

A photograph showing several people in a computer lab or training room. They are seated at desks with multiple computer monitors. Some are looking at the screens, while others are in discussion. The room has a casual, educational atmosphere.

# GTPB

The Gulbenkian Training Programme in Bioinformatics  
(Since 1999)

Pedro Fernandes, Organiser



# IB16S

## Introductory Bioinformatics

12-16 December 2016

(Second 2016 run of this Course)

## Basic Bioinformatics Sessions

### Practical 1: Databases and Tools

Tuesday 6 December 2016

## Investigating gene(s) associated with Aniridia

As a starting point for this exercise, imagine you have a vital interest in discovering and investigating the main human gene responsible for the terrible disease of the eye, **Aniridia**. There are many ways (including **google!**) you could discover what this gene might be. I choose to delve into the vast seas of knowledge so generously proffered by the **The National Center for Biotechnology Information (NCBI)**.

So, begin by going to the **Home Page** of the **The National Center for Biotechnology Information (NCBI)** ("<http://www.ncbi.nlm.nih.gov/>").

You will arrive at a page offering access to the many **NCBI** resources available to you. Currently, you only require to search for genes, specifically those that relate to **Aniridia**, so first set the database selection field of the **Search** facility at the top of your page to **Gene**, set the **Search** field to **Aniridia** and click on the **Search** button.

A fine list of genes will emerge, including those sought. However, our interest is specific to Human, so the search should really be organism specific. To do this, one needs to execute an **Advanced** search. So, click on the **Advanced** button of the **Search** tool.

Now you can specify the precise field(s) of the annotation you wish to interrogate. In this case, set the **Disease/Phenotype** field to **Aniridia** and the **Organism** field to **Human**. As the two conditions are linked by **AND**, both must be true for any gene to be listed.

Name/Gene ID	Description	Location	Aliases	MIM
<input type="checkbox"/> <a href="#">PAX6</a> ID: 5080	paired box 6 [ <i>Homo sapiens</i> (human)]	Chromosome 11, NC_000011.10 (31784792..31817961, complement)	AN, AN2, D11S812E, FVH1, MGDA, WAGR	607108
<input type="checkbox"/> <a href="#">WT1</a> ID: 7490	Wilms tumor 1 [ <i>Homo sapiens</i> (human)]	Chromosome 11, NC_000011.10 (32387775..32435535, complement)	AWT1, EWS-WT1, GUD, NPHS4, WAGR, WIT-2, WT33	607102
<input type="checkbox"/> <a href="#">ITPR1</a> ID: 3708	inositol 1,4,5-trisphosphate receptor type 1 [ <i>Homo sapiens</i> (human)]	Chromosome 3, NC_000003.12 (4493348..4847840)	ACV, CLA4, INSP3R1, IP3R, IP3R1, PPP1R94, SCA15, SCA16, SCA29	147265
<input type="checkbox"/> <a href="#">ELP4</a> ID: 26610	elongator acetyltransferase complex subunit 4 [ <i>Homo sapiens</i> (human)]	Chromosome 11, NC_000011.10 (31509729..31784525)	AN, C11orf19, PAX6NEB, PAXNEB, DJ68P15A.1, hELP4	606985
<input type="checkbox"/> <a href="#">DEL11P13</a> ID: 100528024	Wilms tumor, aniridia, genitourinary anomalies and mental retardation syndrome [ <i>Homo sapiens</i> (human)]		C11DELp13, WAGR	194072

Just a few genes survive. OK, this is just an exercise, so trust me ... it is **PAX6** that is the most interesting gene, in this context. This is the one to follow up by clicking on the link to its details.

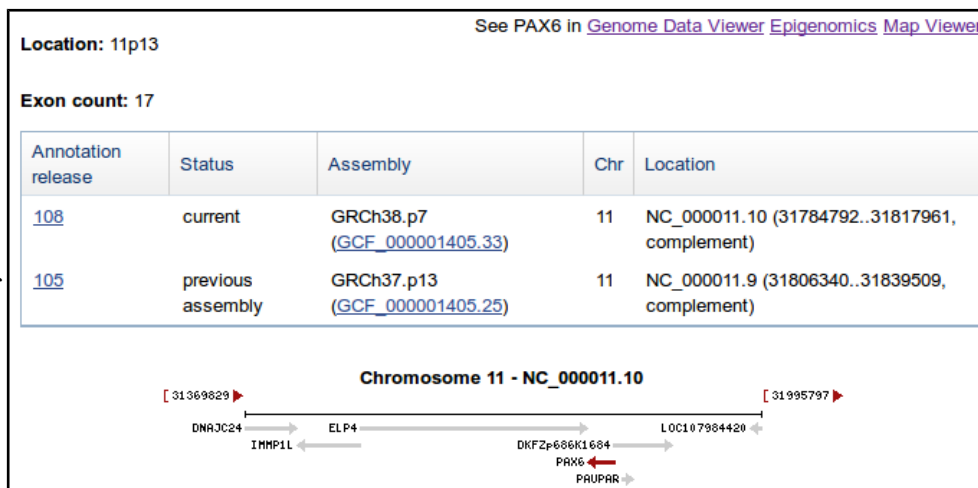
From the Summary section one can conclude (sticking to the features that pertain to this exercise) that:

- there are two major domains, a paired domain and a homeobox, both of which bind DNA
- the gene regulates transcription (is a transcription factor)
- there is more than one protein isoform, and thus more than one transcript variant.

**Summary** This gene encodes a homeobox and paired domain-containing protein that binds DNA and functions as a regulator of transcription. Activity of this protein is key in the development of neural tissues, particularly the eye. This gene is regulated by multiple enhancers located up to hundreds of kilobases distant from this locus. Mutations in this gene or in the enhancer regions can cause ocular disorders such as aniridia and Peter's anomaly. Use of alternate promoters and alternative splicing result in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jul 2015]

From the Genomic context section it can be seen that:

- **PAX6** is situated on **Chromosome 11**, band **p13**
- **PAX6** is on the complementary strand relative to that chosen by **Map Viewer** to represent **Chromosome 11**
- **ELP4** (another gene in the list of human genes associated with **Aniridia**) is exceedingly close, on the opposite strand to **PAX6**. This might be worthy of investigation, at another time?
- There are **17** exons for **PAX6**. Jolly good, but I really wanted to know how many transcripts there were according to the **NCBI**? That is, how many different ways it is thought that nature spliced the **17** exons together. I would also like to discover how many distinct **isoforms** the **NCBI** imagines to result from however many **transcripts**. I proceed with impatience!



Click either the [Genome Data Viewer](#) or the [MapViewer](#) link. Both offer you essentially the same story, the choice really is cosmetic. Do you like your genomes vertical or horizontal. I am a horizontal man myself, so I prefer the [Genome Data Viewer](#). The data is from the [Map Viewer Genome Database](#), whichever choice you make.

I reproduce both views here. The [Genome Data Viewer](#) picture is included in the **PAX6** gene page for free, so maybe the [MapViewer](#) link is the best one for you to choose? Or both, of course! First consider the marginally clearer and simpler [Genome Data Viewer](#) picture.



So, if I tell you the region displayed is the entire **PAX6** region of **Chromosome 11** and the green lines labelled on the right as something beginning with **NM\_** represent the different transcripts, **can you now say how many transcripts there are according to this view?** In passing, the blobs along each line represent the exons. Dark blobs are coding exons. Light blobs represent the exons that form the **3'/5' UTR** regions of each transcript. The Introns are the pale green lines joining the blobs together.

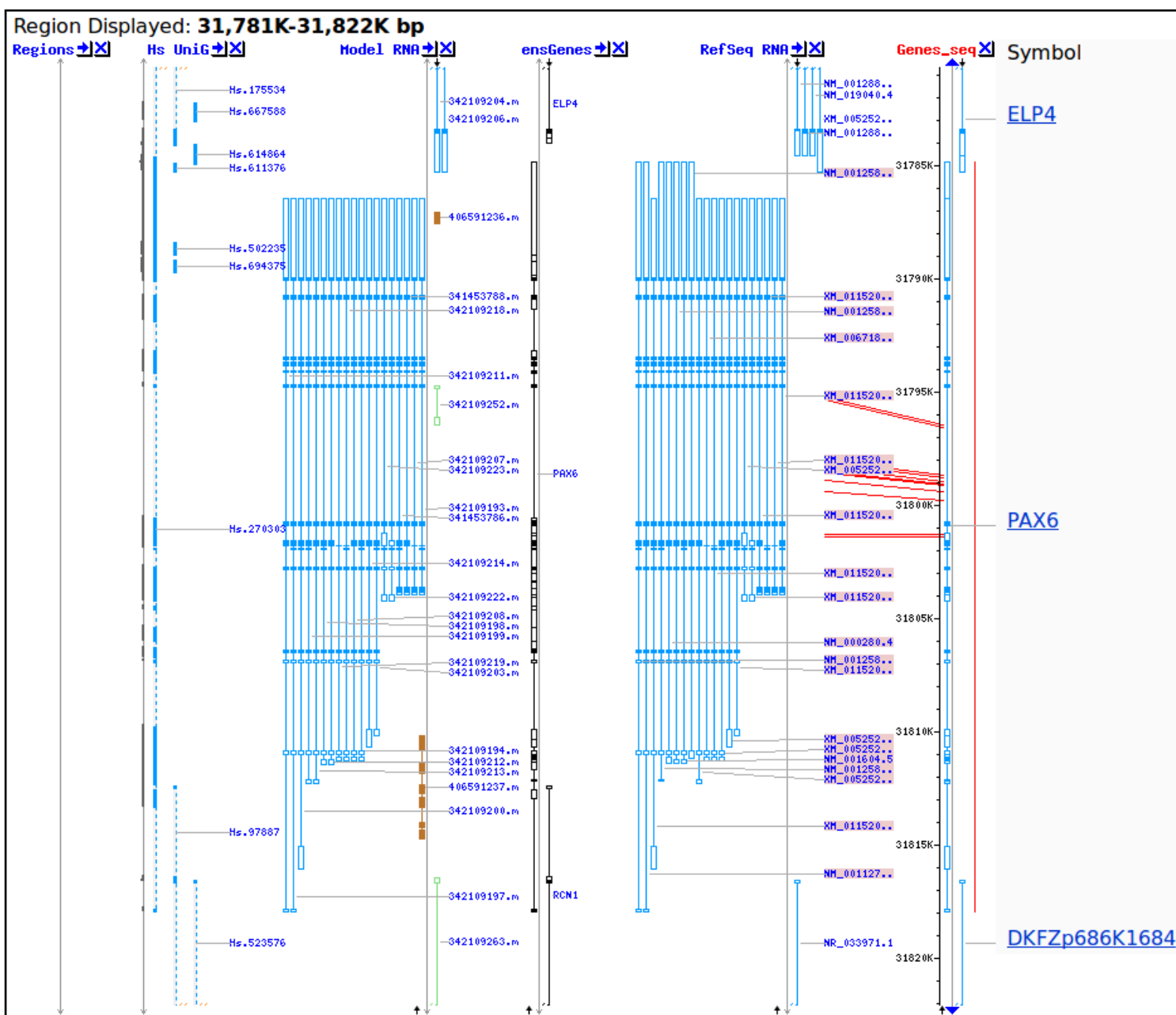
You need also to realise that the prediction of the transcripts shown here are based on database searches of all Human mRNA sequences stored in **RefSeq** against this region of the genome. The theory is that every human mRNA

sequence must match (nearly) perfectly somewhere in the human genome. Where it matches, there must be the genomic DNA from which the mRNA was transcribed. How charmingly simple!

To differentiate between coding and non-coding exons of a transcript, why not compare all human proteins with the genome (after suitable translation to amino acid codes in all six reading frames). They too must match near perfectly somewhere, identifying the **CoDing Sequence (CDS)** of each transcript. Transcript fully located. Job done! Of course, it does not always work so very neatly, but we need not admit that for the moment at least.

Comparing proteins with the genome is clumsy, compute intensive, slow. For major organisms (currently just Human and Mouse), specially comprehensive **databases of extremely reliable DNA Coding Sequences** have been constructed. Searching with these databases is so much more efficiently searching against proteins, serves exactly the same purpose and is thus very much preferred.

And so to the **Map Viewer** version of exactly the same region of **Chromosome 11**.




OK, times up, **how many transcripts are predicted for PAX6 then?** It would appear the answer depends which viewer you chose to view the **PAX6** region? I count **11** in the **Genome Data Viewer** but **20** in the **MapViewer** picture. There is an explanation, but really!! Sometimes I wonder if there is ever any straight answers out there at all.

The explanation? Well, clearly, there should be one transcript predicted for every **PAX6** mRNA sequence in **RefSeq**. At the time the **Map Viewer** illustration was computed, there were **20** mRNA sequences in **RefSeq**, **9** of these were not as well evidenced as the other **11**. You can tell the difference as the “good” ones have names beginning with **NM\_** and the “less good” ones begin **XM\_**. If you have very good eyesight, you can confirm that there are **11** **NM\_s** and **9** **XM\_s** in the **Map Viewer** picture.



The **Genome Data Viewer** declines to show the less worthy matches. The discrepancy is due to **Map Viewer** using a slightly older version of **RefSeq** than the **Genome Data Viewer** to compute its message<sup>1</sup>. There are no **XM\_mRNAs** for **PAX6** in the current **RefSeq**. They appear to all have been recently dismissed as “unworthy”. Unfortunately, this kind of inconsistency due to version differences and similar are not uncommon. One needs to be a little cautious at times.

Even without database version conflicts, seemingly trivial inquiries such as “how many transcripts are there” might be simple. Move back to the page describing the **PAX6** gene. In the familiar graphic at the top of the **Genome regions, transcripts and products** section you will find routes to corresponding information from the **Ensembl Genome Database**. Hover over the **PAX6** (also known as **ESNG00000007372**, by **Ensembl** and other close friends) green line in the bottom half of the picture. You will be rewarded by cheery gray box full of links to **Ensembl** and other exciting places.

 ENSG00000007372  
 Gene: ENSG00000007372  
 Title: PAX6  
 Location: complement(31,784,792..31,817,961)  
 Length: 33,170  
  
 gene\_biotype: protein\_coding  
 gene\_id: ENSG00000007372  
 gene\_name: PAX6  
 gene\_source: ensembl\_havana  
 gene\_version: 20  
 havana\_gene: OTTHUMG00000041447  
 havana\_gene\_version: 12  
 Merged features: 46  
 Links & Tools  
 View ENSEMBL: [ENSG00000007372](#)  
  
 GenBank View: [NC\\_000011.10 \(31,784,792..31,817,961\)](#)  
 FASTA View: [NC\\_000011.10 \(31,784,792..31,817,961\)](#)  
 BLAST Genomic: [NC\\_000011.10 \(31,784,792..31,817,961\)](#)

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	RefSeq
PAX6-201	<a href="#">ENST00000419022</a>	6922	<a href="#">436aa</a>	Protein coding	<a href="#">CCDS31452</a>	<a href="#">F1T0F8</a> <a href="#">P26367</a>	<a href="#">NM_001258462</a> <a href="#">NM_001310158</a> <a href="#">NM_001310161</a> <a href="#">NP_001245391</a> <a href="#">NP_001297087</a> <a href="#">NP_001297090</a>
PAX6-202	<a href="#">ENST00000606377</a>	6860	<a href="#">436aa</a>	Protein coding	<a href="#">CCDS31452</a>	<a href="#">F1T0F8</a> <a href="#">P26367</a>	<a href="#">NM_001258463</a> <a href="#">NM_001310161</a> <a href="#">NP_001245392</a> <a href="#">NP_001297090</a>
PAX6-009	<a href="#">ENST00000379129</a>	2616	<a href="#">436aa</a>	Protein coding	<a href="#">CCDS31452</a>	<a href="#">F1T0F8</a> <a href="#">P26367</a>	-
PAX6-011	<a href="#">ENST00000379107</a>	2591	<a href="#">436aa</a>	Protein coding	<a href="#">CCDS31452</a>	<a href="#">F1T0F8</a> <a href="#">P26367</a>	-
PAX6-008	<a href="#">ENST00000379132</a>	2574	<a href="#">422aa</a>	Protein coding	<a href="#">CCDS31451</a>	<a href="#">P26367</a> <a href="#">Q66SS1</a>	<a href="#">NM_001127612</a> <a href="#">NP_001121084</a>
PAX6-003	<a href="#">ENST00000379123</a>	2160	<a href="#">422aa</a>	Protein coding	<a href="#">CCDS31451</a>	<a href="#">P26367</a> <a href="#">Q66SS1</a>	<a href="#">NM_000280</a> <a href="#">NM_001258464</a> <a href="#">NP_000271</a> <a href="#">NP_001245393</a>
PAX6-001	<a href="#">ENST00000379115</a>	1763	<a href="#">436aa</a>	Protein coding	<a href="#">CCDS31452</a>	<a href="#">F1T0F8</a> <a href="#">P26367</a>	<a href="#">NM_001604</a> <a href="#">NP_001595</a>
PAX6-002	<a href="#">ENST00000241001</a>	1631	<a href="#">422aa</a>	Protein coding	<a href="#">CCDS31451</a>	<a href="#">P26367</a> <a href="#">Q66SS1</a>	-
PAX6-005	<a href="#">ENST00000379111</a>	1627	<a href="#">422aa</a>	Protein coding	<a href="#">CCDS31451</a>	<a href="#">P26367</a> <a href="#">Q66SS1</a>	<a href="#">NM_001258465</a> <a href="#">NP_001245394</a>
PAX6-004	<a href="#">ENST00000379109</a>	2157	<a href="#">422aa</a>	Protein coding	-	<a href="#">P26367</a> <a href="#">Q66SS1</a>	-
PAX6-020	<a href="#">ENST00000525535</a>	677	<a href="#">2aa</a>	Protein coding	-	-	-
PAX6-021	<a href="#">ENST00000524853</a>	574	<a href="#">57aa</a>	Protein coding	-	<a href="#">E9PKM0</a>	-
PAX6-016	<a href="#">ENST00000423822</a>	567	<a href="#">61aa</a>	Protein coding	-	<a href="#">B1B1I9</a>	-
PAX6-012	<a href="#">ENST00000455099</a>	497	<a href="#">124aa</a>	Protein coding	-	<a href="#">B1B1J0</a>	-
PAX6-013	<a href="#">ENST00000438681</a>	455	<a href="#">38aa</a>	Protein coding	-	<a href="#">B1B1I8</a>	-
PAX6-029	<a href="#">ENST00000533156</a>	847	No protein	Processed transcript	-	-	-
PAX6-014	<a href="#">ENST00000471303</a>	782	No protein	Processed transcript	-	-	-
PAX6-027	<a href="#">ENST00000531910</a>	643	No protein	Processed transcript	-	-	-
PAX6-015	<a href="#">ENST00000481563</a>	613	No protein	Processed transcript	-	-	-
PAX6-028	<a href="#">ENST00000530373</a>	572	No protein	Processed transcript	-	-	-
PAX6-025	<a href="#">ENST00000530714</a>	567	No protein	Processed transcript	-	-	-
PAX6-024	<a href="#">ENST00000534353</a>	540	No protein	Processed transcript	-	-	-
PAX6-019	<a href="#">ENST00000533333</a>	6173	No protein	Retained intron	-	-	-
PAX6-006	<a href="#">ENST00000470027</a>	2842	No protein	Retained intron	-	-	-
PAX6-007	<a href="#">ENST00000494377</a>	2460	No protein	Retained intron	-	-	-
PAX6-010	<a href="#">ENST00000464174</a>	979	No protein	Retained intron	-	-	-
PAX6-017	<a href="#">ENST00000474783</a>	702	No protein	Retained intron	-	-	-
PAX6-030	<a href="#">ENST00000532916</a>	627	No protein	Retained intron	-	-	-
PAX6-026	<a href="#">ENST00000534390</a>	578	No protein	Retained intron	-	-	-
PAX6-023	<a href="#">ENST00000532175</a>	524	No protein	Retained intron	-	-	-
PAX6-022	<a href="#">ENST00000527769</a>	487	No protein	Retained intron	-	-	-

Use the link labelled **View ENSEMBL**. A view of the region of **Chromosome 11** similar to those you have already considered will leap forth. As before, the exons for each transcript are represented by blobs (filled for coding, empty for UTR regions). Introns are represented by wiggly lines joining the blobs. The colour scheme we might consider later. For now, notice is that there are considerably more than **20** transcripts represented here! **31** to be precise, as it says near the top of the page in tiny letters. For a clearer view, click on the [Show transcript table](#) link.

Indeed **31**. But see that not all are protein coding. One reason for the difference in transcript count is due to the fact that **Ensembl** includes non-coding transcripts, whereas the **NCBI** list only transcripts that code for proteins.

Also, and more importantly, **Ensembl** and **MapViewer** use different strategies to predict transcripts (and just about everything else!). Both use database searches in roughly the manner described above and (for the human genome at least) the same basic assemblies of the genome and sequence databases, but clearly **Ensembl** has less blind faith in the accuracy of the **RefSeq** mRNAs. As you can see, in the **RefSeq** column of the table, the matches with **RefSeq** mRNAs are taken into consideration. However, they do not appear to be considered individually

sufficient to define a transcript. Some **Ensembl** transcripts reference more than one **RefSeq** mRNA. The lesser quality **XM\_mRNAs** are ignored (possibly because they no longer exist in the version of **RefSeq** used by **Ensembl**?). Not all **11** better quality are used (just **1** ignored). Counting just the protein coding transcripts predicted by **Ensembl**, I see a total of **15**.

We could go on, other sources (not necessarily **Genome Databases**) would count differently again. Perhaps the best answer to the question “How many transcripts are there for the **PAX6** gene” is “**Several**”.

<sup>1</sup> The **MapViewer** information is computed from assembly **GRCh38.p2** and the version of the **RefSeq** database current at the time. The **Genome Data Viewer** uses the newer assembly **GRCh38.p7** with a newer version of **RefSeq**. Things might have changed by now?

Before leaving **Ensembl**, it would be good to save the genomic sequence of this region for analysis later on.

To do this, first click on the  **Export data** link on the left hand side of the page.

Ask for **500** base pairs of extra sequence at either end of the **PAX6** gene. That is, set both

5' Flanking sequence (upstream):

3' Flanking sequence (downstream):

to **500**.

Gene to export:	ENSG00000007372 (PAX6)
Output:	FASTA sequence *
Strand:	Feature strand *
5' Flanking sequence (upstream):	500 * (Maximum of 1000000)
3' Flanking sequence (downstream):	500 * (Maximum of 1000000)
<b>Next &gt;</b>	

Deselect all the extra **PAX6** related sequences on offer. You just want the one genomic sequence for the entire **PAX6** region.

Click on the **Next >** button.

Please choose the output format for your export

- [HTML](#)
- [Text](#)
- [Compressed text \(.gz\)](#)

Choose **Text** as the **output format** for the sequence to be saved.

Options for FASTA sequence	
Genomic:	Unmasked *
cDNA:	<input type="checkbox"/>
Coding sequence:	<input type="checkbox"/>
Peptide sequence:	<input type="checkbox"/>
5' UTR:	<input type="checkbox"/>
3' UTR:	<input type="checkbox"/>
Exons:	<input type="checkbox"/>
Introns:	<input type="checkbox"/>

In your browser you should now have the genomic region of the **PAX6** gene, with **500** base pairs of flanking sequence on either end, in **FASTA** format.

Do whatever it takes to download this to a file called:

**pax6\_genomic.fasta**

on your **Desktop**. If you end up with a big blank bit at the top of your file, as I did, it might be nice (but not essential) to delete it.

```
>11 dna:chromosome chromosome:GRCh38:11:31784292:31818461:-1
GGCCAGGTTGAGGGTACTCATCGAGCCTCGAACTCTCCTAAAAATGATTCTGCCAAA
GCGCCTCTCCATCCCGGCGGGCTTCGGGTCTCTCCGATGAAGGAGTCCCTTGGGGAT
CGGAGGAGGGGACAGGGTGATTACCCAGAGAGGTAGCTGGCCAGCCTAAGGGCAGAGATC
TTGGGGCCCTAGTGCCCGAAGGTGCGGAGGAGCGCACTCGGCAAGACTAGTTTCTGGGG
ATCGACTCTACGCCATACAGGACGGCGGCCAGGCTGGACCGGGCCGGGCTAGAGCAGTC
ACAGGCCGGGCAAGGAAGGCCAAAGCAGGGGTTGGAGCCGGCCGGACCTGGGTGGGA
GAAGCAGGCTCCCGCCGGCCGGAAGTCTGCGGCGAGAGCTGTGCCAACTCTAGCC
GCCATGACGTCACGCGGGCCGGGACGCCAATGAGGACGGCGCTGGCGTGATATTAAGGA
AAGTTAGCGCTGCTGAGCACCTCTTTTCTTATCATTTGACATTTAACTCTGGGGCAG
GTCCTCGCGTAGAACGCGGCTGTGAGATCTGCCACTTCCCTGCGAGCGGCGGTGAGAA
GTGTGGGAACCGGCTGCGAGGCTCACCTGCCCTCCCGCCCTCCGCTCCAGGTAACCG
CCCGGGCTCCGGCCCGGGCCGGCTCGGGGCCCGGGGCTCTCCGCTGCCAGCGACTG
CTGTCCCCAAATCAAAGCCGCCCAAGTGGCCCCGGGCTTGATTTTGTCTTTAAAG
GAGGCATACAAAGATGGAAGCGAGTTACTGAGGGAGGGATAGGAAGGGGGTGGAGGAGG
GACTTGCTTTTCCGAGTGTGCTCTTCTGCAAAAGTAGCAAAATGTTCACTCTAAGAG
TGGACTTCCAGTCCGGCCCTGAGCTGGGAGTAGGGGGCGGAGTCTGCTGCTGTCTG
CTAAAGCCACTCGCGACCGCGAAAAATGACAGGAGTGGGGACGCACTTTGCATCCAGAC
TCCTCTGCATCGAGTTCACGACATCCAGCTTGGGAAAGTCGTACCCGCGCTGGAGC
GCTTAAAGACACCTGCGCGGGTCGGGCGAGGTGCAGCAGAAAGTTCCCGCGGTTGCAA
AGTGACAGATGGCTGACCGCAACAAAGTCTAGAGATGGGGTTCGTTTCTCAGAAAGACGC
GGAGTACGAAAGATGCGGCGCAGAGCTGGGACGCGCTAAAGCTCCAGCGTGTGAT
TTGAGCTTCACTTCGGAAGACCTAATAATTAGCGATTCTCACTGAGCTAGAAACGCGGCT
CCGGTTACTCGGGCGCTGCGCTGCTGCTCGCGGGAAGCGCGGGCGCCATGGGAG
```

The next question might be “How many protein isoforms might there be for **PAX6**?”.

Well, whilst the **Ensembl** transcript list is still in view, glance down the **Protein** column which displays the size of the protein products for each transcript. Clearly insufficient evidence for a serious **isoform** count, but enough to set a lower limit, as the same **isoform** cannot be more than one length! I conclude the **Ensembl** predicts a minimum of **7 isoforms**. Most are either **422** or **436** amino acids long. Some of the others might cause a raised eyebrow or two, especially the one that is **2** amino acids long? But, who are we to question! **At least 7** is the informal **Ensembl** total.

Click your way back to the **NCBI PAX6** gene entry. Next I would like to discover the number of protein products (**isoforms**) that the **NCBI** predicts. This view makes this simple question clumsy to answer as the protein products of each transcript are reported separately, even when they are identical (as does the equivalent **Ensembl** view)???

However, it can be done. Click on the **NCBI Reference Sequences (RefSeq)** link in the **Table of contents** on the right hand side of the page. Focus on the **mRNA and Protein(s)** sub-section. Skim down the entries for every transcript (only the **11** in the current release of **RefSeq**. The “good” **NM\_** ones). Check the different **isoform** names.

01 - NM_000280.4	→ NP_000271.1	paired box protein Pax-6 isoform a
02 - NM_001127612.1	→ NP_001121084.1	paired box protein Pax-6 isoform a
03 - NM_001258462.1	→ NP_001245391.1	paired box protein Pax-6 isoform b
04 - NM_001258463.1	→ NP_001245392.1	paired box protein Pax-6 isoform b
05 - NM_001258464.1	→ NP_001245393.1	paired box protein Pax-6 isoform a
06 - NM_001258465.1	→ NP_001245394.1	paired box protein Pax-6 isoform a
07 - NM_001310158.1	→ NP_001297087.1	paired box protein Pax-6 isoform b
08 - NM_001310159.1	→ NP_001297088.1	paired box protein Pax-6 isoform c
09 - NM_001310160.1	→ NP_001297089.1	paired box protein Pax-6 isoform d
10 - NM_001310161.1	→ NP_001297090.1	paired box protein Pax-6 isoform d
11 - NM_001604.5	→ NP_001595.2	paired box protein Pax-6 isoform b

I count **4**, imaginatively named **Isoform a**, **Isoform b**, **Isoform c** and **Isoform d**. One associated with each transcript description. Look carefully at the annotations and there is more information. In particular:

**Description** field: **Isoform b** is also known as **Isoform 5a**. Why this is interesting will become apparent in a page or so.

**Isoform b** is also reported to be longer than **Isoform a**.

### Conserved Domains.

Conserved Domains (2) summary		
	<a href="#">smart00351</a> Location:4 → 128	PAX; Paired Box domain
	<a href="#">pfam00046</a> Location:214 → 266	Homeobox; Homeobox domain

Both **Isoform a** and **Isoform b** are recorded as having two domains. A **Paired Box Domain** at the beginning, and a **Homeobox Domain** further along.

Conserved Domains (2) summary		
	<a href="#">smart00351</a> Location:4 → 142	PAX; Paired Box domain
	<a href="#">pfam00046</a> Location:228 → 280	Homeobox; Homeobox domain

Both **Paired Box Domains** are primarily indicated by a hit with the relevant entry in the **SMART** database. Both **Homeobox Domains** are supported by matches with **Pfam** database entries. Other domain

databases will almost certainly provide supporting evidence, but reference to just one match is sufficient here.

From the location information, the **Paired Box** of **Isoform a** appears to include an extra **14** amino acids.

**UniprotKB** offers yet another version of this story. Just for a for a few clicks, let us intrude into the **UniProt** session of your course.

Protein Accession	Links	
	GenPept Link	UniProtKB Link
P26367.2	<a href="#">GenPept</a>	<a href="#">UniProtKB/Swiss-Prot:P26367.2</a>

At the very bottom of the current page, you will find a link to **UniprotKB**. Use it.

Lo! the **PAX6** human protein as seen and understood by **UniProtKB**. Click on the [Sequences \(3\)](#) button on the left hand side of the page. **UniProtKB** declares **3** isoforms! At least, **3** that it is willing to admit to publicly.

Sequences (3)	
Sequence status <sup>i</sup> :	Complete.
This entry describes <b>3</b> isoforms <sup>i</sup> produced by <b>alternative splicing</b> .	



There is **isoform 1**, also known as **isoform a** in America. Note that this is the “*canonical sequence*” for this protein. That is, this is the isoform used to represent this protein. The sequence(s) of all other isoform(s) are recorded as elements of the annotation.

**Isoform 1** (identifier: **P26367-1**) [UniParc] [FASTA](#) [Add to basket](#)  
*This isoform has been chosen as the 'canonical' sequence. All positional information in this entry refers to it. This is also the sequence that appears in the downloadable versions of the entry.*

Also we have **Isoform 5a** (or **PAX6-5a**), also known as **isoform b** in America (where it also answers to **Isoform 5a** when pressed). Note that the entry declares the sequence difference to be:

**47-47: Q → QTHADAKVQVLDNQ**

Literally:

“The amino acid at **position 47** is a **Q** in the canonical sequence. In **isoform 5a** this is replaced by the **15** amino acids **QTHADAKVQVLDNQ**”.

More coherently this amounts to:

“**isoform 5a** differs from the canonical **isoform 1** in that it has an insertion of **14** amino acids after the **47<sup>th</sup>** amino acid of the canonical protein”.

It is significant to note that position **47** is right in the middle of the **Paired Box Domain** that occurs in both isoforms. This confirms that which was discovered at the **NCBI**.

Finally **UniProtKB** proudly presents the somewhat ephemeral **isoform 3** (or **PAX6-5A,6\*** for those who enjoy formality). But, this one has no known sequence? Not much that Bioinformatics can offer here methinks.

**Isoform 3** (identifier: **P26367-3**)  
 Also known as: Pax6-5A,6\*  
 Sequence is not available

So I hope you will agree that the **UniProtKB** count stands at a very modest **2**, plus a ghost.

To visualise the differences between the **2** isoforms with sequence, click on the [Align](#) button at the top of the **Sequences** section. After deep thought and much fumbling, **UniProtKB** will multiply align all the isoform sequences for you. As there are only **2** in this case, this will appear very similar to a **Pairwise** alignment. Highlight the **DNA binding regions** and the **Domains**.

I leave the interpretation of this splendid display to you.

**Highlight**

**Annotation**

- ☐ Sequence conflict
- ☐ Helix
- ☐ Beta strand
- ☐ Turn
- ☐ Chain
- ☐ Compositional bias
- ☒ DNA binding
- ☒ Domain
- ☐ Alternative sequence
- ☐ Natural variant

The extra **14** amino acids of **isoform 5a** are due to the inclusion of a tiny extra (**42** base pair) exon in some transcripts.

Can you see the evidence for this assertion in the two regional genomic maps of a few pages back?

Alignment				
<a href="#">How to print an alignment in color</a>				
P26367	PAX6_HUMAN	1	MONSHSGVNQLGGVFVNGRPLPDSTROKIVELAHSGARPCDISRILO-----	47
P26367-2	PAX6_HUMAN	1	MONSHSGVNQLGGVFVNGRPLPDSTROKIVELAHSGARPCDISRILOTHADAKVQVLDNQ	60
P26367	PAX6_HUMAN	48	-VSNGCVSKILGRYYETGSIRPRAIGGSKPRVATPEVVS KIAQYKRECPSIFAW EIRDL	106
P26367-2	PAX6_HUMAN	61	NVSNGCVSKILGRYYETGSIRPRAIGGSKPRVATPEVVS KIAQYKRECPSIFAW EIRDL	120
P26367	PAX6_HUMAN	107	LSEGVCTNDNIPSVSSINRVLRLNASEKQMGADGMYDKLRMLNGQTGSWGRPGWYPGT	166
P26367-2	PAX6_HUMAN	121	LSEGVCTNDNIPSVSSINRVLRLNASEKQMGADGMYDKLRMLNGQTGSWGRPGWYPGT	180
P26367	PAX6_HUMAN	167	SVPGQPTQDGCQQQEGGENTNSISSNGEDSDEAQMRLQLKRKLORNRSTFTEQIEALE	226
P26367-2	PAX6_HUMAN	181	SVPGQPTQDGCQQQEGGENTNSISSNGEDSDEAQMRLQLKRKLORNRSTFTEQIEALE	240
P26367	PAX6_HUMAN	227	KEFERTHYPDVFAERLAAKIDLPEARIQVWFNRRAKWRREEKLNRORQASNTPSHIP	286
P26367-2	PAX6_HUMAN	241	KEFERTHYPDVFAERLAAKIDLPEARIQVWFNRRAKWRREEKLNRORQASNTPSHIP	300
P26367	PAX6_HUMAN	287	ISSSFSTSVYQIPQPTTPVSSFTSGSMLGRD TALNTYSALPPMPSFTMANNLPQPP	346
P26367-2	PAX6_HUMAN	301	ISSSFSTSVYQIPQPTTPVSSFTSGSMLGRD TALNTYSALPPMPSFTMANNLPQPP	360
P26367	PAX6_HUMAN	347	VPSQTSSYSCLPTSPSVNGRSYDITYTPPHMQTHMNSQPMGTSGTTSTGLISPGVSPVQ	406
P26367-2	PAX6_HUMAN	361	VPSQTSSYSCLPTSPSVNGRSYDITYTPPHMQTHMNSQPMGTSGTTSTGLISPGVSPVQ	420
P26367	PAX6_HUMAN	407	VPGSEPDMSQYWPRLQ	422
P26367-2	PAX6_HUMAN	421	VPGSEPDMSQYWPRLQ	436

We need to save a some protein sequences for future analysis. This is easiest from **UniProtKB** so now is good. To declare your intention to save the entire canonical version of the **PAX6** protein to a file, move back from your alignment. Move to the top of the page where you will find the bizarre invitation to [Add to basket](#) ? Just do it.

You also need to download the sequences of both domains in separate files, via your basket. First the **Paired Box**.

Click the [Family & Domains](#) button on the left of the page. Then use the [Add](#) button adjacent to the **Paired** entry. Its now in your basket you will be ecstatic to know.

Feature key	Position(s)	Length	Description	Graphical view	Feature identifier	Actions
Domain <sup>i</sup>	4 – 130	127	Paired PROSITE-ProRule annotation			<a href="#">Add</a> <a href="#">BLAST</a>



As they are so conveniently in view, take note of the **Compositional bias** features. They will be of interest when we look at database searching.

Feature key	Position(s)	Length	Description	Graphical view	Feature Identifier	Actions
Compositional bias <sup>i</sup>	131 – 209	79	Gln/Gly-rich			Add BLAST
Compositional bias <sup>i</sup>	279 – 422	144	Pro/Ser/Thr-rich			Add BLAST

Natural variant	(VAR_008694)	29	I → S in AN.	1 Publication
Natural variant	(VAR_003811)	29	I → V in AN.	1 Publication
Natural variant	(VAR_008695)	33	A → P in AN.	1 Publication
Natural variant	(VAR_008696)	37 – 39	Missing in AN.	1 Publication
Natural variant	(VAR_008697)	42	I → S in AN; mild.	1 Publication
Natural variant	(VAR_008698)	43	S → P in AN.	1 Publication
Natural variant	(VAR_003812)	44	R → Q in AN.	1 Publication

Then take an excursion to glance at the **Pathol./Biotech** section. Note the many **Natural variants** recorded as responsible for AN (ANiridia, that is). Particularly those around amino acid positions 29 to 44 and specifically that at position 33.

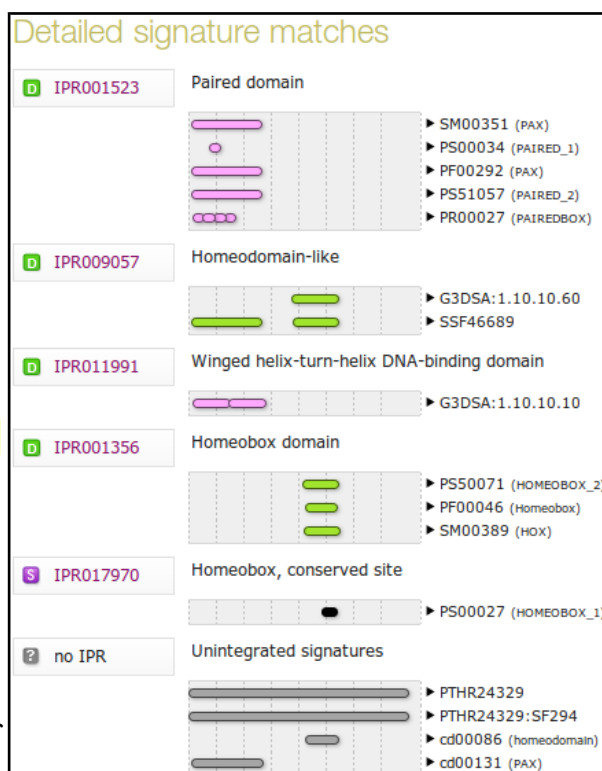
Looking at **PCR Primer Design** later, you will be attempting to create a **PCR** products from patients that, when sequenced, will determine the presence or absence of this variant.

Finally, skip nimbly to the **Family & Domains** section. Concentrate on the Family and domain database sub-section. Here are stored the results of comparing the **PAX6** protein with many of available **Domain/Motif Databases**, including those of the **Interpro Consortium** collectively.


Are the results broadly as you might expect?




For the best summary, click on the **Graphical view** for the **Interpro** results. If the detail is not entirely transparent, hopefully there will be time to discuss this graphic at some point.

The results you are looking at are computed, largely automatically, by the **UniProtKB** annotation system. However, running exactly the same analysis manually is trivial. Maybe you will do this as part of these exercises?




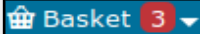
Back to saving sequences for later! To get to the **Homeobox** domain, you need to move back from the **Interpro**

graphic and click on the  **Function** button on the left hand side of the page.


Feature key	Position(s)	Length	Description	Graphical view	Feature Identifier	Actions
DNA binding <sup>i</sup>	210 – 269	60	Homeobox PROSITE-ProRule annotation			 Add  BLAST










A valid question at this point might be “Why is the **Homeobox** domain a **Function** (specifically a **DNA binding** feature), but the **Paired** domain is a **Domain** feature?” To which the answer is “*History, dear boy, history*” to paraphrase a disputed quote of dear Harold (Macmillan that is).

The fact both are **Domains**, and both are **DNA binding**. The illogicality of them being recorded in different places is accepted, however, to fix this early mistake now would not be trivial. So, we live with it. So doing, click on the appropriate  **Add** button and head for the checkout desk (Good Grief! I am beginning to get used to this!).

Shimmy back to the top of the page. You should have  things in your basket.

Click on the basket to view your booty.

For each of the 3 items in turn (not all at once or you get all sequences in one file), select and .

UniProtKB (3)		UniRef (0)	UniParc (0)	(max 400 entries) ✕
<input type="checkbox"/>	Entry	Entry name	Organism	Remove
<input type="checkbox"/>	P26367	PAX6_HUMAN	Homo sapiens (Human)	
<input type="checkbox"/>	P26367[4-130]	PAX6_HUMAN	Homo sapiens (Human)	
<input type="checkbox"/>	P26367[210-269]	PAX6_HUMAN	Homo sapiens (Human)	
     				


Download selected (1) ✕


☐ Download all (3)

**Format:**

FASTA (canonical) ▾

☐ Compressed ☒ Uncompressed



Each time ensure the download parameters are set to **Uncompressed** and **FASTA (canonical)**. Then click the  button.

The next few steps, as before, are very browser/OS dependant. Just do whatever it takes to save the three sequences in files called, as appropriate:

**pax6\_human.fasta**

**pax\_domain.fasta**

**homeobox\_domain.fasta**

Now move back to America to the NCBI view of the **PAX6** gene. If you have problems getting there ... [click here](#).

At the bottom of the page, there is a section called **Related sequences**. Click on the last entry, the mRNA called **AB209177.1**. You will be rewarded by a **GenBank** entry in **GenBank format**. Formats are tedious, but we will discuss them briefly at some point. You have already witnessed **FASTA format**. I expect we will bump into **EMBL format** at some point. The other **137** or so formats I suggest be ignored!

Can you see the official gene name **PAX6**, mentioned in this entry? The **Gene Name** field (where **PAX6** should most certainly be mentioned) is entirely missing! If you searched **GenBank** (or **EMBL** come to that) for this sequence using the most obvious search **Keyword**, that is **PAX6**, do you think you would find this **PAX6 mRNA**? You clearly should! A case for more consistent annotation? Perhaps something to consider further when we superficially mention the **Gene Ontology** project later.

Next, search the nucleotide databases, by textual **Keyword**, for **PAX6** related sequences and download one or two for investigation. To achieve this worthy goal, move to the top of the current page and note that the database selection has changed from **Gene** to **Nucleotide**. Click on the **Advanced** search option button.

Then in the **Nucleotide Advanced Search Builder**, change **All Fields** to **Title** in the pull down menu associated with the first search field and type in the keywords:

chromosome 11

In the second search field, again change **All Fields** to **Title** and type in the keyword:

pax6

You are asking **Entrez** to search for all **Nucleotide** database entries that contain the terms **chromosome 11** and **pax6** in the section of their annotation intended to be a succinct brief description (I.e. **Title**) of the entry. Click on the **Search** button to start the search going.

There is just one matching entry which is arrayed before you in **Genbank** format, very neat!! It was the **DEFINITION** line that you searched by selecting the **Field** value **Title**. I needed a few tries to get the right search to find just what was needed, and was a bit surprised at the simplicity and accuracy of the final search. You are looking at a **RefSeqGene** (a subset of the **RefSeq** database) entry. As such, it represents a genomic sequence for a “well-characterised gene”, in this case **PAX6**.

Take a look at the **FEATURES** for this entry. You will see that there are **two** genes mentioned. **PAX6**, of course, and **ELP4** on the strand that is the complement of that represented here.

```

join(16551..16560,20128..20258,21186..21401,22106..22271,
28174..28332,28848..28930,29160..29310,29409..29524,
32102..32252,32943..33028)
/gene="PAX6"
/gene_synonym="AN; AN2; D11S812E; FVH1; MGDA; WAGR"
/note="isoform a is encoded by transcript variant 1;
paired box protein Pax-6; paired box homeotic gene-6;
oculorhombin; aniridia type II protein"
/codon_start=1
/product="paired box protein Pax-6 isoform a"
/protein_id="NP_000271.1"
/db_xref="CCDS:CCDS31451.1"
/db_xref="LRG:p1"
/db_xref="GeneID:5080"
/db_xref="HGNC:HGNC:8620"
/db_xref="MIM:607108"
/translation="MQNSHSGVNLGGVFNVRPLPDSTROKIVELAHSGARPCDISR
ILQVSNCGVSKILGRYYETGSIIRPRAIGGSKPRVATPEVVSQIAQYKRECPISFAWEI
RDRLLEGVCTNDINIPSVSSINRVLRLNLASEKQMGADGMYDKLRMLNGQTGSWGTTP
GWYPGTSVPGOPTQDGCQQQEGGGENTNSISSNGEDSDEAQMRLQLKRKLQRNRTSFT
QEIQEIALEKEFERTHYPDVFAERARLAAKIDLPEARIQVWFSNRRAKWRREKLRNQR
QASNTPSHIPISSSFSTSVYQIPQPTTPVSSFTSGSMLGRDALTNTYSALPPMPS
FTMANNLPMPQPPVPSQTSYSYCMPLTSPSVNGRSYDTYTPPHMQTHMNSQPMGTSGTT
STGLISPGVSVPVQPGSEPMDSQYWPRLQ"

```

```

gene
5001..38170
/gene="PAX6"
/gene_synonym="AN; AN2; D11S812E; FVH1; MGDA; WAGR"
/note="paired box 6"
/db_xref="GeneID:5080"
/db_xref="HGNC:HGNC:8620"
/db_xref="MIM:607108"

gene
complement(38437..>40170)
/gene="ELP4"
/gene_synonym="AN; AN2; C11orf19; dJ68P15A.1; hELP4;
PAX6NEB; PAXNEB"
/note="elongator acetyltransferase complex subunit 4"
/db_xref="GeneID:26610"
/db_xref="HGNC:HGNC:1171"
/db_xref="MIM:606985"

```

At the top of your page, **Analyze this sequence** by clicking on the **Highlight Sequence Features** option. The **CoDing Sequence (CDS)** feature for **PAX6** is displayed for you by highlighting the relevant parts (the coding **exons**) of the sequence and displaying the **CDS** details including the DNA regions that need to be **joined** to form the **CDS** and the **translation** of the **CDS**.

Use the controls at the bottom of your page to look at the other features of this entry (select feature **number** and then click on the **Feature** button).

What were the features that you found? \_\_\_\_\_

Why might you have expected more features than there were? \_\_\_\_\_

COMMENT REVIEWED [REFSEQ](#): This record has been curated by NCBI staff in collaboration with Isabel Hanson, David FitzPatrick. The reference sequence was derived from [Z95332.1](#) and [Z83307.1](#). This sequence is a reference standard in the [RefSeqGene](#) project.

PRIMARY	REFSEQ_SPAN	PRIMARY_IDENTIFIER	PRIMARY_SPAN	COMP
	1-18852	Z95332.1	2023-20874	
	18853-40170	Z83307.1	105-21422	

Take a look at the **COMMENT** and **PRIMARY** sections just above the **FEATURES**. This entry is suggested to be constructed from two sequences from **GenBank**. That is, the products of two sequencing projects.

Take a quick look at the **GenBank** entries by entering their **ACCESSION** numbers into the **Search** box at the top of your page. Click on the **Search** button.

Nucleotide  [Limits](#) [Advanced](#)

Summary ▾ 200 per page ▾ Sort by Accession ▾

Items: 50

☐ [Homo sapiens F](#) [e, exon 7](#)

1. 166 bp linear DNA  
Accession: AY34235  
[GenBank](#) [FASTA](#)

☐ [Homo sapiens n](#) [6 and partial cds](#)

Sort by

- ☐ Default order
- ☒ Accession
- ☐ Date Modified
- ☐ Date Released
- ☐ Organism Name
- ☐ Taxonomy ID

- ☐ [Human DNA sequence from clone CFAT5 on chromosome 11, complete sequence](#)
1. 20,874 bp linear DNA  
Accession: Z95332.1 GI: 2190397  
[GenBank](#) [FASTA](#) [Graphics](#)
- ☐ [Human DNA sequence from clone A1280 on chromosome 11, complete sequence](#)
2. 22,253 bp linear DNA  
Accession: Z83307.1 GI: 1730464  
[GenBank](#) [FASTA](#) [Graphics](#)

Lo and behold, the two **GenBank** entries are summoned forth. Take a look at one or both. Not particularly illuminating I think<sup>2</sup>. These are clones sequenced as part of the **Human Genome Project (HGP)**. They served to cover regions of **Chromosome 11** and have little biological significance in themselves.

Move back to the list, as illustrated. Elect to **Analyze these sequences**, selecting from the extensive range of possibilities **Run BLAST**. We will look at **blast** properly later, the idea here is to simply prove that these two sequencing clones really do overlap in the fashion suggested by the evidence so far. So, elect to **Align two or more sequences**<sup>3</sup>. Cut and paste one of the sequencing clone **accession numbers** from the **Enter Query Sequence** box to the **Enter Subject Sequence** section of the form. Elect to **Show results in a new window**<sup>4</sup>. Firmly address the **BLAST** button.

Enter Query Sequence BLASTN programs search nucleotide subjects using a nucleotide query. [more...](#)

Enter accession number(s), gi(s), or FASTA sequence(s) [Clear](#) [Query subrange](#)

Z95332.1 From  To

Or, upload file  [Browse...](#)

Job Title

Enter a descriptive title for your BLAST search

☒ Align two or more sequences

Enter Subject Sequence

Enter accession number, gi, or FASTA sequence [Clear](#) [Subject subrange](#)

Z83307.1 From  To

Or, upload file  [Browse...](#)

Program Selection

Optimize for

- ☒ Highly similar sequences (megablast)
- ☐ More dissimilar sequences (discontiguous megablast)
- ☐ Somewhat similar sequences (blastn)

[Choose a BLAST algorithm](#)

**BLAST** Search nucleotide sequence using Megablast (Optimize for highly similar sequences)

☒ Show results in a new window

Just one region of overlap should be identified.

```
Query 20771 GATCCGGAGCGACTTCCGCCTATTTCCAGAAATTAGCTCAAACCTTGACGTGCAGCTAGT 20830
Sbjct 1 GATCCGGAGCGACTTCCGCCTATTTCCAGAAATTAGCTCAAACCTTGACGTGCAGCTAGT 60

Query 20831 TTTATTTTAAAGACAAATGTCAGAGAGGCTCATCATATTTTCCC 20874
Sbjct 61 TTTATTTTAAAGACAAATGTCAGAGAGGCTCATCATATTTTCCC 104
```

How does the alignment you generated match up with the annotation of the original **RefSeq** entry you discovered? \_\_

- The annotation is very sparse which makes these entries very hard to find directly. The **EML-Bank** versions include some links to **Ensembl** codes. These would have been helpful but are not part of the official **International Nucleotide Sequence Database Collaboration (INSDC)** annotation that should be consistent between **GenBank**, **European Nucleotide Archive (ENA)**, which includes **EML-Bank**, and **DNA Data Bank of Japan (DDBJ)**.
- As opposed to comparing each of the two clones against an entire sequence database.
- Just because its neater. In my, significantly less then humble, opinion anyway.



Now for an entirely new search. The easiest way to get a fresh start is to move back to your browser tab displaying the **GenBank Search results**, and then click on the **Advanced** option of the **Search** facility at the top of the page. You should arrive back at the **Nucleotide Advanced Search Builder** offering a fresh start.

Set up a new search as illustrated and set it going. Ultimately simple this time. You have requested all **Human** sequences that are centrally associated with the gene **PAX6**.

A list of **50** or so sequences, all clearly claiming **PAX6** association and announcing their humanity loudly in Latin, will tumble forth.

You will have more hits than are displayed in one go, by default. Also, the hits are arranged in a “**Default**” order which has thus far defied all my attempts to associate with any definition of logic!

To deal with both of these issues, use the display control pull down menus at the top of your page to set the items **per page** to something big and the **Sort by** option to something sane.

The list shows matches between the terms entered and the **annotation** of DNA sequences. Not all relevant sequences will be present. For example, the **mRNA** with accession number **AB209177** was justifiably referenced in the **PAX6 Gene** entry but will not be in this list. **PAX6** appears nowhere in the annotation of **AB209177** including its **DESCRIPTION** (or **Title**) field.

Move far down the list, you will come to the **RefSeq PAX6** mRNAs of a few pages back. Just before these entries is **M77844.1**. Save this one for later analysis. I choose **M77844.1** as it includes a few variations that will add interest. Select the target sequence.

You could now use the diminutive **Send to:** button which is near the bottom of your page to download all the selected sequences into a single file.

However, as there is only one sequence, and it would be so nice to be introduced properly before such intimacies as “downloading”. Click on the link to the database entry to see it in all its **GenBank Format** glory.

The sequence is for analysis rather than decoration, so use the format menu at the top of the page (currently set **GenBank**), and ask for **FASTA** format.

Now click the tiny **Send:** button and **Choose Destination** to be **File**.

Strike the **Create File** button with a firm resolve. With irritating presumption, the choice of file name is made for you. Your sequence will be stored in a file named:

**sequence.fasta**

The **NCBI** is justifiably not famed for its understanding of poetry! Do whatever it takes to rename this file to be called:

**pax6\_mrna.fasta**

Back to **Ensembl**. More with the objective of looking at more sources of information via **Ensembl** than becoming expert **Ensembl** users.

Go to the **Ensembl** home page ([www.ensembl.org](http://www.ensembl.org)). Choose to **View full list of all Ensembl species** using the link just under the **Select a species** menu.

Note that **Ensembl** (and **MapView**, of course) offers far more than just the Human Genome.

In particular, note the links to **EnsemblPlants**, **EnsemblFungi**, **EnsemblBacteria** etc. **Ensembl** databases at the bottom of the list.

During this exercise, you will only look at the Human genome, by far the most completely recorded. However, all the other **Ensembl** genomes are behind the same interface. The techniques required to examine the Human genome are broadly those required to examine any **Ensembl** genome.

Move back to the home page and go straight to the Human **PAX6** gene information by setting up the **Search** fields as shown and clicking the **Go** button boldly.

The target gene is at the top of the hit list.

Click on the link to the **PAX6 (Human Gene)**.

You should recognise the view you now see. The list of transcripts and a view of the genomic region roughly similar to those offered by the **NCBI**.

There is much to investigate here, but maybe that should wait for a specialised **Ensembl** course. They are run regularly in **Cambridge** and elsewhere.








To make a bit more space, elect to [Hide transcript table](#).

First a look at how **Ensembl** sees the **Homologues** of **PAX6**. First the **Orthologues** and then the **Paralogues**. Click on the **Othologues** link in the left hand side of your browser page.

Take a look at some of the alignments providing support for the homologous relations. The protein alignments are the more informative (from the **View Sequence Alignments** menu, select **View Protein Alignment**).

Ensembl Species

Note: to find out which species were in previous releases, please see the [table of assemblies](#)

Show <span>All</span> entries <span>Show/hide columns</span> <span>Filter</span>								
Common name	Scientific name	Taxon ID	Ensembl Assembly	Accession	Variation database	Regulation database	Pre assembly	
 <b>Aardvark</b> <a href="#">Ⓔ</a> (Pre)	<i>Orycteropus afer</i>	1230840	-	-	-	-	-	<a href="#">OryAfe1</a> <a href="#">Ⓔ</a>
 <b>Alpaca</b>	<i>Vicugna pacos</i>	30538	vicPac1	-	-	-	-	
 <b>Amazon molly</b>	<i>Poecilia formosa</i>	48698	Poecilia_formosa-5.1.2	GCA_000485575.1	-	-	-	
 <b>Anole lizard</b>	<i>Anolis carolinensis</i>	28377	AnoCar2.0	GCA_000090745.1	-	-	-	
 <b>Armadillo</b>	<i>Dasypus novemcinctus</i>	9361	Dasnov3.0	GCA_000208655.2	-	-	-	
 <b>Zebra Finch</b>	<i>Taeniopygia guttata</i>	59729	taeGut3.2.4	-	Y	-	-	
 <b>Zebrafish</b>	<i>Danio rerio</i>	7955	GRCz10	GCA_000002035.3	Y	Y	-	

[Credits page for species images](#)

**Other Metazoa**

Additional metazoan genomes (initially insect vectors and nematodes) are available from [EnsemblMetazoa](#) [Ⓔ](#)

**Plants and Fungi**

Plant and fungal genomes can be found at [EnsemblPlants](#) [Ⓔ](#) and [EnsemblFungi](#) [Ⓔ](#)

**Protists, Bacteria and Archaea**

Unicellular eukaryotic and prokaryotic genomes can be found at [EnsemblProtists](#) [Ⓔ](#) and [EnsemblBacteria](#) [Ⓔ](#) respectively.

Search:  for

e.g. [BRCA2](#) or [rat 5:62797383-63627669](#) or [rs699](#) or [coronary heart disease](#)

**PAX6 (Human Gene)**  
**ENSG00000007372** 11:31784792-31817961:-1  
 Paired box 6 [Source:HGNC Symbol;Acc:HGNC:8620]

LRG\_720 (LRG display in Ensembl record; description: Locus Reference Genomic record for **PAX6**) is an external reference matched to Gene ENSG00000007372

Variant table • Phenotypes • Location • External Refs. • Regulation • Orthologues • Gene tree

**PAX6-011 (Human Transcript)**  
**ENST00000379107** 11:31789182-31810305:-1  
 Paired box 6 [Source:HGNC Symbol;Acc:HGNC:8620]

P26367 (UniProtKB/Swiss-Prot record; description: Paired box protein **Pax-6**), with synonyms of Q99413, Q6N006, is an external reference matched to Translation ENSP00000368401

Location • External Refs. • cDNA seq. • Exons • Variant table • Protein seq. • Population • Protein summary

**PAX6-004 (Human Transcript)**  
**ENST00000379109** 11:31789936-31810667:-1  
 Paired box 6 [Source:HGNC Symbol;Acc:HGNC:8620]

P26367 (UniProtKB/Swiss-Prot record; description: Paired box protein **Pax-6**), with synonyms of Q99413, Q6N006, is an external reference matched to Translation ENSP00000368403

Location • External Refs. • cDNA seq. • Exons • Variant table • Protein seq. • Population • Protein summary

**PAX6-005 (Human Transcript)**  
**ENST00000379111** 11:31789922-31811045:-1  
 Paired box 6 [Source:HGNC Symbol;Acc:HGNC:8620]

P26367 (UniProtKB/Swiss-Prot record; description: Paired box protein **Pax-6**), with synonyms of Q99413, Q6N006, is an external reference matched to Translation ENSP00000368406

Location • External Refs. • cDNA seq. • Exons • Variant table • Protein seq. • Population • Protein summary

Armadillo  
 (*Dasyurus novemcinctus*)

1-to-1  
[View Gene Tree](#)

PAX6 ([ENSDNCG00000000761](#))  
[Compare Regions](#) (JH561443)

[View Sequence Alignments](#)

[Orthologue Alignment](#)  
[View Protein Alignment](#)  
[View cDNA Alignment](#)

How many **PAX** protein paralogues are there for human? Suggest a prettier naming scheme than **PAX1**, **PAX2**, ...

Now click the **Exons** link (from **Transcript-based displays** → **Sequence**). **Exons**, **Introns** and **Variations** within **Exons** are clearly displayed.

Intron	Start	End	Start	End	Length	Sequence
<a href="#">Intron 2-3</a>	31,810,827	31,806,926			3,902	gtgagtcgcgtctctctctctctctgct.....ttttctcctctgtttgtcttag
<a href="#">ENSE00001098662</a>	31,806,925	31,806,849	-	-	77	GGGGAAGACTTTAACTAGGGGCGCGCAGATGTGTGAGGCCCTTTATTGTGAGAGTGGACA GACATCCGAGATTTCAG
<a href="#">Intron 3-4</a>	31,806,848	31,806,463			386	gcaagtctctgtggtggtgcttcttg.....ttaactcctatctttctgtctaacag
<a href="#">ENSE00002523992</a>	31,806,462	31,806,402	-	1	61	AGCCCATATTCGAGCCCGTGGAATCCCGCGGCCCCAGCCAGAGCCAGCATCGAGAAC A
<a href="#">Intron 4-5</a>	31,806,401	31,802,835			3,567	gtaagtgcctctggtctctctctgga.....tttctctctctctctctctctctcag
<a href="#">ENSE00003602163</a>	31,802,834	31,802,704	1	0	131	GTCACAGCGGAGTGAATCAGCTGGTGGTCTTTTCACAGGCGGCCACTGCCGGACT CCACCGGGCAGAAGATTGTAGAGCTAGCTCAGAGCGGGGCGGGCGGTGCGACATTCCG GAATTCTCCAG
<a href="#">Intron 5-6</a>	31,802,703	31,801,913			791	gtgatcctcccgcgccgccccact.....ttgaaggtatatctttgtgttatag
<a href="#">ENSE00003512677</a>	31,801,912	31,801,871	0	0	42	ACCCATGCAGATGCAAAAGTCCAAGTGTGGAACAATGAAAC
<a href="#">Intron 6-7</a>	31,801,870	31,801,777			94	gtaagcttgctattgtttaatgcat.....ttttctgtccacttccccatgcat
<a href="#">ENSE00003523920</a>	31,801,776	31,801,561	0	0	216	GTTGTCAAACGATGTGTAGTAAATTCTGGGAGGTATTACGAGACTGCTCCATCACA CCAGGGCAATCGGTGCTGTAAACCGAGAGTGGGACTCCAGAGTTGTAGCAAAATA GCCCACTATAAGCGGAGTGCCCGTCCATCTTTCTTGGGAAATCGGAGACAGATTAATG TCCGAGGGGGTCTGTACCAACGATAACATACAAAC

How long is the sixth exon and why would this concur with your expectations? \_\_\_\_\_

Click on some of the colourful variation locations. The colours are explained in the legend at the top of the display.

Exons/ Introns	Translated sequence	Flanking sequence	Intron sequence	UTR			
Variants	3 prime UTR	5 prime UTR	Coding sequence	Frameshift	Inframe deletion	Missense	Splice region
	Stop gained	Stop lost	Synonymous				

The variations come from a number of variation databases, including **dbSNP**. The **dbSNP** entries are those whose names begin with “rs”. **dbSNP** can be investigated directly at the **NCBI**, of course, but it very handy to have all the variation information built into **Genome Databases** such as **Ensembl**.

The figure displays three screenshots of the Ensembl Variant Effect Predictor (VEP) interface, each showing a different variant. Each screenshot includes a table with the following columns: Position, Alleles, cDNA position, Protein position, Amino acids, Codons, and Consequences. The first screenshot shows variant rs750195797, the second shows C1080974, and the third shows rs755018027. Each screenshot also includes a link to 'Explore this variant' and a link to 'Gene/Transcript Locations'.

Variant	Position	Alleles	cDNA position	Protein position	Amino acids	Codons	Consequences
rs750195797	11:31801684	T/C	745	92	V/V	gtA/gtG	Synonymous variant
C1080974	between 11:31801701 & 11:31801702	HGMD_MUTAT...	728	87			Coding sequence variant
rs755018027	11:31794652	C/T	1171	234	E/E	gaG/gaA	Synonymous variant

Click on the **Domains & features** link (from **Transcript-based displays** → **Protein Information**).

Domain source	Start	End	Description	Accession	InterPro
PANTHER	1	411	-	PTHR24329	-
PANTHER	1	411	-	PTHR24329:SF294	-
Prosite_profiles	222	282	Homeobox domain	<a href="#">PS50071</a>	<a href="#">IPR001356</a> <a href="#">[Display all genes with this domain]</a>
Smart	224	286	Homeobox domain	<a href="#">SM00389</a>	<a href="#">IPR001356</a> <a href="#">[Display all genes with this domain]</a>
Pfam	226	281	Homeobox domain	<a href="#">PF00046</a>	<a href="#">IPR001356</a> <a href="#">[Display all genes with this domain]</a>
Prosite_patterns	257	280	Homeobox, conserved site	<a href="#">PS00027</a>	<a href="#">IPR017970</a> <a href="#">[Display all genes with this domain]</a>
Superfamily	6	143	Homeodomain-like	<a href="#">SSF46689</a>	<a href="#">IPR009057</a> <a href="#">[Display all genes with this domain]</a>
Gene3D	201	284	Homeodomain-like	1.10.10.60	<a href="#">IPR009057</a> <a href="#">[Display all genes with this domain]</a>
Superfamily	205	283	Homeodomain-like	<a href="#">SSF46689</a>	<a href="#">IPR009057</a> <a href="#">[Display all genes with this domain]</a>
Pfam	4	142	Paired domain	<a href="#">PF00292</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
Smart	4	142	Paired domain	<a href="#">SM00351</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
Prosite_profiles	4	144	Paired domain	<a href="#">PS51057</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
PRINTS	8	23	Paired domain	<a href="#">PR00027</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
PRINTS	26	44	Paired domain	<a href="#">PR00027</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
PRINTS	60	77	Paired domain	<a href="#">PR00027</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
PRINTS	78	95	Paired domain	<a href="#">PR00027</a>	<a href="#">IPR001523</a> <a href="#">[Display all genes with this domain]</a>
Gene3D	7	86	Winged helix-turn-helix DNA-binding domain	1.10.10.10	<a href="#">IPR011991</a> <a href="#">[Display all genes with this domain]</a>
Gene3D	87	150	Winged helix-turn-helix DNA-binding domain	1.10.10.10	<a href="#">IPR011991</a> <a href="#">[Display all genes with this domain]</a>

Are you shocked and dismayed that the precise location of the **PAX6** Homeobox domain is not identically predicted by the **SMART** and **Pfam Domain Databases**? If not, why not?

How is that all the predictions, of different domain databases, for a **Paired domain** have the same **Interpro** identifier?

Why does **PRINTS** appear to predict four **Paired\_domains**?

Click on the link to the **SMART** entry for the **Paired domain** (**SM00351**).

Here you will find (quoted from **Interpro**) a **Description** of a **Paired domain**.

Where would you expect a **Paired domain** to occur in a protein?

What expectations do you have concerning what typically follows a **Paired domain**?

The paired domain is an approximately 126 amino acid DNA-binding domain, which is found in eukaryotic transcription regulatory proteins involved in embryogenesis. The domain was originally described as the 'paired box' in the Drosophila protein paired (prd) [(PUBMED:2877747), (PUBMED:3123319)]. The paired domain is generally located in the N-terminal part. An octapeptide [(PUBMED:10811620)] and/or a homeodomain can occur C-terminal to the paired domain, as well as a Pro-Ser-Thr-rich C terminus.

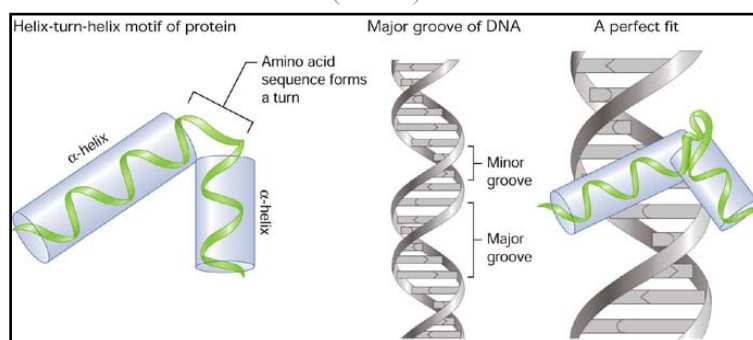
Paired domain proteins can function as transcription repressors or activators. The paired domain contains three subdomains, which show functional differences in DNA-binding. The crystal structures of prd and Pax proteins show that the DNA-bound paired domain is bipartite, consisting of an N-terminal subdomain (PAI or NTD) and a C-terminal subdomain (RED or CTD), connected by a linker. PAI and RED each form a three-helical fold, with the most C-terminal helices comprising a helix-turn-helix (HTH) motif that binds the DNA major groove. In addition, the PAI subdomain encompasses an N-terminal beta-turn and beta-hairpin, also named 'wing', participating in DNA-binding. The linker can bind into the DNA minor groove. Different Pax proteins and their alternatively spliced isoforms use different (sub)domains for DNA-binding to mediate the specificity of sequence recognition [(PUBMED:11103953), (PUBMED:15148315)].

The reason for these two questions will become apparent later.



The second paragraph of the **Description** claims, in gross summary:

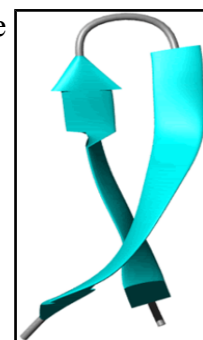
- A paired domain is a DNA binding domain that has 2 binding regions each of which involves a helical triplet
- The second and third helices of each helical triplet form **Helix-Turn-Helix (HTH)** motifs



- The **HTH** regions bind the DNA major groove<sup>s</sup>

- The first helical triplet is preceded by a  **$\beta$ -turn** and  **$\beta$ -hairpin** (“wing”) that participate in the DNA binding

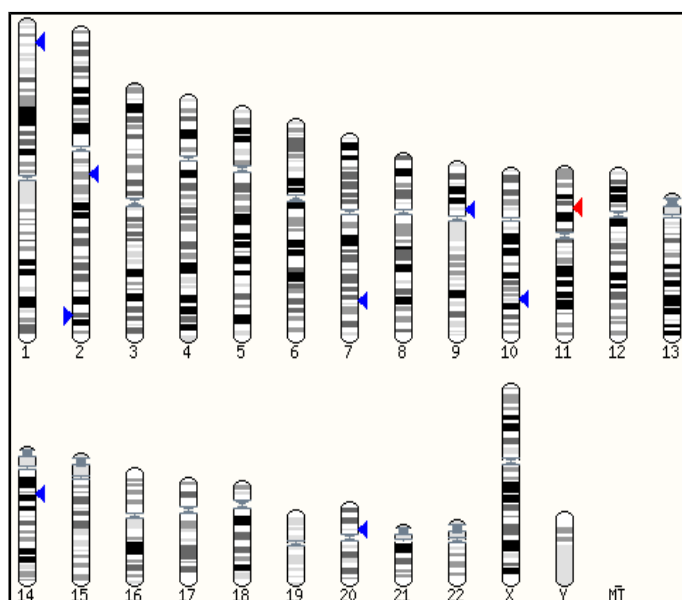
- The linker region between the two helical triplets can bind the **DNA minor groove**



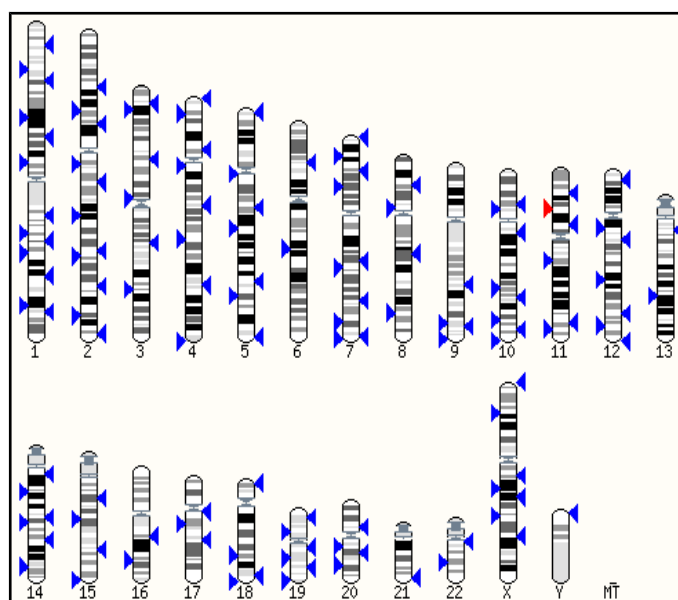
Bear this in mind when looking at the 3D structures a couple of pages on.

Click on **Display all genes with this domain** for the **Paired domain** and **Homeobox domain** InterPro families. The locations of all genes including each domain will be displayed graphically and textually. **PAX6** is shown in red.

#### Paired domain - IPR001523



#### Homeobox domain - IPR001356

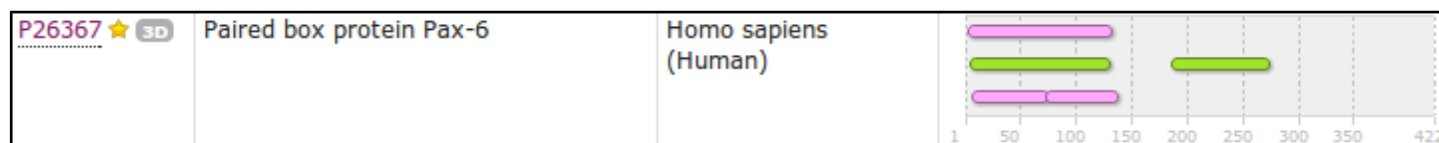


Which domain, **Paired domain** or **Homeobox domain** is more common in humans? \_\_\_\_\_

How many human **PAX** genes are there? \_\_\_\_\_

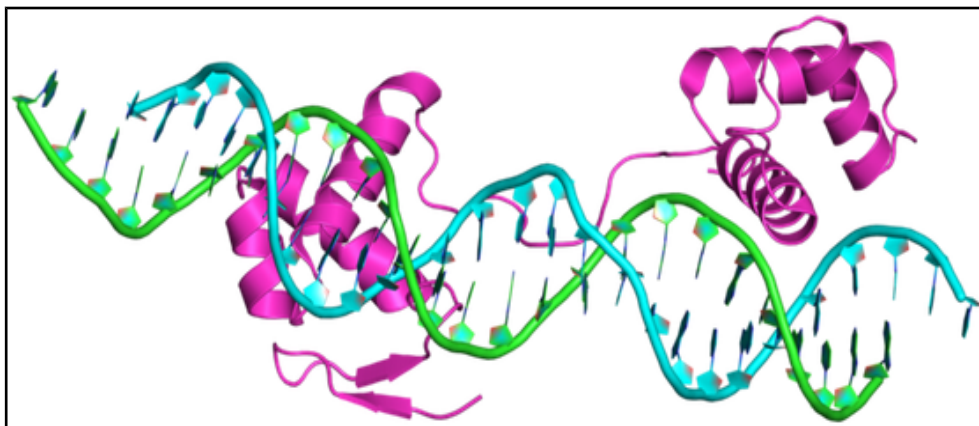
Are all the **PAX** genes on **Chromosome 11**? \_\_\_\_\_

Move back to the **Domains & features** display. Link to the **InterPro** database entry for **Paired domain**, also known as **IPR001523**. Here you will find the origins of the **SMART** documentation. Click on the **Proteins matched** link. You will see listed a number of representations of proteins that, according to **InterPro**, include a **Paired domain**. Amongst these will be the human **PAX6** protein, also known as **P26367**<sup>6</sup>. There are links provided to entries in a number of relevant databases for each listed protein.



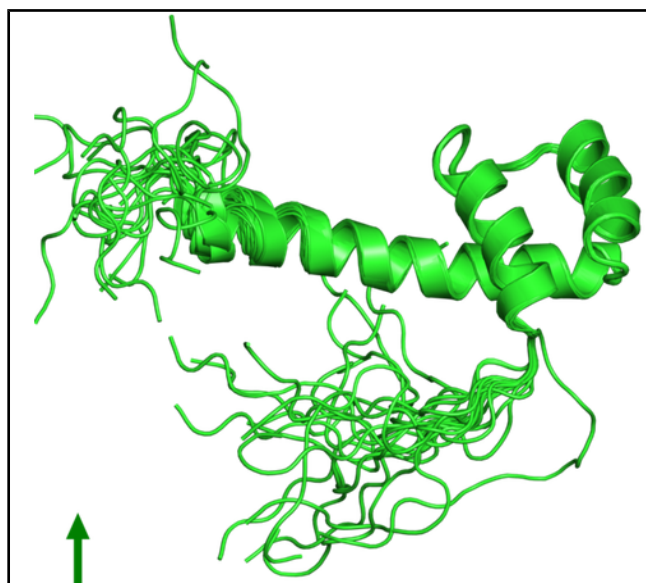
What type of **Helix-Turn-Helix (HTH)** is predicted by **InterPro** for all the **Paired domains** in the list?

Click on the **Structures** link in the top left hand corner of the page. **InterPro** will offer links to relevant entries in the **PDB**, **SCOP** and **CATH**<sup>7</sup> databases. Click on the link to the **6pax** entry in the **PDB** database. You will arrive at the entry for **6pax** in **PDB**, the European version of **PDB** maintained at the **EBI**. Views of this structure are offered on the right hand side of the page. Click on the largest image which shows the paired box protein domain binding DNA rather beautifully. Once you have admired this image sufficiently, move back to the **6pax PDB** entry. From the **Quick links** on the right of the page, select the **3D Visualisation** option.



The **SMART** documentation you read earlier suggested two paired box subdomains, each of which “... form a three-helical fold, with the most C-terminal helices comprising a **helix-turn-helix (HTH)** motif that binds the **DNA major groove**”. Move your image around to confirm this assertion.

The same **SMART** documentation claims the subdomain nearer the N terminal “... encompasses an N-terminal **beta-turn** and **beta-hairpin**, also named '**wing**', participating in DNA-binding. The linker can bind into the **DNA minor groove**”. Manipulate your image to investigate the veracity of these assertions.



Once you have seen all there is to see of **6PAX**, move back to the **Ensembl Domains & features** display. Try the same tricks with the **InterPro Homeobox domain**. This time, it is difficult to find **P26367** in the huge list<sup>8</sup> **Proteins matched**, but you do not need to in order to link to the **Structures**. There are many more structures to choose from this time. I suggest you go for **2cue**. You have to imagine the DNA this time.

It looks rather as if the **Homeobox domain** also includes a helical triplet including a **Helix-Turn-Helix**. You could have confirmed this by reference to the relevant **SMART** documentation (as you did for the **Paired box** domain). It is the **HTH** that the **Homeobox** uses to bind to DNA.

**InterPro** did not detect the **Homeobox HTH** as it did the **Paired box HTH**. Have you any thoughts as to why this might be?

Can you explain the strangely frayed ends displayed in some of the representations of the **2cue** 3D structure? \_\_\_\_\_

6 Third from the bottom of the first page, last time I counted.

7 **PDB** is the main database for **3D** protein structures. **SCOP** and **CATH** are also **3D** structure related databases.

8 If you really wanted to, the best approach is to search for **P26367** in the search box at the top of the page and then look for the **Homeobox domain** entry in the **Detailed signature matches** list.

To end, a gesture towards demonstrating that you could quite easily have computed most of the information you have been accessing, ready packed, from various databases. There are many way this objective could be achieved, I choose to search for the features of the **PAX6** protein.

As has been discovered from several information sources, the **PAX6** human protein has two DNA binding domains. A paired box at the **N terminal** and a homeobox a little further along. Both of the domains include **Helix-Turn-Helix (HTH)** motifs. In this exercise, you will investigate how you might discover these domains and motifs using the various freely available domain databases (discussed previously) and other feature prediction programs. Clearly, this is superfluous for this particularly, well documented protein, but a valuable option in other circumstances.

One approach would be to consider each relevant domain database in turn. Each major domain database has its own Home web site and customised software to take **Query** protein sequences, compare those sequences with domain representations (typically based on **Hidden Markov Models**) and to report convincing matches. This would work, but would be tedious as there are many viable databases to consider. It would be dangerous to rely on too few of the databases available as none is perfect. You need a consensus prediction to be sure you miss nothing.

Also, you would need to know which databases are particularly appropriate for each domain you considered might be present. All databases cannot be optimised for all types of domain (for example, the **SMART** database specialises in domains that occur in signalling proteins).

So, let us not search individual domain databases in the main part of these exercises. Instead, I offer a supplementary exercise investigating a representative selection of the available searches. I selected the **Prosite**, **Pfam** and **PRINTS** domain databases, If you do this exercise, consider particularly the **PRINTS** section. It illustrates how and why **PRINTS** just fails to see one of the two **PAX6** domains (as you already discovered when looking at **UniProt**).

Here, use just **Interpro** to do the whole job. **Interpro** will search for all domains using the appropriate domain databases, thus removing the tedium of interrogating a miscellany of domain searching resources individually.

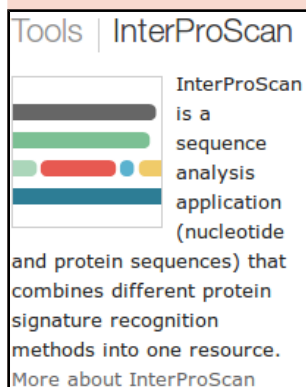


InterPro defines protein families according to the way that proteins match elements of a wide range of protein family databases, including all those we have discussed thus far. **Interpro** provides a search tool that will search all or any of the major protein family databases and assign **Interpro** family associations to the query protein(s) accordingly. To have a look at some of the possibilities offered by **Interpro**, Go to:

<http://www.ebi.ac.uk/interpro/>

If you were to enter the **PAX6** human protein into the obvious place on the **InterPro** home page, you would produce almost exactly the results you saw many pages back, when you were looking at **GeneCards**. Do this if you have the time and inclination.

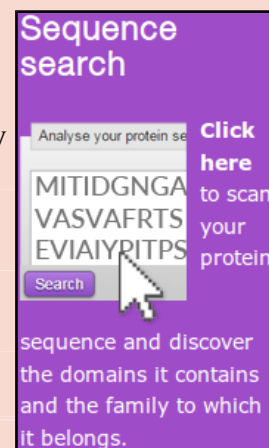
By implication, **InterPro** offers a fuller experience via the **InterProScan** search tool. Other than the opportunity not to search **ALL** the domain databases, and having the results arranged slightly differently, I am unsure what the extra effort brings? Never mind, there are many things of which I am unsure, so, from the **InterPro** Home page ...



Select the **InterProScan** link. Here you will be offered the opportunity to download the **InterProScan** program.

I am not sure this is too useful an offer for most? But it is there.

For now, chose the online **Sequence search**.



You will arrive at a page that looks very similar to that from which you started, as far as the offer to run a domain search is concerned? Except! We now have **Advanced options**. Click on the **Advanced options**.

The **Advanced options** only allow you to choose which databases you wish to search and which feature prediction programs you wish to run. The default is to use all the databases and to run all the predictor programs. I struggle to imagine an occasion I would want to save the **EBI** servers a few cycles by considering which options to deselect, but it so nice to know I could if I wished to.

In passing, the offer to run the feature predictor programs in the **Other sequence features** section is relatively new. Of course, all these programs could be run individually from their home websites (follow the links behind the program names), in the same way as the domain databases can be searched individually. **Interpro** just aims to make thing easy for the user. The programs currently offered are:

- **Coils** is a program for predicting **coiled coils**.
- **Phobius** & **TMHMM** are programs to predict **Transmembrane regions** (essentially **hydrophobic, uncharged** regions). There is no reason to expect any **Transmembrane regions** in this protein.
- **SignalP** predicts the presence and location of **signal peptide cleavage sites** in amino acid sequences from different organisms. I am pretty certain that there is no reason to expect signal peptides in this protein.

Do you think it a good idea for **Interpro** to offer feature prediction programs as well as domain database searches?\_\_

Analyse your protein sequence

>sp|P26367|PAX6\_HUMAN Paired box protein Pax-6 OS=Homo sapiens GN=PAX6 PE=1 SV=2  
 MQNSHSGVNQLGGVFVNGRPLPDSTRQKIVEL  
 AHSGARPCDISRIQVSNCGVSKILGRY  
 YETGSIRPRAIGGSKPRVATPEVVSIAQYKRECP  
 SIAFWIIRDRII SEGVCCTNDNIPSV

Advanced options

Select the applications to run: Uncheck all Select all

**Member databases**

Families, domains, sites & repeats

☒ CDD ☒ HAMAP ☒ PANTHER ☒ PfamA ☒ PIRSF ☒ PRINTS  
☒ ProDom ☒ Prosite-Profiles ☒ SMART ☒ TIGRFAM ☒ Prosite-Patterns

Structural domains

☒ Gene3d ☒ SFLD ☒ SUPERFAMILY

**Other sequence features**

☒ Coils ☒ MobiDB Lite ☒ Phobius ☒ SignalP ☒ TMHMM

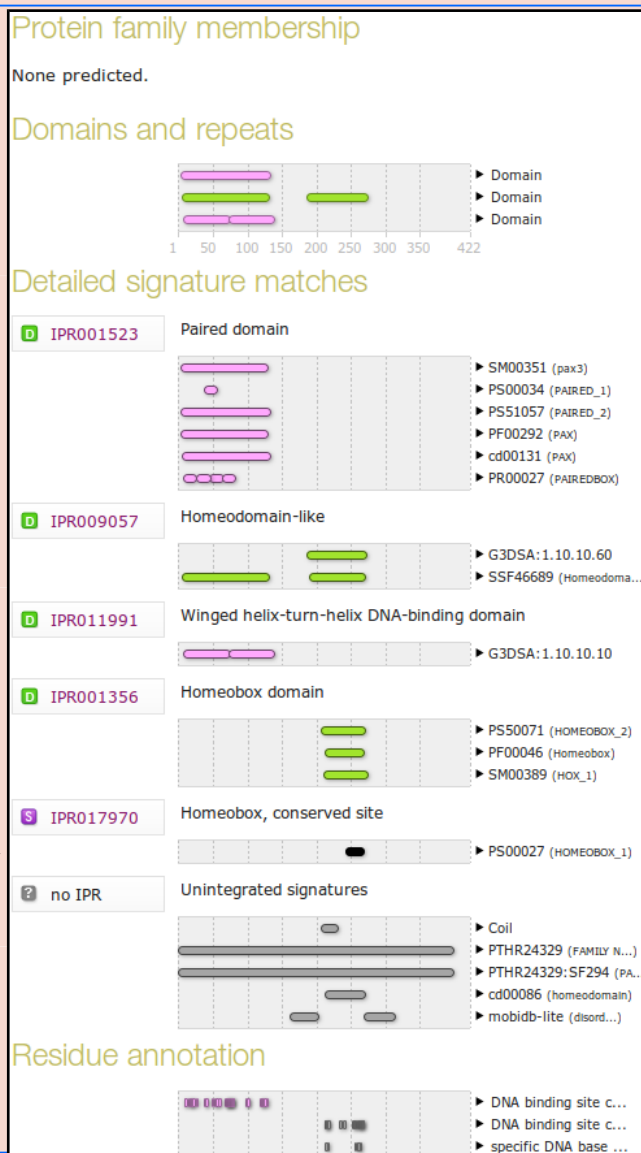
Search Clear Example protein sequence

Paste the human **PAX6** sequence into the patiently waiting box (from the file you made earlier called **pax6\_human.fasta**). Accept the "**do everything**" default. Click on the **Search** button.

After several moments of deep thought, filtering and validating, you will be presented with a table of results looking very much like the one your saw earlier when looking around **UniProtKB**.

There is, however, one significant difference. In the **Unintegrated signatures** section, you will see that a **coiled coil** has been detected by the program **Coils**. This was not included in the **UniProtKB** information, maybe as **Interpro** has only recently included analysis using **Coils**? **UniProtKB** might catch up next time it is updated.

Do you think the Coil prediction might be correct?\_\_





Notice that **Interpro** assigns both the **PAX** domain and the **Homeobox** domain of human **PAX6** to the **Interpro** family **Homeodomain-like**. Both of these associations are based on the hit behind the link **SSF46689**.

**SCOP classification**

Root: [SCOP hierarchy in SUPERFAMILY \[SCOP\\_0\]](#) (11)

Class: [All alpha proteins \[SCOP\\_46456\]](#) (284)

Fold: [DNA/RNA-binding 3-helical bundle \[SCOP\\_46688\]](#) (14)

**Superfamily:** [Homeodomain-like \[SCOP\\_46689\]](#) (19)

Families: [Homeodomain \[SCOP\\_46690\]](#) (40)

[Recombinase DNA-binding domain \[SCOP\\_46728\]](#) (5)

[Myb/SANT domain \[SCOP\\_46739\]](#) (15)


[SLIDE domain \[SCOP\\_100998\]](#)

[GARP response regulators \[SCOP\\_81683\]](#)

[DNA-binding domain of telomeric protein \[SCOP\\_46745\]](#) (2)


[Paired domain \[SCOP\\_46748\]](#) (3)

Follow this link and you will see it leads to the **Homeodomain-like**

**superfamily** of the  database that specialises in very general (SCOP<sup>9</sup> **superfamily** level) protein classifications. One **Superfamily** entry will typically correspond to a number of more specific domain definitions in other domain databases. Here you can see that the **Superfamily** domain **Homeodomain-like** includes both the **Homeodomain** & **Paired domain** Families.

Return to your **Interpro** results page. The **Gene3D** database is similar to **superfamily** but based on the **CATH** database<sup>10</sup>. It suggests the two **HTH** motifs of the paired box are both **Winged helix-turn-helix**. The **HTH** in the **Homeobox domain** is not detected?


Why might you suppose **Interpro** predicts only 2 of the 3 **helix-turn-helix domains** that might be expected?\_\_\_\_\_

Follow the link to the **Interpro** family **Homeodomain-like** (**IPR009057**). Click on the  button in the **Domain relationships** section to show the full list of **Homeodomain-like Interpro** domains.


**Contributing signatures**

Signatures from InterPro member databases are used to construct an entry.

**GENE3D**

 **G3DSA:1.10.10.60**  
(G3DSA:1.10.10.60)

**SUPERFAMILY**

 **SSF46689** (SSF46689)

Note also the **Contributing signatures** in the top right hand corner of the page. Here is listed the domain databases that are searched to determine the presence of an **Interpro Homeodomain-like** domain.

Essentially, if **Gene3D** finds a match with its **Homeodomain-like superfamily** and/or **Superfamily** finds a match with its **Homeodomain-like superfamily**, then **Interpro** acknowledges a match with its **Homeodomain-like** domain (**IPR009057**).

None of the other domain databases **Interpro** searches are used to determine membership of **IPR009057**.

**Domain relationships****D Homeodomain-like (IPR009057)**

- ... **D** DNA binding HTH domain, Fis-type (IPR002197)
- ... **D** DNA binding HTH domain, AraC-type (IPR018060)
- ... **D** DNA binding HTH domain, Psq-type (IPR007889)
- ... **D** DNA-binding HTH domain, TetR-type (IPR001647)
- ... **D** HTH CenpB-type DNA-binding domain (IPR006600)
- ... **D** Homeo-prospéro domain (IPR023082)
- ... **D** Homeobox domain (IPR001356)
- ... **D** Homeodomain, ZF-HD class (IPR006455)
- ... **D** Homeodomain, phBC6A51-type (IPR024978)
- ... **D** Mor transcription activator (IPR014875)
- ... **D** Rap1 Myb domain (IPR015010)
- ... **D** Resolvase, HTH domain (IPR006120)
- ... **D** SANT/Myb domain (IPR001005)
- ... **D** SLIDE domain (IPR015195)
- ... **D** SWIRM domain (IPR007526)
- ... **D** Transposase IS30-like HTH domain (IPR025246)
- ... **D** Transposase, Synechocystis PCC 6803 (IPR002622)
- ... **D** TyrR family, helix-turn-helix domain (IPR030828)

<sup>9</sup> Structural Classification Of Proteins.

<sup>10</sup> CATH is similar to SCOP in that it is another Structural classification database.

To obtain an impression of how widely spread throughout nature is this domain. Click on the **Species** button on the left hand side of the page.

As you can see, this is a very popular domain. You can make this list enormous by injudicious employment of the expansion buttons (the **Number of protein** links). Why not? It amused me for a few moments anyway.

Proteins matched: Homeodomain-like (IPR009057)

Filtered by species: **Schizosaccharomyces pombe (strain 972 / ATCC 24843) (Fission yeast)** (excludes child species) ([change species](#))

Showing 1 to 20 of 28 results

Accession	Protein name	Species	Domain architecture
<a href="#">Q13719</a> ★	SWIRM domain-containing protein laf1	Schizosaccharomyces pombe (strain 972 / ATCC 24843) (Fission yeast)	
<a href="#">Q13788</a> ★	SWI/SNF and RSC complexes subunit ssr1	Schizosaccharomyces pombe (strain 972 / ATCC 24843) (Fission yeast)	
<a href="#">Q13877</a> ★	DNA-directed RNA polymerases I, II, and III subunit RPABC5	Schizosaccharomyces pombe (strain 972 / ATCC 24843) (Fission yeast)	
<a href="#">Q14013</a> ★	RNA polymerase I-specific transcription initiation factor rrm5	Schizosaccharomyces pombe (strain 972 / ATCC 24843) (Fission yeast)	

Domain

Species: [Homeodomain-like \(IPR009057\)](#)

Key Species

Key species	Number of proteins	FASTA	Protein IDs
<a href="#">Arabidopsis thaliana (Mouse-ear cress)</a>	<a href="#">1268</a>		
<a href="#">Homo sapiens (Human)</a>	<a href="#">1134</a>		
<a href="#">Oryza sativa subsp. japonica (Rice)</a>	<a href="#">1023</a>		
<a href="#">Danio rerio (Zebrafish)</a>	<a href="#">961</a>		
<a href="#">Mus musculus (Mouse)</a>	<a href="#">893</a>		
<a href="#">Drosophila melanogaster (Fruit fly)</a>	<a href="#">487</a>		
<a href="#">Caenorhabditis elegans</a>	<a href="#">208</a>		
<a href="#">Escherichia coli (strain K12)</a>	<a href="#">104</a>		
<a href="#">Saccharomyces cerevisiae (strain ATCC 204508 / S288c) (Baker's yeast)</a>	<a href="#">36</a>		
<a href="#">Schizosaccharomyces pombe (strain 972 / ATCC 24843) (Fission yeast)</a>	<a href="#">28</a>		

Taxa

- ☒ **cellular organisms** [810656 proteins](#) | [FASTA](#) | [Protein IDs](#)
- ☒ **Archaea** [3269 proteins](#) | [FASTA](#) | [Protein IDs](#)
- ☒ **Bacteria (eubacteria)** [650577 proteins](#) | [FASTA](#) | [Protein IDs](#)
- ☒ **Eukaryota (eucaryotes)** [156810 proteins](#) | [FASTA](#) | [Protein IDs](#)
- ☒ **unclassified sequences** [4253 proteins](#) | [FASTA](#) | [Protein IDs](#)
- ☒ **Viruses** [1255 proteins](#) | [FASTA](#) | [Protein IDs](#)
- ☒ **other sequences** [19 proteins](#) | [FASTA](#) | [Protein IDs](#)

By clicking on the appropriate button, you can get to either the protein sequences in **Fasta** format or list their accessions codes. Try a few, but be careful! It really does get you **ALL** the sequences, and that is often quite a lot, which can take time.

THE END

DPJ – 2016.12.06

## Model Answers to Questions in the Instructions Text.

### Notes:

For the most part, these “**Model Answers**” just provide the reactions/solutions I hoped you would work out for yourselves. However, sometime I have tried to offer a bit more background and material for thought? Occasionally, I have rambled off into some rather self indulgent investigations that even I would not want to try and justify as pertinent to the objective of these exercises. I like to keep these meanders, as they help and entertain me, but I wish to warn you to only take regard of them if you are feeling particularly strong and have time to burn. Certainly not a good idea to indulge here during a time constrained course event!

Where things have got extreme, I am going to make two versions of the answer. One starting:

### Summary:

Which has the answer with only a reasonably digestible volume of deep thought. Read this one.

The other will start:

### Full Answer:

Beware of entering here! I do not hold back. Nothing complicated, but it will be long and full of pedantry.

This makes the Model answers section very big. **BUT**, it is not intended for printing or for reading serially, so I submit, being long and wordy does not matter. Feel free to disagree.

From your investigations using **Entrez**:

What were the features that you found?

**Summary:**The first feature was the **CoDing Sequence (CDS)** for a **PAX6** isoform.The other three features were the coding sequences for three **ELP4** isoforms.

```
complement(39424..>39569)
/gene="ELP4"
/gene_synonym="AN; AN2; C11orf19; dJ68P15A.1; hELP4;
PAX6NEB; PAXNEB"
/inference="similar to AA sequence (same
species):RefSeq:NP_001275654.1"
/exception="annotated by transcript or proteomic data"
/feature="isoform 2 is encoded by transcript variant 2;
elongator complex protein 4; PAX6 neighbor gene protein;
elongation protein 4 homolog"
/codon_start=3
/product="elongator complex protein 4 isoform 2"
/protein_id="NP_001275654.1"
/db_xref="CCDS:CCDS73271.1"
/db_xref="GeneID:26610"
/db_xref="HGNC:HGNC:1171"
/db_xref="MIM:606985"
/translation="MAAVATCGSVAASTGSAVATASKSNVTSFQRRGPRASVTNDSGP
RLVSIAGTRPSVRNGQLLVSTGLPALDQLLGGGLAVGTVLLIEEDKYNISPLLFKYF
LAEGIVNGHTLLVASAKEDPANILQELPAPLLDDCKCKEFDVYNHKTPESNIMKI
AWRYQLLPKMEIGPVSSSRFGHYDASKRMPQELIEASNMHGGFLPEKISSTLKVEPC
SLTPGYTKLLQFIQNIIEEGFDGNSPQKQKRNILRIGIQNLGSPWGGDICCANGG
GNHSLTKFLYVLRGLRLTSLACIITMPTHLIQNKAIARVTTLSDDVVGLSFISGE
ERETNPLYKYDHYGLHIRIQIPRLNLLICDESDVKDLAFKLKRLFTIEVQDNYLRQE
RNIYPPGFSYLLKQKDSAWGEGSLQHSFTLMSFLAKATAFASRLVRHSEPLKQNGSGR
IRQAAGPRLWHBGRQRQEPAGLLGIPP"
```

```
complement(39438..>39569)
/gene="ELP4"
/gene_synonym="AN; AN2; C11orf19; dJ68P15A.1; hELP4;
PAX6NEB; PAXNEB"
/inference="similar to AA sequence (same
species):RefSeq:NP_061913.3"
/exception="annotated by transcript or proteomic data"
/feature="isoform 1 is encoded by transcript variant 1;
elongator complex protein 4; PAX6 neighbor gene protein;
elongation protein 4 homolog"
/codon_start=1
/product="elongator complex protein 4 isoform 1"
/protein_id="NP_061913.3"
/db_xref="CCDS:CCDS7875.2"
/db_xref="GeneID:26610"
/db_xref="HGNC:HGNC:1171"
/db_xref="MIM:606985"
/translation="MAAVATCGSVAASTGSAVATASKSNVTSFQRRGPRASVTNDSGP
RLVSIAGTRPSVRNGQLLVSTGLPALDQLLGGGLAVGTVLLIEEDKYNISPLLFKYF
LAEGIVNGHTLLVASAKEDPANILQELPAPLLDDCKCKEFDVYNHKTPESNIMKI
AWRYQLLPKMEIGPVSSSRFGHYDASKRMPQELIEASNMHGGFLPEKISSTLKVEPC
SLTPGYTKLLQFIQNIIEEGFDGNSPQKQKRNILRIGIQNLGSPWGGDICCANGG
GNHSLTKFLYVLRGLRLTSLACIITMPTHLIQNKAIARVTTLSDDVVGLSFISGE
ERETNPLYKYDHYGLHIRIQIPRLNLLICDESDVKDLAFKLKRLFTIEVQDNYLRQE
RNIYPPGFSYLLKQKDSAWGEGSLQHSFTLMSFLAKATAFASRLVRHSEPLKQNGSGR
IRQAAGPRLWHBGRQRQEPAGLLGIPP"
```

```
complement(39533..>39569)
/gene="ELP4"
/gene_synonym="AN; AN2; C11orf19; dJ68P15A.1; hELP4;
PAX6NEB; PAXNEB"
/inference="similar to AA sequence (same
species):RefSeq:NP_001275655.1"
/exception="annotated by transcript or proteomic data"
/feature="isoform 3 is encoded by transcript variant 3;
elongator complex protein 4; PAX6 neighbor gene protein;
elongation protein 4 homolog"
/codon_start=2
/product="elongator complex protein 4 isoform 3"
/protein_id="NP_001275655.1"
/db_xref="CCDS:CCDS73272.1"
/db_xref="GeneID:26610"
/db_xref="HGNC:HGNC:1171"
/db_xref="MIM:606985"
/translation="MAAVATCGSVAASTGSAVATASKSNVTSFQRRGPRASVTNDSGP
RLVSIAGTRPSVRNGQLLVSTGLPALDQLLGGGLAVGTVLLIEEDKYNISPLLFKYF
LAEGIVNGHTLLVASAKEDPANILQELPAPLLDDCKCKEFDVYNHKTPESNIMKI
AWRYQLLPKMEIGPVSSSRFGHYDASKRMPQELIEASNMHGGFLPEKISSTLKVEPC
SLTPGYTKLLQFIQNIIEEGFDGNSPQKQKRNILRIGIQNLGSPWGGDICCANGG
GNHSLTKFLYVLRGLRLTSLACIITMPTHLIQNKAIARVTTLSDDVVGLSFISGE
ERETNPLYKYDHYGLHIRIQIPRLNLLICDESDVKDLAFKLKRLFTIEVQDNYLRQE
RNIYPPGFSYLLKQKDSAWGEGSLQHSFTLMSFLAKATAFASRLVRHSEPLKQNGSGR
IRQAAGPRLWHBGRQRQEPAGLLGIPP"
```

**Full Answer:**

Note that only the final coding exon of **ELP4** is within this **RefSeq** sequence, which is defined as the genomic region for **PAX6**. This is clear from the length of the **translations** offered. The exon referenced is only long enough to code for just over **40** amino acids which is far short of any of the three isoform sequences offered here.

Note also that this final coding exon of **ELP4** (stretching from **39424/39438/39533** to **39569** of this **RefSeq** entry) does **not** overlap the coding region of the **PAX6** gene itself (stretching from **16551** to **33028** of this **RefSeq** entry)".

In fact, the two entire genes do not overlap according to the evidence here. The entire **PAX6** gene extends from **5001** to **38170**. The portion of the **ELP4** gene that is included in this entry extends from **40170** (the end) to **38437** (in the opposite direction). This give a gap between the two genes stretching from **38171** to **38436**.

**RefSeqGenes**, comprise the entire gene plus **5,000** "extra" base pairs in either direction. The overlap here is entirely within the "extra" base pairs.

```
join(16551..16560,20128..20258,21186..21401,22106..22271,
28174..28332,28848..28930,29160..29310,29409..29524,
32102..32252,32943..33028)
/gene="PAX6"
/gene_synonym="AN; AN2; D11S812E; FVH1; MGDA; WAGR"
/feature="isoform a is encoded by transcript variant 1;
paired box protein Pax-6; paired box homeotic gene-6;
oculorhombin; aniridia type II protein"
/codon_start=1
/product="paired box protein Pax-6 isoform a"
/protein_id="NP_000271.1"
/db_xref="CCDS:CCDS31451.1"
/db_xref="LRG:p1"
/db_xref="GeneID:5080"
/db_xref="HGNC:HGNC:8620"
/db_xref="MIM:607108"
/translation="MQNSHSGVNLQGGVFNVRPLPDSTRQKIVELAHSGARPCDISR
ILQVSGNCVSKILGRYYETGSIIRPRAIGGSKPRVATPEVVSIAQYKRECPSIFAWEI
RDRLLSEGVCNTNDIPSVSSINRVLRLASEKQMGADGMYDKLRMLNGQTGSWGTRP
GWYPGTSVPGPQPTDQGCQQEGGENTNMISSNGEDSDEAQMRLQLKRLQRNRTSFT
QEQTAELEKEFERTHYPDVFAERLARAAIDLPARIQVWFSNRRAKWRREEKLRNQRN
QASNTSPSHIPISSSFSTSVYQIPQPTTPVSSFTSGSLMGRDOTALNTYTSALPPMPS
FTMANNLPMPQPPVPSQTSSYSCLPTSPSVNGRSYDTYTPPHMQTHMNSQPMGTS GTT
STGLISPGVSVFVQVPGSEPDMSQYVWRLQ"
```

```
gene 5001..38170
/gene="PAX6"
/gene_synonym="AN; AN2; D11S812E; FVH1; MGDA; WAGR"
/feature="paired box 6"
/db_xref="GeneID:5080"
/db_xref="HGNC:HGNC:8620"
/db_xref="MIM:607108"
```

As you will see later, **Ensembl** will confirm the lack of overlap between these two genes graphically as well as their relative positions.

```
gene complement(38437..>40170)
/gene="ELP4"
/gene_synonym="AN; AN2; C11orf19; dJ68P15A.1; hELP4;
PAX6NEB; PAXNEB"
/feature="elongator acetyltransferase complex subunit 4"
/db_xref="GeneID:26610"
/db_xref="HGNC:HGNC:1171"
/db_xref="MIM:606985"
```

The annotation (specifically the **gene\_synonyms**) of **ELP4** associate this gene with **PAX6**. However, I believe only because of its proximity.



Why might you have expected more features than there were?

### Summary:

All the evidence has suggested that **PAX6** has at least **2** isoforms. This would lead me to expect at least **2** CDS features here related to **PAX6**?

### Full Answer::

The explanation from the **NCBI** is that this sort of **RefSeq** entry is intended to be used as a template against which sequences from an individual can be mapped to seek variations. Only a token **CDS** feature is included to indicate the position of the gene. For such an entry, recording every isoform is not essential.

This sounded convincing to me, Until I began to wonder why there were three **CDS** features for **ELP4** which is not even the gene primarily represented by this entry? Maybe I will ask more questions if and when I ever have the strength. In the meantime, mostly for my information, I record their exact explanation here.

“ ... note that **RefSeqGene** defines genomic sequences to be used as reference standards for well-characterized genes. These sequences serve as a stable foundation for reporting mutations, for numbering exons and introns, and for defining the coordinates of other variations. We normally select one **RefSeq** transcript to serve as a reference standard. The goal is not to record all introns and exons of all isoforms, but just to choose one representative to help define the locus. Therefore, most of our **RSG** records have only a single **RefSeq** as reference standard. If an **LSDB** manager or other stakeholder requests that other **RefSeqs** be added as alternate standards, this can easily be done (with the complication that, if a public **LRG** exists, the **RefSeqGene** record is fixed). We receive requests from stakeholders to include **RefSeqs** that represent all known exons, or **RefSeqs** that have become community standards. Often, after creating an **RSG** using our own internal criteria, we receive stakeholder requests to change or add transcripts. Many of these requests come from the **LRG** project regarding transcripts to be included on the **LRG** records.

Generally, **RefSeq** accessions can be added or removed without reversioning, unless a transcript is upgraded or a new one defined that extends beyond the bounds of the **RSG**, or matches a new build of the genome, in which case the **RSG** will be extended and reversioned as needed.

Regarding the chromosomal locus, our standard range is 5 kb upstream from the 5' end and 2 kb downstream from the 3' end of the mRNAs with the greatest extent. For this calculation, we do indeed use all available **RefSeq** (NM\_) accessions. If the database manager or stakeholder has information on promoters or other upstream or downstream regulatory regions, we can certainly extend the **RefSeqGene** locus to accommodate these.

Regarding mismatches, the goal is to exactly match the current build of the genome, unless there is overwhelming transcript and EST evidence that a mismatch should be retained.

Regarding the confusing subject of exon numbering, exon numbers are currently provided only on **RSG** genomic records based on a subset of available transcript **RefSeqs** for the gene. These are often those selected by locus-specific databases as reference sequence reporting standards. You can find an explanation of how exons are numbered here:

<http://www.ncbi.nlm.nih.gov/refseq/rsg/faq/#exon>

You will find links to more information on **RefSeqGenes** on the home page for the **RefSeqGene** project:

<http://www.ncbi.nlm.nih.gov/refseq/rsg/>

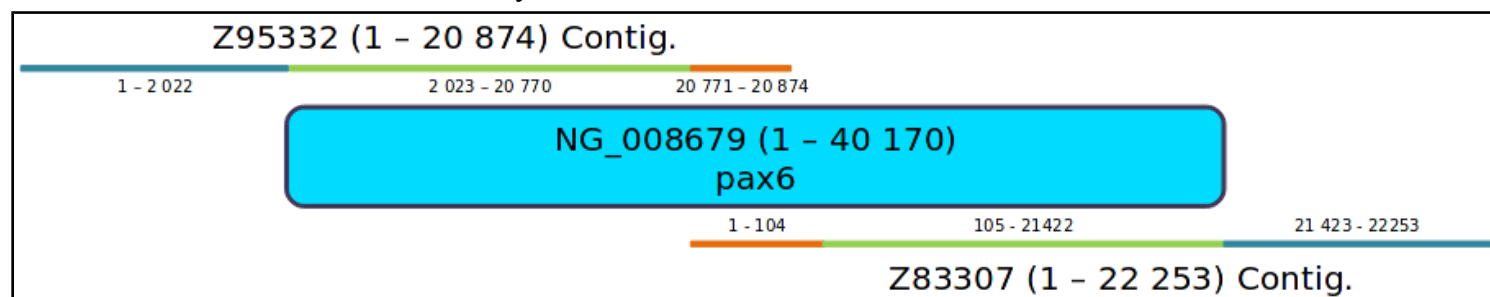
Regarding the **PAX6 RSG** sequence, only difference I see between **NG\_008679.1** and the current build of the genome (**GRCh38**) is an extra 'G' beyond the 3'-UTR of the **PAX6** transcripts (at **NC\_000011.10:g.31,819,125**). ... “

Yes, well I think I followed most of that? and that my interpretation is broadly correct? In summary, there are no fixed rules.

How does the alignment you generated match up with the annotation of the original **RefSeq** entry you discovered?

### Summary:

The most intuitive way of encapsulating graphically the way these two sequencing clones overlap was donated by **Cecilia Pinto (Oeiras, 2013.12.09-12)**. Much better than my rambling attempts, that I keep for sentimental reasons in the “Full Answer”. Thank you Cecilia.



### Full Answer:

Do not spend too much time working this one out, the picture above should be more than sufficient. I just needed to see it all balanced ... then I can sleep soundly?

If you do want to read on, I strongly suggest you look at the picture contributed by Cecilia (now promoted to the “**Summary Answer**”) first. So simple! I have to admit I cannot follow my own wonderful table at all now ... at least, not without bleeding! Although, it did feel good at the time?

- So ...
- ☐ [Human DNA sequence from clone CFAT5 on chromosome 11, complete sequence](#)
  - 1. 20,874 bp linear DNA  
Accession: Z95332.1 GI: 2190397  
[GenBank](#) [FASTA](#) [Graphics](#)
  - ☐ [Human DNA sequence from clone A1280 on chromosome 11, complete sequence](#)
  - 2. 22,253 bp linear DNA  
Accession: Z83307.1 GI: 1730464  
[GenBank](#) [FASTA](#) [Graphics](#)

```
Query 20771 GATCCGGAGCGACTTCGCCTATTTCCAGAAATTAAGCTCAAACCTTGACGTGCAGCTAGT 20830
Sbjct 1 GATCCGGAGCGACTTCGCCTATTTCCAGAAATTAAGCTCAAACCTTGACGTGCAGCTAGT 60
Query 20831 TTTATTTTAAAGACAAATGTCAGAGAGGCTCATCATATTTTCCC 20874
Sbjct 61 TTTATTTTAAAGACAAATGTCAGAGAGGCTCATCATATTTTCCC 104
```

The Query sequence is **Z95332** (Length **20,874**)

The Subject sequence is **Z83307** (Length **22,253**)

PRIMARY	REFSEQ_SPAN	PRIMARY_IDENTIFIER	PRIMARY_SPAN	COMP
1-18852	Z95332.1	2023-20874		
18853-40170	Z83307.1	105-21422		

NG_008679 Range Start	NG_008679 Range End	NG_008679 Range	Z95332 Range Start	Z95332 Range End	Z95332 Range	Z83307 Range Start	Z83307 Range End	Z83307 Range
-	-	-	1	2022	2022	-	-	-
1	18748	18748	2023	20770	18748	-	-	-
18749	18852	104	20771	20874 (end)	104	1	104	104
18853	40170 (end)	21319	-	-	-	105	21422	21318
-	-	-	-	-	-	21423	22253 (end)	831
		40171			20874			22253

### Legend:

Not used in construction of **RefSeq** entry **NG\_008679**

Non-overlapping **GenBank** entry used in construction of **RefSeq** entry **NG\_008679**

Overlapping **GenBank** entry used in construction of **RefSeq** entry **NG\_008679**

Total entry lengths

The **RefSeq** entry was thus constructed by overlapping the two **Genbank** entries and then manually trimming away the edges to form a biologically meaning region. If I was a bit brighter, I think I might have come to that conclusion without the fuss above? Oh well, one has to use what one has.

I refer you again to the far more intuitive way of encapsulating the same message graphically, donated by **Cecilia Pinto** that is now the “**Summary Answer**” above. Much better! Thank you once more Cecilia.

Using the evidence of the protein alignments, which **PAX6** isoforms do the fruitfly orthologues most resemble?

Fruitfly ( <i>Drosophila melanogaster</i> )	1-to-many <a href="#">View Gene Tree</a>	ey (FBgn0005558) <a href="#">Compare Regions</a> (4,697,589-721,173-1)
		<a href="#">View Sequence Alignments</a> <a href="#">Orthologous Alignment</a>
Fruitfly ( <i>Drosophila melanogaster</i> )	1-to-many <a href="#">View Gene Tree</a>	toy (FBgn0005559) <a href="#">View Protein Alignment</a> <a href="#">View cDNA Alignment</a> <a href="#">Compare Regions</a> (4,389,725-1,007,322-1)
		<a href="#">View Sequence Alignments</a>

There are two **Fruitfly** orthologues recorded here, with the gene names **ey** and **toy**. Looking at the first few lines of the protein alignments for these genes, it is clear that that **14** amino acid insert that defines **isoform 5a** (**THADAKVQVLDNQN**) is not present in either. It is therefore reasonable to conclude that the representative fly proteins are both closest to the canonical protein sequence of **PAX6** human (**isoform 1**).

```

ENSP00000404100/1-436      -MQN-----SHSGVNQLGGVFNVRPLPDSTR
FBpp0099810/1-898          GKPSPTMEAVEASTASHPHSTSSYFATTYYHLTDDCEHSGVNQLGGVFVGGRPLPDSTR
                               * :                               , ***** : *****

ENSP00000404100/1-436      KIVELAHSGARPCDISRILQTHADAKVQLDNQNVNSGCVSKILGRYYETGSIRPRAIG
FBpp0099810/1-898          KIVELAHSGARPCDISRILQ-----VSNGCVSKILGRYYETGSIRPRAIG
                               *****                               ,*****

ENSP00000404100/1-436      SKPRVATPEVVS KIAQYKRECP SIFAW EIRDRLLEGVCTNDNI PSVSSINRVLRNLAS
FBpp0099810/1-898          SKPRVATAEVVSKI SQYKRECP SIFAW EIRDRLLEQNVCTNDNI PSVSSINRVLRNLAA
                               *****,*****,* ,*****

ENSP00000404100/1-436      -MQN-----SHSGVNQLGGVFNVRPLPDSTRQKIVELAH
FBpp0088249/1-543          MMLTTEHIMHGHPHSVQG STLFGC STAGHSGINQLGGVVYNVRPLPDSTRQKIVELAH
                               * ,*****

ENSP00000404100/1-436      GARPCDISRILQTHADAKVQLDNQNVNSGCVSKILGRYYETGSIRPRAIGGSKPRVAT
FBpp0088249/1-543          GARPCDISRILQ-----VSNGCVSKILGRYYETGSIKPRAIGGSKPRVAT
                               *****                               ,*****

ENSP00000404100/1-436      EVVSKIAQYKRECP SIFAW EIRDRLLEGVCTNDNI PSVSSINRVLRNLASEKQMMGAD
FBpp0088249/1-543          PVVQKIADYKRECP SIFAW EIRDRLLESEQVCNSDNI PSVSSINRVLRNLASEKQEQQAQ
                               **,***,*****

```

### Protein alignment for toy

Well, maybe also it is not that simple? I would not be surprised If there were isoforms for **ey** and/or **toy** that were roughly equivalent to human **isoform 5a**. The alignment displayed could well reflect the relatively arbitrary choice of **Ensembl** as to which isoform it decides to use for the alignments, rather than any deep and meaningful biological truth. Already you can see that **Ensembl** prefers the (presumably) less important human isoform, merely because it is longer (more letters to match). Again, useful though these displays are, caution is required before reading too much “biology” into them.

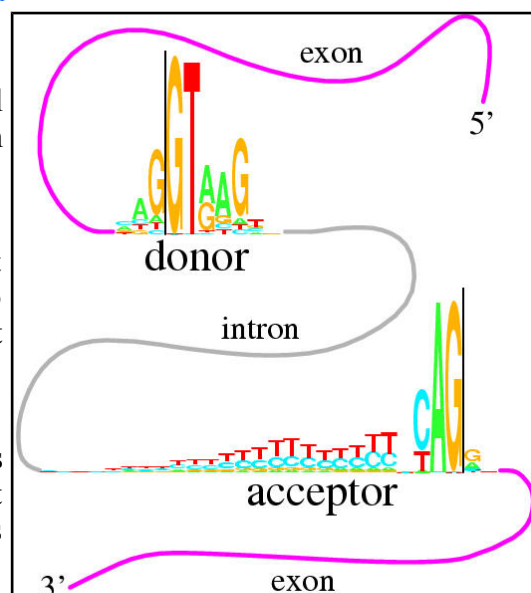
**Ensembl** does not pick up all fruitfly homologues of **PAX6**? Again, I wonder why. Mind you, **Ensembl** does only claim “**Selected orthologues**”? Still **prd**, in particular, is a pretty important one to pass over!

What are the first two bases and what are the last two bases of nearly every intron?

As you are probably well aware, introns are highly conserved at each end. They typically begin with **GT** and end with **AG**. This rule is obeyed by all but one of the introns of this transcript (**intron 3-4** starts **GC** rather than **GT**).

As the cartoon suggests, the conservation does not apply just to the first and last two bases, but that is where the conservation is most strict. So strict that when exceptions from this rule were sought in the databases, it was thought most of the deviations were due to annotation error!

The cartoon also suggests that introns have **C/T rich regions** towards their ends (the **Polypyrimidine tract**). This too is clearly evident in most of the introns of this transcript, even though only small parts of the introns are displayed.



How long is the sixth exon and why would this concur with your expectations?

It is **42** base pairs long, so it codes for **14** amino acids. Specifically, it codes for the **14** extra amino acids that define **isoform 5a**.

Explain the **Start Phase** and **End Phase** columns?

An exon/intron boundary can occur anywhere in a codon. The **Start** and **End Phases** record how an intron has been inserted into a coding region with respect to the coding reading frame.

If an exon ends at the end of a codon, then its **End Phase** is **0**.

Clearly, the next exon must begin at the start of a codon. Its **Start Phase** is also **0**.

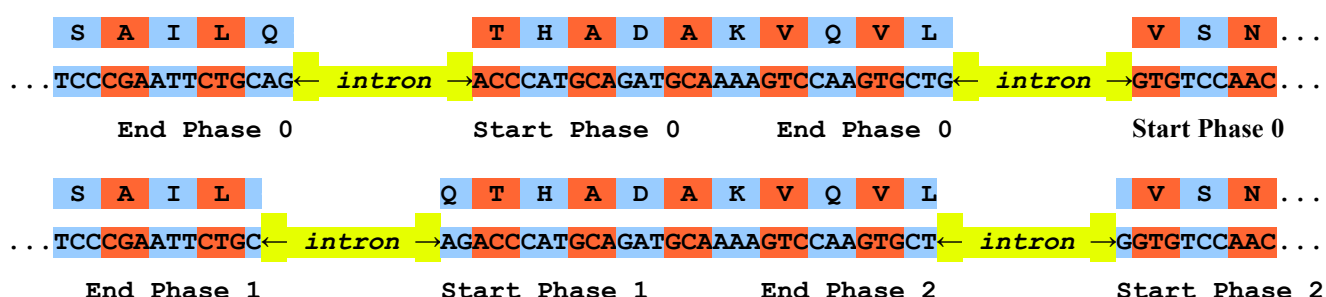
If an exon ends after the first base of a codon, then its **End Phase** is **1**.

Clearly, the next exon must begin after the first base of a codon. Its **End Phase** is also **1**.

If an exon ends after the second base of a codon, then its **End Phase** is **2**.

Clearly, the next exon must begin after the second base of a codon. Its **End Phase** is also **2**.

I attempt a picture, though I am sure that is clear? I just like pictures, and lots of colours.



Why does **Prints** appear to predict four **Paired\_domains**?

**Prints** does not find the **Homeobox\_domain** at all. If you were to investigate by using the **Prints search** carefully, you will find it nearly does, but the evidence is not quite strong enough. As has been discussed, none of these systems are perfect. They all occasionally fail. That is why it is always best to use **Interpro** to consult them all and deliver a consensus answer.

**Prints** appears to find **FOUR Paired\_domains**. This is only because of the way **Prints** works. **Prints** finds **FOUR** signatures (or **motifs**) that together indicate **ONE Paired domain**. **Prints** searches for ordered series of **motifs** that together indicate **domains**. Here it reports each of four motifs separately, but it is only claiming one **Paired domain**.

Which domain, **Paired domain** or **Homeobox domain** is more common in humans?

How many human **PAX** genes are there?

As you will have expected, there are but **9 Paired domains** in the Human genome. There are many more **Homeobox domains**.

Are all the **PAX** genes on **Chromosome 11**?

Of course not? What a stupid question!

Well, I suppose they could all be on **Chromosome 11**? By chance ... or maybe design ... who knows, the lack of predictable pattern in all this business never ceases to astound me.

But, philosophy aside, the answer is **NO**.



How does **Interpro** match with the **PAX6 Paralogues** reported by **Ensembl/GeneCards** earlier?

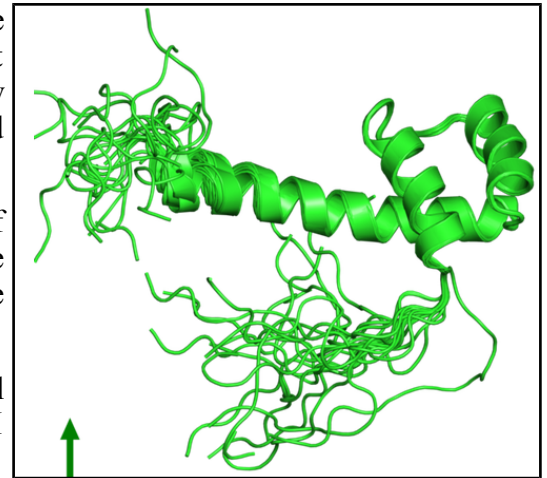
The evidence from both **GeneCards** and **Ensembl** is that there are **9 PAX** paralogues in Human. Yep, we all agree and ... these questions are becoming a trifle repetitive one feels!

Can you explain the strangely frayed ends displayed in some of the representations of the **2cue** 3D structure?

**2cue** is a 3D structure determined by Nuclear Magnetic Resonance (NMR). This is a process that does not involve immobilizing the target as a crystal (as is the case with structures determined by **X-ray crystallography**). Parts of the protein will still be moving around whilst its structure is being determined.

I think of **NMR** as analogous to taking a long exposure photograph of a group of children. Each child will appear in many different places! The frayed ends represent various positions in which the ends of the **homeobox** were detected during the **NMR** process.

In some views, including the one you were offered to move around, all the possible positions are averaged out before the structure is stored. I prefer the fuzzy view ... much more fun.



I broadly believe that which I have just typed, however, I must stress that my understanding of **NMR** is tragically incomplete. If anyone would like to offer a better explanation, I am very willing to hear it.

From your investigations of Domain & Motif identification using Interpro

Do you think it a good idea for **Interpro** to offer feature prediction programs as well as domain database searches?

Well ... why not? The purpose of **InterProScan** is to associate regions of query proteins with **Interpro** domains. This was originally achieved, exclusively, by simply comparing a query sequence with all entries of relevant individual domain databases. These entries being representations of alignments of examples of specific domains constructed by homology searching (i.e. **blast** and similar).

I would suggest including a few predictor programs would provide extra evidence gathered from more general, more theoretical definitions of domains. I would imagine the inclusion of these programs has improved and widened the picture provided by **InterProScan**.

Searching domain databases, typically composed of **HMM profiles**, such as **Pfam**, **Prosite** and **PRINTS** is quite different to running the predictor programs. As I cannot improve on the justification of this claim offered to me by Geoff Barton (Head of the group responsible for **Jalview**, **Jpred**, **Jnet** and much more), I will just reproduce his explanation here:

“ ... The main difference is that with an **HMM profile** you have a "specific" example of a domain or motif whereas with something like **COILS**, you have something trained across all examples.

For example, for secondary structure prediction, you could (a) do predictions of alpha-helix and beta-strand just by aligning a sequence to a protein of known structure, or an **HMM** from a family of aligned proteins of known structure. This is a specific case of secondary structure in the context of one protein family. Or (b) you can train a predictor from **ALL** protein families and then apply this. The advantage of (a) is it is very specific to the individual protein family and so should be more accurate for that family. The disadvantage is that it does not generalise to proteins that are not very like the specific example. The advantage of (b) is that it will work with any protein but will likely be less accurate than (a) for proteins that fit into the (a) category. ... “

Do you think the Coil prediction might be correct?

I do not recall anything in what we have discovered thus far that would directly suggest there should be a **coiled coil** here, in the middle of the **HTH**. However, **wikipedia** does suggest **coiled coils** are associated with **transcription factors** (which **pax6\_human** is).

" ... Many **coiled coil**-type proteins are involved in important biological functions such as the regulation of **gene expression**, e.g. **transcription factors**. ... "

I think I would not be overly convinced by this prediction, but I would not make that judgement with any great confidence. The all knowing **wikipedia** says:

“ ... **Coiled coils** usually contain a repeated pattern, **hxxhxc**, of **hydrophobic (h)** and **charged (c)** amino-acid residues, referred to as a **heptad repeat**. ... “

Geoff Barton comments:

“ ... Sometimes the pattern that is particular to **coiled-coils** also turns up in other helices that pack against each other. You would need to look at some examples of coiled-coil structures to see if the example you are using fits structurally. ... “

Which seems very reasonable. The **heptad repeat** pattern could easily occur just by chance. **COILS** surely cannot predict the structure of the helices well enough to make an assured judgement? **COILS** offers a suggestion the user must follow up with other resources.

There is also the evidence that **Jpred** (a system for secondary structure prediction), possibly using the **COILS** program disguised as **LUPAS**, does not detect any coiled coils. This could be for a number of reasons. Possibly **LUPAS** is not the same program as **COILS**, or it is a different version, or **Jpred** runs **COILS**, but with different parameters.

Not many clear and confident answers in Bioinformatics are there!

Why might you suppose **Interpro** predicts only **2** of the **3 helix-turn-helix domains** that might be expected?

**2 Winged helix-turn-helix (wHTH) DNA-binding domains** are predicted coincident with the helical triplets of the **Paired domain**. This should broadly match your expectations.

No **helix-turn-helix (HTH) domain** is detected coincident with the **Homeobox domain**, where one might also have been expected?

I am not entirely certain why this might be, so I speculate.

**Pfam** attempts to classify a variety of types of **HTH**, and offers a range of **HTH** domain models (**HTH\_17**, **HTH\_38**, **HTH\_39** and **HTH\_40** to name but a few) and a number of **wHTH** domain models (including **HTH\_33** and **HTH\_24**).

**Interpro** also has a considerable number of **HTH** entries (**IPR017895**, **IPR032877**, **IPR007394**, **IPR013197** and more) and **wHTH** entries (**IPR005104**, **IPR023120** to name but 2).

**Contributing signatures**

Signatures from InterPro member databases are used to construct an entry.

- SUPERFAMILY**
  - SSF46785** (SSF46785)
- Pfam**
  - PF12840** (HTH\_20)
  - PF14502** (HTH\_41)
- GENE3D**
  - G3DSA:1.10.10.10** (G3DSA:1.10.10.10)

**Interpro** does use **Pfam** models to detect its various flavours of **HTH/wHTH** domain, but it does so selectively. For example, to detect the **wHTH** domains discovered here, only two **Pfam** families were used (**HTH\_20** and **HTH\_41**, see illustration). These appear not to have matched in this instant as only a **G3DSA** entry is quoted.

All the above suggests that no one model exists to pick up all **HTH** domains? Possibly also, the fact that **HTH** domains come in such a variety of forms makes them difficult to detect reliably?

There is a simple **EMBOSS** program to detect **HTHs**. It easily detected the **Homeobox domain HTH** but essentially failed to detect the **wHTHs** recorded here. This must be because the, very simple, model (based on a **Weight Matrix** built from about **100** examples) used by the program only reliably applies to a specific range of **HTH** domains/motifs that includes the one in the **Homeobox domain** of the human **PAX6** protein?

I am very open to better explanations. I am not completely convinced by the discussion above.



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