	1	0.38	1.00	0.87	0.69	--	--	--	--	--
68803_at ²	U95-E	1	0.31	1.00	-0.36	0.27	--	--	--	--	--
200892_s_at ²	U133-A	1	1.00	1.00	--	--	--	--	--	--	--
210180_s_at ²	U133-A	1	1.00	1.00	--	--	--	--	--	--	--
200893_at ²	U133-A	1	1.00	1.00	--	--	--	--	--	--	--
239447_at ²	U133-B	1	0.27	1.00	--	--	--	--	--	--	--
200892_s_at ²	U133Plus2	1	1.00	1.00	--	--	--	--	--	--	--
210180_s_at ²	U133Plus2	1	1.00	1.00	--	--	--	--	--	--	--
200893_at ²	U133Plus2	1	1.00	1.00	--	--	--	--	--	--	--
239447_at ²	U133Plus2	1	0.27	1.00	--	--	--	--	--	--	--

[About this table](#)

GeneDecks TRA2B for expression [About GeneDecksing](#)

Data from GeneNote (Publications) and GNF BioGPS
[About these images](#)

Experimental tissue vectors: Duplicate measurements were obtained for twelve normal human tissues (out of 28 tissues shown) hybridized against Affymetrix GeneChips HG-U95A-E (GeneNote data) and for 22 normal human tissues hybridized against HG-U133A (GNF data). [more...](#)

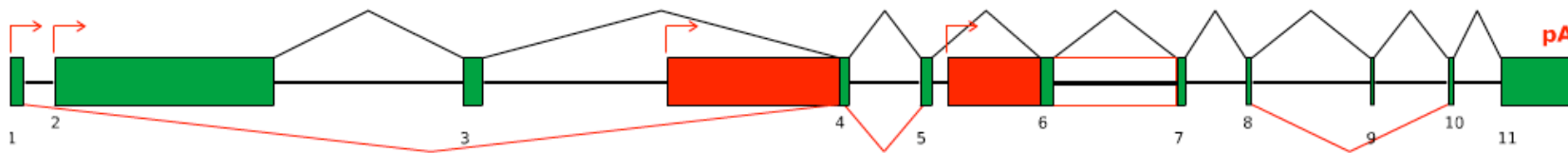
Fast, comprehensive overview for a given gene (expression, names, publications...)

FAST-DB

splicing factor, arginine/serine-rich 10 (transformer 2 homolog, Drosophila)
(ID 15379)

Human mRNAs Human ESTs Mouse mRNAs

Human mRNAs MAIN PAGE NAMES / SYMBOLS PUBMED TRANSCRIPTS VIEW TISSUE-SPECIFICITY IN SILICO PCR PROBE ALIGN. PDF SEQUENCES



[21284 bp]

[chromosome (strand): 3(-):187117238-187138521]

[11 EXON(S)]

[legend]

TRANSCRIPTION INITIATION & FIRST EXON(S)

FIRST EXON(S)	Number of evidence(s)
EXON 1	9
EXON 2	1
EXON 4	1

>exon 5 163 bp (12406-12568)
GTCTAGATCCAGAAGAAGCTCCCGAAGGCATTATACCC
GGTCACGGTCTCGCTCCCGCTCCCATAGACGATCACGT
AGCAGGTCTTACAGTCGAGATTATCGTAGACGGCACAG
CCACAGCCATTCTCCCATGTCTACTCGCAGGCGTCATG
TTGGGAATCGG

<http://www.fast-db.com>

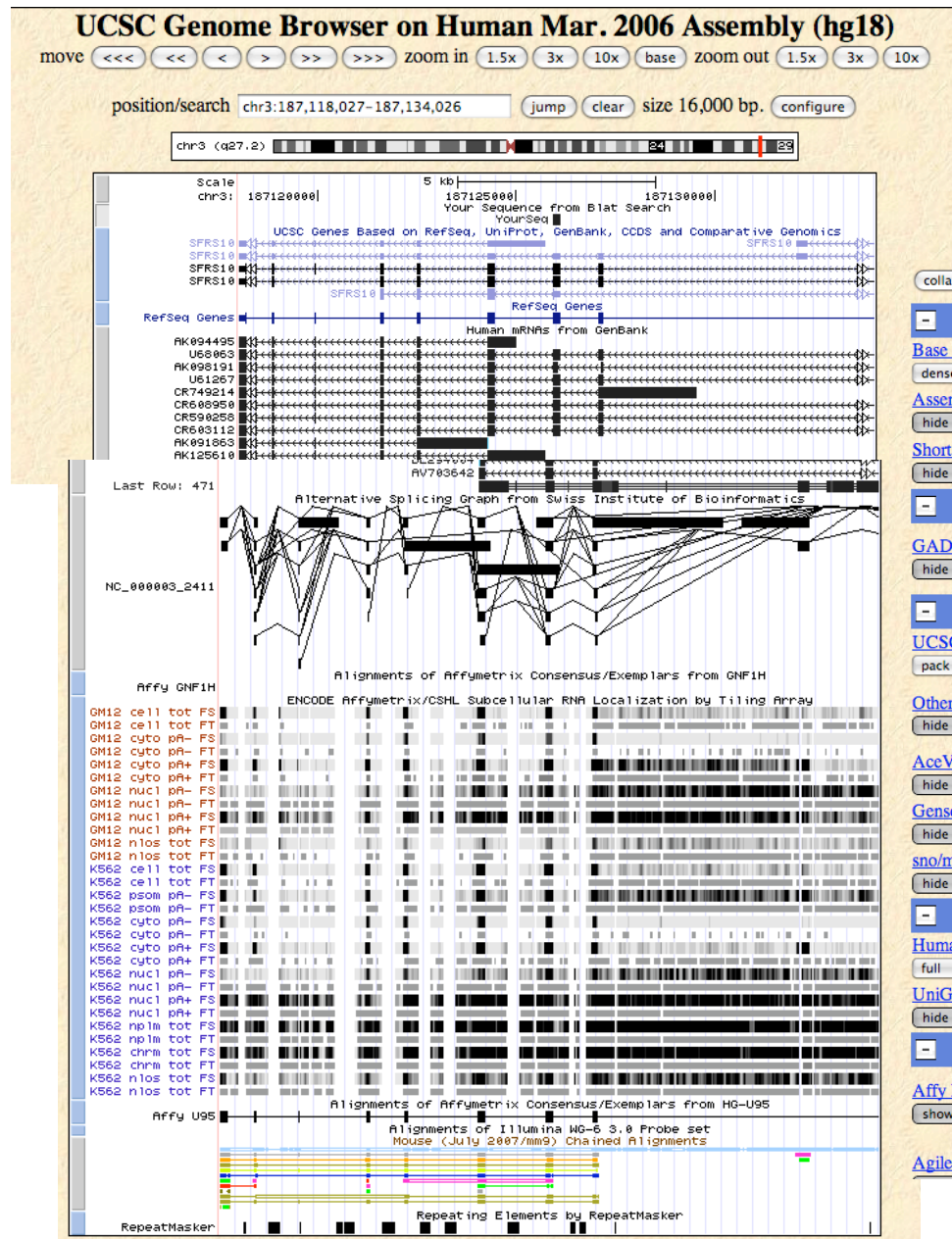
User-friendly

Stefan@stamms-lab.net

PW: OUSMqllerX

UCSC genome browser

UCSC blat



Use drop-down controls below and press refresh to alter tracks displayed. Tracks with lots of items will automatically be displayed in more compact modes.

collapse all expand all

Mapping and Sequencing Tracks

refresh

Base Position	Chromosome Band	STS Markers	FISH Clones	Recomb Rate	Map Contigs
dense	hide	hide	hide	hide	hide
Assembly	Gap	Coverage	BAC End Pairs	Fosmid End Pairs	GC Percent
hide	hide	hide	hide	hide	hide
Short Match	Restr Enzymes	Wiki Track	Blat Sequence	Mapability	
hide	hide	hide	pack	hide	

Phenotype and Disease Associations

refresh

GAD View	OMIM Genes	RGD Human QTL	RGD Rat QTL	MGI Mouse QTL
hide	hide	hide	hide	hide

Genes and Gene Prediction Tracks

refresh

UCSC Genes	Old UCSC Genes	Alt Events	Gencode Genes	CCDS	RefSeq Genes
pack	hide	hide	hide	hide	dense
Other RefSeq	MGC Genes	ORFeome Clones	TransMap...	Vega Genes	Ensembl Genes
hide	hide	hide	hide	hide	hide
AceView Genes	SIB Genes	N-SCAN	CONTRAST	SGP Genes	Geneid Genes
hide	hide	hide	hide	hide	hide
Genscan Genes	Exoniphy	Augustus	RNA Genes	ACEScan	EvoFold
hide	hide	hide	hide	hide	hide
sno/miRNA	Pos Sel Genes				
hide	hide				

mRNA and EST Tracks

refresh

Human mRNAs	Spliced ESTs	Human ESTs	Other mRNAs	Other ESTs	H-Inv
full	full	full	hide	hide	hide
UniGene	Gene Bounds	SIB Alt-Splicing	Poly(A)	CGAP SAGE	
hide	hide	full	hide	hide	

Expression

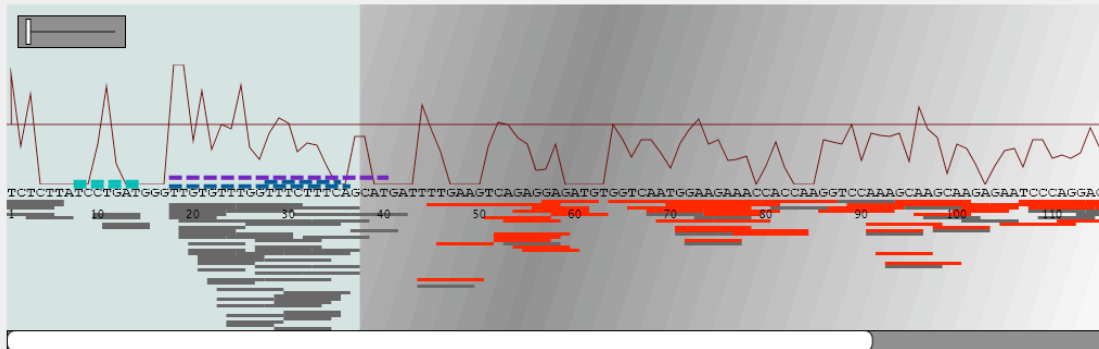
refresh

Affy Exon...	Affy GNF1H	Affy RNA Loc	Affy U133	Affy U133Plus2	Affy U95
show	dense	dense	hide	hide	dense
Agilent Array	Allen Brain	Bertone Yale TAR	CSHL Long RNA-seq	GIS PET RNA	GNF Atlas 2

Gil Ast homepage

Splicing Regulation Online Graphical Engine

[help](#) [download](#)



Regulatory Sets

[SF2](#)
[SRP40](#)
[SRP55](#)
[SC35](#)
[Fairbrother](#)
[Chasin ESE](#)
[Wang](#)
[Chasin ESS](#)
[Goren](#)
[Yeo up](#)
[Yeo down](#)
[Voelker up](#)
[Voelker down](#)

Splice Site Scores

Represented by dashed annotations above the sequence.

Element	Start	End	Score	Score Percentile (Const Exons)	Score Percentile (Alt Exons)
Branch site	8	14	3.45 (Kol et al)	0.8	0.82
	8	14	3.00 (Schwartz et al)	0.05	0.034
Polypyrimidine tract	18	36	0.78 (Kol et al)	0.41	0.43
	28	35	8.00 (Schwartz et al)	0.31	0.33
3' splice site	18	40	9.36 (Max entropy)	0.61	0.66
			87.50 (PSSM)	0.66	0.71
			-5.40 (Delta-G)	0.42	0.46
5' splice site	119	127	9.06 (Max entropy)	0.57	0.65
			82.85 (Senepathy)	0.5	0.57

Display Preferences

☐ Display Matches For Mutations (place cursor over nucleotide to view mutations)

Graphs Display:

☒ Mutability Index (☒ Use only selected sets)

☐ NJ Values

Mutability Index - the ratio between regulatory sequences overlapping each position and the sequences that would be found if the position was mutated.

Splicing regulatory sequences: Annotated beneath the sequence.

Name	Type	Upstream Intron	Exon	Downstream Intron	Density	Density Percentile (Const Exons)	Density Percentile (Alt Exons)
SF2	E	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.17	0.28	0.31
SRP40	E	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.42	0.94	0.92
SRP55	E	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.00	0	0
SC35	E	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.49	0.94	0.91
Fairbrother	E	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.69	0.99	0.99
Chasin ESE	E	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.49	0.91	0.92
Wang	S	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.00	0	0
Chasin ESS	S	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.00	0	0
Goren	R	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	0.71	0.97	0.98
Yeo up	R	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	0.46	0.5	0.5
Yeo down	R	<input type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	0.49	0.62	0.62
Voelker up	R	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	0.31	0.14	0.18
Voelker down	R	<input type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	0.49	0.47	0.51

(Type: E - enhancer S - silencer R - regulator)

Density: (# bases overlapped by regulatory sequence / sequence length)

SCROOGLE

Christopher Burge lab

EXONSCAN

ExonScan results page

locus: HUMATPGG length: 12900 cutoff: 192
Predicted exons:

Begin	-	End	(3'ss	5'ss	ESE	ESS	GGG	total)
35	-	178	(92	37	102	6	39 276)
294	-	353	(50	96	12	0	51 209)
925	-	1128	(97	52	3	13	39 204)
2631	-	2744	(101	85	19	-1	9 213)
2948	-	3029	(78	58	75	10	6 227)
4371	-	4569	(90	63	36	-3	27 213)
5066	-	5200	(79	91	9	-2	21 198)
5715	-	5907	(109	85	25	5	9 233)
7751	-	7887	(117	74	15	-2	15 219)
7973	-	8123	(97	80	34	3	21 235)
8487	-	8641	(89	85	66	0	30 270)
9615	-	9856	(116	43	27	-4	39 221)
10380	-	10525	(132	72	33	-3	6 240)
12450	-	12551	(95	85	25	5	24 234)
12636	-	12727	(104	60	16	14	39 233)

Key:

- locus: gene name (taken from line beginning with > if present)
- length: number of bases in given sequence
- cutoff: minimum total exon score to be considered for prediction
- Begin: index (starting at one) of first base in predicted exon
- End: index (starting at one) of last base in predicted exon
- 3'ss: maximum entropy score of 3' splice site (beginning of exon)
- 5'ss: maximum entropy score of 5' splice site (end of exon)
- ESE: sum of scores for all RESCUE-ESEs in predicted exon
- ESS: sum of scores for all FAS-hex3 ESSs in predicted exon
- GGG: sum of scores for all GGGs in certain intronic regions flanking predicted exon
- total: sum of all other scores

All scores are in tenths of bits.

[Return to ExonScan Web Server](#)

SUMMARY AND OUTLOOK

Lots of information available about gene expression that can be visualized using browsers

Problem with databases is their upkeep, UCSC and ENSEMBLE up to date for nucleic acids

Multiple analysis tools are available on the web

For your pet gene:

determine all names using genecards

how many known exonic SNPs are there

find two commercial antisera for the protein

get the complete sequence for your pet gene using the UCSC database and

compare the exon intron-organisation with the one predicted by exonscan