Two independent and interactive DNA-binding subdomains of the Pax6 paired domain are regulated by alternative splicing

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Vertebrate Pax proteins share a conserved 128-amino-acid DNA-binding motif, the paired domain. The PAX6 gene, which is mutated in the murine Small eye and human aniridia developmental defects, also encodes a second protein with a 14-amino-acid insertion in the paired domain. This protein, which arises by alternative mRNA splicing, exhibits unique DNA-binding properties. Unlike other paired domains, which bind DNA predominantly by their amino termini, the extended Pax6 paired domain interacts with DNA exclusively through its carboxyl terminus. This property can be simulated by deletion of 30 amino-terminal residues from the Pax6 or Pax2 paired domains. Thus, the insertion acts as a molecular toggle to unmask the DNA-binding potential of the carboxyl terminus. The functional nonequivalence of the two Pax6 proteins is underscored by a T → C mutation at position -3 of the alternative splice acceptor site that changes the ratio of the two isoforms and causes a distinct human ocular syndrome.

Key Words: Pax genes; transcription factor; alternative mRNA splicing; DNA binding; ocular development; aniridia

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Transcription factors responsible for embryonic patterning and tissue differentiation regulate the expression of target genes by specific DNA sequence recognition. In large part, their unique functional and DNA-binding characteristics have evolved through the modular assembly of conserved domains [Kessel and Gruss 1990]. These domains define classes of transcription factors, such as the basic leucine zipper (bZIP), basic helix-loop-helix (bHLH), zinc finger, and homeo box families. Mechanisms that refine DNA-binding specificity within these families are of particular interest, as they generate target gene diversity.

The Pax (paired box) family of developmentally regulated transcription factors share an amino-terminal DNA-binding motif known as the paired domain [for review, see Chalepakis et al. 1992]. This domain consists of 128 amino acids and has been identified in a number of metazoan species, including Drosophila and humans [Noll 1993]. The nine mammalian Pax genes are divided into four groups on the basis of their amino acid sequence, intron–exon boundaries, and the presence of other conserved motifs [Walther et al. 1991; Wallin et al. 1993]. The paired domain is predicted to contain at least three α-helical regions [Bopp et al. 1989] and circular dichroism spectral data suggest that a stable conformation with α-helical characteristics is adopted upon specific DNA binding [Epstein et al. 1994]. Within the paired domain, an evolutionary discontinuity divides the relatively conserved 74-amino acid amino terminus from the more divergent 54-amino-acid carboxyl terminus and coincides with an exon boundary [Walther et al. 1991].

The PAX6 gene is mutated in Small eye mice [Hill et al. 1991] and humans with the congenital eye disease aniridia [Ton et al. 1991; Glaser et al. 1994b]. Because these phenotypes are semidominant and the mutations cause a loss of function, the Pax6 proteins appear to control rate-limiting steps in eye development. The PAX6 gene spans 22 kb, is divided into 14 exons, and encodes a 422-amino-acid protein that is identical in humans and mice. In addition to the paired domain, Pax6 contains a homeo domain and a carboxy-terminal region capable of activating transcription [Glaser et al. 1994a]. The paired domain (amino acids 3–130) is encoded by exons 5–7. An alternatively spliced exon [exon 5a] has been identified in several vertebrate PAX6 genes and is included variably.

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in the mature mRNA transcript, resulting in the insertion of a 14-amino-acid peptide in the paired domain after Gln47 [Walther and Gruss 1991; Glaser et al. 1992; Puschel et al. 1992]. Both splice products are expressed in the developing eye, brain, spinal cord, and olfactory epithelium, with the exon 5a-containing transcripts at slightly lower levels [Ton et al. 1991; Walther and Gruss 1991; T. Glaser and R.L. Maas, unpubl.].

We have identified previously a high-affinity binding site for the Pax6 paired domain using an in vitro-binding site selection (Epstein et al. 1994). A similar sequence has been identified by stepwise replacement of positions in a potential genomic DNA-binding site for Pax5 [Czerny et al. 1993]. Although Pax6 and Pax5 [Barberis et al. 1990; Czerny et al. 1993; Kozmik et al. 1993; Plaza et al. 1993] bind related DNA sequences such that up to 28 bp are protected from DNase I digestion, other paired domains exhibit different DNA-binding characteristics. The Drosophila Paired protein recognizes the e5 sequence located in the promoter region of the even-skipped gene. In this case, the paired domain footprints only 15 bp and deletion of the carboxyl terminus has no effect on e5 binding [Treisman et al. 1991]. In contrast, the Pax1 paired domain contacts modified e5 sequences along a 24-bp distance, with the amino-terminal portion responsible for sequence specificity and the carboxy-terminal portion required for high-affinity binding or stability of the protein–DNA complex [Chalepakis et al. 1991]. The Pax8 paired domain recognizes sequences within the promoter regions of two thyroid-specific genes that are not obviously related to e5 and footprints 15 bp at each site [Zannini et al. 1992]. Pax5 appears to contact DNA in a bipartite manner in which both regions of contact contribute to the overall affinity of binding [Czerny et al. 1993]. Together, these results suggest that the amino terminus of the paired domain is required for specific DNA recognition. However, the role of the carboxyl terminus is unclear, and the function of the 5a insertion in Pax6 is unknown.

In this report we describe unique properties of the alternatively spliced form of Pax6, which have allowed us to examine the DNA-binding characteristics of the paired domain in detail. We demonstrate that (1) the paired domain functions as two discrete subdomains with distinct binding specificities, (2) insertion of the 5a peptide into the amino-terminal subdomain enables the carboxy-terminal subdomain to recognize its binding site, and (3) this binding is an important determinant of transcriptional activity. Finally, we describe a human PAX6 mutation that changes the ratio of the alternatively spliced forms and causes an unusual eye phenotype, indicating that the two Pax6 paired domains function differently during embryogenesis.

Results

A splice acceptor mutation in exon 5a

To investigate the biological function of the extended Pax6 paired domain, we tested DNA samples from patients with aniridia and related ocular disorders for point mutations in exon 5a and flanking genomic DNA. Heteroduplex and single-strand conformational analysis revealed a single-base substitution in one family [Bh] with a distinct clinical phenotype (Fig. 1). The disease affects a mother and daughter with bilateral juvenile cataracts, peripheral corneal opacification, glaucoma, and pendular nystagmus, an unsteady oscillating gaze attributed to a congenital sensory deficiency [see Materials and methods]. However, in contrast to the typical manifestations of aniridia, such as iris aplasia or severe hypoplasia [Nelson et al. 1984], the irides in these two patients were relatively intact, the pupils were round, and the maculas, the central region of the retinas with the greatest density of photoreceptors, appeared normal. Abnormalities were noted in the anterior aspect of the iris derived from neural crest, including small radial defects, decreased vasculature and crypt density, and thinning of the iris stroma. The posterior pigmented iris layer derived from neural ectoderm was essentially normal.

The Bh family mutation is a T → C transition at position −3 in the exon 5a splice acceptor (Fig. 2A). The mutation appeared as a de novo change in the mother’s DNA and creates an MaeIII restriction site, which was used to follow its transmission in the pedigree. The mutation is present only in the mother and daughter, correlating completely with the ocular phenotype, and was not observed in any of >50 unrelated DNA samples.

To determine the effect of the Bh mutation, we compared mutant and wild-type genomic fragments in an in vivo splicing assay (Fig. 2B). Transcripts containing ex-
Figure 2. Mutational analysis of PAX6 in family Bh. (A) Exon 5a splice acceptor mutation. (Top) MaeIII digest of radiolabeled PCR products spanning exon 5a. These products were amplified from genomic DNA of indicated family members and pSPL1 plasmid clones containing mutant or wild-type sequences. (DC) Normal, (●●) affected. (Bottom) Genomic map of PAX6, showing the 5' end of exon 5a and the T→C mutation at position −3. Coding regions are shaded. A consensus splice acceptor site (Senapathy et al. 1990) is aligned above for comparison. The mutation creates an MaeIII restriction site, which divides the 236-bp PCR product into 109- and 127-bp fragments. The sequence shown is identical to mouse (data not shown) and quail (Dozier et al. 1993) PAX6 genomic DNA. The bracket indicates the genomic fragment used in the in vivo splicing assay. (B) The Bh mutation significantly increases the use of alternative exon 5a. (Top left) In vivo splicing assay. COS7 cells were transfected with expression plasmids containing mutant or wild-type (wt) PAX6 genomic sequences, no DNA (mock), or pSPL1 DNA alone (vector). The relative expression of the two PAX6 transcripts was assessed by RT PCR analysis. The addition or omission of RT enzyme from these reactions is indicated above the lanes. (Bottom) Schematic representation of PCR products detected in the in vivo splicing assay. (Solid bar) PAX6; (stippled bar) HITat sequences. The upstream primer is located in exon 5 and the downstream primer spans the tat-exon 6 junction. (Top right) RT PCR of PAX6 transcripts in lymphoblasts derived from two healthy individuals and a heterozygous patient with the Bh mutation. The primers are located in exons 3 and 6. Fragment sizes are indicated in bp.

were expressed in Escherichia coli as glutathione S-transferase (GST) fusion proteins, and their DNA-binding characteristics were examined in an electrophoretic mobility shift assay (EMSA). The extended paired domain [Pd-5a] does not bind P6CON (Fig. 3A), which is the consensus sequence identified previously using the shorter P6x5a paired domain, Pd (Epstein et al. 1994). Pd-5a also does not bind other paired domain recognition sites, including e5 and modified e5 (PRS4) sequences [data not shown], even under low stringency conditions; therefore, we performed a DNA-binding site selection with Pd-5a protein under the same conditions used to select P6CON. Similar results were obtained with oligonucleotide pools containing a core of 25 or 35 random positions (see Materials and methods). After nine rounds of selection using oligonucleotides containing 35 random positions, 75% (18/24) of the selected oligonucleotides contained a 22-bp consensus sequence (Fig. 3A) with no more than two mismatches. This sequence [denoted 5aCON] binds more tightly to Pd-5a by EMSA than any other sequence tested and was therefore chosen as an appropriate binding site for molecular analysis of the extended paired domain.

Identification of a high-affinity DNA-binding sequence for Pd-5a

The two Pax6 paired domains (denoted Pd and Pd-5a)
5aCON contains two copies of an imperfect 11-bp direct repeat and has no significant homology to P6CON. It forms several complexes when incubated with Pd-5a protein (Fig. 3A), indicating that one, two, or more protein molecules can bind the DNA probe. Bands representing higher order complexes predominate even under conditions where DNA is in excess. The Pd protein also binds the 5aCON probe, although the pattern of shifted bands is different (Fig. 3A), and there is no such preference for higher order complexes, although a low intensity second band is sometimes observed at high protein concentrations (see Fig. 6B, below). The formation of DNA–protein complexes is not affected by the GST moiety, as Pd and Pd-5a proteins containing an amino-terminal polyhistidine tail instead of GST give identical patterns of shifted complexes in EMSA experiments (data not shown).

To compare the protein–DNA ratios directly within the complexes formed upon Pd-5a binding to 5aCON, we performed double-label experiments in which Pd-5a was expressed as a GST fusion protein with a single cAMP-dependent protein kinase phosphorylation site. This protein, in EMSA with 5aCON, gave an identical pattern of shifted complexes to the fusion protein lacking the phosphorylation site (data not shown). After phosphorylation with [γ-32P]ATP, the protein was incubated with 5aCON end-labeled with [γ-32P]ATP. The relative protein to DNA content of shifted complex C2 compared with C1 was $1.95 \pm 0.14$ (mean ± s.e.), suggesting that C2 is a dimer of Pd-5a binding to 5aCON. Previous studies of Pd binding to P6CON (Epstein et al. 1994) suggest that C1 likely represents monomeric binding. For technical reasons (see Materials and methods) the protein/DNA ratio for the most slowly migrating Pd-5a/5aCON complex seen in Figure 3A could not be determined.

An equilibrium dissociation constant ($K_d$) was estimated for binding of both paired domain forms to the 5aCON probe by performing EMSA reactions with a
fixed, limiting amount of DNA in the absence of non-specific DNA competitor [data not shown]. We obtained $K_d$ values of $5.5 \times 10^{-9}$ M for Pd and $3 \times 10^{-9}$ M for Pd-5a. These values are comparable to the $K_d$ observed for binding of Pd to P6CON ($2.5 \times 10^{-9}$ M) [Epstein et al. 1994], and represent an upper limit, as they were calculated assuming bimolecular, monomeric binding reactions with 5aCON [Fried 1989].

**Pax6 activates transcription upon binding 5aCON**

To assess the functional significance of the 5aCON consensus, we tested the ability of full-length Pax6 and Pax6-5a proteins to activate transcription from this sequence in vivo, in comparison with murine Pax2, which has been shown to activate transcription in murine P19 embryonic carcinoma cells [Fickenscher et al. 1993]. We cotransfected P19 cells with Pax6, Pax6-5a, or Pax2 expression plasmids and a reporter plasmid in which the chloramphenicol acetyltransferase (CAT) gene was placed downstream of a minimal Elb promoter and two to six copies of the P6CON- or 5aCON-binding sites. We also tested a reporter containing six 5aCON half-sites, each separated by 6–14 bp [see Materials and methods]. The Pax6-5a protein efficiently stimulates CAT expression when 5aCON is present, whereas Pax6 is effective when either P6CON or 5aCON is present [Fig. 3B]. Transcriptional activation correlates with the in vitro ability of the Pax6 paired domain, but not Pd-5a, to bind P6CON [Fig. 3A]. Pax2 also stimulates CAT expression from P6CON, but to a lesser extent than Pax6, in accord with its lower affinity for this site [Epstein et al. 1994]. Thus, this activation is independent of the presence of a homeo domain, as Pax2 lacks a homeo domain. In accord with DNA-binding data, Pax6-5a can activate transcription from separated 5aCON half-sites [5aCON[1/2] in Fig. 3B] whereas Pax6 and Pax2 cannot.

**Pd-5a binds to adjacent half-sites**

The tendency of Pd-5a to form higher order complexes upon binding 5aCON suggests that individual protein molecules bind to each of the two predicted half-sites of the direct repeat composing 5aCON. Pd-5a binds a probe containing one copy of this 11-bp half-site [5aCON[1/2]], yielding a single band in EMSA experiments [Fig. 3A]. This probe is not bound by the Pd protein, indicating that the two forms of the Pax6 paired domain bind 5aCON by different mechanisms.

Methylation interference assays were performed to identify specific nucleotides contacted by Pd and Pd-5a proteins in each complex. The residues contacted by Pd-5a in the faster migrating complex [C1] cluster predominantly around the right-hand half-site, whereas residues contacted in the more slowly migrating complex [C2] span the entire length of the direct repeat [Fig. 4E]. The two half-sites are not identical, therefore, differences in sequence may account for the preference for the right-hand half-site. The Pd-5a protein, therefore, binds 5aCON by recognizing one of the two half-sites preferentially and subsequently occupying the second half-site. In contrast, the interference pattern for Pd reveals contacts along the entire length of the direct repeat in 5aCON, although the migration of the Pd/5aCON complex in the EMSA suggests that only one protein molecule is bound.

DNase I footprinting experiments are also consistent with Pd-5a binding to 5aCON by consecutively filling individual half-sites, whereas Pd recognizes 5aCON as one long binding site [Fig. 4A,E]. The Pd-5a domain protects ~33 bp of 5aCON from DNase I digestion but only 18 bp of the 5aCON[1/2] half-site probe [Fig. 4C]. When Pd binds 5aCON, ~28 bp is protected, as was shown previously for Pd binding to P6CON [Epstein et al. 1994]. Pd-5a consistently protected a slightly longer sequence from DNase I digestion, as compared with Pd, when incubated with 5aCON [Fig. 4A].

We tested the importance of the relative position of the two half-sites with respect to each other by synthesizing DNA probes with two [5aCON+2] or four [5aCON+4] A-T base pairs inserted between the half-sites of 5aCON. Pd-5a binding and the formation of higher order complexes is reduced as the distance between half-sites is increased [Fig. 5]. The effect of the separation on higher order complex formation can best be appreciated by comparing lanes 2, 7 and 12, in which Pd-5a protein forms approximately equal amounts of C2 complex with each of the three probes, respectively. As the half-sites are separated, the relative amount of probe in the C1 complex, and the ratio of C1 to C2, increases.

The 2- and 4-nucleotide insertions not only physically separate the two half-sites from each other but change progressively their rotational relationship in the DNA helix. Pd also binds very poorly to the 5aCON+4 probe [Fig. 6C].

**Different protein subdomains determine unique DNA-binding characteristics of Pd and Pd-5a**

To define the parts of each paired domain that are required for binding DNA, we prepared a set of Pd and Pd-5a proteins with amino- or carboxy-terminal deletions and tested their ability to bind P6CON and 5aCON by EMSA [Fig. 6]. Deletion of the amino-terminal 23 amino acids from Pd abolishes binding to P6CON, whereas deletion of the carboxy-terminal 33 amino acids decreases binding to P6CON only slightly and reduces the DNase I footprint from 28 to 16 bp [Fig. 6A]. This 16-bp footprint of the amino-terminal subdomain contains the region of homology among several paired domain binding sites [Czerny et al. 1993; Epstein et al. 1994]. We conclude that the amino subdomain determines the majority of Pd–DNA-binding affinity, whereas the carboxyl subdomain makes less important contacts with adjacent nucleotides.

Binding of Pd and Pd-5a to 5aCON requires the carboxy-terminal subdomain. Deletion of the carboxy-terminal 33 amino acids from Pd or Pd-5a abolishes 5aCON binding [Fig. 6B]. In contrast, amino-terminal deletions of Pd confer the ability to bind 5aCON, form higher or-
Alternative binding by paired box subdomains

Figure 4. The two Pax6 paired domains recognize 5aCON by different mechanisms. (A) DNase I footprint analysis. Approximately 28 bp of the 5aCON sequence is protected by Pd protein, and a 33-bp region is protected by Pd-5a protein. The reverse strand is shown, labeled at its 3' end. The visible band at the upper extent of the footprint of 5aCON with Pd-5a was reproducibly less intense than the corresponding band in control or Pd lanes, whereas other subtle differences between the footprints were not reproducible. (B) Methylation interference analysis. 5aCON probes were labeled at their 5' ends and modified with dimethylsulfate (DMS) or hydrazine (Hyd). (C) DNA–protein complex; (F) free probe; (T) total probe. The rapidly (C1) and slowly (C2) migrating complexes formed with Pd-5a were analyzed separately. The indicated residues correspond to numbered positions in E. In complex C1, contacts are clustered in the righthand half-site of 5aCON. In complex C2, contacts are formed with both half-sites. (D) DNase I footprint of Pd-5a binding to half-site probe 5aCON(1/2). An 18-bp segment is protected. (E) Methylation interference of Pd-Sa binding to 5aCON(1/2). (F) Summary of footprint and interference data. The protein and probe in each complex is indicated. The numbers refer to the 22-bp 5aCON consensus sequence. Dark lines indicate regions of the reverse strand protected from DNase I digestion. \( \Psi \) residues where methylation interferes with protein binding; \( \Delta \), bottom, an augmentation of binding when this thymidine is modified by hydrazine.

...der complexes, and recognize 5aCON + 4 [Fig. 6C], properties that are characteristic of Pd-5a. We conclude that the 5a peptide does not participate directly in half-site recognition or coordinate binding to adjacent half-sites but functions by inactivating the amino-terminal portion of the paired domain, thus exposing the latent DNA-binding potential of the carboxyl subdomain.

In accord with these observations, we found that amino-terminal deletions of the Pax2 and Pax3 paired domains also abolish binding to P6CON and enhance high order complex formation of the Pax2 paired domain with 5aCON [Fig. 6C]. The Pax3 paired domain differs significantly from Pax6 in the carboxy-terminal region [Walther et al. 1991] and does not bind 5aCON.

Discussion

Our findings provide evidence for two autonomous but interacting subdomains that reside in the amino- and carboxy-terminal portions of the paired domain. This
Alternative splicing changes the population of sequences recognized by the paired domain, potentially allowing Pax6-5a to regulate an expanded or restricted set of genes in comparison to Pax6. Although we do not know whether half-sites or tandem repeat 5aCON-like sequences are used in nature, it is clear that some DNA sequences are recognized by both forms of Pax6, whereas others are recognized specifically by one form only. The use of variously oriented half and full DNA-binding sites has been demonstrated in nature among members of the nuclear hormone receptor superfamily, which can bind DNA as monomers or dimers [Näärä et al. 1991; Umesono et al. 1991; Kato et al. 1992; Parker 1993].

An alternative splicing mutation in PAX6 causes a distinct ocular phenotype

The importance of the disparate DNA-binding properties of the alternative paired domains is underscored by our finding of a PAX6 mutation that disturbs the ratio of the two splice products and causes an ocular phenotype. The T→C mutation at position −3 of the exon 5a splice acceptor is unusual in that it increases splicing efficiency significantly. Although not absolutely essential [Fu et al. 1988], the presence of a pyrimidine at position −3 is thought to be important for effective interaction between the splice acceptor site of the pre-mRNA and small nuclear ribonucleoproteins [snRNPs] U1 and U5 of the spliceosome [Green 1991; Guthrie 1991]. In a survey of 3724 splice acceptor sites, cytosine was found 3.5 times more frequently than thymidine at this position (75% vs. 21%; Senapathy et al. 1990), suggesting that cytosine may be enzymatically preferred. In the model proposed by Steitz (1992), the conserved AG dinucleotide of the splice acceptor site pairs with the C9Ulo dinucleotide in the 5′ tail of U1 snRNA during assembly of the mammalian spliceosome. In this configuration, the cy-
Figure 6. Paired domain DNA binding is mediated by two distinct subdomains. (A) DNase I footprint of a Pd-selected sequence (related to P6CON, see Epstein et al. 1994). The 5' end of the forward strand was radiolabeled. The Pd protein protects a 28-bp segment. Deletion of 33 [PdΔC1] or 48 [PdΔC2] amino acids from the carboxyl terminus of Pd reduces the footprint to 16 bp. The 5' boundary of the footprint is unchanged. [B] EMSA analysis of paired domains with carboxy-terminal deletions. Both carboxy-terminal deletions prevent binding of Pd and Pd-5a to 5aCON but do not prevent binding of Pd to P6CON. Pd-5a does not bind P6CON. (C) EMSA analysis of paired domains with amino-terminal deletions. Deletion of 24 [PdΔN1] or 34 [PdΔN2] amino acids from the amino terminus of Pd prevents binding to P6CON. The Pax2 and Pax3 paired domains also bind P6CON and deletion of 30 amino acids from the amino terminus of each prevents binding. Amino-terminal deletions of Pd promote the formation of higher order complexes with 5aCON, similar to the pattern observed for Pd-5a binding to 5aCON. Likewise, Pax-2Pd binds 5aCON and Pax-2PdΔN binds 5aCON forming high order complexes, whereas Pax-3Pd and Pax-3PdΔN do not bind this sequence. Amino-terminal deletions enable Pd to bind a half-site probe 5aCON(1/2) and probe 5aCON + 4, in which the two half-sites are separated by 4 bp. [D] Summary showing Pax6 paired domain deletion constructs and DNA-binding data. Relative binding to P6CON and 5aCON is indicated at right. Amino acids are numbered starting from the initiation methionine. The paired domain extends from amino acid 3 to 130. The solid bar indicates a 16-amino-acid segment that is highly conserved among paired domains. Open bars indicate α-helical regions predicted from primary sequence data [Bopp et al. 1989]. The 5a peptide is inserted after amino acid 47.

tosine at position −3 is aligned with G11 of the U1 snRNA. If pairing of these complementary bases also occurs, it could increase splicing efficiency and explain the effect of the Bh mutation. The few human mutations reported in splice acceptors at position −3 introduce purines at this position, and all appear to decrease splicing efficiency [Krawczak et al. 1992]. A 195-bp fragment encompassing exon 5a is 97.0% identical in human and mouse genomic DNA [data not shown], and the T at position −3 is invariant among quail [Dozier et al. 1993], human, and mouse PAX6 genes. The exact sequence of the acceptor site, in particular the pyrimidine at position −3, therefore plays an important role in modulating the relative expression of the two paired domain forms.

Mammalian ocular and central nervous system development is exquisitely sensitive to PAX6 gene dosage. Mutant phenotypes range from anophthalmia with profound olfactory and brain malformations in PAX6 homozygotes [Schmahl et al. 1993; Glaser et al. 1994a] to relatively mild findings in the present case. Aniridia is caused by heterozygous null mutations, either cytogenetic deletions of 1lp13 in individuals with the WAGR syndrome [Wilms tumor, aniridia, genitourinary anomalies, mental retardation] [Ton et al. 1991] or intragenic nonsense or frameshift mutations [Glaser et al. 1994b]. In both groups of patients, the phenotype is characterized by severe hypoplasia of the iris, with anterior segment and retinal defects [Nelson et al. 1984]. All mutations described thus far are predicted to affect both Pax6 and Pax6-5a transcripts [Glaser et al. 1994b]. The Bh mu-
The 5a insertion functions as a molecular toggle

The exon 5a insertion has been conserved during vertebrate evolution, with only a single amino acid substitution distinguishing human, mouse, quail, axolotl, and zebrafish 5a peptides (Glaser et al. 1994b). This implies that the 5a insertion itself has an important functional role. As indicated by aminoterminal deletion analysis, the 5a peptide is not required for dimerization upon DNA binding or recognition of the 5aCON site. Instead, we suggest it functions as a molecular toggle, preventing the amino subdomain from binding DNA and allowing the carboxy terminus to bind.

Figure 7. A model for DNA binding by the two alternative Pax6 paired domains. The paired domain is shown consisting of amino and carboxyl subdomains. The Pd protein is folded so that the amino subdomain binds its preferred sequence, P6CON. The carboxyl subdomain makes limited contacts with P6CON but is shielded partially from or conformationally incapable of binding its preferred sequence, 5aCON(1/2). Insertion of the 5a peptide deforms the amino subdomain, preventing it from binding DNA and exposing the independent binding capability of the carboxyl subdomain. This conformational change allows the carboxyl subdomain to bind 5aCON(1/2), and promotes binding of Pd-5a molecules to adjacent half-sites in 5aCON.

Structural studies of the Drosophila Paired protein reveal the existence of a HTH motif in the amino subdomain that is not predicted from the amino acid sequence (W. Xu, M. Rould, S. Jun, C. Desplan, and C. Pabo, unpubl.). The 5a insertion site is located at the end of the first α-helix of the HTH motif. There are several prece-
gaze was unsteady and neither eye could fixate on objects for an extended period. The corneas were small, measuring 10 mm in horizontal diameter and were opacified at the superior and inferior margins by corneal pannus. The intraocular pressures were slightly elevated while receiving topical glaucoma medication. A moderately broad iris leaf was present bilaterally with a surgical coloboma above in each eye. The optic discs showed evidence of glaucomatous cupping bilaterally. There was no macular abnormality.

The daughter had poor vision through early childhood. Cataracts were first reported at age 12, consisting of cortical equatorial opacities and diffuse posterior pole opacities that progressed during 3 years of observation. At age 13, the visual acuity was 20/80 (right) and 20/400 (left). After bilateral cataract extractions at age 16, unsteady pendular fixation persisted, suggesting a congenital sensory nystagmus. The corneas measured 9.5 mm in horizontal diameter. Pannus defects extended 2 mm inward from the superior and inferior corneal margins and stained well with fluorescein. The pupils were round and dilated normally, but were displaced nasally. The irides were blue stained well with fluorescein. The pupils were round and di-

PAX6 mutational analysis

Genomic DNA was extracted from venous blood samples obtained with informed consent. Fragments spanning PAX6 exons were amplified by the polymerase chain reaction (PCR) using previously described primers and conditions (Glarner et al. 1992) and screened for mutations by heteroduplex (White et al. 1992) and single-strand conformational (Orita et al. 1989) analysis. The Bb mutation was detected by altered migration of exon 5a heteroduplex molecules. Mutant and wild-type PCR products were cloned into pGEM3Z and sequenced. To confirm the exon 5a mutation, radiolabeled PCR products were digested with Msel. The presumed genetic relationships within the family were confirmed using seven highly polymorphic DNA markers on chromosome 11.

Reverse transcriptase PCR analysis

RNA was extracted from Epstein-Barr virus-transformed lymphoblast cell lines by the phenol–guanidinium thiocyanate method (Chomczynski and Sacchi 1987). Twenty microliters reverse transcriptase [RT] reactions were performed at 37°C as described (Kawasaki 1990) using 2 μg RNA as template and 0.2 μg pd[N6] as primer. Fragments spanning the exon 5a insertion site were amplified from the resulting cDNA by PCR, using upstream primer 5’-GGAGAGTTATGAGTCAGGTT-3’ (exon 3) and downstream primer 5’TTCGGTATGATGTTATCG-3’ (exon 6) as follows: 1 min at 95°C, 2 min at 60°C, and 2 min at 72°C, for 45 cycles. The expected products are 416 and 458 bp long and extend between nucleotides 292 and 707 of PAX6 clone AHPX-2 (Glarner et al. 1992).

In vivo splicing assay

A 1.3-kb genomic fragment encompassing PAX6 exons 5, 5a, and 6 was amplified by PCR from a FIX-3 (Glarner et al. 1992) and cloned in the sense orientation into the BamHI site of exon amplification vector pSPL1 (Buckler et al. 1991). This fragment begins 64 bp upstream of exon 5 and ends 54 bp downstream of exon 6. The cloning site is located within a 2.3-kb intron derived from the human immunodeficiency virus (HIV-1) tat gene. The intron forms part of a transcription unit that begins with the SV40 ori promoter and is flanked by tat splice donor and acceptor sites. An isoformic derivative containing the Bh point mutation was created by site-directed mutagenesis (Deng and Nickoloff 1992). The DNA sequence surrounding exon 5a was confirmed in both plasmid clones. COS7 cells were plated in 60-mm Petri dishes and transfected with 1.5 μg of plasmid DNA coated with a cationic lipopolyamine (transfectam, Promega, Madison, WI). Total cellular RNA was extracted after 72 hr and PAX6–tat chimeric transcripts were amplified by RT PCR using an upstream primer in exon 5 (5’-GTACACGCC-

DNA-binding site selection

A modified binding-site selection (Wilson et al. 1993) was performed using a degenerate oligonucleotide with a core of 25 random positions (see Epstein et al. 1994). However, because the region of Pd-5a binding defined by DNase I protection analysis extended into the flanking sequences, a longer oligonucleotide (93 bp) with a core of 25 random positions was used for all further experiments. Approximately 100 ng of purified GST–Pd-5a protein was attached to glutathione–agarose beads and incubated with 5 μg of oligonucleotide in a buffer containing 20 mm Hepes (pH 7.4), 150 mm KCl, 250 μg/ml of poly[d(I-C)], 20% glycerol, 0.5 mg/ml of bovine serum albumin (BSA), and 1 μM DTT. Samples were incubated at room temperature for 45 min with gentle agitation and washed extensively with cold binding buffer. Pd-5a/DNA complexes were released from the GST moiety by thrombin cleavage and used for subsequent PCR amplification. After nine rounds of selection, bound oligonucleotides were cloned into pGEM3Z and sequenced. The results of the selection performed with 35 random positions are presented.
The consensus obtained from the experiment using 25 random positions [24 selected sequences analyzed] contained the same 22-bp imperfect direct repeat except that position 4 (as numbered in Fig. 4E) was T instead of C, and position 14 was A instead of G.

**EMSA**

EMSA reactions were performed essentially as described [Epstein et al. 1994], in a buffer consisting of 10 mm Tris (pH 7.4), 50 mm KCl, 100 µg/ml of poly[d(I-C)], 20% glycerol, 2.5 mg/ml of BSA, 1 mm β-mercaptoethanol, and 1 µm protein, unless otherwise noted. For measurement of Kₐp, the poly[d(I-C)] competitor was omitted. Double-stranded probes were made by annealing an oligonucleotide template with a short complementary primer and filling in the remaining sequence using Klenow DNA polymerase. Probes were radiolabeled by kinasing the oligonucleotide with [γ-32P]ATP or including [α-32P]dCTP in the polymerase reaction, and were purified on polyacrylamide gels. Binding reactions were performed at 4°C for 30 min followed by electrophoresis through 6% polyacrylamide gels in 0.5x TBE. DNA-protein complexes and free probes were excised separately, electroeluted in 0.2x TBE, and precipitated with ethanol, resuspended in water, and analyzed by autoradiography with a PhosphorImager (Molecular Dynamics). The probes were P6CON, 5'-TGGAAATCCGAGAAAAATTTTACGCCTGTAGCTACGCTGAGAATGGTACGCTC-3'; 5aCON, 5'-AATAAATTTTACCCGTTACGAGTGCTACGCTGAGAATGGTACGCTC-3'; 5aCON[1/2], 5'-GATCCAATTTTACCGTTACGAGTGCTACGCTGAGAATGGTACGCTC-3'.

**Double-label binding experiments**

To calculate the relative ratio of protein to DNA in the complexes formed upon Pd-Sa binding to 5aCON, we cloned the oligonucleotide with [γ-32P]ATP or including [α-32P]dCTP in the polymerase reaction, and were purified on polyacrylamide gels. Binding reactions were performed at 4°C for 30 min followed by electrophoresis through 6% polyacrylamide gels in 0.5x TBE. DNA-protein complexes and free probes were excised separately, electroeluted in 0.2x TBE, and precipitated with ethanol, resuspended in water, and analyzed by autoradiography with a PhosphorImager. The consensus obtained from the experiment using 25 random positions [24 selected sequences analyzed] contained the same 22-bp imperfect direct repeat except that position 4 (as numbered in Fig. 4E) was T instead of C, and position 14 was A instead of G.

**CAT assays**

P19 murine embryo carcinoma cells [5x10⁴] were plated in 60-mm petri dishes and transfected with plasmid DNA coated with cationic lipopolymine (transfectam). The pGECAT reporter plasmid contains the adenovirus E1bTATA promoter upstream of the bacterial CAT gene [Lillie and Green 1989]. Derivative plasmids with six copies of the P6CON sequence, two copies of a 5aCON sequence, or six copies of the 5aCON[1/2] sequence upstream of the promoter were prepared by inserting two copies of PRDCONx3 (5'-TCGACAAAAATTTTACCGTTAGTTCACGCTGAGAATGGTACGCTC-3'), two copies of 5aCON-ava (5'-CTAGTGAACATGGCTAGTTCACGCTGAGAATGGTACGCTC-3'), or three copies of 5aCON[1/2]-6A (5'-CTAGTGAACATGGCTAGTTCACGCTGAGAATGGTACGCTC-3') into the SalI site of pGECAT after the addition of SalI linkers. Expression plasmids were derived by replacing the NorI-bgal cassette of pCMV8 [MacGregor et al. 1987] with full-length coding sequences for Pax6, Pax6-5a, or Pax2, and expression of Pax proteins was confirmed by Western analysis. The 1.6-kb human Pax6 DNA fragment was amplified by PCR from pHXP-2 [Glarer et al. 1992] using an upstream primer in the 5'-untranslated region (5'-AAACCATTGCAGCTGGCCCAAGC-ACCAACAGCCGACGCA-3') and a downstream primer in flanking vector DNA (5'-AATTACGGCGCGCCATACGGC-TCATATAAGGGCC-3'). Mouse Pax2 sequences were recombined as a 2.6-kb NorI fragment from two partial cDNA clones [Dressler et al. 1990]. The Pax6-5a plasmid was derived by replacing the 371-bp XhoI fragment of pHXP-2 with the corresponding 359-bp XhoI fragment containing exon 5a. Each dish was transfected with 2 µg of reporter and 1 µg of expression plasmid DNA. Cell extracts were prepared after 48 hr and assayed for CAT activity as described by Gorman et al. [1982]. Samples were diluted as necessary to measure activity in the linear range and equalize the amount of protein added. The percent acetylation was quantitated from thin-layer chromatography plates using a PhosphorImager.

**DNase I footprinting and methylation interference**

Footprinting and interference experiments were performed as described [Epstein et al. 1994]. For DNase I footprint analysis, double-stranded oligonucleotides P6CON, 5aCON, and 5aCON[1/2] were cloned in the polylinker of plasmid pGEM3Z or pGEM4Z. Restriction fragments containing these sequences were end-labeled by kinasing the oligonucleotide with [γ-32P]ATP or including [α-32P]dCTP in the polymerase reaction, gel purified, and mixed with 1 µM GST-Pd or GST-Pd-5a protein in 20 µl of EMUSA buffer including 5 µg/ml of poly[d(I-C)], 10 mm CaCl₂, and 2 mm MgCl₂. After DNase I digestion, samples were precipitated with ethanol and run on denaturing 6% or 10% polyacrylamide gels. For interference assays, complementary oligonucleotides were end-labeled by kinasing with [γ-32P]ATP, annealed, and extended with Klenow DNA polymerase. These probes were gel purified, modified with dimethylsulfate or hydrazine, precipitated with ethanol, and resuspended in 20 µl of EMUSA buffer with 0.1 µM GST-Pd or GST-Pd-5a. After 30 min at 4°C, samples were electrophoresed through a denaturing 6% polyacrylamide gel in 0.5x TBE and exposed to X-ray film. DNA-protein complexes and free probes were excised separately, electroeluted in 0.2X TBE, and precipitated with ethanol. Samples were then resuspended in 1 µl pipidine, cleaved for 30 min at 95°C, lyophilized three times, and run on a denaturing 10% polyacrylamide gel.

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