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Inhibition of the DNA-binding activity of *Drosophila* Suppressor of Hairless and of its human homolog, KBF2/RBP-Jκ, by direct protein–protein interaction with *Drosophila* Hairless

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We have purified the sequence-specific DNA-binding protein KBF2 and cloned the corresponding cDNA, which is derived from the previously described RBP-Jκ gene, the human homolog of the *Drosophila* Suppressor of Hairless [Su(H)] gene. Deletion studies of the RBP-Jκ and Su(H) proteins allowed us to define a DNA-binding domain conserved during evolution. Because Su(H) mutant alleles exhibit dose-sensitive interactions with Hairless [H] loss-of-function mutations, we have investigated whether the RBP-Jκ or Su(H) proteins directly interact with the H protein in vitro. We show here that H can inhibit the DNA binding of both Su(H) and RBP-Jκ through direct protein–protein interactions. Consistent with this in vitro inhibitory effect, transcriptional activation driven by Su(H) in transfected *Drosophila* S2 cells is inhibited by H. These results support a model in which H acts, at least in part, as a negative regulator of Su(H) activity. This model offers a molecular view to the antagonistic activities encoded by the H and Su(H) genes for the control of sensory organ cell fates in *Drosophila*. We further propose that a similar mechanism might occur in mammals.

[Key Words: RBP-Jκ; KBF2; Su(H); H; protein–protein interaction; transcriptional interaction]

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The human protein KBF2 specifically binds as a monomer to various κB sites and is present in the nuclei of all cells analyzed so far [Israel et al. 1989]. We have purified this factor and isolated the corresponding cDNA. Data base searching revealed a perfect homology with proteins belonging to the RBP-Jκ family. RBP-Jκ was originally purified based on its specific binding to the recombination signal of the κκ immunoglobulin gene [Hamaguchi et al. 1989; Matsunami et al. 1989]. However, recent data strongly suggest that RBP-Jκ acts as a transcriptional regulator [Dou et al. 1994] by binding to specific DNA sites, whose consensus sequences have been determined [Tun et al. 1994], rather than as a recombinase in V(D)J recombination [Hamaguchi et al. 1992].

A better understanding of RBP-Jκ function may benefit from the genetic analysis of the *Drosophila* RBP-Jκ homolog Suppressor of Hairless [Su(H)]. The murine RBP-Jκ and *Drosophila* Su(H) proteins share 82% identity and 94% similarity over most of their length [Furukawa et al. 1991; Schweisguth and Posakony 1992]. Su(H) mutations were first isolated as dominant suppressors of the Hairless [H] dominant, haploinsufficient loss-of-function mutations [Ashburner 1982]. Detailed phenotypical studies have shown that H and Su(H) act in an opposite manner to affect various developmental decisions [Bang et al. 1991; Bang and Posakony 1992; Schweisguth and Posakony 1992, 1994; Posakony 1994]. In particular, loss-of-function mutations of Su(H) and H have opposite effects on two successive alternative cell fate decisions during the formation of the adult sense organs. H and Su(H) activities are first required in the larva for the determination of single sensory organ precursor (SOP) cells from groups of equipotent cells called proneural...
clusters (for review, see Ghysen and Dambly-Chaudière 1989). $H$ and $Su(H)$ activities are also required later in the development of the pupa to determine the alternative cell fates adopted by two accessory cells of the mechanosensory organs (Bang et al. 1991; Schweisguth and Posakony 1994). Each of these two cell fate decisions also requires the activity of the neurogenic genes, including Notch [for reviews, see Simpson 1990; Posakony 1994]. The role of the $H$ and $Su(H)$ gene products in these two Notch-mediated cell fate decisions remains to be elucidated.

$H$ encodes a novel basic 109-kD protein (Bang and Posakony 1992, Maier et al. 1992). The biochemical activity and subcellular localization of the $H$ protein are so far unknown. The lack of a clear epistatic relationship between $H$ and $Su(H)$ for SOP determination has suggested that $H$ and $Su(H)$ act antagonistically in the same genetic operation (Schweisguth and Posakony 1994), though the molecular mechanism underlying the genetic suppression of the $H$ phenotypes by $Su(H)$ mutant alleles has not yet been investigated.

We report here that $Su(H)$ may act as a transcriptional activator in a transient transfection assay and that $H$ down-regulates transcriptional activation by $Su(H)$ in this assay. Negative regulation by $H$ appears to result from the formation of $H$–$Su(H)$ complexes that can prevent the in vitro binding of $Su(H)$ to the DNA. Finally, we show that the DNA-binding activity of KBF2/RBP-JK can likewise be regulated negatively by Drosophila $H$ and that the proteins interact in vitro, as well as in vivo using the yeast two-hybrid system.

Results

Cloning and DNA-binding properties of different splicing products of the human RBP-Jk gene and Drosophila $Su(H)$

KBF2 protein was purified from HeLa cells on the basis of binding to NF-κB sites, microsequence, and cloned [see Materials and methods for details]. Sequencing analysis of the obtained cDNA fragment indicated that it encodes part of the conserved region of RBP-Jk proteins (Amakawa et al. 1993).

The human RBP-Jk gene appears to be composed of 14 exons [named N1, 1, 1', N2, and 2–11]. Four classes of cDNAs, that differ in their first exon but share exons 2–11, have been described (Amakawa et al. 1993; Dou et al. 1994; see Fig. 1). KBF2 could be encoded by any one of the four alternatively spliced RBP products already described. In this paper the cDNAs and the proteins encoded by the aPCR1, aPCR2, and aPCR3 transcripts of the RBP-Jk gene [Amakawa et al. 1993] are called RBP1, RBP2, and RBP3, respectively (see Fig. 1).

To compare the properties of purified KBF2 to RBP1, RBP2, and RBP3, we tried to isolate their corresponding cDNAs by PCR amplification. The RBP1 and RBP3 cDNAs were obtained from a human lymphocyte cDNA library. However, we failed to isolate the RBP2 cDNA from two different libraries or by reverse transcription–polymerase chain reaction (RT–PCR) from various sources of RNAs [see Materials and methods for details]. A novel isoform, named RBP4, was isolated from the same library and was present in various RNA preparations [data not shown]. Its sequence is shown in Figure 1. It differs from RBP2 by the use of an alternative splice donor site located within exon N1 [see Amakawa et al. 1993 for exon nomenclature].

The DNA-binding activity of KBF2/RBP-Jk was examined using a new putative RBP-binding site found in the regulatory region of the mouse Hairy Enhancer of split (HES1) gene (Akazawa et al. 1992; Sasai et al. 1992). HES1 is one of the five mammalian proteins that display structural homology with a family of Drosophila bHLH proteins encoded by neurogenic genes located in the Enhancer of split [E(spl)] complex. HES1 appears to be involved in neural [Ishibashi et al. 1994] as well as in muscle differentiation [Sasai et al. 1992]. Our finding of a motif identical to the RBP consensus binding site, 85 bp upstream of the transcription initiation site of the HES1 gene, was guided by the previous description of a strong in vitro RBP-binding site in the regulatory region of the Drosophila E(spl) m8 gene (Tun et al. 1994). A double-stranded oligonucleotide corresponding to this putative KBF2/RBP-Jk-binding site [probe HES] was used in gel retardation experiments. We show here that purified
H inhibits DNA binding of Su(H) and RBP-J/KBF2

Figure 2. DNA-binding properties of the KBF2/RBP and Su(H) proteins. [A] The RBP proteins have the same DNA-binding specificity as purified KBF2. The proteins encoded by the RBPl, RBP3, and RBP4 splicing products were tested for binding to the HES probe: RBP3 (lanes 1, 5–10), RBPl (lane 2), and RBP4 (lane 3) were obtained by in vitro transcription/translation in a rabbit reticulocyte lysate. Binding of RBP3 was compared with that of purified KBF2 (lane 4). As cold competitor, an excess of 300-fold of probes 2122 (lane 6), KBF (lane 7), IgK (lane 8), p2m (lane 9), and Kβ2 (lane 10) was used. The arrows indicate the specific retarded band of each splicing product. [B] Mapping of the DNA-binding domain of RBP3. Deletion derivatives of RBP3 were obtained by in vitro transcription/translation and tested in a gel shift experiment with the HES probe: RBP3 [1–486] (lane 1), RBP3 [1–435] (lane 2), RBP3 [1–329] (lane 3), RBP3 [1–271] (lane 4), RBP3 [175–486] (lane 5), RBP3 [7–486] (lane 6), and RBP3 [98–486] (lane 7) [see Fig. 5A for a schematic representation of these constructs]. Note that the upper band in lane 2 corresponds to the full-length RBP3 translation product, because of an incomplete digestion of the template DNA. [C] Mapping of the DNA-binding domain of Su(H). Full-length and carboxy-terminal deletion derivatives of the Su(H) protein were produced in vitro in a rabbit reticulocyte lysate (lanes 1–5), whereas the amino-terminal deletion mutants were analyzed as GST fusion proteins (lanes 6–10). In all lanes, the 2122 probe was used. The relative amount of radioactively labeled Su(H) proteins was determined by Phosphorlager quantification following SDS-PAGE. A similar amount of wild-type and deleted Su(H) proteins was used in each lane: Su(H) [10–594] (lane 2), Su(H) [10–525] (lane 3), Su(H) [10–457] (lane 4), Su(H) [10–399] (lane 5). In lanes 6–10, ~250 ng of purified GST fusion proteins were assayed: GST-Su(H) [10–594] (lane 6), GST-Su(H) [10–525] (lane 7), GST-Su(H) [10–457] (lane 8), GST-Su(H) [10–399] (lane 9). As a control for specificity, a competition experiment with a 10-fold excess of cold 2122 probe is shown in lane 10.
KBF2 protein (Fig. 2A, lane 4) and in vitro-translated RBP proteins (Fig. 2A, lanes 1–3, 5–10) strongly bind to the HES probe (Fig. 2A). These results suggest that the HES1 promoter is a potential target for RBP binding. The KBF2 and RBP proteins also recognize the RBP-JK promoter is a potential target for RBP binding. The KBF2 proteins (Fig. Btou et al. lane 4) or in vitro-translated products (lanes 1-3) can be GAAagtccct) used in the purification of KBF2. Formation of complexes involving either purified KBF2 (Fig. 2A, lane 4) or in vitro-translated products (lanes 1–3) can be competed by several but not all NF-kB-binding sites, as well as by the consensus sequence probe (Fig. 2A, lanes 5–10, data not shown). These results are consistent with data described previously for purified RBP-Jk and KBF2 proteins (Israel et al. 1989; Tn et al. 1994).

We then compared the relative DNA-binding affinities of the three isoforms, RBPl, RBP3, and RBP4, for the HES site (Fig. 2A, lanes 1–3). Whereas RBPl and RBP3 bind with a similar affinity to the HES probe, the RBP4 product seems to bind this site poorly (equivalent quantities of each translation product were used in each lane). This indicates that the isoform-specific amino-terminal regions may contribute to RBP DNA-binding specificity.

The importance of the variable amino-terminal regions in DNA binding was investigated using a deletion construct that starts at the first amino acid common to all isoforms, the first K residue encoded by exon 2 (see Fig. 1) corresponding to amino acid 7 in RBP3 (see Fig. 5, below, for a schematic representation of all deletion constructs studied). As shown in Figure 2B (lane 6), RBP3 [7–486] still binds to the HES probe. We conclude that the variable amino-terminal regions of RBP are not strictly required for DNA binding. Still, the poor binding ability of RBP4 suggests that sequences encoded by the isoform-specific first exons may modulate the affinity and/or specificity of RBP binding to the DNA.

Further amino-terminal deletions within the protein sequence common to all RBP isoforms were also tested for their ability to bind the HES probe: RBP [98–486] and RBP [175–486] appear unable to bind DNA in our gel retardation assay (Fig. 2B, lanes 7 and 5, respectively). The effects of carboxy-terminal deletions, introduced in the RBP3 cDNA, were also analyzed: The RBP [1–435] construct recognized the probe (Fig. 2B, lane 2), whereas RBP [1–329] did not (Fig. 2B, lane 3). Thus, the region extending from exon 2 [amino acid 7 in RBP3] to amino acid 435 appears sufficient for RBP binding to the HES probe.

The Drosophila Su[H] protein was subjected to a similar deletion analysis. The nearly full-length in vitro-translated product (Su[H] [10–594]) binds specifically to the RBP consensus-binding site (probe 2122) and more weakly to the E(spl) binding site identified by Tun et al. (1994) (Fig. 2C, lane 2; data not shown). Whereas the truncated Su[H] [10–525] and Su[H] [10–457] proteins were still able to bind DNA (Fig. 2C, lanes 3,4), Su[H] [10–399] (Fig. 2C, lane 5) did not. Amino-terminal deletions were analyzed as glutathione S-transferase [GST] fusion proteins. A fusion protein between GST and Su[H] starting at the amino acid position 110 (corresponding to amino acid 22 in human RBP3) gave a specific gel-retarded complex (Fig. 2C, lane 6), whereas further deletions starting at amino acid 139 (corresponding to amino acid 50 in RBP3) abolished GST–Su[H] DNA binding (Fig. 2C, lanes 7–9). Thus, a region extending from amino acid positions 110–457 (22–369 in RBP3) appears necessary for Su[H] binding to the 2122 probe. These results are summarized in Figure 5A, below.

Thus, part of the Su[H]/RBP-Jk-conserved region, defined as the 425-amino-acid region, which is 82% identical between the Drosophila and human proteins (Schweisguth and Posakony 1992; see Fig. 5A, below), does not appear to be required for DNA binding. One may speculate that this carboxy-terminal region [amino acids 458–528 and 369–439 in Su[H] and RBP3, respectively], which has been strongly conserved during evolution, provides an interaction surface for molecules other than DNA.

Direct protein–protein interaction between Drosophila Su[H] or human RBP3 and Drosophila H

The H and Su[H] activities have been proposed to act antagonistically in the same genetic operation (Schweisguth and Posakony 1994). It is thus possible that H and Su[H] control common downstream functions in an opposite manner. Alternatively, the H and Su[H] proteins may interact directly, resulting in the inhibition of either H or Su[H] activity. We therefore investigated whether the H and Su[H] proteins may interact in vitro. First, an Escherichia coli GST–Su[H] fusion protein, which contains the conserved region [GST–Su[H] [110–594]], was tested for its ability to interact with the full-length H protein synthesized in vitro in a reticulocyte lysate (Fig. 3A, lanes 1–3). As shown in Figure 3A, the H protein was efficiently retained on GST–Su[H] [110–594] (lane 2) but not on control GST beads (lane 3). A number of other Drosophila nuclear proteins were tested as controls and were not retained onto the GST–Su[H] [110–594] beads in this assay [AEF-1 (Falb and Mannatis 1992), Daughterless (Caudy et al. 1988), Groucho (Tata and Hartley 1993), Neuralized (Price et al. 1993) [data not shown]. Thus, the H and Su[H] proteins appear to be able to specifically interact in vitro under these conditions.

We then tested various H-deletion constructs for their ability to interact in vitro with GST–Su[H] [110–594] (Figs. 3A, and 5B for a summary of the results). The first 293 amino acids appear sufficient to efficiently bind the GST–Su[H] [110–594] protein (Fig. 3A, lanes 4–6). In contrast, the H [1–236] in vitro translated product did not interact with GST–Su[H] [110–594] (Fig. 3A, lanes 7–9). This indicates that the H protein contains an interaction surface from amino acid 236 to 293, which is required for interaction with Su[H]. Furthermore, this region is not sufficient for the interaction because the H [1–84/236–
presence of \(^{35}\text{S}\)Met and tested for their interaction with H, using the GST–H [1–292] fusion protein immobilized on glutathione–agarose beads [see also Fig. 5A]. GST is used here as a negative control. Input lanes [1,4,7,10,13] show the different Su[H] in vitro-translated products, prior to incubation with the beads. The corresponding H constructs are indicated above the lanes (see also Fig. 5B for a schematic representation of these constructs). The amount of H proteins bound to the GST–Su[H] [110–594] beads, shown in lanes 2, 5, 8, 11, and 14, is compared with the control lanes 3, 6, 9, 12, and 15, which correspond with nonspecific binding of the same H samples to GST beads. Note that identical amounts of in vitro-translated proteins are shown in the input lanes or incubated with the beads. Between 10% and 25% of the H [1–1076] or H [1–293] fraction is reproducibly retained onto the GST–Su[H] [110–594], compared with the 0.5–1% background level on GST beads [lanes 1–6]. Weaker binding [5–10% of retention] is observed with the H [212–293] construct [lanes 13–15], whereas no specific binding is detectable with the H [1–236] and [1–84/236–293] constructs [lanes 7–12]. (B) Deletion mapping of the HID of Su[H]. (Lanes 1–15) Carboxy-terminal truncation derivatives of Su[H] were in vitro-translated in the presence of \(^{35}\text{S}\)Met and tested for their interaction with H, using the GST–H [1–292] fusion protein immobilized on glutathione–agarose beads [see also Fig. 5A]. GST is used here as a negative control. Input lanes [1,4,7,10,13] show the different Su[H] in vitro-translated products, as indicated above the lanes. The fraction of each Su[H] protein retained onto GST–H [1–292] beads is shown in lanes 2, 5, 8, 11, and 14, and nonspecific binding of Su[H] proteins to GST beads is indicated in lanes 3, 6, 9, 12, and 15. Between 50% and 70% of Su[H] [10–594] or [10–525] is reproducibly retained onto GST–Su[H] [110–594], compared with the 0.5–2% background levels observed with the GST beads alone [lanes 3, 6]. No specific binding is detectable for the Su[H] [10–457], [10–399], and [10–287] constructs [lanes 7–15]. Molecular mass markers are indicated at the left of the 10% polyacrylamide gel. (Lanes 16–22) \(^{35}\text{S}\)Met-labeled H [1–1076] was tested for its ability to interact with GST–Su[H] fusion proteins corresponding to progressive amino-terminal deletions [see also D]. For each GST–Su[H] fusion protein, the bound fraction [lanes 17–21] was compared with the input [lane 16] and with the background interaction with GST alone [lane 22]. Molecular mass markers are indicated at the left of the 8% polyacrylamide gel. (C) Deletion mapping of the HID of RBP3. \(^{35}\text{S}\)Met-labeled in vitro-translated derivatives of RBP3 [as indicated above the lanes] were assayed for their ability to associate with GST–H [1–292] bound to glutathione–agarose beads. Controls were GST alone and GST–p50, which is a fusion protein between GST and the p50 subunit of NF-κB, as indicated. Molecular mass markers are indicated at the left of the 10% or 12% polyacrylamide gels. Asterisks [*] mark the positions of the in vitro-translated products.

293] protein was not retained on GST–Su[H] [110–594] beads [Fig. 3A, lanes 10–12]. Weak Su[H] binding could be detected using the short H [212–293] in vitro-translated product [Fig. 3A, lanes 13–15], suggesting that this region of the H protein contributes in part to the interaction with Su[H].

To map the H interaction domain within Drosophila Su[H], we used a bacterially produced GST–H [1–292] fusion protein that contains the domain of interaction with Su[H] defined above. The nearly full-length in vitro-translated Su[H] [10–594] protein was specifically and efficiently retained by the GST–H [1–292] fusion protein, compared with the control GST beads [Fig. 3B, lanes 1–3]. The analysis of carboxy-terminal deletion constructs of Su[H] revealed that [1] Su[H] [10–525] still interacts with GST–H [1–292] [Fig. 3B, lanes 4–6], indicat-
RBP3–H protein interaction was then tested in vivo using the yeast two-hybrid system (Fields and Song 1989; Durfee et al. 1993; Harper et al. 1993). Two series of expression plasmids were constructed: One encodes segments of the RBP3 protein fused to the GAL4 DNA-binding domain [GAL[DB] plasmids in Table 1], the other encodes the GAL4 activation domain, either alone [plasmid GAL[AD]] or fused to the first 292 amino acids of H [plasmid GAL[AD]–H [1–292]]. Each pair of plasmids was cotransformed into a yeast strain carrying a GAL4 upstream activating sequence (UAS)–lacZ reporter construct integrated into its genome (Harper et al. 1993). RBP3–H interactions would be expected to mediate the formation of a complex between the GAL[DB]–RBP3 and GAL[AD]–H [1–292] proteins, resulting in the restoration of transcriptional activity. This would be detected as a blue yeast colony because of induced β-galactosidase activity. First, the GAL[DB]–RBP3 fusion constructs were not sufficient to induce detectable transcriptional activation of the UAS–lacZ gene when cotransformed with GAL[AD], with the exception of the GAL[DB]–RBP3 [330–438] fusion protein that gave light blue colonies (Table 1). Similarly, the H [1–292] domain fused to the GAL4 transcription activation domain did not induce lacZ reporter gene transcription when cotransformed with the GAL[DB]. In contrast, easily detectable lacZ expression was observed when the GAL[DB]–RBP3 [1–486], GAL[DB]–RBP3 [175–486], and GAL[DB]–RBP3 [330–438] constructs were cotransformed with the GAL[AD]–H [1–292] plasmid. As a negative control, lacZ expression was not restored when using GAL[DB]–RBP3 [1–271], in which the HID is deleted. Thus, activation of the lacZ reporter gene transcription most likely results from specific protein–protein interaction.

Table 1.  RBP3 and H interact in vivo in yeast

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Two series of plasmids were constructed, one encoding the GAL[DB] fused to various fragments of RBP3 (amino acid numbering refers to RBP3), and the other encoding a hybrid of the GAL4 activation domain GAL[AD] fused to the first 292 amino acids of H [GAL[AD]–H [1–292]]. The plasmids were introduced into a yeast strain containing the lacZ reporter gene under the control of GAL4 elements. β-Gal activity was monitored semi-quantitatively by the colony lift assay. [+] Colonies turned blue in <15 min; [+] in 15–60 min, [–] colonies remained white.

In vivo interaction between RBP3 and H in yeast

Table 1 shows that the carboxy-terminal sequences that have not been conserved throughout evolution are dispensable for interaction with H, and further deletions of conserved sequences abolish the interaction with GST–H [1–292] (constructs Su[H] [10–457], Su[H] [10–399], and Su[H] [10–287]; Fig. 3B, lanes 7–15; Fig. 5A). Amino-terminal deletion constructs were analyzed using a different experimental strategy: GST–Su[H] fusion proteins carrying progressive amino-terminal deletions were tested for their ability to retain the H [1–1076] in vitro translated product. Negative controls include H binding to GST alone (Fig. 3B, lane 22) and binding of the Drosophila AEF1 nuclear protein on these GST–Su[H] beads (not shown). Deleting the amino-terminal half of the conserved region appears to have no effect on H binding in vitro (see the GST–Su[H] [288–594] construct in Fig. 3B, lane 20). The GST–Su[H] [444–594] deletion construct, which leaves only 84 amino acids from the conserved region upstream of the evolutionary divergent carboxy-terminal tail shown above to be dispensable for H binding, still exhibits a weak H-binding activity in vitro (see Fig. 3A, lane 21). Similar results were obtained using the H [1–293] in vitro-translated product (not shown). We conclude that the H interaction domain (HID) is included within the conserved region of Su[H], between amino acids 288 and 525 (see Fig. 5A, below).

The genetic interactions observed between H and Su[H] are thought to unravel a fundamental aspect of Su[H] function. Because of the very strong conservation of Su[H]/RBP primary structure during evolution, we believe that both proteins share similar biochemical activities. We therefore investigated the ability of the RBP3 isoform to interact with Drosophila H. In vitro-translated H [1–1076] and H [1–293] interacted with GST–RBP [1–486] (data not shown), and, conversely, the RBP3 in vitro-translated protein [RBP3 [1–486]] was retained onto the GST–H [1–292] fusion protein bound to glutathione–agarose beads (Fig. 3C, lane 3). This interaction appears specific because no significant signal was detected upon binding onto GST or GST–P50, a fusion protein that consists in the p50 subunit of NF-kB fused to GST (Fig. 3C, lanes 1,2). A deletion analysis of RBP3 was carried out to independently define the HID. The RBP3 [1–435] and RBP3 [175–486] mutant proteins were still specifically retained onto GST–H [1–292] beads (Fig. 3C, lanes 7–9,13–15). This indicates that sequences upstream of amino acid 175 and downstream of amino acid 435 are dispensable for interaction with H in this assay. However, the RBP3 [1–329] and RBP3 [1–271] carboxy-terminal truncated proteins were unable to interact with H (Fig. 3C, lanes 10–12, and 4–6, respectively). We then assayed a short internal region (RBP3 [330–438]) for its capacity to interact with GST–H [1–292], as shown in Figure 3C (lanes 16–18), this 108-amino-acid in vitro-translated product is specifically retained onto GST–H [1–292] beads. Thus, RBP3 can interact in vitro with Drosophila H. Both Su[H] and RBP3 interact with a similar region of H, located between amino acid positions 1 and 292 (Figs. 3A and 5B; data not shown). Likewise, a similar HID can be defined in both Su[H] and RBP3 proteins (Fig. 5A, below). Note also that the HID partly overlaps the DNA-binding domain (see Fig. 5A, below).
action in the yeast nucleus between the HID of RBP3 and H [1–292]. These results fully confirm our conclusions based on the in vitro study presented in Figures 3 and 5.

**H inhibits the DNA-binding activity of Su(H) and RBP proteins**

The antagonistic nature of H and Su(H) activities, together with the direct protein–protein interaction between H and Su(H), suggest that H could possibly control the activity of Su(H). This could be at the level of DNA binding, subcellular localization, or transcriptional activity on DNA binding of Su(H). We first examined whether the in vitro-translated H protein affects the DNA-binding activity of Su(H) [1-292]. Increasing amounts of H, added to a constant amount of Su(H) proteins and DNA probes [Fig. 4A, lane 3; data not shown], correlates with a progressive reduction in the intensity of the retarded complex. Adding a twofold molar excess of the H protein, relative to Su(H), decreases the amount of the retarded complex by about threefold (data not shown). H alone did not bind to the 2122 nor HES probes (Fig. 4A, lanes 1,11; data not shown), and the formation of new, supershifted complexes was not observed. Thus protein–protein interactions between H and Su(H) apparently lead to the formation of complexes that are unable to recognize the Su(H) target site. The specificity of this inhibitory effect could be demonstrated further using two Su(H) truncated proteins. Su(H) [10–525] was shown to both bind DNA (Fig. 2C, lane 3) and interact in vitro with H (Fig. 3B, lane 5). Consistent with these data, we show here that H inhibits the DNA binding of Su(H) [10–525] (Fig. 4A, lanes 4,5). In contrast, Su(H) [10–457] is known to bind DNA but not H (Fig. 2C, lane 4; Fig. 3B, lane 8; Fig. 5A, below). Accordingly, H had no detectable effect on its DNA-binding ability (Fig. 4A, lanes 6,7). This result confirms that a region between amino acids 457 and 525 in Su(H) is necessary for interaction with H but not for DNA binding (Fig. 5A). We then investigated whether *Drosophila* H may also inhibit KBF2/RBP-Jκ DNA-binding activity. As shown in Figure 4A, H effectively decreases binding of purified KBF2 (Fig. 4A, lanes 8,9), as well as of the in vitro-translated proteins RBP1 (lanes 13,14), RBP3 (lanes 10,12), and RBP4 (data not shown), to the HES probe. A similar inhibitory effect was also observed with recombinant GST–RBP3 fusion protein (Fig. 4B, cf. lanes 11 and 7) or with the endogenous KBF2 activity present in nuclear extracts from several cell lines [human kidney 293T and neuroblastoma NGP, mouse myoblast C2; data not shown]. As a negative control we verified that H does not affect the DNA-binding activity of the p50 subunit of NF-κB [data not shown]. Finally, several H-deletion mutants were tested for their capacity to inhibit Su(H) or RBP3 DNA-binding activity. A similar fivefold molar excess of H proteins relative to Su(H) was used for all H-deletion constructs (Fig. 4B, lanes 1–6). The amount of H proteins compared with bacterially produced GST–RBP3 proteins was kept constant in all lanes (Fig. 4B, lanes 7–11). Full-length H and H [1–710] exhibited strong

**Figure 4.** H inhibits the DNA-binding activity of Su(H), KBF2, and RBP isoforms. (A) A fivefold molar excess of in vitro-translated H [lanes 1,3,5,7,9,11,12,13] or unprogrammed rabbit reticulocyte lysate [lanes 2,4,6,8,10,14] was preincubated with in vitro-translated Su(H) [lanes 2,3], Su(H) [10–525] [lanes 4,5], Su(H) [10–457] [lanes 6,7], RBP3 [lanes 10,12], RBP1 [lanes 13,14], or purified KBF2 [lanes 8,9] before addition of 2122 [lanes 1–7] or HES probe [lanes 8–14] and assayed by gel shift. (B) Deletion mapping of the H region required to inhibit the binding of Su(H) to the 2122 probe [lanes 1–6] or of recombinant GST–RBP3 to the HES probe [lanes 7–11]. Identical amounts of in vitro-translated H [lanes 2,7], H [1–710] [lane 8], H [1–293] [lanes 3,9], H [1–236] [lanes 4,10], H [1–84/236–293] [lane 5], H [212–293] [lane 6], or unprogrammed lysate [lanes 1,11] were incubated with Su(H) or GST–RBP3 as indicated, before addition of probes 2122 [lanes 1–6] or HES [lanes 7–11].
DNA-binding inhibition activity, whereas H [1–293] gave a weaker inhibitory effect [Fig. 4B lanes 2,3,7–9]. The H [1–236], H [1–84/236–293], and H [212–293] mutant proteins had no inhibitory effects upon Su[H] DNA binding [Fig. 4B, lanes 4–6,10]. These results, summarized in Figure 5B, are all consistent with the in vitro protein interaction results [Figs. 3A and 5B], with the exception of the H [212–293], which was able to weakly

**Figure 5.** Schematic representation of the functional domains of the Su[H], RBP3, and H proteins. (A) Deletion mapping of Su[H] (top) and RBP3 (bottom) DNA-binding domain and HID. Su[H] and RBP3 are drawn at the same scale and are aligned following their highly conserved region represented as a solid box (Su[H], amino acids 103–528; RBP3, amino acids 15–440). The results of the gel shift experiments of Fig. 2, B and C, are summarized in the column labeled DNA binding (+, DNA binding; -, no DNA binding), the results of the in vitro interaction experiments of Fig. 3 in the column labeled in vitro interaction with H (+, interaction with H; -, no detectable interaction; + / -, weak interaction; ns, not shown; nd, not determined), the results of gel shift experiments of Fig. 4 in the column labeled inhibition of DNA binding (−, H inhibits the DNA binding of this protein; –, H has no effect on DNA binding). In the central part, the DNA-binding domain and the HID of Su[H] and RBP3 are compared. (B) Deletion mapping of the H protein. The results presented in Figs. 3 and 4 are summarized here as in A.

<table>
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<tr>
<th><strong>DNA-binding</strong></th>
<th><strong>in vitro interaction with H</strong></th>
<th><strong>inhibition of DNA binding</strong></th>
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<tr>
<td><strong>Su[H]</strong></td>
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<tr>
<td>Su[H] [10–994]</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Su[H] [10–525]</td>
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<td>+</td>
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<tr>
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<td>-</td>
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<tr>
<td>Su[H] [10–287]</td>
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<td>-</td>
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<tr>
<td>Su[H] [110–594] GST</td>
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<td>+</td>
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<tr>
<td>Su[H] [139–594] GST</td>
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<td>Su[H] [444–594] GST</td>
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<tr>
<th><strong>Human RBP3</strong></th>
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<tr>
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<td>+</td>
<td>+</td>
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<tr>
<td>RBP3 [1–329]</td>
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<tr>
<td>RBP3 [1–271]</td>
<td>-</td>
<td>-</td>
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<tr>
<td>RBP3 [175–486]</td>
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<td>+</td>
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<tr>
<td>RBP3 [330–438]</td>
<td>+</td>
<td>+</td>
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<tr>
<td>RBP3 [7–486]</td>
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<td>RBP3 [330–438]</td>
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**DNA binding domain**

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<thead>
<tr>
<th>Su[H]</th>
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<tr>
<td>RBP3</td>
<td>7–435</td>
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**H interaction domain**

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<tr>
<th>Su[H]</th>
<th>288–525</th>
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<tr>
<td>RBP3</td>
<td>330–418</td>
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**Hairless**

| H [1–1076] | +     |
| H [1–716]  | + (ns) |
| H [1–293]  | +     |
| H [1–236]  | -     |
| H [1–84/236–293] | -     |
| H [212–293] | +/-   |
interact in vitro with GST–Su(H) [110–594] but did not inhibit Su(H) DNA-binding activity or supershift the retarded complex.

_H inhibits Su(H)-dependent transcriptional activation in transfected S2 cells_

The importance for transcription regulation of the inhibition by H of the Su(H) DNA-binding activity was examined in cotransfection experiments using the Drosophila cell line S2 [Fig. 6]. Two direct repeats of the 2122 DNA-binding sites were introduced upstream of the minimal Adh promoter [−86, +53] [Englund et al. 1990] driving CAT gene expression [[2122]2 Adh-86 CAT]. CAT activity was measured following cotransfection of the above with expression plasmids producing the Su(H) and/or H gene products under the control of the actin 5C promoter.

First, although no effect was observed using the Adh-86 CAT reporter plasmid alone, CAT expression from the [2122]2 Adh-86 CAT construct was increased upon cotransfection with Su(H). Average stimulation was found to be 7.4-fold, indicating that Su(H) acts as a transcriptional activator in this assay. Second, cotransfection of the Su(H) and H expression vectors resulted in a dose-dependent reduction of the stimulated CAT expression. The Su(H)-dependent stimulation was 2.9-fold with a H/Su(H) ratio of 1:1, and 1.8-fold with a 4:1 ratio. As a control, H did not, on its own, affect the level of expression of the two reporter genes used. These results indicate that H may inhibit in vivo the transcriptional activation of a synthetic promoter mediated by Su(H).

**Discussion**

We report here the cloning of a cDNA encoding the KBF2 protein. Peptide and cDNA sequencing revealed that KBF2 is identical to RBP-Jk [Matsumani et al. 1989; Amakawa et al. 1993]. The genomic organization of the human RBP-Jk gene has been reported [Amakawa et al. 1993]. Three different splicing products, RBP1, RBP2, and RBP3 [according to the nomenclature proposed in this paper], have been described by Amakawa et al. [1993]. We describe here a fourth isoform, RBP4, that contains part of exon N1 linked to exon 2. A fifth protein that corresponds to the previously described mouse splicing product RBP2N [Kawaichi et al. 1992] was reported recently [Dou et al. 1994]. It is possible that other potential splicing products may exist. The purified KBF2 activity may thus correspond to a single splicing product or a combination of specific RBP splicing products. The DNA-binding domain of RBP maps to a large, highly conserved region common to all isoforms [from amino acid 7 to 435 in RBP3, and from 110 to 457 in Su(H)]. The possible functional diversity associated with these various RBP isoforms remains to be investigated. Similarly, although the RBP-Jk gene products are expressed in many different cell types [Hamaguchi et al. 1992; data not shown], the detailed expression pattern of each isoform has not been described and awaits the use of isoform-specific probes.

**Direct protein–protein interaction with Drosophila**

_H prevents Su(H)/RBP3 binding to DNA_

Further insights into the possible regulation of RBP activity were gained from studies of Su(H) function in fly development. Su(H), together with H, controls at least two alternative cell fate decisions during peripheral nervous system (PNS) formation of the adult fly [Bang et al. 1991; Bang and Posakony 1992, Schweisguth and Posakony 1992, 1994]. We show here that Su(H) binds the consensus-binding site defined for the mouse RBP protein and acts upon transfection as a transcriptional activator of an artificial promoter carrying two copies of this sequence. These two assays for Su(H) function allowed us to investigate the molecular basis of the antagonistic activity of the H protein, for which no biochemical activity is known so far. First, H acts as a repressor by decreasing in a dose-dependent manner the transcriptional activation mediated by Su(H) in transfected cells. We show further that H specifically prevents Su(H) binding to its target site in vitro and that H interacts directly with Su(H) in vitro. The ability of H to inhibit Su(H) DNA binding extends to its vertebrate cognates, RBP1, RBP3, and RBP4, as well as to purified KBF2. Likewise, H and RBF3 were shown to interact in vivo using the yeast

**Figure 6.** Transcriptional activities of Su(H) and H in Drosophila S2 cells. CAT assays of S2 cell extracts following transfection with 2 μg of the reporter genes Adh-86 CAT (open bars) or [2122]2Adh-86 CAT (shaded bars) along with variable amounts of the Su(H) and/or H expression vectors as indicated. Histograms correspond to the average of at least three independent experiments. Each activity is expressed as the fold increase of CAT activity over the value from 0.3 μg of the reporter gene plasmid alone, which was set at 1.

<table>
<thead>
<tr>
<th>Expression plasmids (μg)</th>
<th>0.5</th>
<th>0.5</th>
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<tr>
<td>Su(H)</td>
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<td>H</td>
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**GENES & DEVELOPMENT 2499**
two-hybrid system and in vitro by direct protein–protein interactions. The functional domain required for interaction with H (HID) lies within the region highly conserved between Su(H) and RBP-κ. The HID displays a high percentage of prolines (10%) and hydrophobic amino acids (28% for RBP3 and 26% for the corresponding region of Su(H)). We note that the H domain required for the interaction with Su(H) also includes a potential α-helix with hydrophobic residues on one side of the helix. The role of these hydrophobic residues in the formation of H–Su(H) complexes remains to be investigated. The PRD repeat located at the carboxy-terminal end of the H protein, which may regulate protein–protein interactions, plays no apparent role in the H–Su(H) interaction.

The mechanism by which H prevents Su(H)/RBP DNA binding is at unclear present. Considering that the HID partly masks the DNA interaction surface of Su(H) and RBP, H and Su(H)/RBP would then form stable complexes unable to bind DNA. This hypothesis is reminiscent of the mechanism by which calreticulin prevents the glucocorticoid receptor from binding to its DNA response element (Burns et al. 1994; Dedhar et al. 1994) or IP1 inhibits AP1 DNA binding (Auwerx and Sassone-Corsi 1991). An alternative mechanism that cannot be formally excluded is one in which the H–Su(H)/RBP complex could still bind DNA but with an altered specificity.

Could the formation of H–Su(H) heteromeric complexes account for the antagonistic genetic interactions observed between H and Su(H)?

The results described in this paper would fully support a model in which H acts as a negative regulator of Su(H) activity. However, the dose-sensitive genetic interactions associated with partial loss of function in these two genes initially led Ashburner (1982) to propose that Su(H) acts as a negative regulator of H function. This conclusion stems from the observation that a reduction in Su(H) gene dosage suppresses the partial H loss-of-function phenotype, whereas an increase in gene dosage enhances it (Ashburner 1982). However, as first pointed out by Locke et al. (1988), these dose-dependent effects could also be explained by the formation of multiprotein complexes involving the Su(H) protein. We thus propose that Su(H) acts on DNA binding to favor the adoption of an epithelial, that is, epidermal and socket-producing, cell fate and that H inhibits its DNA-binding activity to promote a subepithelial, that is, neuronal and shaft-secreting, cell fate. This proposal is entirely consistent with a role for H in protecting the future SOP cell from residual negative signaling that would be mediated by Su(H) (Bang and Posakony 1992). The observation that the neurogenic Su(H) null phenotype is partially suppressed in Su(H);H double mutant larvae (Schweisguth and Posakony 1994) may also suggest that H acts independently of Su(H) to negatively regulate additional neurogenic activities. Thus, it is possible that H, when not associated with Su(H), may still participate in other regulatory interactions. We propose that direct protein–protein interactions between H and Su(H) control both the activity of Su(H) and the amount of H protein unbound to Su(H). The dominant loss of H function phenotype predicts that H is present in limiting amount. A threefold increase in Su(H) gene dosage results in an adult phenotype very similar to the dominant H phenotype observed in H heterozygote flies (Schweisguth and Posakony 1994; data not shown). Although the subcellular distribution of the H protein is not yet known, the nuclear localization of both Su(H) (F. Schweisguth and J.W. Posakony, unpubl.) and RBP (Hamaguchi et al. 1992) suggests that the H–Su(H) complex probably forms in the nucleus.

Functional conservation between Su(H) and RBP-κ

Our results suggest a possible interpretation for the remarkable degree of structural similarity between Su(H) and RBP. First, these proteins have a large DNA-binding domain. Second, the highly conserved sequences that are not required for DNA binding appear involved in protein–protein interactions. Considering that regulation by H is a fundamental aspect of Su(H) activity in flies and that the human RBP proteins interact with Drosophila H in a manner identical to Su(H), we propose that a human H homolog should exist that has the same ability to interact with the RBP proteins. Experiments are in progress to isolate this human homolog of H. This study also suggests another level of functional conservation between Su(H) and RBP. Genes from the E(spl) complex are thought to act at the last step in a genetic cascade of gene activities restricting neural fate and to be the genetic target of the H gene activity (de la Concha et al. 1988). Consistent with this proposition, the promoter of the m8 gene of the E(spl) complex contains a potential binding site for Su(H) (Tun et al. 1994; M. Lecourtois and F. Schweisguth, unpubl.). We further report here that the promoter of the HES1 gene, a mammalian homolog of the E(spl) genes, also contains a strong RBP-binding site. This raises the exciting possibility that not only the specificity of DNA binding and the negative regulation by H, but also their direct downstream target genes have been conserved throughout evolution. Whether RBP-κ may also act as Su(H) in the Notch-mediated signaling pathway during mammalian development remains to be determined.

Materials and methods

Purification and cloning of KBF2

KBF2 was purified from 250 grams of HeLa cells. Its activity was followed by bandshift assay as described in Israel et al. (1989). Nuclear extract preparation, ammonium sulfate precipitation, and hydroxylapatite chromatography have been described in Kieran et al. (1990). The fraction eluted from the hydroxylapatite
column, which contains both KBF1 and KBF2, was dialyzed against a buffer containing 0.1 M NaCl and then loaded onto a DNA affinity column containing the H-2K\textsuperscript{b} X\textsuperscript{b} site in the presence of poly[d(I-C)] [Kieran et al. 1990]. This step allowed separation of KBF1, retained on the affinity column, from KBF2 that does not bind to the H-2K\textsuperscript{b} X\textsuperscript{b} site under the conditions used. The flowthrough was subsequently loaded onto a heparin-agarose column. KBF2 activity was eluted at 0.3 M NaCl, dialyzed, and loaded onto a sulfolipid ion-exchange column. Active fractions eluting at 0.6 M NaCl were pooled and dialyzed against a buffer containing 50 mM NaCl. A second DNA affinity column containing the P2m site [Israel et al. 1989] was then used. Most of the KBF2 activity bound to this column, and the KBF2 protein was eluted at 0.2 M NaCl. The active fraction was dialyzed and loaded once again onto the P2m affinity column. SDS gel analysis detected a single 58-kD protein. This 58-kD protein could be specifically cross-linked by UV irradiation to RBP2 cDNA is 1.6 kb. The HeLa RT–PCR product was subcloned and sequenced: It corresponds to RBP4 [see Fig. 1].

Synthesis of in vitro-translated products and bandshift assays

In vitro-translated proteins were all synthesized in a reticulocyte lysate-coupled transcription/translation system [Promega], from cDNA fragments inserted into the P7T\textsuperscript{b}link vector. The cDNA fragment encoding the first 293 amino acids of H was subcloned following PCR amplification, using a composite full-length cDNA clone as template [plasmid KH, a gift from A. Preiss, Biozentrum, Basel, Switzerland] [Maier et al. 1993]. The full-length H was obtained by inserting an EcoRI fragment purified from plasmid KH into the unique EcoRI site of P7T\textsuperscript{b}linkH [1–293]. The amino-terminal deletion mutants RBP3 [1–157], RBP3 [7–286], RBP3 [98–286], and RBP3 [1–271] were obtained by PCR amplification from a template containing the full-length RBP3 cDNA. The carboxy-terminal deletion mutants RBP3 [1–435], RBP3 [1–329], and H [1–710] were obtained by digestion with restriction enzymes Sial, Apol, and Xhol, respectively, of the corresponding P7T\textsuperscript{b}link constructs. The carboxy-terminal deletion mutants Su[H] [10–525], Su[H] [10–547], Su[H] [10–399], and Su[H] [10–287] were obtained from internal deletions of the P7T\textsuperscript{b}link Su[H] [10–594] construct, using the restriction enzymes Eagl–Xbal, Pstl, Narl–Smal, and HincII–Smal, respectively. The following H constructs—H [1–236], H [212–236], and H [1–84/236–293]—were generated by internal deletions of the P7T\textsuperscript{b}link H [1–293], using the DraI–EcoRI, NeoI–HindIII, and NotI–Dral restriction enzymes.

Bandshift assays were performed as described in Israel et al. [1989] and Antoniewski et al. [1994]. The following double-stranded oligonucleotides were used as probes or competitor:

\begin{align*}
2122: & 5'\text{-gaccGCGGATCCGAGGAGAGGGACCTTTCAGGGGACag-5'} \\
HES: & 5'\text{-gaccGTTACTGTCGAGGAAAGATCTAATGCACCCCTTTCTTCCATGCTG...-5'} \\
KBF: & 5'\text{-gacGATGCGCAAGCTCGCGCGCTCGGCCTGGG...-5'} \\
\beta2: & 5'\text{-gaccGCTGGGAAACCCCGAGCCACCTTGGTCGCCGCT-5'} \end{align*}

GST fusion proteins and interaction assays

For GST–RBP3 protein expression in E. coli, a restriction fragment from P7T\textsuperscript{b}link RBP3 construct was subcloned into the pGEX-KT vector as a fusion with GST [Smith and Johnson 1988; Hakes and Dixon 1992]. The amino-terminal deletion derivatives of Su[H] were obtained by subcloning the NeoI–Ndel, XmnI–Ndel, Scal–Ndel, HincII–Ndel, and MscI–Ndel cDNA fragments purified from SK12 [Schweiguth and Posakony 1992] into the filled-in BamHI site of the pGEX-KG vector [Hakes and Dixon 1992].

In the in vitro-translated proteins (\(3 \mu\)L) were mixed with glutathione–agarose beads (10 \(\mu\)L) coated with GST protein and preadsorbed for 1 hr at 4°C in buffer A (40 mM HEPEs pH 7.5, 5 mM MgCl\(_2\), 0.2 mM EDTA, 1 mM DTT, 0.5% NP-40, 100 mM KCl). The supernatant was incubated with glutathione–agarose beads carrying either GST or GST fusion proteins (RBP3, Su[H], or H [1–293]) for 1 hr at 4°C. The beads were washed extensively with buffer A, then with buffer A containing 200 mM KCl. Associated proteins were eluted by boiling in Laemmli buffer and subjected to SDS-PAGE analysis, followed by fluorography.
Y190 strain (Harper et al. 1993) contains a GAL-lacZ reporter integrated into the genome. Its genotype is [MATa, leu2-3,112, ura3-52, trp1-901, his3-D200, ade2-101, gal4a gal80Δ URA3:: GAL-lacZ, LYS::GAL-HIS3, cyh2]. The pAS1–CYNH2-derived plasmids contain the first 147 amino acids of GAL4, which correspond to its DNA-binding domain [GAL[DB]] fused to various fragments of RBP3, obtained by Ncol and Xhol digestion of the pT7link-derived vectors and insertion at the Ncol and SalI sites of pAS1–CYNH2. The GAL[AD]H–[I–292] encodes the activation domain of GAL4 [plasmid pACTII (Dufree et al. 1993)] fused to the amino-terminal 292 amino acids of H. The pT7link–H plasmid was digested by Xbal, filled-in by the Klenow enzyme, and redigested by Ncol. The appropriate fragment was inserted into pACTII vector digested by Ncol and SmaI.

Y190 was transformed by the pACTII- and PAS1-derived plasmids using the lithium acetate method (Gietz et al. 1992). The appropriate fragment was inserted into pACTII vector digested by Ncol and SmaI.

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C Brou, F Logeat, M Lecourtois, et al.

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