

Nova Regulates GABA_A Receptor γ 2 Alternative Splicing via a Distal Downstream UCAU-Rich Intronic Splicing Enhancer

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Nova is a neuron-specific RNA binding protein targeted in patients with the autoimmune disorder paraneoplastic opsoclonus-myoclonus ataxia, which is characterized by failure of inhibition of brainstem and spinal motor systems. Here, we have biochemically confirmed the observation that splicing regulation of the inhibitory GABA_A receptor γ 2 (GABA_AR γ 2) subunit pre-mRNA exon E9 is disrupted in mice lacking Nova-1. To elucidate the mechanism by which Nova-1 regulates GABA_AR γ 2 alternative splicing, we systematically screened minigenes derived from the GABA_AR γ 2 and human β -globin genes for their ability to support Nova-dependent splicing in transient transfection assays. These studies demonstrate that Nova-1 acts directly on GABA_AR γ 2 pre-mRNA to regulate E9 splicing and identify an intronic region that is necessary and sufficient for Nova-dependent enhancement of exon inclusion, which we term the NISE (Nova-dependent intronic splicing enhancer) element. The NISE element (located 80 nucleotides upstream of the splice acceptor site of the downstream exon E10) is composed of repeats of the sequence YCAU, consistent with previous studies of the mechanism by which Nova binds RNA. Mutation of these repeats abolishes binding of Nova-1 to the RNA in vitro and Nova-dependent splicing regulation in vivo. These data provide a molecular basis for understanding Nova regulation of GABA_AR γ 2 alternative splicing and suggest that general dysregulation of Nova's splicing enhancer function may underlie the neurologic defects seen in Nova's absence.

The regulation of eukaryotic gene expression involves an ordered sequence of events within various subcellular compartments. RNA binding proteins (RBPs) play a critical role in this process, acting at multiple levels to regulate and diversify gene expression, and such functions are likely to be of particular importance to neurons. RBPs regulate processes such as alternative splicing, which is an important mechanism for generating complexity in specialized cells, particularly in neurons. For example, all of the major neurotransmitter receptors contain subunits that are alternatively spliced, and these variants can alter receptor localization, ligand binding, signal transduction, and electrophysiological properties (13, 20, 44, 51). Moreover, RBPs and aberrant splicing of mRNAs encoding proteins critical for proper functioning of neurons have been associated with a number of neurological diseases (reviewed in reference 19). To understand the role RBPs play in regulating gene expression in neurons, it would be necessary to identify their complement of RNA targets and to understand how they are recognized amidst the complexity of the cellular environment.

Regulation of alternative splicing occurs through recognition of *cis*-acting sequences in pre-mRNA by *trans*-acting factors. *cis*-acting repressor elements that mediate the exclusion of the neuron-specific exons in nonneuronal tissues have been identified through studies of neuron-specific splicing in transcripts such as *c-src*, the GABA_A receptor γ 2 (GABA_AR γ 2) subunit, the *N*-methyl-D-aspartate receptor NR1 subunit, and clathrin light-chain B mRNAs (3, 9, 10, 60, 63, 64). These repressor sequences are mostly pyrimidine rich and, when

present *in trans* as RNA competitors in vitro, stimulate neuronal splicing patterns (10, 64). A number of generally expressed *trans*-acting factors, including hnRNP H, hnRNP F, KSRP, and PTB (3, 10, 12, 46), have been found to bind such sequences and are believed to play important roles in regulating alternative splicing.

In addition to the contribution of generally expressed factors, it has long been suspected that tissue-specific factors regulate alternative splicing in the brain (1). Study of the paraneoplastic neurologic disorders (PNDs) has resulted in the identification of Nova-1 (7), a neuron-specific protein involved in splicing regulation (32). PNDs are rare immune-mediated disorders associated with ectopic expression of neuron-specific proteins in tumors (reviewed in references 15 and 48). Patients harbor characteristic high titers of antibodies specific for these antigens in their sera, and immune targeting of these antigens in the central nervous system (CNS) is thought to result in the neuronal death and neurological symptoms seen in PNDs. Nova proteins are the target antigens in paraneoplastic opsoclonus-myoclonus ataxia, a PND characterized by loss of inhibitory control of motor neurons in the spinal cord and brainstem.

The Nova proteins are RBPs containing 3 KH-type (for hnRNP K homology) RNA binding domains. Widely expressed KH domain-containing proteins have been shown to play roles in a number of aspects of RNA processing, including mRNA stability and translational control (hnRNP K and hnRNP E1/E2) (35, 49), subcellular localization of RNAs to specific sites in the cell cytoplasm (ZBP-1 and Vera) (17, 55), and RNA splicing (KSRP [46] and SF1 [2] in mammals, PSI [57] in *Drosophila*, and MER-1 [21] in *Saccharomyces cerevisiae*). Nova-1 is expressed exclusively in neurons of the CNS, including

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the brainstem and spinal motor neurons targeted in paraneoplastic opsoclonus-myoclonus ataxia. Nova-1 null mice display a severe phenotype. Although they appear normal at birth, they are markedly smaller than their wild-type littermates at as little as 3 to 4 days postbirth and display symptoms of motor impairment such as motor weakness and an action-induced tremor (32).

Nova proteins act as sequence-specific RNA binding proteins. A consensus binding sequence determined by *in vitro* RNA selection consists of a stem-loop RNA containing (UCAUY)₃ in the loop region (6). This binding is sequence specific, as a number of discrete nucleotide substitutions, such as mutagenesis of this core to (UCUUY)₃, abolish RNA binding. Studies using the third Nova KH domain in isolation identified high-affinity targets containing a single UCAY element, again within a hairpin loop context (33), and allowed solution of the X-ray structure of a Nova KH3/RNA cocrystal. The crystal structure reveals that the RNA stem-loop presents the UCAY sequence to a "molecular vise" formed by a cleft between the variable and invariant protein loops of the Nova KH3 domain (42). In particular, the CA dinucleotide forms contacts with the protein to create pseudo-Watson-Crick base pairs and predicts that these residues are absolutely necessary for binding to occur. The structure agrees precisely with Nova KH3 domain-RNA binding studies in which RNA mutations still conforming to the consensus HYCAY (where H denotes A, C, or U and Y denotes C or U) are tolerated (33).

These studies led to the identification of GlyR α 2 inhibitory neurotransmitter receptor subunit pre-mRNA as a Nova target RNA. Nova binds this transcript 80 nucleotides (nt) upstream of an alternatively spliced exon, E3A, which is spliced in a mutually exclusive manner with a downstream exon, E3B. These exons encode part of the extracellular domain of the receptor and thus may be involved in ligand binding (39). Nova protein in mouse brain lysates coimmunoprecipitates with GlyR α 2 pre-mRNA (6) and can be UV cross-linked to an RNA encoding a region of the GlyR α 2 intronic sequence (32). Cotransfection of GlyR α 2 minigene constructs with a Nova-1 expression plasmid demonstrated that Nova-1 is able to act directly on GlyR α 2 pre-mRNA to enhance the inclusion of GlyR α 2 E3A and that this action is dependent on the integrity of the intronic (UCAU)₃ Nova binding site (32).

Analysis of GlyR α 2 alternative splicing *in vivo* provided genetic support for these conclusions, as Nova-1 null mice showed a decrease in utilization of GlyR α 2 E3A relative to wild-type littermates (32). This defect in alternative splicing regulation was restricted to regions that normally express Nova-1 and was specific. Regulation of a number of alternatively spliced RNAs was unaffected in Nova-1 null mice, with one notable exception: alternative splicing of exon 9 in the inhibitory neurotransmitter receptor subunit pre-mRNA, GABA_A-R γ 2, was disrupted in Nova-1 null mice in a dose-dependent manner, and these defects were again restricted to regions which normally express Nova-1.

Alternative splicing of GABA_A-R γ 2 cassette exon E9 generates two mRNA isoforms, γ 2L (long [E9 included]) and γ 2S (short), which show differential spatial and temporal expression characteristics (24, 27, 59). Inclusion of E9 adds 8 amino acids to an intracellular loop of the protein which contain a consensus site for phosphorylation by protein kinase C (37,

61). Phosphorylation of this site has been shown to modulate GABA_A-R function (38). Mice engineered to express only the short form of the GABA_A-R γ 2 subunit display a higher level of anxiety than control mice (30) and increased sensitivity to benzodiazepines (53). Studies aimed at trying to understand the mechanism behind regulation of GABA_A-R γ 2 splicing have shown that nucleotides within the exon and the adjacent splice sites are essential for regulation in a neuronal versus a non-neuronal cell line (63). PTB has been shown to inhibit γ 2L splicing, leading to the hypothesis that splicing is regulated by a mechanism involving derepression of splicing at this site in neurons of the CNS (3, 64). However, it is unknown whether these observations are relevant to γ 2L splicing in neurons and/or whether they involve Nova protein.

To understand how Nova acts to regulate GABA_A-R γ 2 splicing, we have determined the sequence requirements for Nova-1 regulation of E9 inclusion. We find that previously mapped elements important for the regulation of splicing in tissue culture are not involved in Nova regulation of γ 2L. Remarkably, a broad and unbiased screen for regulatory elements over 3.7 kb of GABA_A-R γ 2 pre-mRNA revealed that a single intronic YCAY repeat element is necessary and sufficient to mediate Nova-dependent regulation. These studies define Nova as a neuron-specific splicing regulator that functions through action on a newly identified intronic splicing enhancer which lies far downstream of the regulated exon.

MATERIALS AND METHODS

RNA preparation, RNase protection assays, and RT-PCR. Using a modified guanidine-acid phenol protocol (11), total RNA was extracted from mouse tissues. RNA from tissue culture cells was purified using the RNeasy Mini kit system (Qiagen). RNase protection assays were performed using the RPA III kit (Ambion) according to the manufacturer's instructions and 5 μ g of total RNA isolated from mouse spinal cords. Hybridization was performed at 56°C, and digests were performed with RNase T1. Using T7 RNA polymerase from gel-purified PCR products generated with primers gaba-rpa-F (5'-GAAATTAATACGACTCACT ATAGGGAGTTGAATGGTTGCTGATCTGGGACG), gaba-rpa-R (5'-TC CCGCTCGTCTGCTGTTATGGCACCCCTGCATTATTTTGTC), cla-rpa-F (5'-GAAATTAATACGACTCACTATAGGGAGGGGGTCTCCTCTGG ATTCT), and cla-rpa-R (5'-TCCCGCTCGTCTGACCGAACAGGAG TGGCGGGAG) or from plasmid pTRI- β -Actin-Mouse antisense control template (Ambion), antisense probes were *in vitro* transcribed. Probes were labeled by incorporation of [α -³²P]UTP, and specific activities were adjusted by incorporation of unlabeled UTP to give protected fragments of approximately equal intensities for each message. Reverse transcription-PCR (RT-PCR) was performed as previously described (Jensen et al. [32]). PCR primers to human β -globin were E1F (5'-CTGAGGAGAAGTCTGCCGTTA CTG) and E3R (5'-CAGCACACAGACCAGCAGTTG).

Plasmid constructs. The GABA_A-R γ 2 minigene containing full-length introns was generated by long-range PCR using primers γ 2 GABA F and γ 2 GABA R (32) and a subclone derived from mouse BAC 608F8 (Research Genetics). The product was cloned into pGem-T-Easy (Promega) and transferred to pcDNA3 (Invitrogen) via *Eco*RI. Mutations of minigene constructs were made with a QuikChange site-directed mutagenesis kit (Stratagene) and specifically designed primers (Operon).

Using plasmid sp64-H β Δ 6 (a kind gift from A. R. Krainer) as the template, the β -globin minigene was generated by PCR using primers glo-KpnF (5'-ATGGT ACCTGACTCCTGAGGAGAAG) and glo-Rc (5'-CCACTTTCTGATAGGC AGCC) and cloned into pcDNA3 (Invitrogen) via *Kpn*I and *Eco*RI. Exon 2 was shortened by PCR. Chimeric minigene constructs were generated by PCR.

cDNAs encoding T7-tagged Nova-1 including alternative exon H and T7-tagged hnRNP E1 were cloned into pCI-neo (Promega). All DNA constructs generated by PCR or site-directed mutagenesis were sequenced in their entirety. Plasmid DNA for transfection into cell lines was prepared by a modified cesium chloride method (56) and dialyzed with 1 \times Tris-EDTA buffer.

Cell transfection. Using Fugene6 (Roche) as described by the manufacturer, N2A and 293T cells (American Type Culture Collection) were transfected with 2.25 μ g of total DNA in a mixture composed of 0.25 μ g of the appropriate minigene and variable amounts of pNova-1 and empty pcneo vector. After 40 h, the cells were washed with 1 \times phosphate-buffered saline, collected by scraping, and halved for RNA extraction and protein extraction. Western blotting was performed using horseradish peroxidase-conjugated anti-T7 tag monoclonal antibody (Novagen) and detected by chemiluminescence (NEN).

Fusion protein synthesis. Full-length recombinant T7/His-tagged Nova-1 fusion protein was produced and purified as previously described (41). Full-length Nova-1 cDNA including alternative exon H was cloned into pET21a (Novagen) and transformed into BL21-competent cells, followed by standard IPTG (isopropyl- β -D-thiogalactopyranoside) induction and purification by nickel chelation chromatography.

Nitrocellulose filter binding assays and boundary mapping. Labeled RNA (100 to 200 fmol) was incubated with the indicated concentrations of protein. Protein-RNA mixes were incubated in a total volume of 50 μ l of 1 \times BB (200 mM KOAc, 10 mM TrisOAc [pH 7.7], 1 mM MgOAc, 10 mM dithiothreitol, 1 mg of heparin/ml) for 15 min at room temperature. Binding solutions were passed through nitrocellulose filters (0.45- μ m pore size; Millipore) and washed with 5 ml of 1 \times BB without dithiothreitol or heparin. Data were plotted as the percentage of total bound RNA versus the log of the protein concentration, and $K_{d,s}$ were determined with Kaleidograph software (Synergy Software). SB2 RNA was synthesized as previously described (6). GABA wild-type and mutant RNAs were transcribed using T7 RNA polymerase and gel-purified PCR templates. Glo RNA was transcribed using T7 RNA polymerase directly from minigene pGlo2 Δ digested with *Bbs*I to yield a 175-nt RNA. Boundary mapping was performed essentially as described previously (14), with binding conditions identical to those used for nitrocellulose filter binding assays.

RESULTS

Splicing defects in Nova-1 null mice. To confirm and extend the previous analysis of Nova-1 null mice by Jensen et al. (32), RNase protection assays were performed to assess the relative abundance of GABA_AR γ 2-spliced mRNAs. RNA isolated from the spinal cord of Nova-1 null mice displayed a decrease of roughly 2.5-fold in the ratio of E9-inclusive (γ 2L) to E9-exclusive (γ 2S) splice forms of GABA_AR γ 2 (Fig. 1 B and C). Moreover, no difference was detected in the total abundance of GABA_AR γ 2 mRNA when compared to that of actin (Fig. 1D), indicating that the decrease in γ 2L is accompanied by a complementary increase in the γ 2S spliced form. As a control for the specificity of Nova action, we examined the alternative splicing and steady-state levels of clathrin light chain B (Cl_B), which were found to be unchanged in the Nova-1 null mice, as expected (Fig. 1B to D). These results demonstrate specific, Nova-dependent alterations in alternative splicing, but not stability, of the γ 2L isoform of GABA_AR γ 2 pre-mRNA in the CNS of mice.

Nova regulates GABA_AR γ 2 alternative splicing in cell lines. To assess the sequence requirements necessary for Nova-1 action on GABA_AR γ 2 alternative splicing, a minigene construct was generated which contains the entire mouse genomic region surrounding the alternatively spliced exon, E9, extending into the adjacent exons, E8 and E10 (pGABA; Fig. 2A). This construct was transiently transfected into N2A or 293T cells in the presence or absence of pNova-1 plasmid encoding full-length Nova-1 protein, and spliced products were analyzed by semi-quantitative RT-PCR. Consistent with the analysis of GABA_AR γ 2L splicing in Nova-1 null mice, cotransfection of pNova-1 caused a dose-dependent increase in E9 inclusion, resulting in an up to fivefold increase in the ratio of E9-inclusive (γ 2L) to E9-exclusive (γ 2S) splice forms in 293T cells (Fig. 2B). Similar results were seen with a mouse neuroblas-

toma cell line, N2A, for which a fourfold increase in exon 9 utilization was observed (Fig. 2C). No products were detected in control cells (Fig. 2BI and CI, lanes M), indicating that spliced products were derived from pGABA. Notably, the baseline level of E9 inclusion without the addition of exogenous Nova-1 was approximately 10-fold higher in N2A cells than in 293T cells, perhaps reflecting the fact that N2A cells express endogenous Nova proteins whereas 293T cells do not (data not shown).

In contrast to the action of Nova-1, addition of exogenous hnRNP E1 caused a modest decrease in the utilization of GABA_AR γ 2 E9 (Fig. 2B and C). Western blotting and immunofluorescence microscopy confirmed that using the plasmid concentrations indicated, comparable levels of transfected Nova-1 and hnRNP E1 were expressed and both proteins were localized to the nucleus (Fig. 2BII and CII and data not shown). Since hnRNP E1 and Nova contain three KH-type RNA binding domains whose sequences and spacings are similar, these results are consistent with the hypothesis that sequence-specific interactions between Nova-1 and GABA_AR γ 2 RNA are necessary for splicing regulation.

Defining the region of GABA_AR γ 2 pre-mRNA necessary for Nova-1 regulation. Genomic sequences surrounding GABA_AR γ 2 E9 from humans and mice were examined to identify putative conserved Nova-1 binding sites. Within approximately 250 nt on either side of E9, only two YCAY elements conserved between humans and mice were found, one upstream of E9 in intron 8 and one downstream in intron 9. These elements plus an adjacent CA dinucleotide were mutated (Fig. 3), and the resulting pGABA mutants were assessed for Nova-1-dependent splicing. No changes in E9 splicing were seen upon mutation of intron 8 in 293T or N2A cells (Fig. 3A and Fig. 3B, panel m8). In contrast, a 5- or 15-fold decrease in E9 splicing was seen upon mutation of intron 9 in 293T cells and N2A cells, respectively, reducing γ 2L to barely detectable levels in the absence of pNova-1. Despite this, cotransfection of pNova-1 led to an increase in E9 inclusion in both cell lines (Fig. 3B and C, panels m9), although quantitation of these effects in 293T was difficult due to very low abundance of the γ 2L species. A construct harboring both mutations (Fig. 3B and C, panel m8+9) behaved in a manner identical to that of the intron 9 mutation, indicating that the two mutated sites do not act synergistically. We conclude that neither the intron 8 or 9 mutation disrupts the major site of Nova-1 action. Although we have not further explored the effect of the intron 9 mutation, its action on basal E9 splicing might be due to disruption of a TGCATG element, a common component of downstream intronic enhancer elements. Various numbers of repeats of this sequence have been shown to be important for alternative splicing of fibronectin (31), fibroblast growth factor receptor 2 (16, 43), *src* (47), calcitonin/CGRP (28), nonmuscle myosin II heavy-chain B (34), and possibly AMP deaminase (22).

To identify sites capable of mediating Nova-dependent GABA_AR γ 2 E9 inclusion, we used exonuclease III to generate pGABA deletion constructs harboring introns of differing lengths. These experiments were used as an initial means to screen the 3.7 kb of intronic sequence for potential Nova-dependent elements, bearing in mind that introns are known to harbor widely scattered elements that might affect splicing (43). Screening of over 20 constructs in transfected N2A cells

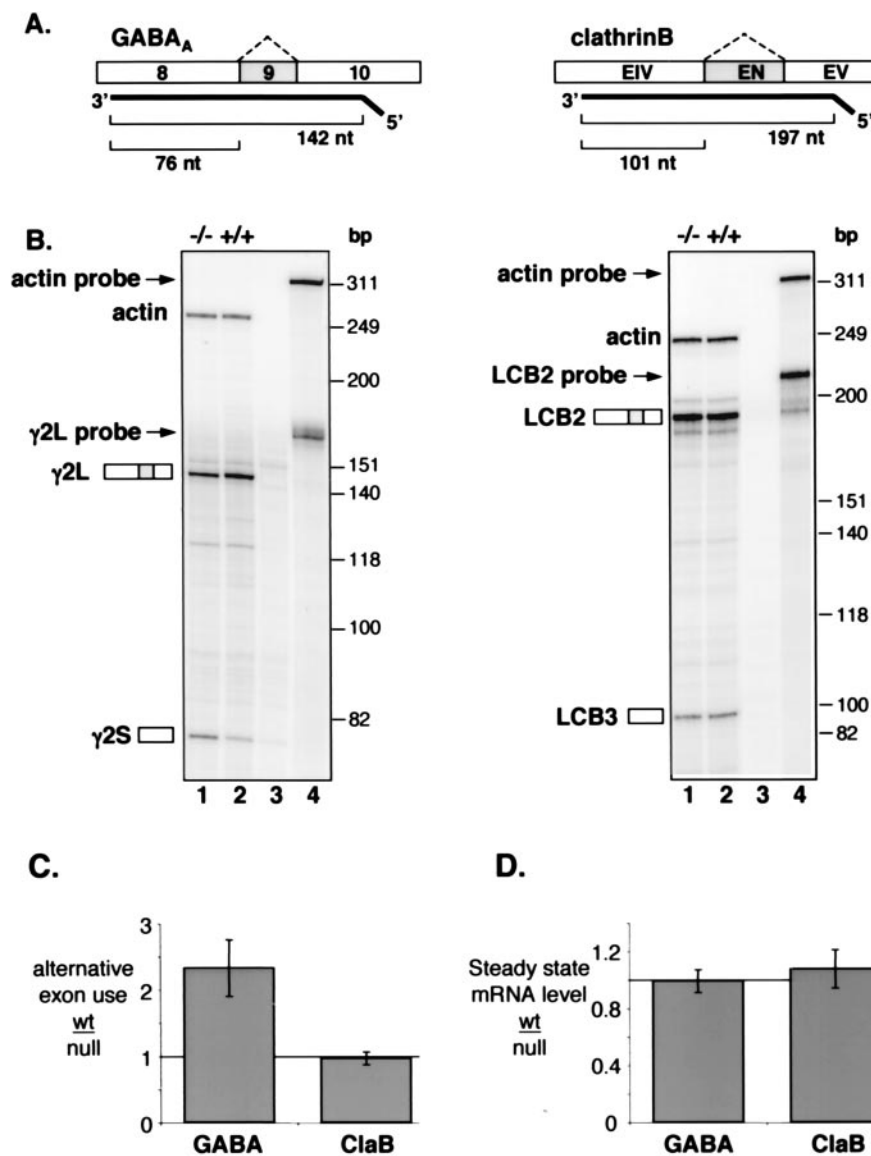


FIG. 1. Changes in alternative splicing but not steady-state levels of $GABA_A\gamma 2$ transcripts in Nova-1 null mice. (A) Schematic representation of body-labeled antisense RNA probes used for RNase protection assays and the resulting product sizes used for quantitation. The cassette exons shown in grey are alternatively spliced, producing $\gamma 2L$ (contains exon 9) and $\gamma 2S$ (lacks exon 9) forms of the $GABA_A\gamma 2$ mRNA and LCB2 (contains exon EN) and LCB3 (lacks exon EN) forms of the *ClaB* mRNA (60). An antisense RNA probe specific for β -actin mRNA was also made. (B) Spinal cord RNA isolated from P9 ($GABA_A\gamma 2$ analysis) and P11 (*ClaB*) Nova-1 null mice ($-/-$) and their wild-type ($+/+$) littermates was analyzed by an RNase protection assay. Lanes 3, yeast RNA controls; lanes 4, undigested probes. (C and D) Quantitation of the data presented in panel B together with additional RNA samples ($n = 5$ littermate pairs). Products were quantitated by phosphorimager and corrected for the expected number of ^{32}P -labeled uridines. (C) The ratio of long to short alternatively spliced products measured in control animals (wt) relative to the respective ratio for the corresponding Nova-1 null littermate. (D) For steady-state comparisons, the total of both alternatively spliced products was normalized to the amount of actin message detected in the same lane and compared between control animals (wt) and the corresponding Nova-1 null littermate. The solid bars represent the average changes in wild-type–null exon usage (C) or mRNA level (D); error bars represent standard deviations.

demonstrated that most of introns 8 and 9 can be deleted without eliminating Nova-dependent splicing regulation (data not shown). Using this deletion data, we generated a truncated $GABA_A\gamma 2$ minigene, pGABA Δ , derived from the shortest intron 8 and intron 9 deletions that retained Nova-responsive E9 splicing. pGABA Δ recapitulated the Nova-dependent inclusion of E9 seen with the full-length construct, showing a 6- to 14-fold increase in E9 utilization upon the addition of No-

va-1 in N2A and 293T cells, respectively, and a lower level of basal E9 inclusion in 293T cells compared to that in N2A cells (Fig. 4A). Taken together, these data indicate that the size of introns 8 and 9, and most of their sequence content, is not critical for Nova-dependent regulation of $GABA_A\gamma 2$ E9 splicing.

To further delineate Nova-responsive elements, a heterologous minigene was constructed using the human β -globin gene

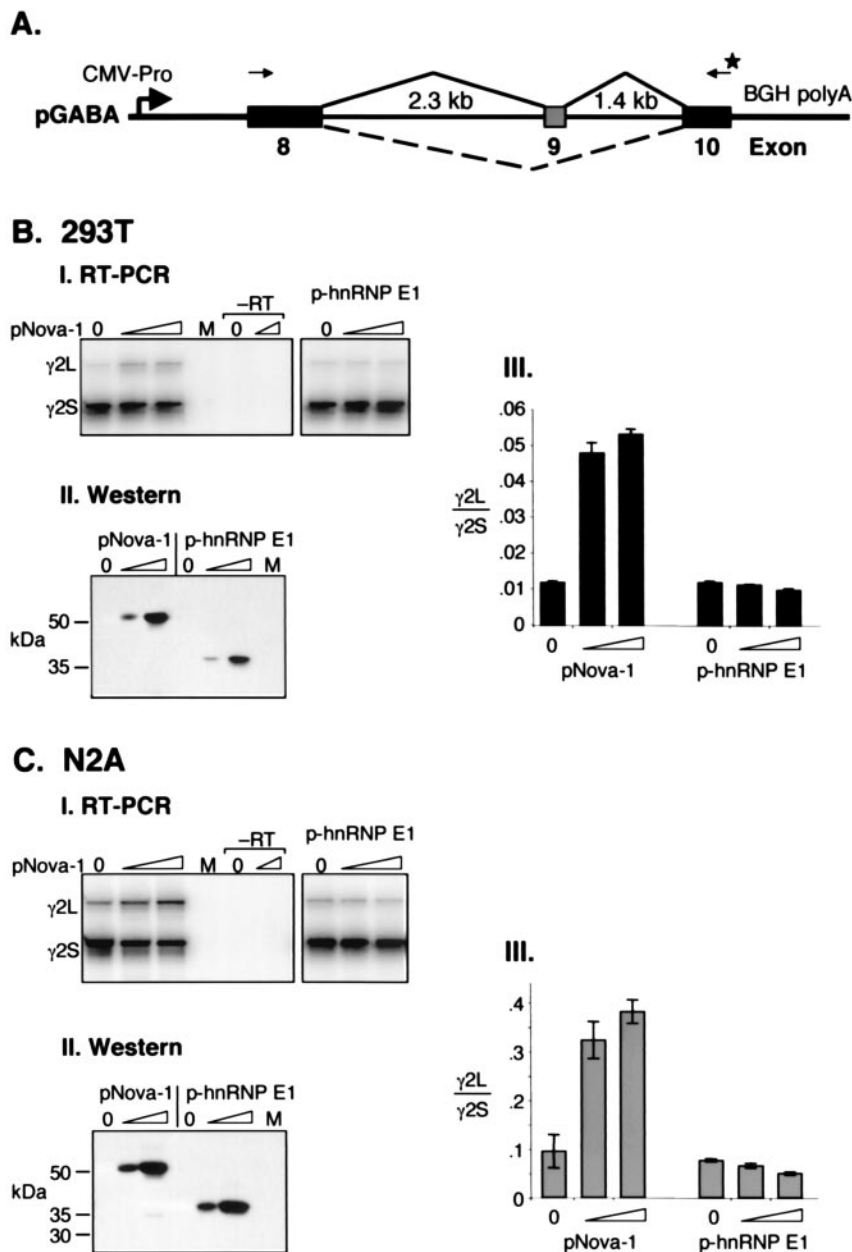


FIG. 2. GABA_AR γ 2L splicing is enhanced specifically by Nova-1 in heterologous cell lines. (A) Schematic representation of the GABA_AR γ 2 (pGABA) minigene containing the full mouse intronic regions surrounding and including exon 9 plus shortened exons 8 and 10 constructed for transient transfection assays. The primers used for RT-PCR analysis are depicted as arrows; the star denotes the ³²P-labeled primer. Figure is not to scale. (B and C) RNA was isolated from 293T (B) or N2A (C) cells transiently transfected with pGABA and increasing amounts of pNova-1 (0, 0.5, or 2.0 μ g) or p-hnRNP E1 (0, 0.05, or 0.2 μ g) mammalian expression plasmid and analyzed by RT-PCR. Mock transfections were performed using no plasmid DNA (lanes M). (Panels I) RT-PCRs were performed in duplicate using controls without RT, representatives of which are shown (-RT). (Panels II) Western blot using anti-T7-tagged antibody showing titration of T7-tagged Nova-1 and hnRNP E1 protein levels after transfection. (Panels III) Quantitation of the data presented in panels I plus two additional independent transfections (*n* = 3). The bars represent the ratios of γ 2L/ γ 2S spliced products \pm standard deviations.

as a backbone into which elements from pGABA Δ can be inserted (Fig. 4A). Since exon size can directly influence splicing efficiency (4, 18), the central region of β -globin exon 2 was deleted to give a central exon size of 28 nt (pGlo2 Δ) (Fig. 4A), making it comparable in size to GABA_AR γ 2 E9 (24 nt). pGlo2 Δ showed a level of alternative exon inclusion (29%

inclusion of exon E2 Δ) similar to that seen with pGABA Δ (24% inclusion of exon E9) in N2A cells, although the exon was less well utilized in 293T cells (9% inclusion of exon E2 Δ). However, pGlo2 Δ was not regulated upon the addition of pNova-1 in either cell line (Fig. 4A).

A chimeric construct in which 121 nt (surrounding and in-

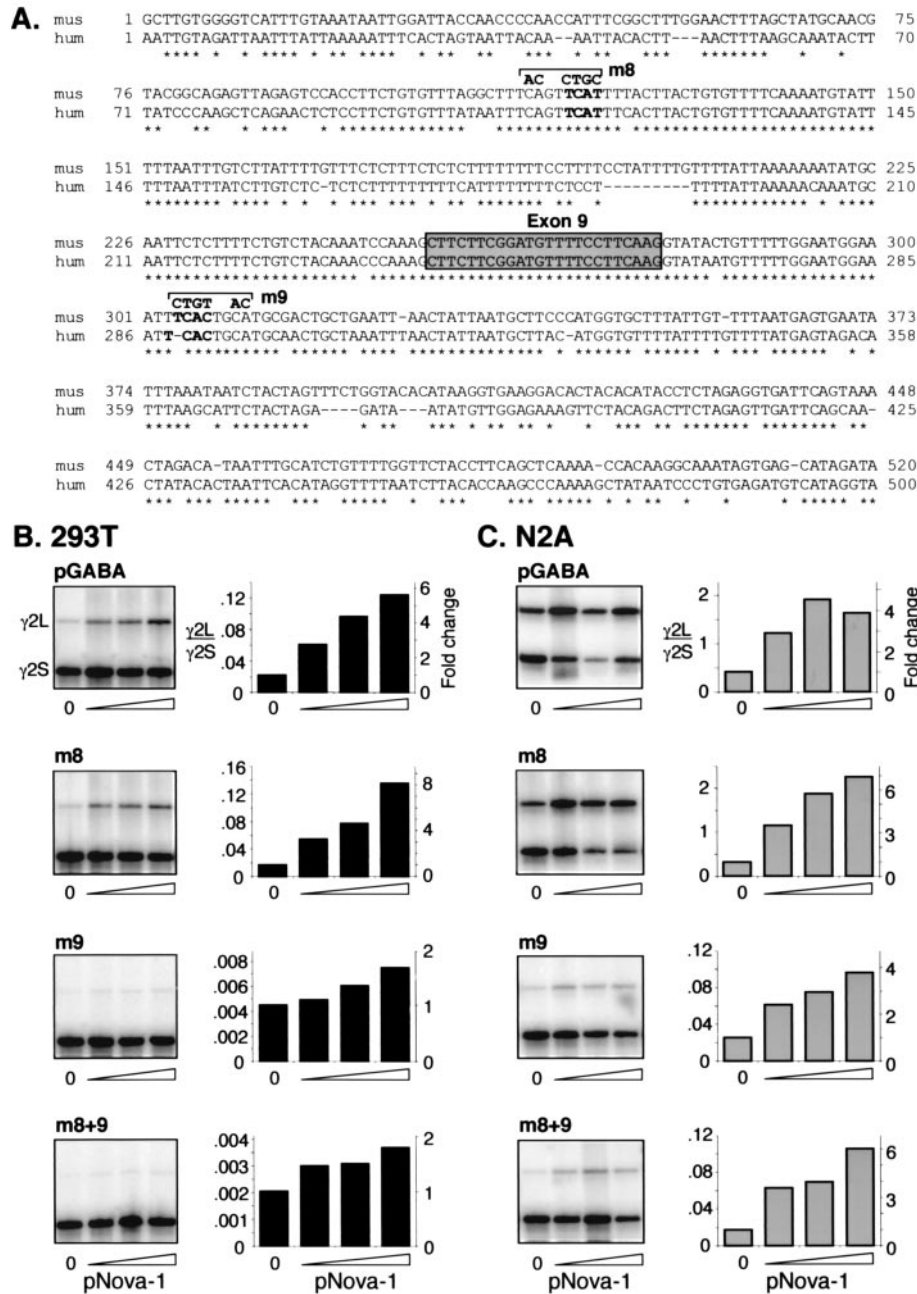


FIG. 3. Mutations within the adjacent intronic regions do not interfere with Nova's ability to regulate GABA_ARγ2 exon 9 alternative splicing. (A) Sequence comparisons of intronic regions surrounding GABA_ARγ2 exon 9. Genomic sequences surrounding and encompassing GABA_ARγ2 E9 (gray box) from human (hum) and mouse (mus) were aligned by ClustalW alignment (MacVector). Conserved YCAY elements are shown in boldface characters. m8 and m9 represent mutations made in the mouse minigene within introns 8 and 9, respectively. (B and C) The mutations depicted in panel A were made in pGABA, either individually or in combination, and the resulting minigenes were cotransfected with increasing amounts of pNova-1 (0, 0.5, 1.0, or 2.0 μg) into 293T (B) or N2A (C) cells. Spliced products were analyzed as described for Fig. 2. The graphs represent quantitation of the autoradiograph data shown in the left column of each panel; the left y axes indicate the ratios of γ2L/γ2S spliced products, and the right y axes indicate the severalfold changes in these ratios upon addition of pNova-1. Titration of transfected pNova-1 expression was monitored by Western blotting (data not shown). Similar results were obtained in at least two independent transfection experiments.

cluding E9) were inserted into pGlo2Δ was not able to mediate Nova-dependent splicing (Fig. 4Ba). In addition, inclusion of additional GABA_ARγ2 sequence derived from E8 and its downstream intron was not sufficient to regain Nova-dependent alternative splicing of E9 (Fig. 4Bb). However, inclusion

of the region derived from the constitutively spliced exon E10 and its upstream intron did restore Nova-dependent splicing regulation, indicating that this region is necessary for Nova's effect (Fig. 4Bc). Moreover, the 211-nt sequence upstream of exon E10 was itself able to mediate Nova-dependent regula-

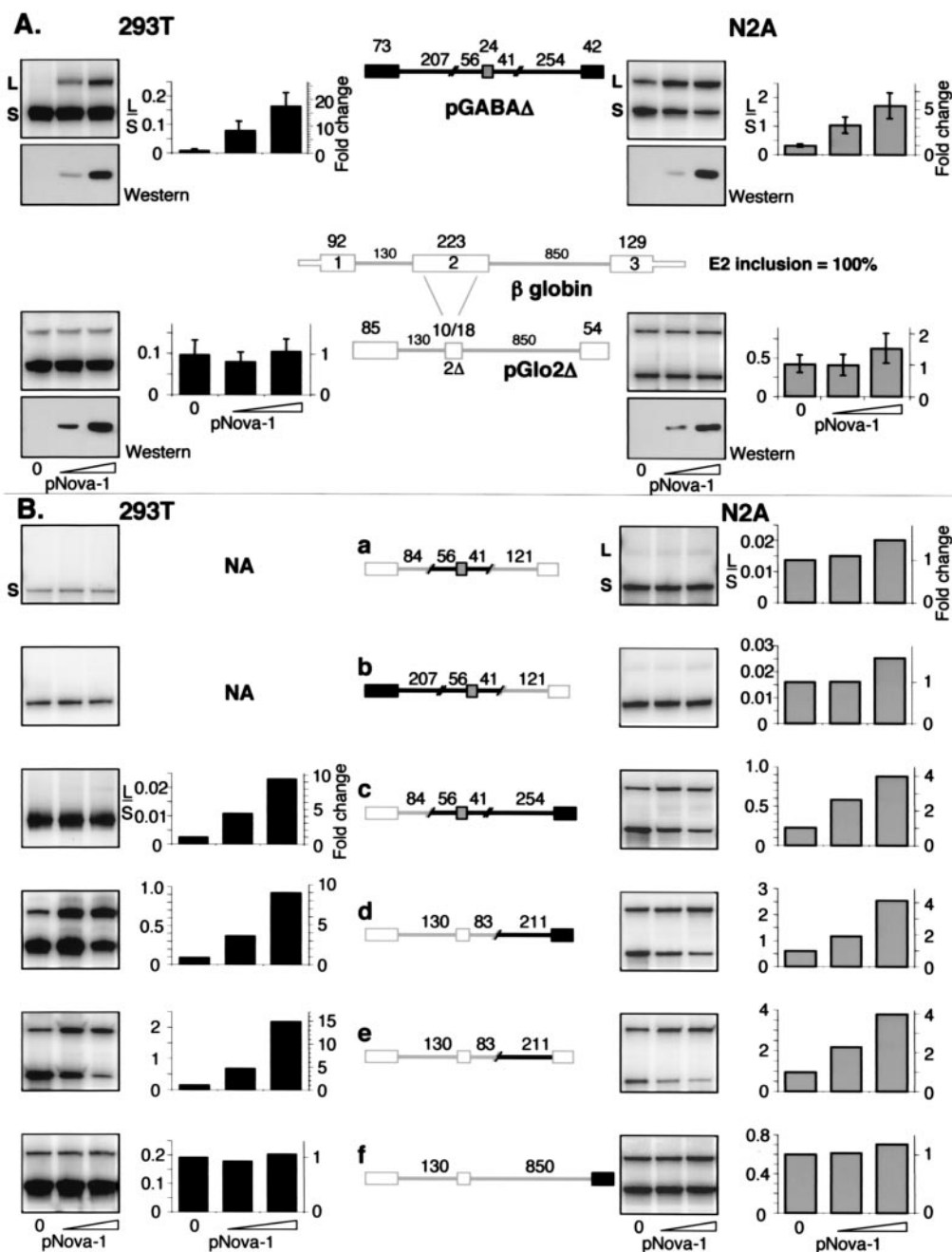


FIG. 4. The intronic sequence upstream of GABA_AR γ 2 E10 is sufficient for regulation of alternative splicing by Nova-1. (A) A truncated GABA_AR γ 2 minigene (pGABA Δ) was generated by fusing intron deletion constructs (data not shown). A β -globin minigene (pGlo2 Δ) was derived from the human gene, with the central region of exon 2 deleted such that exon 2 Δ retains 10 bp 5' and 18 bp 3' of the deletion site. The sizes (in base pairs) of the exonic and intronic sequences contained in the minigenes are noted above the schematics. Minigenes were cotransfected into 293T or N2A cells with increasing amounts of pNova-1 (0, 0.5, or 2.0 μ g), and spliced products were measured by RT-PCR and phosphorimage analysis as described for Fig. 2. Titration of transfected pNova-1 expression was monitored by Western blotting using α -T7-tagged antibody (lower panel of each pair of panels). Quantitation of the data plus that of additional experiments ($n = 4$ for pGABA Δ , $n = 5$ for pGlo2 Δ) is shown at the right side of each pair of panels. The bars represent L/S exon use ratios \pm the standard deviations. (B) Chimeric constructs were made by ligating portions of the GABA_AR γ 2 (black lines and filled boxes) and β -globin (gray lines and empty boxes) constructs shown in panel A. Minigenes were cotransfected into 293T or N2A cells with increasing amounts of pNova-1 (0, 0.5, or 2.0 μ g), and spliced products were measured by RT-PCR using primers specific for the outside exons and phosphorimage analysis as described for Fig. 2. The graphs represent quantitation of the autoradiograph data shown in the left column of each panel; the left y axes indicate the ratios of L/S spliced products, and the right y axes indicate the severalfold changes in these ratios upon addition of pNova-1. NA, not applicable (no long form detected). Titration of transfected pNova-1 expression was monitored by Western blotting (data not shown). Similar results were obtained in at least two independent transfection experiments.

A.

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mus 1 TTTTGATACATAAAATAAATTTTTGTGCATAAAAATCAACGAACCTCTTTATTTCTCCATGAGTACTAGCATTGTGCTACTAAAATTTGTCAGTATAGTTG 100
hum 1 CTTTAAAACATAAGCCATACCATTGGGTATACAATCAGGAAAATTCGTGATTAT--ATATATTTAACATTTCTATTAAACATT---TAATACATTG 95
    * * * * *

mus 101 TTCAGTATGATGGGAC--CTATTTGCCTCTAAGTTAGCATGACATGCAATTTGATTGAAAATCAATGCATTTTATTTCTCTATTTTACTCATTGGCAG 198
hum 96 CTCAGAGTAATGGGTTTTCTATTTGCCTATAACTAAGCATGACACAATATTTGATTTCAGAATTGATGCATTTTATTTCTCTGTCCACTCACATGCAG 195
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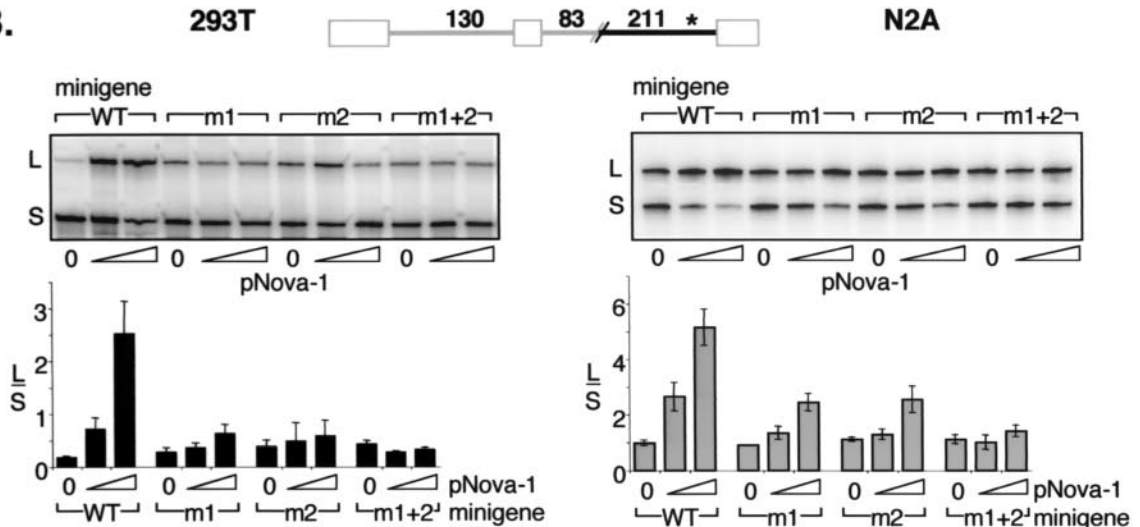
mus 199 CATCATTAAGTACTGCTTCATGTAAGTGGCTTAACATTTAATAAATTACTAGAATGAGATTCCATGTAGTTTGATTAAATCTTTATACCTTTGATTCTTTTG 298
hum 196 CACCATCAAAATGCCTTCATGTAACCAATGTCACATCTAATACTTACTAGAATAAGATTCCATGTAATTTGATTAAATATTA-ACCTTGAGTCTTTTG 294
    * * * * *

mus 299 GTGGTCAATAGTCTTCTGCTGTAGCTATTAGCTTGATGACATTATTTCCAGCTGCACTGCTTAAGCGCAAAAATCTGAAACCTGCAACTGTGCTATCTTTCTG 398
hum 295 GTGTTCAATCTCCTTGTCTATAGCTATTAGCTTGATGATATTATTTCCAGCTGCACTGCTTAAGCTCAAAAATTTGAAATCTGCAAAATGTGCTATCTTTCTA 394
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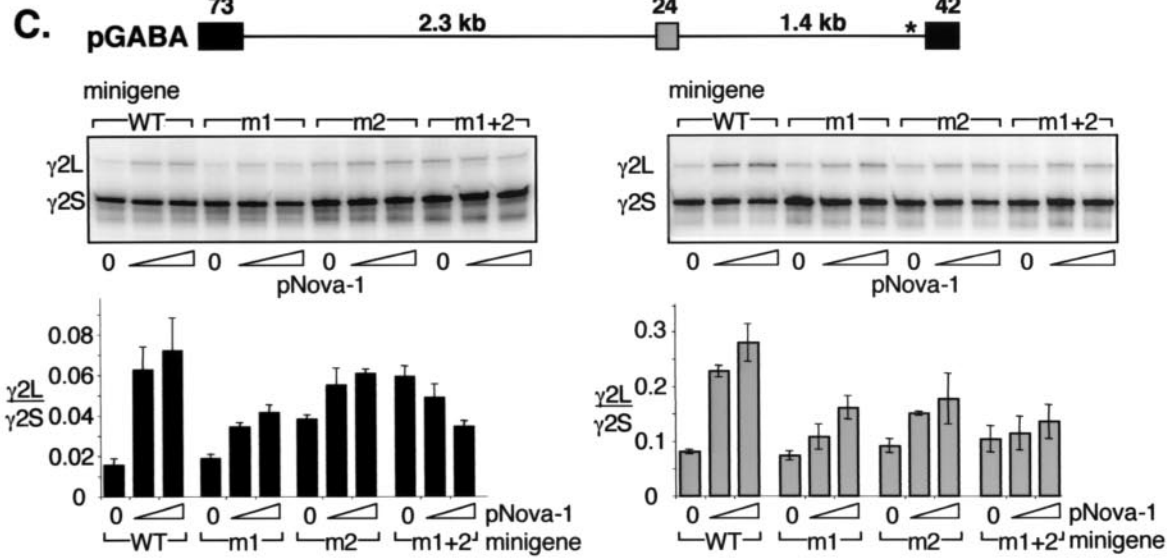
mus 399 AGTTCAATTTTACCATCGTAAGATCACGATGTCATAAGCAATTTTCTGAATACTCATTTTCAGATTCATCATTCATTTGATGACATTGTGGTAAGTACTGT 498
hum 395 AGTTCAATTTTACCATGTA-GATCATGATGTCATA-GCAATTTCTGAGTACCCATTTTCAGATTCATCATCACATTGGTGACATTGTGGTAAACACGC 492
    * * * * *

mus 499 GTTATATAGCTCCAGTATAACTAACTTTGCTCTCTCTCTTACCTACCTT-GTCTCAGAGCCCTACCATTTGATATTCGTCCAGATCAGCAACCATTCAT 597
hum 493 CTAGGATCTCTCGAGAACAACAACT--GATCCCTCTCTCTCCCTACCCTCGTCCAGAGCCCTACCATTTGATATTCGTCCAGATCAGCAACCATTCAT 590
    * * * * *
  
```

B.



C.



tion of GABA_AR γ 2 alternative splicing when placed into the heterologous pGlo2 Δ minigene, and E10 by itself did not contribute to the effect (Fig. 4Bd to f). These results indicate that the 211-nt intronic element immediately upstream of E10 in GABA_AR γ 2 pre-mRNA is sufficient to mediate Nova-dependent regulation of alternative splicing (Fig. 4B).

Defining a Nova-dependent intronic splicing enhancer (NISE). This 211-nt intronic element contains seven YCAAY repeats (Fig. 5A) and is highly conserved between mouse and human sequences. Mutations were made in the first three (Fig. 5B, lanes m1), last four (lanes m2), or all seven (lanes m1+2) YCAAY elements, mutating each to YAAY, a sequence which does not bind to Nova (6, 32, 42). In 293T cells, these mutations increased the baseline level of E9 inclusion in the absence of cotransfected pNova-1 up to 2.4-fold, possibly due to inadvertent generation or disruption of a binding site for another protein factor. Nonetheless, mutation of the first three or last four YCAAY elements markedly reduced the effect of Nova in 293T cells from a 14.5-fold effect to a 2.5-fold (Fig. 5B, lanes m1) or 1.8-fold (lanes m2) effect on splicing, respectively, and mutation of all seven repeats abolished the effect of Nova on alternative splicing (lanes m1+2).

In N2A cells, which express endogenous Nova proteins, mutation of the Nova binding site would be expected to decrease the baseline level of E9 inclusion in the absence of cotransfected pNova-1. However, no change was seen, likely due to a competing increase in E9 inclusion identical to that observed in 293T cells. Nonetheless, the mutations showed identical behavior with respect to Nova-dependent exon inclusion in N2A cells (Fig. 5B), and taken together, these data demonstrate that YCAAY repeats within a conserved 211-nt intronic element are absolutely necessary for Nova-dependent regulation of GABA_AR γ 2 alternative splicing in the context of the chimeric minigene. Since these repeats are both necessary and sufficient to mediate Nova-dependent enhancement of exon inclusion, we refer to this region as a NISE element.

To confirm the role of the NISE element in mediating Nova action in the context of the native GABA_AR γ 2 pre-mRNA, identical YCAAY-to-YAAY mutations were made in pGABA, which harbors the intact GABA_AR γ 2 genomic sequence between E8 and E10 (Fig. 5C). Cotransfection of these plasmids with pNova-1 into 293T or N2A cells yielded results similar to those seen with the chimeric pGlo2 Δ constructs. In N2A cells, mutation of the first three or last four YCAAY repeats reduced Nova-dependent regulation from a 3.5- to a 2.2- or 1.9-fold effect, respectively (Fig. 5C). Mutation of all seven YCAAY

repeats (Fig. 5C, lanes m1+2) markedly diminished the effect of Nova-1 to a 1.3-fold change, a change comparable to the background effect of Nova on pGlo2 Δ in N2A cells (compare Fig. 4A).

In 293T cells, mutation of the YCAAY repeats altered the baseline level of E9 inclusion up to 3.9-fold; however, the magnitude of the Nova effect was markedly reduced from a 4.7-fold to a 2.2-fold (Fig. 5B, lanes m1) or 1.6-fold (lanes m2) effect. Mutation of all seven YCAAY repeats (Fig. 5B, lanes m1+2) resulted in a paradoxical 1.7-fold decrease in the inclusion of E9 upon the addition of Nova-1 in 293T cells. Similar baseline and paradoxical Nova-dependent effects were also seen upon mutation of YCAAY repeats to YAAY in GlyR α 2 minigenes (32). While not fully understood, the aberrant effect of Nova might reflect a dominant-negative action of the protein under these conditions. Taken together, these results indicate that the YCAAY repeats within the NISE are necessary for the regulation of GABA_AR γ 2 E9 alternative splicing by Nova-1 in the context of the wild-type pre-mRNA.

In vitro analysis of Nova binding to GABA_AR γ 2 RNA. To assess whether Nova protein directly interacts with the NISE element defined above, we performed nitrocellulose filter binding assays with purified RNA and Nova-1 protein. Purified full-length Nova-1 fusion protein (NFP) (Fig. 6A) is capable of binding a 159-nt RNA encompassing the NISE element (Fig. 5A) with high affinity (K_d , 7 nM) similar to the affinity with which Nova binds to RNA selection winners (SB2; K_d , 5 nM) (6) (Fig. 6B). Mutation of the first three YCAAY repeats to YAAY results in a modest decrease of approximately 4-fold in NFP binding affinity (K_d , 29 nM), while mutation of the last four reduces binding affinity by >14-fold (K_d , 100 nM). Mutation of all seven repeats reduced the binding affinity >100-fold to a level similar to that of NFP binding to an irrelevant β -globin RNA of similar length (Fig. 6B). These results are consistent with the conclusion that Nova binds directly to the NISE element in vivo to mediate GABA_AR γ 2 E9 alternative splicing.

To map the RNA sequence necessary for high-affinity Nova-1 binding, the 5' and 3' boundaries of the smallest RNA that retains high affinity to Nova-1 were determined (Fig. 6C). In vitro-transcribed RNA was labeled with ³²P at either the 5' or 3' end and subjected to mild alkaline hydrolysis. This RNA was then incubated with NFP, and bound RNAs were captured by filtration through nitrocellulose filters, eluted, and analyzed by denaturing polyacrylamide gel electrophoresis (PAGE). The 5' boundary mapped just upstream of the fourth YCAAY

FIG. 5. Mutagenesis of the YCAAY repeats close to E10 abolishes Nova-dependent regulation of GABA_AR γ 2 minigene alternative splicing. (A) Genomic sequences upstream of GABA_AR γ 2 exon 10 (gray box) from mouse (mus) and human (hum) were aligned by ClustalW alignment (MacVector). Conserved YCAAY elements are shown in boldface characters. Brackets indicate the region corresponding to the in vitro transcribed RNA used as described for Fig. 6B and are numbered relative to exon 10. m1 and m2 represent mutations of three and four YCAAY repeats to YAAY, respectively. (B) Chimeric minigenes containing the mutations depicted in panel A were cotransfected into 293T or N2A cells with increasing amounts of pNova-1 (0, 0.5, or 2.0 μ g), and spliced products were measured by RT-PCR and phosphorimage analysis as described for Fig. 2. The transfected pNova-1 expression was monitored by Western blotting (data not shown). Quantitation of the data is shown below representative autoradiographs and is presented as L/S exon use ratios \pm standard deviations for three independent transfections. WT, wild type. (C) GABA_AR γ 2 minigenes containing the mutations depicted in panel A were cotransfected into 293T or N2A cells with increasing amounts of pNova-1 (0, 0.5, or 2.0 μ g), and spliced products were measured by RT-PCR and phosphorimage analysis as described for Fig. 2. The transfected pNova-1 expression was monitored by Western blotting (data not shown). Quantitation of the data is shown below representative autoradiographs and is presented as γ 2L/ γ 2S exon use ratios \pm standard deviations for three independent transfections. WT, wild type.

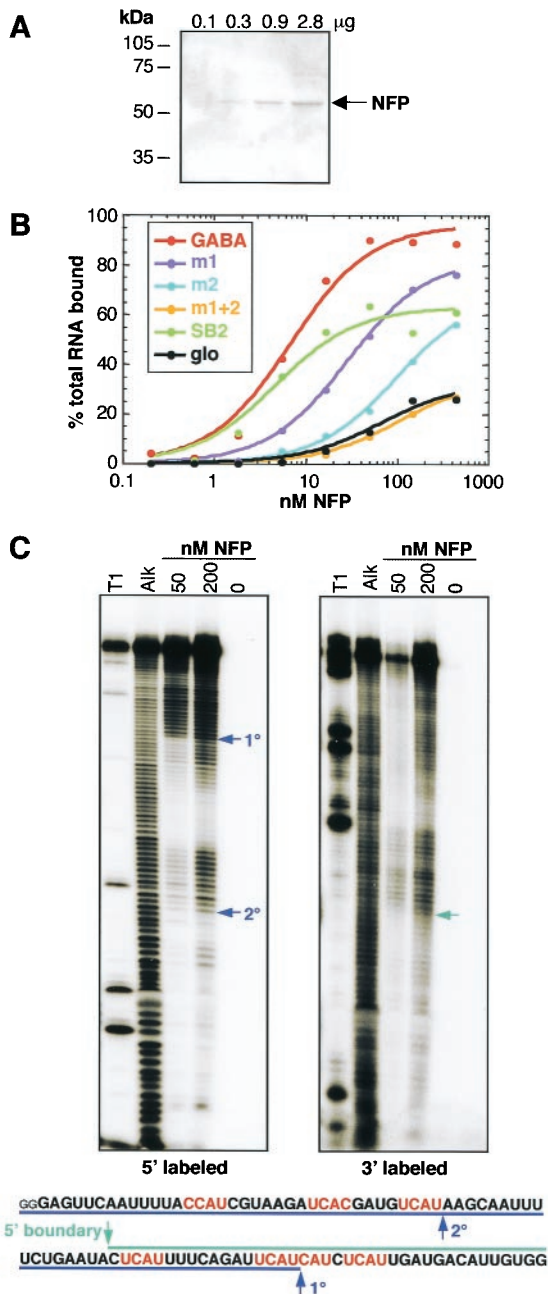


FIG. 6. Nova-1 binds with high affinity to GABA_AR γ 2 RNA in vitro. (A) His-tagged NFP was expressed in *Escherichia coli* and purified over a nickel column. Aliquots of increasing concentrations were analyzed by sodium dodecyl sulfate-PAGE followed by staining with Coomassie stain. (B) Nitrocellulose filter binding assays were performed using the NFP shown in panel A and six RNAs transcribed in vitro. GABA, RNA corresponding to the region of GABA_AR γ 2 intron 8 highlighted in Fig. 5, with the addition of the sequences GGGAG at the 5' end and CUAGCAA at the 3' end derived from the PCR primers used to amplify the template prior to in vitro transcription; m1 and m2, mutations of three and four YCAY repeats to YAAY in GABA, respectively; SB2, RNA obtained by RNA selection with Nova-1 (6); glo, RNA derived from human β -globin which spans regions of exon 1 and intron 1 and contains no YCAY elements. (C) Mapping the boundaries of Nova-1 binding to GABA_AR γ 2 intron 9 RNA. RNA was labeled with ³²P at either the 5' or 3' end and subjected to mild alkaline hydrolysis (Alk). The RNA was then incubated with NFP at a final concentration of 50 or 200 nM, and protein:RNA complexes were

tetranucleotide such that four YCAY repeats were contained within the smallest bound RNA. The position of the boundary at 1 nt upstream of a YCAY sequence agrees with structural data indicating that this nucleotide makes stacking interactions with the Nova KH domain in such a way as to contribute to binding affinity (42). We detected two 3' boundaries; the predominant boundary (Fig. 6C) occurred immediately downstream of the fifth YCAY repeat. A secondary boundary (Fig. 6C) immediately downstream of the third YCAY repeat was apparent at a high protein concentration (200 nM), suggesting that NFP binds these RNAs with lower affinity. The results suggest that a minimum of four or five YCAY repeats are necessary for high-affinity binding of the RNA to Nova-1.

To correlate the in vitro binding studies with the function of Nova-1 on the regulation of alternative splicing, additional β -globin-based minigenes harboring GABA_AR γ 2 intronic elements of 114, 68, 39, or 24 nt from intron 9 were assayed by transfection into 293T and N2A cells and were all found to retain Nova-dependent splicing capacity (Fig. 7). Reduction of the element to only 24 nt containing a total of four YCAY repeats was sufficient to mediate Nova-dependent alternative splicing. This contrasts with the relatively small effect of Nova on pGABA retaining only these four repeats (Fig. 5). However, two UCAU repeats separated by a single nucleotide are present in the β -globin pre-mRNA at 21 nt upstream of the site at which GABA_AR γ 2 sequences were inserted (data not shown), which might have contributed to Nova binding and splicing regulation. Mutation of the four GABA_AR γ 2-derived repeats (Fig. 7) abolished Nova-1's effect. Taken together with the in vitro binding studies, these results are consistent with the conclusion that Nova-1 binding to a small set of YCAY intronic repeats downstream of the regulated exon is necessary and sufficient for regulation of alternative splicing and imply that the number of YCAY repeats correlates with Nova binding affinity and the magnitude of splicing regulation.

DISCUSSION

The observation that the human genome contains a smaller number of genes than expected has enhanced the appreciation of alternative splicing as a mechanism for generating proteomic complexity. Comparisons of transcribed sequence databases with genomic data suggest that alternative splicing is particularly abundant in the brain (62), and understanding alternative splicing regulation in the brain is of profound importance (19, 25). While the mechanisms by which tissue-specific regulated alternative splicing decisions are controlled are not fully understood, an overriding theme appears to be the balance between positive and negative signals in pre-mRNA, which interact with constitutive or tissue-specific protein factors to promote or inhibit exon usage.

Nova-1 is a neuron-specific RBP important for neuronal

captured by filtration through nitrocellulose filters. Bound RNAs were eluted and analyzed by denaturing PAGE. RNase T1 digestion (T1) was used to generate size standards. Boundaries are highlighted by arrows and indicated on the transcribed RNA sequence shown at the bottom of each panel. 1°, primary boundary; 2°, secondary boundary. Small letters represent sequences derived from primers.

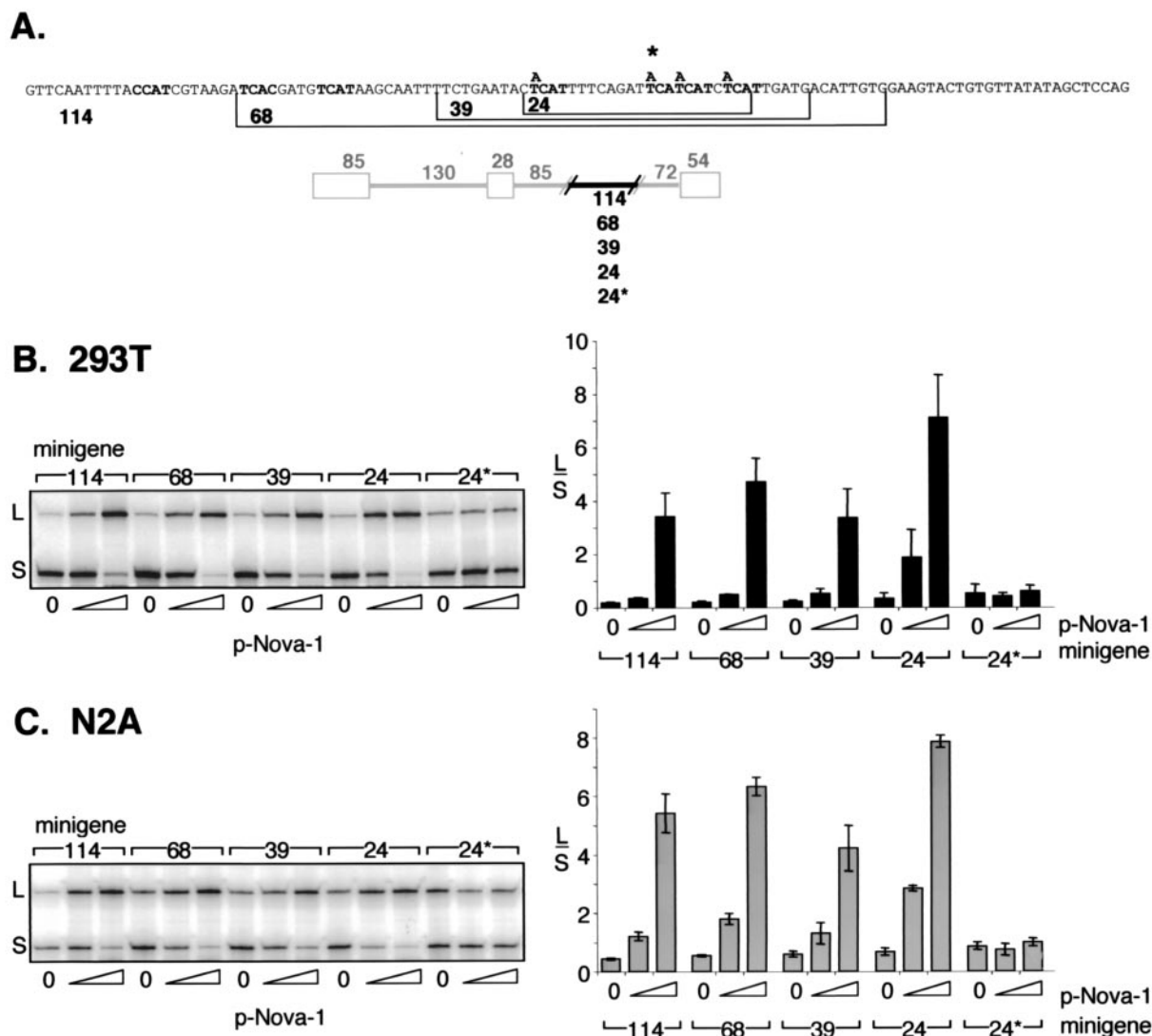


FIG. 7. A 24-nt intronic sequence is sufficient for regulation of alternative splicing by Nova-1 in a heterologous context. (A) DNA sequences derived from GABA_AR γ 2 intron 9 contained within the chimeric minigenes used in panels B and C. Numbers specify the lengths (in base pairs) of the GABA_AR γ 2 intronic regions cloned into the β -globin minigene. The star denotes the presence of four C-to-A mutations within the 24-nt GABA_AR γ 2 sequence. (B and C) The chimeric minigenes depicted in panel A were cotransfected into 293T (B) or N2A (C) cells with increasing amounts (0, 0.5, or 2.0 μ g) of pNova-1, and spliced products were measured by RT-PCR and phosphorimage analysis as described for Fig. 2. The titration of transfected pNova-1 expression was monitored by Western blotting (data not shown). Representative autoradiographs are shown at the left side of each panel; graphs summarize data from three independent transfections. The bars represent the ratios of L/S spliced products \pm standard deviations.

survival and regulation of alternative splicing in mammals (32). We have previously used in vitro methods to identify RNA binding targets for Nova and in turn used those targets to drive toward identification of an in vivo function for Nova binding to YCAY elements in an intron of GlyR α 2 pre-mRNA to regulate exon inclusion. However, the generality of this phenomenon was not previously known. The fortuitous identification of GABA_AR γ 2 as a second transcript whose alternative splicing was affected by Nova allowed us to prospectively examine a mechanism regarding which we had no prior understanding.

A number of factors have been previously reported to influence splicing of GABA_AR γ 2 E9, including the small exon size,

a weak 5' splice site, sequence elements within and upstream of E9 (63), and the RBP PTB (3). The finding that Nova-1 null mice showed a decrease in the level of inclusion of E9 (Fig. 1) (32) indicated that Nova-1 also plays a role in regulation of GABA_AR γ 2 alternative splicing but did not address whether Nova acts directly on GABA_AR γ 2 pre-mRNA, and if so, whether Nova binds to an element resembling the YCAY sequences selected in vitro. Here we show that GABA_AR γ 2 E9 inclusion is regulated by Nova action on a distal downstream intronic YCAY repeat element. These results provide an important verification of Nova function as a sequence-specific splicing factor, underscore the significance of YCAY repeats

as Nova targets *in vivo*, and provide insight into the mechanisms of tissue-specific splicing regulation.

The NISE element. Several lines of evidence demonstrate that Nova regulates GABA_AR γ 2 E9 inclusion by a direct action on clustered YCAY intronic repeats. First, the NISE element contains seven YCAY repeats, which are known to be the preferred RNA sequence for Nova KH domain binding (6, 33, 42), and these are conserved between mice and humans in GABA_AR γ 2 (Fig. 5A). Second, mutation of these YCAY repeats to YAAY completely abolished Nova's effect on splicing (Fig. 5B and C), indicating that these sequences are necessary for Nova-dependent splicing regulation. Third, Nova binds with high affinity to the wild-type but not the mutated sequence corresponding to this region (Fig. 6B). Boundary mapping experiments indicated that this binding abruptly drops when the sequence contains fewer than four YCAY repeats (Fig. 6C). Consistent with this result, we found that a 24-nt element spanning the last four YCAY repeats is sufficient to mediate Nova-dependent E9 inclusion (Fig. 7). These findings do not entirely exclude the possibility that other sequences within the GABA_AR γ 2 pre-mRNA contribute to Nova-dependent splicing of E9; however, our data indicate that any such additional Nova regulatory regions would play a minor regulatory role and would be dependent on the integrity of the NISE element.

The requirement for a number of YCAY elements for high-affinity Nova-1 binding suggests that alternative splicing regulation of GABA_AR γ 2 might be mediated through the binding of multiple KH domains to multiple YCAY repeats within the RNA. A similar mechanism has been reported for Vg1 mRNA localization mediated by VgRBP/Vera. Binding to the Vg1 mRNA requires two KH didomains, and these KH didomains bind RNA cooperatively (23). It is possible that a single Nova molecule can bind up to three YCAY elements within an RNA. Moreover, Nova proteins can multimerize (41, 42, 52), suggesting that several Nova protein molecules cooperatively bind an RNA target. Biophysical evidence suggests that both of these mechanisms can apply to the formation of Nova-RNA complexes (54). Either mechanism can effectively increase the affinity of binding to an RNA target and help to explain the requirement for several YCAY elements in the GABA_AR γ 2 pre-mRNA for regulation of alternative splicing by Nova.

A general principle of RBP action can be seen in this data. RNA targets of high specificity yet low complexity can be recognized *in vivo* via clustering of the target elements. This can be seen in several examples of RNA translational control and localization mediated by RBP-target RNA interactions. These include the action of hnRNPs K and E1 on LOX mRNA translation in erythrocytes which requires clustered differentiation control elements (49, 50), the action of bruno on *oskar* mRNA translation in *Drosophila* which requires clustered bruno response elements (36), and the action of Vg1RBP/Vera on Vg1 and VegT mRNA localization in *Xenopus* which requires clustered VM1 elements (5, 40). Our finding that clustered YCAY repeats are necessary and sufficient to mediate the action of Nova helps confirm this principle and extend it to alternative splicing.

Mechanism of Nova action to regulate alternative splicing.

The mechanism by which Nova-1 enhances E9 inclusion is unclear. In some instances of alternative splicing control, enhancement of splicing is mediated through inhibition of a neg-

ative signal. For example, PTB acts on CT/CGRP through inhibition of a competing pseudoexon (45). PTB has also been shown to inhibit inclusion of GABA_AR γ 2 E9, and nucleotides adjacent to and within E9 are important for this function (3, 64). This observation led to the proposal that inclusion of E9 in neurons of the CNS is regulated by a mechanism involving derepression of splicing by PTB. The finding that GABA_AR γ 2L exon utilization is dysregulated in Nova-1 null mice suggested that Nova is a candidate for mediation of this derepression (32), perhaps through competitive binding to the pre-mRNA. Our findings that the NISE element lies further than 1 kb downstream of E9 and that a small (as short as 24 nt) sequence element corresponding to the region of Nova-1 binding is sufficient to mediate Nova-dependent enhancement of an upstream exon in a heterologous system argue against this hypothesis of Nova action on GABA_AR γ 2L exon utilization.

Alternatively, enhancement can be achieved by positive action such as recruitment of essential splicing factors to an otherwise poorly utilized splice site, as has been proposed for KSRP enhancement of *src* N1 inclusion (46), or the action of the downstream enhancer of (NMHC)-B N30 inclusion (26). Conceptually, Nova-dependent enhancement of GABA_AR γ 2L exon utilization is likely to involve facilitation of splicing of an intron adjacent to E9. It is unclear whether this facilitation involves the intron upstream [in similarity to (NMHC)-B N30] or downstream (in similarity to KSRP/*src* N1) of E9. However, comparison with Nova's action on GlyR α 2 splicing (and the simplest biochemical hypothesis) would predict that Nova enhances splicing of the intron in which it resides. *In vitro* splicing assays utilizing simplified RNA substrates similar to those performed by Guo and Kawamoto (26) would be helpful in assessing this possibility.

The extremely close positioning of the Nova binding site at only 80 nt upstream of the splice acceptor signals of exon 10 is intriguing. The majority of splicing regulatory sequences that have been characterized lie within or close to the exon that they regulate (58), including those reported to affect GABA_AR γ 2 E9. Although our data confirm that mutations within the region surrounding E9 cause a dramatic decrease in the basal level of E9 inclusion, Nova acts at a distance—the NISE element is approximately 1,250 nt downstream of E9. One possibility is that Nova recruits factors that interact with and help assemble the basal splicing machinery in a manner analogous to the distant action of transcriptional enhancer elements. Transcriptional enhancers recruit DNA binding proteins which, in turn, help to recruit the basal transcription machinery to the promoter, which often lies many kilobases away, through protein-protein interactions (reviewed in reference 29). In this context, we hypothesize that Nova binding to the NISE recruits proteins that directly or indirectly interact with either the upstream 5' splice site and/or the downstream 3' splice site. Yeast two-hybrid results have indicated that Nova can bind to spliceosomal elements present at both the 3' and 5' splice sites, namely, U1-70K and U2AF⁶⁵, as well as some SR proteins, including SRp20 (G. Stefani and R. Darnell, unpublished data). Through interactions such as these, Nova may serve a bridging function between the nearby 3' splice site and the next upstream 5' splice site.

Another possibility for the mechanism of Nova action is that Nova slows down the kinetics of spliceosome assembly at E10

by interfering with binding or assembly of essential factors at this site. This might then enhance splicing to E9 by allowing more time for assembly to take place at an otherwise weakly recognized site and thus increase the chances of recognition of this exon. A similar kinetic mechanism has been proposed to explain the influence of transcription on alternative splicing through differences in RNA polymerase II processivity rates (reviewed in reference 8). However, it is difficult to reconcile this hypothesis with what is known about Nova regulation of GlyR α 2 splicing. Nova enhances GlyR α 2 E3A inclusion through binding to an element which lies at the same distance upstream of E3A that the NISE element lies upstream of GABA_AR γ 2 E10 (32). If a similar kinetic mechanism involving interference of spliceosome assembly at the adjacent 3' splice site were involved in regulation of GlyR α 2 splicing, Nova binding would be expected to repress E3A inclusion.

The identification of the NISE element in GABA_AR γ 2 pre-mRNA offers several advantages over the study of GlyR α 2 for the further elucidation of the biochemical mechanism of Nova-dependent alternative splicing regulation. First, alternative splicing of GABA_AR γ 2 involves the skipping or inclusion of a single cassette exon rather than the more complex scenario of mutual exclusion of the two alternative exons E3A and E3B in GlyR α 2. Second, this study demonstrated that the NISE element is necessary for Nova-dependent regulation of E9 splicing in the context of the full-length introns; this has not been shown for GlyR α 2, in part because the Nova binding site lies within an intron approximately 35 kb in length, necessitating the use of truncated introns in the minigenes used to assay GlyR α 2 splicing. Finally, we have extended previous analyses by demonstrating that the NISE element is sufficient to mediate Nova-dependent enhancement of exon inclusion in a heterologous context, enabling us to conclude that the exact nature of the surrounding splice sites and other regulatory elements within the pre-mRNA is not critical for Nova's action. Furthermore, this simplified substrate is useful in *in vitro* splicing assays (Stefani and Darnell, unpublished), providing a potentially powerful tool to help determine the biochemical mechanism by which Nova regulates alternative splicing.

Conclusions. Beginning with the *in vivo* finding that GABA_AR γ 2L exon utilization is perturbed in Nova-1 null mice and working backwards to biochemical function, we have found an intronic Nova binding element—the NISE—that is necessary and sufficient to mediate Nova-dependent splicing regulation. This is the first demonstration that a small RNA element is both necessary and sufficient to mediate the action of a neuron-specific mammalian splicing activator. The NISE element was identified using a systematic approach to screen for Nova-dependent regulatory elements. The finding that the NISE element also conforms to the consensus Nova binding sequences predicted *in vitro* further validates the usefulness of RNA selection as a means to identify biologically relevant targets and leads to the prediction that it might be possible to identify additional Nova targets through bioinformatic screenings. The work presented here, completing a cycle of experiments *in vitro* to *in vivo* and back, validates our identification of Nova binding sites as targets for Nova-dependent regulation and furthers our understanding of how Nova acts as a regulator of alternative splicing in neurons.

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