BEHAB, a New Member of the Proteoglycan Tandem Repeat Family of Hyaluronan-binding Proteins That Is Restricted to the Brain

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Abstract. Hyaluronan (HA) is a ubiquitous component of the extracellular matrix of all tissues. In the mammalian central nervous system (CNS) HA is present throughout development and into adulthood. While the functions of HA are likely to be mediated by HA-binding proteins, no cell or tissue specific HA-binding proteins have been reported. In an effort to characterize the composition of the extracellular matrix of the CNS, we sought to identify neural HA-binding proteins.

We report here the isolation and characterization of a cDNA with a high degree of sequence homology to members of the proteoglycan tandem repeat (PTR) family of HA-binding proteins. Unlike other HA-binding proteins, the expression of this cDNA is restricted to the CNS. We propose the name BEHAB, Brain Enriched HyAluronan Binding protein, for this gene. The expression of BEHAB mRNA is developmentally regulated; expression is first detected in the late embryonic period and peaks during the first two postnatal weeks. In the embryo, BEHAB is expressed at highest levels in mitotically active cells. The sequence of BEHAB has long stretches of identity between rat and cat, suggesting that the encoded protein is functionally important. The size and sequence of BEHAB are consistent with the possibility that it could serve a function like link protein, stabilizing interactions between HA and brain proteoglycans. These observations suggest the existence of other tissue specific HA-binding proteins.

While the presence of an extracellular matrix in non-neural tissues has been universally accepted, only recently has evidence been adduced for an organized extracellular matrix in the mammalian central nervous system (CNS) (Reichardt and Tomaselli, 1991; Hockfield et al., 1990; Sanes, 1989). Because molecules normally associated with the extracellular matrix, such as laminin and collagen, are not present (or present at very low abundance) in the CNS, the extracellular space in neural tissue has sometimes been considered a preparative artifact. However, the application of a variety of staining techniques demonstrated that the neural extracellular space is filled with a matrix-like material (Nakanishi, 1983; Castejon, 1970; Bomdareff, 1967). With the advent of monoclonal antibody technology, it became clear that many components of the extracellular matrix identified in other tissues are indeed present in the CNS (Margolis and Margolis, 1993; Rauch et al., 1991; Zaremba et al., 1990; Hockfield, 1990; Fujita et al., 1989).

The CNS extracellular matrix consists of a heterogeneous mixture of glycoconjugates, many of which are proteoglycans. Proteoglycans are complex macromolecules that consist of a core protein modified with one or more types of glycosaminoglycan chains (Hardingham and Fosang, 1992; Wight et al., 1991). Many functional properties of proteoglycans have been ascribed to glycosaminoglycans. Glycosaminoglycans have been reported to exhibit both adhesive and repulsive properties and, as such, have been suggested to mediate neuronal migration and axon guidance (Prieto et al., 1992; Cole and McCabe, 1991; Snow et al., 1990). Glycosaminoglycans are believed to regulate the local cellular environment primarily by serving as selective filters, facilitating permeability and retention of low molecular weight solutes, including growth factors, while excluding other macromolecules. Hyaluronan (HA) is particularly suited to this function because of its charge density and hydrosopic nature. HA is a ubiquitous component of extracellular matrices of all tissues, including brain, and is believed to organize water and extracellular proteins (Laurent and Fraser, 1992). During development HA plays a role in the regulation of morphogenesis and differentiation of neural tissues (Toole, 1991; Perris and Johansson, 1990).

Because HA is ubiquitously present in extracellular space, cell type specific functions attributed to HA may be mediated through its interaction with HA-binding proteins. Several HA-binding proteins have been reported in the brain, a subset of which have a high degree of sequence similarity to one another, including versican (Zimmermann and Ruos-
lahti, 1989), link protein (Doerge et al., 1986), neurocan (Rauch et al., 1992), glial hyaluronan binding protein (GHAP) (Perides et al., 1989) and CD44 (Culty et al., 1990), and have been called the proteoglycan tandem repeat (PTR) family of HA-binding proteins (Perkins et al., 1989). Additional HA-binding proteins have been isolated from brain, for which sequence is not yet available, including the T1 proteoglycan (Iwata and Carlson, 1993), and the antigen recognized by monoclonal antibody Cat-301 (Fryer et al., 1992).

The spatial distribution and temporal expression of neural extracellular matrix proteoglycans, and HA-binding proteins, in particular, indicate that they may be involved in many events in the development and function of the mammalian CNS. While some HA-binding proteins represent general components of the extracellular matrix (e.g., T1 [Iwata and Carlson, 1993]), others have a restricted pattern of expression on subsets of neurons (e.g., the Cat-301 antigen [Hockfield and McKay, 1983; Hockfield et al., 1983; Hockfield and Sur, 1990; Defoe et al., 1990]). In addition, while some extracellular matrix molecules are transiently expressed during embryogenesis (e.g., laminin [Letourneau et al., 1989; McLoon et al., 1988] and fibronectin [Stewart and Pearman, 1987]), others are first expressed late in the postnatal period (e.g., chondroitin sulfate proteoglycans [Watanabe et al., 1989; Aquino et al., 1984; Hockfield et al., 1983; Maeda et al., 1992]), coincident with the decline in developmental synaptic plasticity.

Our interest in the composition and function of the CNS extracellular matrix, along with the ubiquitous expression of HA in the brain, motivated an investigation of neural HA-binding proteins. Here we describe the identification and characterization of a neural specific cDNA that shows a high degree of sequence similarity to the PTR family of HA-binding proteins. We propose to name this gene BEHAB, for Brain Enriched HyAluronan Binding protein.

**Materials and Methods**

**Isolation of Rat cDNA Clones**

An unamplified postnatal day 12 (P12) rat brain Agt10 cDNA library was screened with rat aggrecan clone pRCP 4 (Doerge et al., 1987). 4 × 10^4 phage (per 150-mm plate) were plated with C600 bacteria, immobilized onto nitrocellulose filters, and prepared for hybridization according to standard techniques (Sambrook et al., 1989). Filters were prewashed for 1 h in 1 M NaCl, 0.1% SDS, 20 mM Tris-HCl (pH 8.0) and 1 M EDTA at 65°C. Filters were then prehybridized for an additional 4–6 h in 50% formamide, 5× SSC (1× SSC = 0.15 M sodium chloride, 0.015 M sodium citrate), 1% SDS, 1× Denhardt's (0.02% Ficoll, 0.02% BSA [Fraction V], 0.02% polyvinylpyrrolidone), 50 mM sodium phosphate (pH 6.7) and 100 μg/ml salmon sperm DNA at 37°C. Hybridization was carried out in the identical solution with the inclusion of 10^6 cpmp cDNA probe for 24 h at 37°C. For all experiments, radiolabeled probes (32P-cDNA, Amer sham) were prepared by random priming (Feinberg and Vogelstein, 1983) (Boehringer Mannheim Corp., Indianapolis, IN) gel purified cDNA clones, followed by the removal of unincorporated radiomucleotides (NICK column, Pharmacia). One post-hybridization wash in 2× SSC, 0.1% SDS and one in 0.2× SSC for 1 h each were performed at room temperature. Phage DNA was isolated using DES2 (Whatman) and the cDNA insert excised by EcoRI digestion. The cDNA was gel purified (Gene-Clean, Bio 101), subcloned into pBluescript KS* (Stratagene, LaJolla, CA) and transformed into DH5α (Mandel and Higa, 1970) (GIBCO BRL, Gaithersburg, MD).

**Isolation of Cat cDNA Clones**

Random nonamers (1.4 mg) were used to synthesize first strand cDNA from 5 μg poly A+ RNA isolated from P39 cat cortex. cDNA synthesis was performed according to manufacturer's instruction for the production of non-directional libraries (Stratagene) and size-fractionated by column chromatography (GIBCO BRL). 50 μg of cDNA was ligated to 1 μg EcoR I cut, phosphatized Lambda Zap II vector and packaged into phage (Gigapack II Gold, Stratagene). This yielded 5 × 10^6 recombinants when transfected into XLI-Blue (Stratagene). The unamplified library was screened with cat clone H1. Hybridization was performed in 6× SSC, 0.1% SDS, 1× Denhardt's and 100 μg/ml salmon sperm DNA at 65°C. Filters were washed twice in 2× SSC, 0.1% SDS and twice in 0.2× SSC at 65°C for 20 min. cDNA inserts of plaque purified positive clones were isolated in pBluescript SK+ by in vivo excision.

**DNA Sequencing and Analysis**

DNA sequencing was performed by the dideoxy chain termination method (Sanger et al., 1977) using Sequenase (U.S. Biochemical, Cleveland, OH). Bluescript SK+KS primers or cDNA specific 20-mers were used. Sequence was verified from overlapping clones or by sequencing both strands of DNA. Sequence compressions were resolved using dITP nucleotides. After labeling, the reactions were incubated at 37°C for 30 min in the presence of 1× reaction buffer, 1 mM GTPs (pH 7.0) and 0.5 U terminal deoxynucleotidyl transferase to prevent premature termination caused by the use of dITP. Sequence analyses were performed using the University of Wisconsin Genetics Computer Group (GGG) programs (Devereux et al., 1984).

**Northern Hybridization**

For Northern analysis, 25 μg total RNA was denatured in 2.2 M formaldehyde, 50% formamide, 1× MOPS (3-N-morpholino) propanesulfonic acid) buffer at 65°C for 15 min. The RNA was electrophoresed on a 1.0% agarose-formaldehyde gel with 1× MOPS buffer at 60°C with buffer recirculation. The gel was briefly neutralized in transfer buffer (2× SSC) and RNA blotted to Zetaprobe (BioRad Labs., Hercules, CA) by capillary transfer. Filters were rinsed briefly in 2× SSC, and RNA was immobilized both by UV cross-linking (Church and Gilbert, 1984) and baking in vacuo (80°C for 1 h). Hybridization in 7% SDS, 1% BSA, 0.5 M phosphate buffer (pH 6.8) (PB), 1 mM EDTA and 0.5–2.5 × 10^6 cpmp rat H1 probe/ml was carried out for at least 8 h at 65°C. Filters were washed twice in 5% SDS, 0.5% BSA, 40 mM PB, 1 mM EDTA and twice in 1% SDS, 40 mM PB, 1 mM EDTA at 65°C for 20 min (Church and Gilbert, 1984), and exposed to film (Hyperfilm, Amersham) at −70°C. Molecular sizes were determined relative to RNA molecular weight standards (GIBCO BRL) and 28S and 18S ribosomal RNA observed during UV illumination. The ubiquitously expressed, non-developmentally regulated gene cyclophilin (Danielson et al., 1988; Lenoir et al., 1986) was used to determine equal loading of lanes. Densitometry was performed using the NIH Image program.

**In Situ Hybridization**

Twelve to fourteen micron thick frozen sections were thaw-mounted onto gelatin-coated slides and postfixed in 0.1 M sodium phosphate buffered 4% paraformaldehyde (pH 7.4). Sections were rinsed in 1× PBS (137 mM NaCl, 2.7 mM KCl, 10 mM Na2HPO4, 1.8 mM KH2PO4, 0.1 M triethanolamine (pH 8.0). Sections were then rinsed in 2× SSC, 1× PBS, dehydrated in ethanol and delipidated in chloroform. Sections were prehybridized in 2× SSC, 50% formamide at 50°C for 1 h, and then hybridized in 0.5 M NaCl, 50% formamide, 1× Denhardt's, 10% dextran sulfate, 30 mM DTT, 10 mM Tris-HCl (pH 7.5), 1 mM EDTA, 100 μg/ml salmon sperm DNA, 0.5 mg/ml yeast tRNA and 10^6 cpm probe per slide at 50°C for 12–15 h. [35S]cCTP (New England Nuclear, Boston, MA) labeled cDNA probes were synthesized using T3 (GIBCO BRL), SP6, and T7 RNA polymerases (New England Biolabs Inc., Beverly, MA). After hybridization, sections were washed in 2× SSC, 50% formamide, 0.1% BME (β-mercaptoethanol) at 50°C for 1 h and treated with 20 μg/ml RNase A in 0.5 M NaCl, 10 mM Tris-HCl (pH 8.0) at 37°C for 30 min. Sections were then washed in 2× SSC, 50% formamide, 0.1% BME at 58°C for 30 min and 0.1× SSC, 0.1% BME at 63°C for 30 min and dehydrated. For initial localization of probe the slides were exposed to film (Hyperfilm, Amersham) for 4 d. Autoradiograms were used as negatives for prints. For higher resolution, the slides were dipped in NTB-2 emulsion (Kodak), developed after 5 d and counterstained with cresyl violet. Neurofilament-middle (NF) antisense (Martin et al., 1992) and rat clone BI sense probes were used as positive and negative controls, respectively.
Results

BEHAB Encodes a Protein with Homology to the PTR Family of HA-binding Proteins

To identify HA-binding proteins involved in neural development, rat aggrecan clone pRC5-4, which encodes the HA-binding region (Doege et al., 1987), was used to screen a postnatal day 12 (P12) rat brain λgt10 cDNA library. A total of 3.2 × 10^6 recombinants were screened resulting in two positives. On Northern blots the two clones recognize the postnatal day 12 (P12) rat brain Xgfl0 eDNA library. A total binding region (Doege et al., 1987), was used to screen a rat aggrecan clone pRCP 4, which encodes the HA-binding region in relation to the coding sequence. The complete composite sequence for BEHAB was obtained from these overlapping clones (Fig. 2). The complete BEHAB coding sequence is 1,113 bp. The nucleotide sequence preceding the first AUG contains a consensus sequence for translation initiation (Kozak, 1991, 1984). In the 3' untranslated region only that sequence verified from clones B1, A2, and A3 is presented. The deduced amino acid composition of the BEHAB protein is comprised of 371 amino acids and includes a putative signal peptide cleavage site at Ala-22 (von Heijne, 1986). The resulting mature protein has a predicted molecular mass of 38,447 kD. Analysis of the deduced amino acid sequence indicates the presence of two NX(S/T) consensus sequences for potential N-glycosylation (Marshall, 1974).

\[ \text{Figure 1. Overlapping cDNA clones encoding BEHAB. (A) cDNA clones for rat BEHAB were isolated from a P12 rat brain λgt10 library and subcloned into pBluescript KS+.} \]

In our previous characterizations of extracellular matrix proteins in brain we focused on expression in the cat visual system. Therefore, we investigated whether the BEHAB gene is also expressed in the cat cortex. Northern hybridization using rat H1 as the probe revealed a single mRNA species in cat visual cortex identical in size to rat BEHAB (see Fig. 6). Rat H1 was used to clone the cat homologue from a P39 cat cortex Lambda Zap II cDNA library. A total of 3.2 × 10^6 recombinants were screened resulting in five positives which were excised in pBluescript SK- (Fig. 1 B). The composite sequence for cat BEHAB was obtained from these overlapping clones (Fig. 3). The complete coding sequence for cat BEHAB is 1,134 bp. The first AUG is preceded by both an in-frame termination codon and the translation initiation consensus sequence. The cat BEHAB sequence encodes 378 amino acids which, like the rat, contains a 22 residue signal peptide. However, cat BEHAB contains six additional amino acids at the carboxy terminus, resulting in a predicted molecular mass of 38,955 kD. In the cat, Trp-373 is encoded by TGG, while the corresponding rat sequence of TAG results in termination. This termination sequence was verified in rat clones B1, A5, and A8 and by sequencing both strands of cat clone Bl. Cat BEHAB also contains one additional site for potential N-glycosylation not present in the rat.

Database analyses at both the nucleic acid and amino acid levels indicated that BEHAB is a previously unreported member of the PTR family of HA-binding proteins. BEHAB has a substantial degree of amino acid identity to the other members of the PTR family, which includes rat aggrecan (48%) (Doege et al., 1987), rat neurocan (48%) (Rauch et al., 1992), human versican (46%) (Zimmermann and Rohs, 1989), and rat link protein (42%) (Doege et al., 1986). The NH2-terminal domain of the cat BEHAB is only that sequence verified from clones B1, A2, and A3. The Ig domain contains two clusters of conserved amino acids around the cysteine residues which generate the disulfide bond of the loop. The consensus sequence YxCxVxH in the COOH-terminal cluster is present in all immunoglobulin and major histocompatibility complex proteins, and is also present in BEHAB (Fig. 4). The most conserved region of the rat family’s HA-binding domain is the sequence CDAGWL(A/S)D(Q/G)(T/S)VRYPI found in PTR 1 and PTR 2. The Ig domains have been shown to mediate binding to HA (Doerge et al., 1991; Foxang and Hardingham, 1989; Gotzinck et al., 1987). The Ig domain contains two clusters of conserved amino acids around the cysteine residues which generate the disulfide bond of the loop. The consensus sequence YxCxVxH in the COOH-terminal cluster is present in all immunoglobulin and major histocompatibility complex proteins, and is also present in BEHAB (Fig. 4). The most conserved region of the rat family’s HA-binding domain is the sequence CDAGWL(A/S)D(Q/G)(T/S)VRYPI found in PTR 1 and PTR 2. Two copies of this sequence are also found in BEHAB. The degree of identity of BEHAB between rat and cat is high (84 % overall), with the greatest conservation in the Ig domain (84 %) (Fig. 5). The relative degree of homology between the PTR 1, PTR 2, and Ig domains observed in rat and cat is also observed between BEHAB and the other members of the PTR family (Table I and Fig. 4).

BEHAB mRNA Is Restricted to the Nervous System

The tissue distribution of rat BEHAB mRNA was determined by Northern blot analysis (Fig. 6). A single 3.9-kb mRNA transcript is detected in adult rat cortex, spinal cord, and cerebellum. This transcript is not detected in liver, kid-
Figure 2. Nucleotide and deduced amino acid sequences of rat BEHAB. The nucleotide sequence was determined from the overlapping cDNA clones shown in Fig. 1A. The deduced amino acid sequence contains a signal peptide, which is underlined, and a putative cleavage site, indicated by an arrow. The sequence also contains two potential N-glycosylation sites, which are boxed. These sequence data are available from EMBL/GenBank/DBJ under accession number Z2836.

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intense hybridization is present in the CA1 subfield, with less intense hybridization in subfields CA2 and CA3, and the dentate gyrus. Examination of emulsion-coated slides counter stained with cresyl violet reveal that the labeling in CA1 is associated with neurons (Figs. 8, A and B). The pattern of NF hybridization in the hippocampus is essentially reciprocal to that of BEHAB (Fig. 7 E); the NF probe hybridizes most intensely in the dentate gyrus. BEHAB hybridization is also seen throughout the inferior colliculus (IC) and less intensely in the superior colliculus (SC). In addition to the hippocampus, BEHAB hybridization in gray matter is most intense in the substantia nigra (SN). The B1 sense probe generates almost no signal in most of the brain, but a low level of hybridization is seen in the hippocampus and dentate gyrus (Fig. 7 I).

In the brainstem, BEHAB is expressed throughout the re-
BEHAB has homology to the PTR family of HA-binding proteins. A database search showed that the deduced amino acid sequence of rat and cat BEHAB have a high degree of homology to several PTR HA-binding proteins. The PTR proteins contain three functional domains: an immunoglobulin fold (A), and two domains thought to be involved in HA binding, PTR 1 (B) and PTR 2 (C). The homology among all of the HA-binding proteins is greatest in PTR 1 and FIR 2, followed by the Ig fold (see Table I). Identical amino acids are shown in black. Intensity of gray counter-shading, indicating amino acid similarity, was determined by the method of Schwartz and Dayhoff (1979). Sequence alignment was produced using the GCG PILEUP function and counter-shaded using PRETTYBOX.

BEHAB expression in the spinal cord is greater in the gray matter than white matter (Fig. 7 C). In the gray matter, BEHAB expression is slightly greater in the ventral than in the dorsal horn. BEHAB hybridization is lacking in the substantia gelatinosa (layer 2), but is present in layer 1. In the ventral horn hybridization is seen over motor neurons (Fig. 8, C and D). In the spinal cord white matter, the size of labeled cells and their distribution indicates that BEHAB is expressed by glial cells. Like BEHAB, NF expression is greater in the ventral horn than in the dorsal horn; however, unlike BEHAB, NF is not detected in the spinal white matter (Fig. 7 G). As observed in the brainstem, no hybridization signal is detected in the spinal cord with the B1 sense probe (Fig. 7 K).

In the cerebellum, BEHAB expression is greatest in the deep cerebellar nuclei (CN) (Fig. 7 D). The cerebellar cortex, labeling is detected in all three cortical layers. The molecular layer distribution of silver grains parallels the distribution of basket and stellate cells. In the Purkinje cell layer labeling is clustered over Purkinje cells and in the granule cell layer it is clustered over Golgi II cells (Fig. 8, E and F). The white matter of the cerebellar cortex also shows hybridization signal. NF is primarily expressed by Purkinje cells.
Rat BEHAB

| MFLPFL AAVLGVTPQAAPAAALDYLKDSEDRAFRVRI.GAAQLRGVLG |

Cat BEHAB

| MAPFLPFL AAVLGVTPAADSALDGSEDRAFRVRI.GSNAPLQGVVLG |

--- Signal peptide ---

**Immunoglobulin Fold**

GWVAIPCHVHLLPPSRRAAPGRXWMTLSGDRVEVLAVGRLRVKV

| GALTISCHVHYLRRPGAVLGSAPRXHTLSGREGAEVLARGRLRVKV |

--- PTR 1 ---

NEAYRFVRVALPAYSLTVLSELRPNDGYSVYRCEVQHGIDDDSDAV

| SEAYRFVRVALPAYSLTVLSELRPNDGYSVYRCEVQHGIDDDSDAV |

--- PTR 2 ---

YAEQLNGELFLGAPPGLVETLKEARDYTGLEHQAINTQGYYAMMNLDR

| YAEQLNGELFLGAPPGLVETLKEARDYTGLEHQAINTQGYYAMMNLDR |

--- PTR 3 ---

CSPSGWLADGSVRYPIIPTSQRCGGGLPGVTLFLFPAQTPGFSPKQNRNFV

| CSPSGWLADGSVRYPIIPTSQRCGGGLPGVTLFLFPAQTPGFSPKQNRNFV |

YCFRDSAHPSAFSEPPAQPLMD 371

| YCFRDSQPSSTPETASDQLGWRPSQ 378 |

**Figure 5.** BEHAB is highly conserved between species. The GCG BESTFIT alignment of rat and cat BEHAB amino acid sequences demonstrates the extent of conservation between species. Overall, the two proteins are 84% identical and possess many conserved substitutions. Based on homology with other HA-binding proteins (see text and Fig. 4), the BEHAB protein contains four domains: a signal peptide, an immunoglobulin fold, and two PTR domains involved in HA binding. The amount of conservation between rat and cat BEHAB is greatest in PTR 1 (95%) and PTR 2 (86%), followed by the immunoglobulin fold (84%) (see Table I).

**BEHAB mRNA Expression Is Developmentally Regulated**

To determine the temporal regulation of BEHAB mRNA expression, Northern blot analysis was performed using total RNA from embryonic and postnatal rat cortex and spinal cord (Fig. 9). The non-developmentally regulated gene cyclophilin was used as a control probe to verify equal loading.

**Table I. Percent Identity of Rat BEHAB to Other Members of the PTR Family of HA-binding Proteins**

|             | Ig | PTR1 | PTR2 |
|-------------|----|------|------|
| Cat BEHAB   | 84%| 95%  | 86%  |
| Aggrecan    | 40%| 59%  | 57%  |
| Neurocan    | 37%| 56%  | 51%  |
| Versican    | 36%| 59%  | 48%  |
| Rat link    | 34%| 93%  | 53%  |
| CD44        | 34%| 93%  | 53%  |

**Figure 6.** BEHAB mRNA expression is restricted to the nervous system. Total RNA (25 μg) from P39 cat cortex and from the indicated adult rat tissues was hybridized with a 32P-labeled probe for rat clone HI (see Materials and Methods). A single 3.9-kb mRNA transcript encoding cat BEHAB is detected. The 3.9-kb mRNA transcript for rat BEHAB, which is identical in size to cat BEHAB, is also present in rat cortex, spinal cord, and cerebellum. This transcript is absent from the non-neural tissues even after prolonged film exposure. The ubiquitously expressed gene cyclophilin was hybridized simultaneously as a positive control. The position of 28S and 18S rRNA is indicated at the right.

and by cells of the deep cerebellar nuclei (Fig. 7 H). The sense B1 probe generates a low level of diffuse hybridization signal throughout the granule cell layer (Fig. 7 L).

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Figure 7. BEHAB mRNA is expressed throughout the central nervous system. In situ hybridization was performed on sections from P21 rat forebrain, brainstem, spinal cord, and cerebellum using a 35S-labeled cRNA probe to rat clone B1 (A-D) (see Materials and Methods). Near adjacent sections were probed with NF (neurofilament, middle) as a marker for neuronal mRNA expression (E-H) and a sense probe of clone B1 as a negative control (I-L). (A–D) BEHAB hybridization is detected in both gray and white matter. A diffuse hybridization signal is detected throughout the gray matter at all levels. In the forebrain (A), the highest levels of hybridization are detected in the hippocampus (H) and substantia nigra (SN). In the hippocampus the most intense hybridization is present in the CA1 subfield (see Figs. 8, A and B). The inferior colliculus (IC) has a higher hybridization signal than the superior colliculus (SC). A high level of hybridization is seen in white matter tracts, including the corpus callosum (CC), the fimbria of the hippocampus (F), and the anterior commissure (AC). (B) BEHAB is expressed in several brainstem nuclei, including the superior olivary nucleus (ON), the vestibular nucleus (VN), the abducens nucleus (AN), and the dorsal column nuclei (DCN). (C) BEHAB expression in the spinal cord is greater in the gray matter than white matter. In the gray matter, BEHAB hybridization is slightly greater in the ventral horn (V) than in the dorsal horn (D). In the dorsal horn, BEHAB is present in layer 1, but is absent in the substantia gelatinosa (see Fig. 8, C and D). (D) BEHAB expression in the cerebellum is greatest in the deep cerebellar nuclei (CN). Labeling is also detected in all three cortical layers and the white matter (see Fig. 8, E and F). (E) NF hybridization is very high in many gray matter regions, but is virtually absent from white matter tracts. In the hippocampus NF localization is reciprocal to that of BEHAB; NF hybridizes most intensely in subfields CA2, CA3, and in the dentate gyrus. (F) NF hybridization in the nuclei of the brainstem is similar to BEHAB, but it is not seen in white matter tracts. (G) In the spinal cord, NF mRNA is restricted to the gray matter, where, like BEHAB, it is present expressed at greater levels in the ventral horn. (H) In the cerebellar cortex NF is primarily expressed by Purkinje cells; it is also expressed at high levels by the neurons in the deep cerebellar nuclei. (I–L) The sense probe generates almost no hybridization signal, except a low level seen in the hippocampus, dentate gyrus, and granule cell layer of the cerebellum. Bar, 1 mm.
Figure 8. BEHAB mRNA is expressed by neurons and glial cells. Emulsion-coated sections were counter stained with cresyl violet and photographed under dark field optics to visualize silver grains (A, C, and E) and with transmitted illumination to visualize cell bodies (B, D, and F). (A and B) In the hippocampus BEHAB hybridization is most intense in the CA1 subfield, and is much less intense in subfields CA2 and CA3, and in the dentate gyrus (DG). Observation of cresyl violet counter stained sections shows that the BEHAB hybridization pattern is not attributable to differences in cell density. (C and D) In the spinal cord BEHAB expression is seen in gray matter (GM) and white matter (WM). In the gray matter, BEHAB expression is slightly greater in the ventral than in the dorsal horn. In the ventral horn hybridization is seen over motor neurons (arrowhead). In the dorsal horn hybridization is present in layer 1 (single arrow), but is absent from the substantia gelatinosa (double arrow). In the white matter the size and distribution of labeled cells suggests BEHAB synthesis by astrocytes. (E and F) In the cerebellar cortex, intense BEHAB hybridization is detected in the Purkinje cell layer (P) and in the white matter (WM). The distribution of silver grains in the molecular (M) and granule cell (G) layer indicates expression by interneurons (arrows). Bars: (A-D) 200 μm; (E and F) 50 μm.
Figure 9. BEHAB mRNA is developmentally regulated. Northern analysis was performed with 25 μg total RNA from E17, P0, P5, P14, P21, and adult (Ad) rat cortex (A) and spinal cord (B). Hybridization was performed simultaneously with 32P-labeled probes of rat clone H1 and the non-developmentally regulated gene cyclophilin (see Materials and Methods). A single 3.9-kb mRNA transcript is detected in both cortex and spinal cord. To determine the developmental expression of BEHAB, densitometry was performed, and standardized by calculating the ratio of BEHAB absorbance relative to cyclophilin at each age (C). In the cortex, BEHAB recognizes a single 3.9-kb mRNA transcript (Fig. 9 A). BEHAB expression is detected at embryonic day 17 (E17) and gradually increases to attain adult levels by P21 (Fig. 9 C). In the spinal cord, BEHAB also recognizes a 3.9-kb mRNA transcript (Fig. 9 B). At all ages, except the adult, BEHAB expression is greater in the spinal cord than cortex. Like the cortex, BEHAB is present in the spinal cord at E17 and gradually increases with age, but reaches maximal levels by P14. Unlike the cortex, BEHAB expression in the spinal cord then declines slightly (Fig. 9 C).

The expression of BEHAB in the embryo, like in the postnatal animal, is restricted to the CNS (Fig. 10 A). BEHAB expression is absent in dorsal root ganglia (DRG), a peripheral nervous system structure (Fig. 10, B and C). Tissues in the embryo that express high levels of closely related genes, such as the cartilage (which expresses aggrecan), also show no hybridization signal for BEHAB (Fig. 10, B and C). The distribution of BEHAB expression in the embryonic CNS differs slightly from the postnatal brain. The highest levels of BEHAB expression are found in regions that contain mitotically active cells, such as the ventricular zone of the medulla, midbrain, and spinal cord (Fig. 10 A). Expression of BEHAB is heterogeneous in the developing brain, the details of which are beyond the scope of this report and will be described in detail elsewhere (unpublished manuscript).

Discussion

BEHAB Encodes a New Member of the PTR Family of Proteins

The complete coding sequence of the BEHAB gene from rat and cat is reported here. The high degree of sequence identity between rat and cat suggests that the protein encoded by BEHAB is a functionally important element of the mammalian CNS. BEHAB shows sequence homology with a subset of HA-binding proteins, the proteoglycan tandem repeat (PTR) family, which includes aggrecan, versican, link protein, neurocan, and CD44.

Among the members of the PTR protein family, the size of the mRNA recognized by BEHAB is most similar to link protein, which has been shown to contain three disulfide-bonded loops. The first loop, formed by a single disulfide bond, has homology to proteins in the immunoglobulin superfamily (Williams and Barclay, 1988; Bonnet et al., 1986). The second and third loops, referred to as PTR 1 and PTR 2 (Perkins et al., 1989), or B and B' (Doege et al., 1990), are each formed by a pair of disulfide bonds. The PTR 1 loop is the most highly conserved of the three loops, followed by the PTR 2 loop, and then the Ig loop. The most conserved regions of BEHAB are, as for the other family members, the PTR 1 loop, and then the PTR 2 loop and the Ig loop. The molecular and histological data presented here clearly show that BEHAB is closely related to, but distinct from, all previously described members of the PTR family.

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In the embryo, as in the adult, BEHAB mRNA is restricted to the central nervous system. In situ hybridization was performed on E17 rat embryos using a 35S-labeled cRNA probe for rat clone BL.

(A) Parasagittal sections of whole embryos demonstrate that BEHAB expression is restricted to the CNS. In the forebrain, BEHAB hybridization is present in the neocortex (NC). BEHAB is also expressed in the developing brainstem and spinal cord. In the midbrain, pons and medulla, the highest levels of expression are found in regions that contain actively mitotic cells, the ventral ventricular zone of the midbrain (MB) and the medial tegmental neuroepithelium (MT) of the medulla. (B and C) Cross sections of the spinal cord photographed as an autoradiogram (B) or a negative image of the same section stained with cresyl violet (C) show in more detail the distribution of BEHAB. BEHAB hybridization is restricted to the ventral portion of the developing spinal cord, with highest levels of expression in the ventral half of the ependymal layer, a region of mitotic activity. There is no detectable BEHAB expression in the peripheral nervous system, as illustrated here for the dorsal root ganglia (DRG). This pair of photomicrographs also demonstrates that cartilage (C), which expresses aggrecan, an mRNA with 60% identity to BEHAB, does not hybridize with the BEHAB probe. The negative cresyl violet stained image was generated by projection printing directly to photographic film. Bars: (A) 1 mm; (B and C) 200 μm.

However, several lines of evidence demonstrate that BEHAB encodes a protein distinct from link protein. First, on Northern blots, at low stringency BEHAB fails to detect a message in chondrosarcoma RNA (data not shown). Second, the reported location of link protein in brain is histologically distinct from that of BEHAB (Ripellino et al., 1989). Third, we have isolated a PCR clone from cat brain that is almost identical to cartilage link protein and is significantly different from cat BEHAB. Therefore, BEHAB is unlikely to encode link protein.

Two HA-binding proteins of approximately the same size as that predicted for BEHAB have been reported in the brain, glial hyaluronate binding protein (GHAP) and hyaluronectin. BEHAB is also distinct from these neural HA-binding proteins. GHAP is a 60-kD glycoprotein isolated from human brain white matter (Perides et al., 1989). The amino acid sequence of GHAP is identical to the N-terminal portion of versican, leading to the suggestion that GHAP may be a proteolytic fragment of versican (Zimmermann and Ruoslahti, 1989). Hyaluronectin is a 68-kD glycoprotein purified from CNS white matter, for which only partial amino acid sequence is available. The NH2-terminal 20 amino acids of hyaluronectin are virtually identical to GHAP (Delpech et al., 1991). The sequence of BEHAB clearly differentiates it from GHAP and hyaluronectin. Also, in contrast to BEHAB, which is only detected in the CNS, hyaluronectin is expressed in the peripheral nervous system and in the kidney, as well as in the CNS (Bignami and Dahl, 1986; Delpech and Delpech, 1984).

The Sequence of BEHAB Predicts an Extracellular HA-binding Protein

The PTR family of HA-binding proteins can be divided into two classes, HA receptors and extracellular HA-binding proteins (Wight et al., 1991). The PTR proteins that have a transmembrane domain, CD44 and TSG-6, act as HA receptors and have only a single PTR loop. Those that lack a transmembrane domain, and are extracellular proteins, have two PTR loops. The BEHAB gene encodes both a PTR 1 and a PTR 2 loop, and lacks the COOH-terminal hydrophobic domain present in transmembrane proteins, indicating that BEHAB encodes an extracellular HA-binding protein (see Fig. 11).

A HA-binding consensus sequence has not been definitively identified. Binding to HA has been demonstrated for aggrecan (Fryer et al., 1992), versican (LeBaron et al., 1992), GHAP (Perides et al., 1991), link protein (Goetinck et al., 1987), CD44 (Miyake et al., 1990), TSG-6 (Lee et al., 1992), Tl (Iwata et al., 1993), and the Cat-301 antigen (Fryer et al., 1992). HA binding by neurocan (Rauch et al., 1992) is predicted from deduced amino acid sequence. Based on sequence conservation, it has been suggested that the HA-binding region resides in the PTR loops (Doerge et
and link protein. Predictions of the structure of the PTR proteins have been described. Those with a single PTR loop (illustrated here for CD44) and a transmembrane domain (shown at bottom right) are called HA receptors. In contrast, extracellular HA-binding proteins (illustrated here for BEHAB) lack a transmembrane domain and contain two PTR loops. Sequence analysis of BEHAB suggests that it is an extracellular HA-binding protein. Binding to HA (shown as a stippled bar) is mediated either through sequences at the tip of the PTR loops (the HAL sequence), or by clusters of basic amino acids, including an arginine (R) in PTR1. The two clusters of basic amino acids which have been shown to be critical for HA binding in CD44 are indicated with arrows. The Ig loop of the extracellular HA binding proteins may permit interaction with other Ig loop containing proteins (see text for details).

Figure 11. Schematic representation of the hypothesized interaction of PTR proteins with HA. Two types of PTR HA-binding protein have been described. Those with a single PTR loop (illustrated here for CD44) and a transmembrane domain (shown at bottom right) are called HA receptors. In contrast, extracellular HA-binding proteins (illustrated here for BEHAB) lack a transmembrane domain and contain two PTR loops. Sequence analysis of BEHAB suggests that it is an extracellular HA-binding protein. Binding to HA (shown as a stippled bar) is mediated either through sequences at the tip of the PTR loops (the HAL sequence), or by clusters of basic amino acids, including an arginine (R) in PTR1. The two clusters of basic amino acids which have been shown to be critical for HA binding in CD44 are indicated with arrows. The Ig loop of the extracellular HA binding proteins may permit interaction with other Ig loop containing proteins (see text for details).

The HAL sequence does not exclusively define proteins that bind HA, because several proteins that bind HA lack the HAL sequence, including T1 (Iwata and Carlson, 1993), CD44 (Stamenkovic et al., 1989), RHAMM (Hardwick et al., 1992), and TSG-6 (Lee et al., 1992). Data from several reports suggest that HA binding may not be conferred by a specific amino acid sequence, but rather, may be determined by a clustering of basic amino acids (Lyon, 1986; Hardingham et al., 1976): peptides that inhibit link protein binding to HA contain clusters of basic amino acids (Goetinck et al., 1987), and the HA-binding region of CD44 contains two clusters of basic amino acids (Peach et al., 1993). Replacement of basic amino acids in CD44 with alanine using site-directed mutagenesis reduces HA binding. Substitution of even a single basic residue reduces HA binding, while multiple substitutions have an additive effect. One substitution, at Arg-41, almost completely abolishes HA binding. An Arg residue is present at an equivalent position in all PTR proteins, and is also present in BEHAB (Arg-166 in Fig. 4).

These sequence analyses show that BEHAB is similar to all members of the PTR family, including CD44. BEHAB is structurally similar to the PTR proteins, contains the HAL sequence, and has a basic residue (Arg) at the position that has been shown to be critical for HA binding by CD44. These data indicate that BEHAB is a new member of the PTR family of proteins and predict that BEHAB encodes an extracellular HA-binding protein.

BEHAB Represents a Neural Specific HA-binding Protein

We show here by Northern analysis that BEHAB is expressed in the mammalian CNS, but is not expressed in liver, kidney, spleen, lung, or muscle. Many of these non-neural tissues contain high levels of HA and express previously characterized HA-binding proteins. Our developmental studies using in situ hybridization demonstrate that BEHAB expression in the embryo is, similarly, restricted to the CNS.

The distribution of BEHAB in the mature CNS demonstrated by in situ hybridization parallels that reported for HA (Fryer et al., 1992; Fryer, H. J. L., G. M. Kelly, R. Kalb, and S. Hockfield, unpublished results). Biochemical assays of HA show that the corpus callosum contains three times the concentration of HA as the cerebral cortex. Also, the brainstem and spinal cord have higher concentrations of HA than the cerebral cortex (Delpech et al., 1989). Similar to the distribution of HA, a higher level of BEHAB hybridization is detected in the corpus callosum than in the cortex, and the brainstem and spinal cord have higher hybridization signals than the cortex (Fig. 7). In the white matter, clusters of silver grains from BEHAB hybridization are found over astrocytes, which synthesize HA (Delpech et al., 1989). This colocalization provides further support for the possibility that BEHAB is a HA-binding protein in brain.

While HA is abundant in the mammalian CNS, no brain specific HA-binding protein has been reported to date. Several HA-binding proteins are expressed in the brain as well as in other tissues. Versican is expressed in endoneurial and endomysial contacts, kidney medulla, precartilagenous mesenchyma, and brain (Bignami et al., 1993). The TI antigen, which is distributed throughout the brain, is also present in the electroplax and at neuromuscular junctions (Carlson et al., 1986). CD44 is expressed in lymphocytes, epithelium, and brain (Picker et al., 1989). The HA-binding protein neurocan, isolated from brain, is not expressed in liver (Rauch et al., 1992), but the presence of neurocan in tissues that contain high levels of HA and HA-binding proteins, such as
kidney, spleen, lung and muscle, has not been investigated. BEHAB represents the first reported, brain specific HA-binding protein.

Hyaluronan is found virtually in all tissues, so the existence of a tissue-specific HA-binding protein, like BEHAB, might not have been predicted. However the high degree of cellular complexity in the mammalian CNS, as well as the heterogeneity in extracellular matrix proteins in brain (see below), might require a greater diversity in HA-binding proteins than that required in other tissues. The identification of this brain specific HA-binding protein could suggest that other tissues also express tissue- or cell-type specific HA-binding proteins.

The Developmental Regulation of BEHAB Parallels the Elaboration of the Mature CNS Extracellular Matrix

The 3.9-kb BEHAB transcript is detected in the rat CNS at embryonic day 17 and its level of expression increases over the first two postnatal weeks. Adult levels of expression are reached at P21 in the spinal cord and cortex. In the spinal cord a transient peak occurs at P14; a similar peak may also occur in the cortex between P21 and adulthood. This would be consistent with the delayed maturation of cortex relative to spinal cord.

The existence of an extracellular matrix in brain has been debated. The extracellular space in brain is electronlucent and contains a very low abundance of laminin and collagen, major components of most extracellular matrices. However, studies over the last several years have shown that some elements of extracellular matrices are found in brain, including HA, heparan sulfate proteoglycans, chondroitin sulfate proteoglycans, keratan sulfate proteoglycans, and fibronectin (Margolis and Margolis, 1993; Fryer et al., 1992; Sheppard et al., 1991; Rauch et al., 1991; Hockfield et al., 1990; Herndon and Lander, 1990; Stewart and Pearlman, 1987; Margolis et al., 1975).

The composition of the brain extracellular matrix changes over the course of development. During the embryonic period, newly born neurons migrate to their final destinations and undergo initial dendritic and axonal outgrowth. The cell motility and synaptic refinement that occurs in this period requires a relatively fluid extracellular environment, consistent with our observations of high levels of BEHAB in regions of mitotic activity. During later stages in development the mature structure and synaptic relationships of neurons are established and are maintained, in large measure, throughout the life of an animal. Over the course of the early postnatal period an insoluble extracellular matrix is elaborated in the CNS. Hyaluronan becomes increasingly associated with a water insoluble fraction of brain during the early postnatal period. Ninety percent of HA is water extractable at P7 while only 15% is water extractable in the adult (Margolis et al., 1975). Several components of this insoluble matrix, including the neuronal cell surface proteoglycan recognized by monoclonal antibody Cat-301, and a variety of other cell surface proteoglycans, are initially detected during the first postnatal week and do not reach adult levels until the third or fourth postnatal week (Maeda et al., 1992; Hockfield et al., 1990; Watanabe et al., 1989; Aquino et al., 1984; Hockfield et al., 1983; Hockfield, S., S. Zaremba, and G. Kelly, unpublished results).

While the CNS contains many extracellular matrix components, the interaction of these components needs to be explored further. The extracellular matrix of cartilage contains large insoluble aggregates, formed by the interaction of aggrecan (a chondroitin sulfate proteoglycan) and HA, which are stabilized by link protein (Neame et al., 1986). The interaction between aggrecan and link protein is mediated by the immunoglobulin loops of the two proteins. As in cartilage, chondroitin sulfate proteoglycans in brain could form large aggregates through multivalent interactions with HA and link protein or a link-like protein. The size and sequence of BEHAB are similar, but not identical, to link protein, suggesting that BEHAB could serve a link-like function in the stabilization of proteoglycan-HA aggregates in CNS extracellular matrix. The cell surface proteoglycans in the CNS appear more molecularly heterogeneous than in cartilage and are expressed on distinct populations of neurons (Bertolotto et al., 1991; Watanabe et al., 1989; Fujita et al., 1989; Hockfield and McKay, 1983). Cell-type specific localization (and possibly function) of these proteoglycans could be mediated through immunoglobulin loops. Many classes of extracellular and transmembrane neural proteins contain immunoglobulin loops, including cell adhesion molecules (e.g., N-CAM) and signal transducing molecules (e.g., tyrosine phosphatases). BEHAB could associate with other immunoglobulin loop containing proteins and participate in cell–cell or cell–matrix interactions.

There is a considerable degree of cellular diversity in the mammalian CNS. The recent identification of chondroitin sulfate proteoglycans present on different populations of neurons has shown that the brain extracellular matrix is heterogeneous and suggest that this matrix may be an important determinant of neuron specific traits. The matrix heterogeneity in the CNS could include the expression of small HA-binding proteins that might serve link-like functions. The gene described here, BEHAB, encodes a neural specific protein that, based on sequence analysis, binds HA and lacks a transmembrane domain. These data, along with the time course of expression, suggest that the BEHAB protein is an important component of the CNS extracellular matrix.

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