Biochemical and genetic evidence for a role of IGHMBP2 in the translational machinery

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The human motor neuron degenerative disease spinal muscular atrophy with respiratory distress type 1 (SMARD1) is caused by loss of function mutations of immunoglobulin \( \mu \)-binding protein 2 (IGHMBP2), a protein of unknown function that contains DNA/RNA helicase and nucleic acid-binding domains. Reduced IGHMBP2 protein levels in neuromuscular degeneration (nmd) mice, the mouse model of SMARD1, lead to motor neuron degeneration. We report the biochemical characterization of IGHMBP2 and the isolation of a modifier locus that rescues the phenotype and motor neuron degeneration of nmd mice. We find that a 166 kb BAC transgene derived from CAST/EiJ mice and containing tRNA genes and activator of basal transcription 1 (Abt1), a protein-coding gene that is required for ribosome biogenesis, contains the genetic modifier responsible for motor neuron rescue. Our biochemical investigations show that IGHMBP2 associates physically with tRNAs and in particular with tRNA\(^{\text{Ty}}\), which are present in the modifier and with the ABT1 protein. We find that transcription factor IIIC-220 kDa (TFIIC220), an essential factor required for tRNA transcription, and the helicases Reptin and Pontin, which function in transcription and in ribosome biogenesis, are also part of IGHMBP2-containing complexes. Our findings strongly suggest that IGHMBP2 is a component of the translational machinery and that these components can be manipulated genetically to suppress motor neuron degeneration.

INTRODUCTION

Spinal muscular atrophy with respiratory distress type 1 (SMARD1) is an autosomal recessive neurodegenerative disease (1,2). The disease is also known as distal spinal muscular atrophy 1 (DSMA1, MIM no. 604320). The primary pathology of SMARD1 consists of degeneration of spinal cord motor neurons and axonal loss leading to muscular atrophy (1,2). Diaphragmatic paralysis is one of the earliest and most severe manifestations of the disease and typically leads to early infant death (1,2). SMARD1 results from mutations in the immunoglobulin \( \mu \)-binding protein 2 (IGHMBP2) (1,2). The human IGHMBP2 is a 993 amino acid protein that contains an RNA/DNA helicase domain (found in the superfamily I helicases), an R3H, single-stranded nucleic acid-binding domain and a zinc finger domain (1–3). IGHMBP2 mRNA is ubiquitously expressed (4). On the basis of in vitro binding assays, IGHMBP2 has been proposed to bind DNA and was given several names in the literature: glial factor 1 \([\text{GF1}]\) (5), immunoglobulin S-\( \mu \)-binding protein-2 \([\text{Smbp2 or IGHMBP2}]\) (4,6), rat insulin-enhancer-binding protein \([\text{Rip1}]\) (7) and cardiac transcription factor 1 \([\text{Catf1}]\) (8). Proposed functions, based on in vitro data, include transcriptional activation, immunoglobulin class switching and pre-mRNA splicing (4–9). In neurons, IGHMBP2 is predominantly found in the cytoplasm, including dendrites and axons (3), arguing that IGHMBP2 has a major function outside the nucleus. The presence of an RNA helicase domain and a single-stranded nucleic acid-binding domain suggests that IGHMBP2 may function in RNA processing, regulation or metabolism.
Neuromuscular degeneration (nmd) mice that are homozygous for mutations of the Igshmbp2 gene develop a phenotype that is very similar to SMARD1 (10). The pathology of nmd mice consists of motor neuron degeneration with axonal loss leading to neurogenic muscle atrophy and death by 8–12 weeks (10–12). We previously reported that the nmd mutation is a single A-to-G transition 23 bp into intron 4 of the mouse Igshmbp2 gene that creates a new favorable splice donor, resulting in a frameshift and premature stop codon (10). In nmd homozygous mice, low levels (~20%) of correctly spliced mRNA are evident in all nmd tissues, indicating that this is a hypomorphic allele and not a complete null mutation (10). We reported that the motor neuron degeneration of nmd mice was completely corrected by transgenic expression of wild-type IGHMBP2 driven by a neuron-specific promoter (11). This finding demonstrates that motor neuron degeneration in nmd mice is cell autonomous and is a direct consequence of reduced IGHMBP2 protein levels in motor neurons. Interestingly, these transgenic mice later developed a primary myopathy affecting both skeletal muscles and the myocardium, suggesting that IGHMBP2 is also essential for myocyte maintenance (11). Such primary myopathy has not been described in human patients with SMARD1, presumably because it is superceded by the severe motor neuronopathy that leads to early death, before manifestation of the primary myopathy (11).

Mice contain 418 tRNA genes and humans contain 505 tRNA genes, corresponding to tRNAs that decode the standard 20 amino acids (13). Approximately 6% of these genes contain introns (13). tRNAs are transcribed in the nucleus by RNA polymerase III as longer pre-tRNA transcripts (14,15). tRNA genes contain intragenic RNA polymerase III promoters whose core elements consist of the A and B boxes (whose sequences roughly correspond to the D- and T-loops of the mature tRNA) (16,17). The A and B boxes are recognized by a multisubunit complex known as general transcription factor IIIC (TFIIC). Mammalian TFIIC220, one of the subunits of TFIIC, binds directly to box B of RNA polymerase III promoters and is essential for tRNA transcription (16,17). Pre-tRNA transcripts contain 5′-leader and 3′-trailer sequences (14,15). Several tRNAs also contain a single, short (14–60 nt) intron which is found in the anticodon loop, typically one nucleotide 3′ to the anticodon (13–15). Processing of pre-tRNAs involves removal of the 5′-leader by RNAses P, removal of the 3′-trailer by various endo- and exo-nucleases, splicing of the intron, addition of the CCA trinucleotide to the 3′-end and numerous tRNA modifications (14,15,18,19). Addition of the CCA trinucleotide typically occurs prior to tRNA splicing and is followed by aminoaetylation and export of the mature tRNA to the cytoplasm (15).

The biogenesis of ribosomes in eukaryotes is a complicated, step-wise process that involves more than 200 proteins and an equal number of small nucleolar RNAs (snoRNAs) (20). RNA polymerase I generates a large pre-rRNA precursor (termed 47S/45S) in mammals that contains the mature 18S, 28S and 5.8S tRNAs. A series of endonucleolytic cleavages and exonucleolytic processing steps leads to generation of the mature rRNAs (21). snoRNAs are involved in the processing and modification of ribosomal RNA (rRNA) and are found in cells complexed with proteins (snoRNPs). U3 snoRNP is a large ribonucleoprotein composed of a 217 nucleotide U3 snoRNA and multiple proteins (22) and directs the first cleavage of the pre-rRNA precursor (22,23).

Spinal muscular atrophy (SMA), although distinct from SMARD1, shares many clinical and pathological features with SMARD1. SMA is caused by deletions or loss of function mutations of the survival of motor neurons (SMN) gene (24). The pathology of SMA consists of degeneration of spinal cord motor neurons, leading to muscular atrophy (25). The SMN protein associates with seven additional proteins, known as Gemins2–8, into a large complex termed the SMN complex (26–28). The SMN complex plays critical roles in the biogenesis of spliceosomal small nuclear RNPs (snRNPs) and possibly other RNP (26,29–32).

Here, we show that IGHMBP2 is not a component of the SMN complex, but rather it interacts with a unique set of small RNAs that includes tRNA Tyr, and with proteins, including helicases, TFIIIC220 and activator of basal transcription 1 (ABT1), that are required for tRNA transcription and ribosome biogenesis and function. These biochemical observations are corroborated in mutant nmd mice by the analysis of a genetic suppressor of motor neuron disease encoding multiple tRNA Tyr genes and ABT1 that we have shown to associate physically with IGHMBP2. Our findings strongly suggest that IGHMBP2 is a component of the translational machinery and that these components can be manipulated genetically to suppress motor neuron degeneration.

RESULTS

IGHMBP2 is not a component of the SMN complex

Because SMARD1 and SMA have similar phenotypes, caused by degeneration of motor neurons, we first wanted to investigate whether IGHMBP2 could interact with SMN or with other components of the SMN complex. We cloned the human IGHMBP2 cDNA in a eukaryotic expression vector in frame with a FLAG epitope tag and generated a stable human 293T cell line, expressing FLAG-IGHMBP2 and another 293T cell line expressing the empty FLAG vector as a negative control. We used anti-FLAG beads to immunoprecipitate FLAG-IGHMBP2 and we probed the immunoprecipitates on western blots using antibodies against components of the SMN complex: SMN, Gemin2, Gemin3, Gemin5 and Gemin6. As shown in Supplementary Material, Figure S1, none of these proteins co-immunoprecipitated with IGHMBP2, indicating that IGHMBP2 does not associate with the SMN complex.

Generation of a monoclonal antibody against IGHMBP2

Next, we generated a monoclonal antibody (mAb11-24) against IGHMBP2 by immunizing mice with an aminoterminal portion of human IGHMBP2 that was generated as recombinant protein in bacteria. As shown in Figure 1, mAb11-24 recognized IGHMBP2 in western blots, in immunoprecipitations (IPs) and by immunofluorescence.

mAb11-24 recognizes a protein band of ~110 kDa corresponding to IGHMBP2 and also a faster migrating band at ~60 kDa, which is inconsistently seen, and likely represents...
None of the antibodies that we have tested was able to efficiently immunoprecipitate Reptin or Pontin, although many of them worked well on western blots. To overcome the lack of immunoprecipitating antibodies for Reptin and Pontin, we used FLAG-tagged constructs for Reptin and Pontin and co-transfected these plasmids along with a plasmid expressing myc-IGHMBP2 in 293T cells. Cells transfected with myc-IGHMBP2 and FLAG-empty vector served as negative controls. We then performed IPs with anti-FLAG antibodies and probed the immunoprecipitates with anti-myc (9E10) antibody. As shown in Figure 2B and C, myc-IGHMBP2 co-immunoprecipitates with FLAG-Reptin and FLAG Pontin. Peptides corresponding to transcription factor IIIC, 220 kDa subunit (TFIIIC220; also known as TFIIICa) were obtained from the P65 band, indicating that TFIIIC220, a large 220 kDa protein, fragmented during cell lysis and IP. To confirm the IGHMBP2–TFIIIC220 association, we performed IPs in 293 cells using an antibody against the endogenous TFIIIC220 and probed the immunoprecipitates with mAb11-24 against IGHMBP2. As shown in Figure 2D (lane 5), IGHMBP2 co-immunoprecipitated with TFIIIC220. Peptides corresponding to tubulin chains were also identified in IGHMBP2 immunoprecipitates. To further assess these findings, we performed IPs using anti-α-tubulin antibody from 293T cells and probed the immunoprecipitates with mAb11-24 against IGHMBP2 but detected only a faint interaction (Fig. 2D, lane 4). These results indicate that IGHMBP2 associates with the helicases Reptin and Pontin and with TFIIIC220. Co-IP experiments showed that IGHMBP2 did not interact with hnRNPH, HSP70 or Zfp161 (Supplementary Material, Fig. S2). Since many RNA or DNA helicases form dimers or homomers, we tested whether IGHMBP2 self-associates. For this, we expressed either FLAG-IGHMBP2, or myc-IGHMBP2, or both FLAG- and myc-IGHMBP2 in 293T cells along with an empty vector as negative control (Fig. 2E). Co-IPs were then performed with either anti-FLAG or anti-myc antibodies, and the immunoprecipitates were probed on western blots with anti-myc (Fig. 2E, top panel) or anti-FLAG (Fig. 2E, bottom panel) antibodies, respectively. The presence of myc-IGHMBP2 in the FLAG-IGHMBP2 immunoprecipitates and the presence of FLAG-IGHMBP2 in the myc-IP immunoprecipitates (Fig. 2E) indicate that IGHMBP2 is able to self-associate in vivo.

Identification of small RNAs that associate with IGHMBP2

We next asked whether IGHMBP2 associates with small RNAs. We generated a stable MN1 cell line, expressing FLAG-IGHMBP2 and another MN1 cell line expressing the empty FLAG vector as a negative control (Fig. 3A). MN1 is a well-characterized hybrid mouse motor neuronal cell line (33–35). We then performed IPs with anti-FLAG antibody from MN1-FLAG-IGHMBP2 and from MN1-FLAG cells. RNA was then isolated from the immunoprecipitates, 3′-end-labeled with [32P]Cp and T4 RNA ligase and resolved on a 10% UREA–PAGE. Two discrete RNA bands (~34 and ~40 nt) were present in the FLAG-IGHMBP2 immunoprecipitates only (Fig. 3B). We then directionally cloned the ~40 nt RNA band from the FLAG-IGHMBP2 immunoprecipitates by employing the same strategy that we used to a proteolytic fragment of IGHMBP2. The predominant localization of IGHMBP2 in human NT2 cells and in mouse MN-1 cells is cytoplasmic but there is also a fraction of IGHMBP2 that is localized in the nucleus. This is consistent with prior reports using a rabbit polyclonal anti-IGHMBP2 antibody (3).

Identification of proteins that associate with IGHMBP2

To identify possible protein partners of IGHMBP2, we used 293T cells stably expressing either FLAG-IGHMBP2 or the FLAG empty vector, in IP experiments. We chose this approach because we can perform IPs by using two different monoclonal antibodies, anti-FLAG and also mAb11-24 against IGHMBP2. Proteins that co-immunoprecipitate with both antibodies are likely to represent authentic IGHMBP2-associated proteins. Furthermore, competitive elution with 3XFLAG peptide from the anti-FLAG immunoprecipitates results in much cleaner co-IP patterns since the antibody chains do not contaminate the samples. As shown in Figure 2, other proteins co-immunoprecipitated with FLAG-IGHMBP2 and were found in both anti-FLAG and mAb11-24 immunoprecipitates. We identified several of these proteins by mass spectrometry and the results are shown in Figure 2A. Interestingly, we identified two closely related ATPase/helicases, Reptin (Tip48, RuvB-like2) and Pontin (Tip49, RuvB-like1) co-immunoprecipitating with IGHMBP2. To confirm the association of IGHMBP2 with Reptin and Pontin, we performed reverse co-IP experiments. None of the antibodies that we have tested was able to efficiently immunoprecipitate Reptin or Pontin, although many of them worked well on western blots. To overcome the lack of immunoprecipitating antibodies for Reptin and Pontin, we used FLAG-tagged constructs for Reptin and Pontin and co-transfected these plasmids along with a plasmid expressing myc-IGHMBP2 in 293T cells. Cells transfected with myc-IGHMBP2 and FLAG-empty vector served as negative controls. We then performed IPs with anti-FLAG antibodies and probed the immunoprecipitates with anti-myc (9E10) antibody. As shown in Figure 2B and C, myc-IGHMBP2 co-immunoprecipitates with FLAG-Reptin and FLAG Pontin. Peptides corresponding to transcription factor IIIC, 220 kDa subunit (TFIIIC220; also known as TFIIICa) were obtained from the P65 band, indicating that TFIIIC220, a large 220 kDa protein, fragmented during cell lysis and IP. To confirm the IGHMBP2–TFIIIC220 association, we performed IPs in 293 cells using an antibody against the endogenous TFIIIC220 and probed the immunoprecipitates with mAb11-24 against IGHMBP2. As shown in Figure 2D (lane 5), IGHMBP2 co-immunoprecipitated with TFIIIC220. Peptides corresponding to tubulin chains were also identified in IGHMBP2 immunoprecipitates. To further assess these findings, we performed IPs using anti-α-tubulin antibody from 293T cells and probed the immunoprecipitates with mAb11-24 against IGHMBP2 but detected only a faint interaction (Fig. 2D, lane 4). These results indicate that IGHMBP2 associates with the helicases Reptin and Pontin and with TFIIIC220. Co-IP experiments showed that IGHMBP2 did not interact with hnRNPH, HSP70 or Zfp161 (Supplementary Material, Fig. S2). Since many RNA or DNA helicases form dimers or homomers, we tested whether IGHMBP2 self-associates. For this, we expressed either FLAG-IGHMBP2, or myc-IGHMBP2, or both FLAG- and myc-IGHMBP2 in 293T cells along with an empty vector as negative control (Fig. 2E). Co-IPs were then performed with either anti-FLAG or anti-myc antibodies, and the immunoprecipitates were probed on western blots with anti-myc (Fig. 2E, top panel) or anti-FLAG (Fig. 2E, bottom panel) antibodies, respectively. The presence of myc-IGHMBP2 in the FLAG-IGHMBP2 immunoprecipitates and the presence of FLAG-IGHMBP2 in the myc-IP immunoprecipitates (Fig. 2E) indicate that IGHMBP2 is able to self-associate in vivo.
clone miRNAs and the (mt)tRNA\textsubscript{Met} from Ago2 immunoprecipitates (36,37) (detailed in 38). Of the 30 clones sequenced, three clones corresponded to the 3\textsuperscript{\text{\textprime}}-end or tRNA Tyr (Fig. 3C and D), one clone corresponded to the 3\textsuperscript{\text{\textprime}}-end of tRNA Met (CAU anticodon), one clone to the 3\textsuperscript{\text{\textprime}}-end of tRNA Arg (UCU anticodon) and the remainder represented clones with adapters only. These results indicated that IGHMBP2 associated with tRNAs and, in particular, tRNA\textsubscript{Tyr}. We were unable to clone the smaller, ~34 nt band, but it likely corresponds to 5\textsuperscript{\text{\textprime}}-ends of tRNAs. Because the RNAs from the FLAG-IGHMBP2 immunoprecipitates were 3\textsuperscript{\text{\textprime}}-end-labeled with a single \([^{32}\text{P}]\)PCp molecule, the stoichiometry of the RNAs may be extrapolated from band intensity. As shown in Figure 3B, the ratio of the ~40 nt to the ~34 nt band is 1:1, suggesting that they may be derived from processing of a tRNA molecule. In experiments described in what follows, we detect association of IGHMBP2 with intact tRNAs (Fig. 3E and F). Cleavage of tRNAs in the anticodon loop has been shown to occur as a conserved response to oxidative stress in yeast, plants and mammalian cells (39) and it is likely
that such fragmentation occurred prior to or during cell lysis and RNA IP. The RNA bands at ~75, ~120 and ~180 nt correspond in size to tRNAs, SS rRNA and 5.8S rRNAs. The SS and 5.8S rRNAs (which we have identified by northern blots; our unpublished data) are likely non-specific as we often observe them in small RNA IPs using various antibodies [such as anti-Ago antibodies (36)], and in the anti-FLAG vector (negative control) RNA immunoprecipitate (Fig. 3B).

Next, we isolated RNAs that co-immunoprecipitated with endogenous IGHMBP2 from mAb11-24 immunoprecipitates from HeLa cells; non-immune mouse serum (NMS) served as a negative control. We 3' end-labeled the RNA with pCp and T4 RNA ligase and resolved the small RNAs on a 10% UREA–PAGE. As shown in Figure 3E, a strong band at ~75 nt, corresponding to tRNAs, was present in mAb11-24 immunoprecipitates but not in the negative (NMS) control. To confirm the association of tRNA-Tyr with endogenous IGHMBP2, we probed RNAs that were isolated from mAb11-24 immunoprecipitates with an oligonucleotide probe against tRNA^{Tyr} on a northern blot.
As shown in Figure 3F, tRNA$^{Tyr}$ was detected specifically in mAb11-24 immunoprecipitates. These findings indicate that IGHMBP2 associates with small RNAs and, in particular, with tRNA$^{Tyr}$.

Positional complementation cloning of the MnMc modifier using BAC transgenic rescue

During the course of our original fine-mapping of the nmd mouse mutation, we observed that $\sim$25% of the (B6-nmd$^{jJ}$ x CAST/EiJ)F2 nmd$^{jJ}$/nmd$^{jJ}$ mice were very mildly affected with regard to the extent of neurogenic atrophy and paralysis (10,11). We mapped the modifier region to mouse Chr 13 and completed construction of a congenic B6.CAST-MnMc line of mice that are homozygous B6 at all loci except around the region containing the MnMc gene. Just as in the original F2 intercross, this single Chr 13 locus suppresses the nmd motor neuron disease in a semidominant fashion but fails to rescue the onset or progression of dilated cardiomyopathy (11). Using the B6.CAST-MnMc congenic strain in a mapping cross, we have obtained additional recombinants to reduce the genetic interval to 0.03 cM between markers CA1 and CAGA. We have constructed a contig across most of the 430 kb MnMc critical interval from CAST-derived (CHORI-26 library, http://bacpac.chori.org) BAC clones (Fig. 4A).

To further localize the MnMc modifier within our genetic interval, we undertook a transgenic rescue strategy using the 166 kb CAST-derived BAC clone (CH26-27K3). As shown in Figure 4A, BAC-27k3 contains two protein-coding genes, Abt1 and Zfp322a. In addition, 1 non-coding EST (ENSMUSESTG00000018860) and 24 tRNA genes (10 tRNA$^{Val}$, 6 tRNA$^{Ala}$, 5 tRNA$^{Tyr}$, 2 tRNA$^{Ile}$ and 1 tRNA$^{Met}$) have been annotated to the region between 23,380,802 and 23,527,157 bp on mouse Chr 13 (NCBI build 37). We have identified eight independent B6-Tg(BAC27k3) transgenic founders and crossed each of them with B6-+/nmd mice to generate B6-Tg(nmd/nmd) transgenic mice. Seven of eight 27k3 founder lines completely recapitulate the MnMc modifier rescue phenotype seen in our B6.CAST-MnMc congenic line though one founder line (line 60) displays a generally mild, yet variable rescue phenotype among littermates. Thus, we have been able to limit the MnMc modifier effect to a single 166 kb CAST-derived BAC clone.

Shotgun sequencing of BAC-27k3 to identify CAST/EiJ sequence polymorphisms was compared with the public mouse C57BL/6J genome sequence. No coding polymorphisms, aberrant alternative splicing events or real-time reverse transcriptase (RT)–polymerase chain reaction (PCR) expression differences have been detected in the CAST-derived Abt1 or Zfp322a sequences that might point to the cause of the genetic modifier effect (data not shown). Apart
from the triplication of an ∼10 kb repetitive element in CAST detected by DNA fingerprint analysis of BAC-27k3, we have been surprised by the low rate of polymorphisms between C57BL/6J and CAST/EiJ in our interval (∼1 SNP/4.1 kb) given that the two subspecies are thought to have diverged approximately 0.5–1 million years ago. However, this is consistent with the mosaic nature of the genome of common laboratory strains with intervals of high SNP density and low SNP density reflecting the recent contributions of *musculus*, *domesticus* and *castaneus*-derived chromosomal segments during inbreeding (40). The ∼10 kb triplication is relatively common among several inbred strains based on Southern blotting (data not shown) and does not appear to be associated with the modifier effect though the sequence contains three tRNA genes (tRNA\textsuperscript{A\textsubscript{ta}} and two tRNA\textsuperscript{Val} genes). Complete sequencing of an 18.1 kb EcoRI fragment of BAC-27k3 containing the *Abt1* gene, six tRNA genes and one noncoding EST revealed only seven polymorphisms in intergenic or intronic regions (summarized in Fig. 4A, GenBank accession number EU880213) that do not immediately suggest a mechanism for the modifier effect of the CAST/EiJ allele versus the C57BL/6J allele in suppressing motor neuron disease. The syntenic genomic area in humans is found on chromosome 6 and contains a cluster of four tRNA\textsuperscript{Tyr} genes and the *ABT1* gene.

The identification of five tRNA\textsuperscript{Tyr} genes in the 27k3 modifier and the physical association of tRNA\textsuperscript{Tyr} with IGHMBP2 are particularly intriguing and indicate that IGHMBP2 may function in tRNA pathways or in translation. This is further supported by the association of IGHMBP2 with TFIIIC220, a general transcription factor for RNA polymerase III, which is essential for transcription of tRNA genes.

**IGHMBP2 associates with the ABT1 protein, whose gene is present in the 27k3 modifier**

The presence of the *Abt1* gene in the 27k3 modifier is particularly interesting because ABT1 is an essential factor required for pre-ribosomal RNA (pre-tRNA) processing and maturation of ribosomes (41). In the initial immunopurification of FLAG-IGHMBP2, we did not detect a clear protein band at ∼31 kDa, corresponding to ABT1, in the Coomassie-blue stained FLAG-IGHMBP2 eluates (Fig. 2A). However, since ABT1 is a small protein and the sensitivity of Coomassie blue is low, we performed co-IP experiments to test whether there is an association of IGHMBP2 with ABT1. We found that there are no anti-ABT1 antibodies suitable for IP. We thus generated a myc-tagged ABT1 construct and co-transfected it along with a plasmid expressing FLAG-IGHMBP2 in 293T cells. We then performed IPs with anti-FLAG or anti-myc antibodies and probed the immunoprecipitates with anti-myc (9E10) or anti-FLAG antibodies to detect co-precipitating ABT1 and IGHMBP2, respectively. As shown in Figure 6, myc-ABT1 co-immunoprecipitated with FLAG-IGHMBP2, indicating that there is a physical association between IGHMBP2 and ABT1. However, co-IP experiments showed that IGHMBP2 did not interact with ZNF322a, the other protein-coding gene found in the 27k3 modifier (Supplementary Material, Fig. S2).

**DISCUSSION**

Dysregulation of pre-mRNA, rRNA or tRNA pathways is emerging as an important feature of neurodegenerative disorders and in particular in degeneration of motor neurons or their axons.

SMA is caused by reduced SMN protein levels, and in the SMA mouse, this reduction results in generalized pre-mRNA splicing defects (42). Most cells, except motor neurons, apparently tolerate such splicing defects caused by reduced SMN levels. It is also possible that in motor neurons, SMN may have additional, as yet unidentified functions that promote axonal and neuronal survival.

Charcot–Marie–Tooth (CMT) neuropathies are common diseases of the peripheral nervous system that are caused by degeneration of axons (either axonal or demyelinating) of motor neurons or sensory neurons. Mutations of the tyrosyl-tRNA synthetase (also known as YARS), the enzyme that aminoacylates tRNA\textsuperscript{Tyr}, cause dominant intermediate CMT neuropathy type C (DI-CMTC) (43). Mutations of glycyl-tRNA synthetase (also known as GARS), the enzyme that aminoacylates tRNA\textsuperscript{Gly}, cause CMT neuropathy type 2D (CMTD2) and distal spinal muscular atrophy type V (dSMA-V) (44,45). Recent studies indicate that dSMA-V and CMTD2 are best characterized as primary motor neuronopathies (45). These findings strongly indicate that disruption of tRNA function may underlie neurodegeneration.

Amyotrophic lateral sclerosis type 4 (ALS4) is caused by mutations of senataxin (46). Senataxin has an RNA helicase domain that is similar to that of IGHMBP2 and shows significant similarity to the yeast helicase Sen1p, which is involved in the processing and maturation of tRNAs and other small noncoding RNAs (47). Senataxin is localized to the cytoplasm.
and in the nucleolus of neurons, suggesting a function in ribosomal RNA biogenesis and/or translation (48).

Two RNA-binding proteins are also major players in the etiopathogenesis of sporadic and familial ALS. Cytoplasmic aggregation of the TARDBP RNA-binding protein [TAR-DNA-binding protein 43 (TDP 43)] is found in ALS motor neurons (49), and mutations of TARDBP cause familial ALS (50–53). Mutations of FUS (fused in sarcoma), another RNA-binding protein, cause familial ALS, and like TARDBP, FUS aggregates in the cytoplasm of affected motor neurons (54,55).

Our study adds SMARD1 as another motor neuron degenerative disease that is likely caused by dysregulation of tRNA and rRNA pathways. Mice contain 10 tRNA Tyr genes, all of which contain introns (13). Intriguingly, we found that this cluster of the five tRNA Tyr genes is present in yeast to humans, they contain introns (13). Another exceptional feature of tRNA Tyr is the presence of a pseudo-uridine modification of the uridine that is found in the center of the GUA anticodon in all tRNA Tyr from yeast to humans (Fig. 3C) and which is required for proper codon recognition (56). Furthermore, this modification is dependent on the presence of the intron and leads to the hypothesis that the strong evolutionary pressure to maintain the intron in virtually all tRNATyr from yeast to humans may be to direct this pseudo-uridine modification of the uridine that is found in the center of the GUA anticodon (56,57). In mice, half of all tRNA Tyr genes are located in an ~8 kb region of chromosome 13 (13). Intriguingly, we found that this cluster of the five tRNA Tyr genes is present in the BAC-27k3 transgene that rescues motor neuron degeneration of nmd mice, and tRNA Tyr physically associates with TATA-binding protein and to stimulate transcription on the basis of its ability to associate with TATA-binding protein and to stimulate weakly the activity of reporter genes (68). However, subsequent protein–protein interaction studies and localization studies indicated that ABT1, like its yeast homolog, functions in ribosome biogenesis (41). The yeast homolog of ABT1 is called Esf2p (for eighteen S factor 2 protein), is an essential protein and biochemical and genetic studies have unequivocally demonstrated that it functions in ribosome biogenesis (41). Specifically, Esf2p associates with U3 snoRNP (small nuclear ribonucleoprotein) and the 5’ ETS (external transcribed spacer) of pre-ribosomal RNA (pre-rRNA) and is essential for early pre-rRNA processing (41). U3 snoRNP contains four proteins that are common to most snoRNPs and also many additional proteins, which are required for its assembly and function. Reptin and Pontin are two of the proteins that associate with U3 snoRNP and are required for its assembly in human cells (22). The yeast homolog of Reptin is also required for the assembly of yeast snoRNPs including yeast U3 snoRNP (69). Our finding that IGHBMP2 also associates with Reptin and Pontin and the fact that the Abt1 gene, whose protein product is required for ribosome biogenesis, is contained in the 27k3 genetic modifier suggest that IGHBMP2 plays a role in ribosome biogenesis or function.

While our manuscript was under revision, Guenther et al. (70) reported that IGHBMP2 associates with ribosomes. Our findings are in support of a role of IGHBMP2 in mRNA translation, and our biochemical and genetic investigations have led...
to a remarkable convergence of the pathways that IGHMBP2 is likely to function, which include ribosome and tRNA function, both of which are essential for mRNA translation. Future work will focus on delineating the molecular mechanism(s) of IGHMBP2 function and its specific requirements for motor neuron maintenance.

MATERIALS AND METHODS

Mice and genotypic selection

Mice were maintained in humidity and temperature-controlled rooms with a 12:12 dark:light cycle. They were given an NIH-mouse/rat diet with 6% fat, except the ALR/LtJ strain, which received 4% diet (SK54, PMI Feeds, Inc., St Louis, MO, USA) ad libitum with free access to water (HCl-acidified, pH 2.8–3.2). All procedures performed on the animals were reviewed and approved by our institutional animal care and use committee. The C57BL/6J–Mm\textsuperscript{C} congenic strain [B6.CAST-(D13Mit17-D13Mit14)/Cx] was constructed by 10 backcross generations of a (B6.BKS--/+\textsuperscript{nm}\textsuperscript{B}/Mm\textsuperscript{C})\textsuperscript{+} mouse to C57BL/6J as described previously (10). For construction of a high-resolution genetic map of the Mm\textsuperscript{C} locus, 3528 (B6.CAST--/+\textsuperscript{nm}\textsuperscript{B}/Mm\textsuperscript{C})\textsuperscript{+} F\textsubscript{2} mice were phenotyped and genotyped to identify recombinants using the flanking SSLP markers CA1 (27k3CA1F CACGGAGGGAGGAGGGAAAC and 27k3CA1R TGGAGCCTGTTCATCA TCCACC) and CAGA (CAGA-F2 ATTTGTCACTCCCC TCCCTGCCC and CAGA-R2 CTAAAACCTTCAAAGCAAA CGTGGG). F\textsubscript{2} mice containing recombinant Mm\textsuperscript{B}/Mm\textsuperscript{C} genotypes in the absence of a homozygous nm\textsuperscript{B}/nm\textsuperscript{B} mutation (+/+ or +/nm\textsuperscript{B}) were mated to known +/nm\textsuperscript{B} heterozygotes until mutant nm\textsuperscript{B} progeny appeared to determine the status of the Chr 13 Mm\textsuperscript{B} modifier allele.

Generation and characterization of transgenic BAC-27K3 mice

Screening of CHORI-26 CAST BAC library by Southern blot hybridization was performed by TIL Molecular Biology Service using five probes derived from B6 sequences across the genetic interval (primer sequences available on request). CHORI-26 BAC 27K3 was microinjected into fertilized C57BL/6J embryos to generate eight independent transgenic CHORI-26 CAST BAC library by Southern blot hybridization was performed by TIL Molecular Biology Service using five probes derived from B6 sequences across the genetic interval (primer sequences available on request). CHORI-26 BAC 27K3 was microinjected into fertilized C57BL/6J embryos to generate eight independent transgenic

DNA sequencing

EcoRI fragments of CHORI-26 CAST BAC clone 27K3 were subcloned into Invitrogen pcR-TOPO vector and were end-sequenced with vector primers T3 and T7 on an Applied Biosystems 3700 DNA sequencer. Complete sequencing of an 18.1 kb EcoRI fragment containing the Abt1 gene was accomplished using a transposon insertion kit, EZ::Tn-Tn5 TET-1 and EZ::TN KAN-2 according to the manufacturers instructions (Epigenic Technologies), and the sequence information was deposited into GenBank (Accession no. EU880213).

DNA constructs

The open-reading frames of IGHMBP2, ABT1, Zfp161 were cloned downstream of the CMV promoter into modified pcDNA3 vectors (Invitrogen) containing either the myc-tag or the FLAG-tag sequences at the amino terminus. All of the constructs were analyzed by DNA sequencing.

Antibodies

The anti-IGHMBP2 monoclonal antibody mAb11-24 was produced by immunization of BALB/c mice and fusion of spleen cells with Sp2/0 myeloma cells. The mouse sera and the hybridoma culture supernatants were both screened by ELISA, western blot, IP and immunofluorescence. Hybridoma cell lines were cloned to homogeneity by limiting dilution. Ig class was determined using an isotyping kit (Roche). Antibodies used were as follows: 11-24 mouse IgG1 monoclonal anti-IGHMBP2; 2B1, mouse IgG1 monoclonal anti-SMN; 2E17, mouse IgG1 monoclonal anti-Gemin2; 12H12, mouse IgG1 monoclonal anti-Gemin3; 17D10, mouse IgG1 monoclonal anti-Gemin4; 9E10, mouse IgG1 monoclonal anti-myc (Developmental Studies Hybridoma Bank); A-14, affinity-purified rabbit polyclonal anti-myc 9E10 epitope (Santa Cruz Biotechnologies); M2, mouse IgG1 monoclonal anti-Flag (Sigma); anti-TFIIC220 antibody (sc-23110, Santa Cruz, Inc.); anti-α-tubulin antibody (Ab4074, Abcam); anti-β-tubulin antibody (Ab6046, Abcam or E-7 from Developmental Studies Hybridoma Bank); anti-hnRNP H (Bethyl); anti-HSP70 (Abcam); anti-ZNF322a (Aviva).

Cell culture, IP, small RNA cloning

Cells were cultured in Dulbecco’s modified Eagle’s medium (Invitrogen) supplemented with 10% fetal bovine serum (Invitrogen). Cells were transfected by the standard calcium phosphate method. Following overnight incubation with DNA, cells were washed and fresh medium was added. Stable clones of MN1 and 293T were obtained by selection in the same fixative and then processed for plastic embedding. For counts of axons, left and right nerves were taken whenever possible, and counts were averaged so that each n represents one mouse and the average count of the left and right nerve. The total number of myelinated axons in each nerve was counted using light microscopy on 0.5 μm sections stained with Toluidine blue. Images were taken on a Nikon E600 microscope using a Leica DFC300 FX digital camera.
presence of G418 (100 μg/ml). IP experiments were performed from total cell extracts followed by western blotting as described previously (36). For small RNA cloning, RNA was isolated from FLAG-IGHMBP2 immunoprecipitates; 3′-end labeling, directional adapter ligation, cloning and sequencing were performed as described previously (38).

**In vitro translation and IP**

FLAG-Zfp161, myc-IGHMBP2 and FLAG-IGHMBP2 proteins were generated by *in vitro* translation using the TNT T7 Quick-Coupled Transcription/Translation System in the presence of 35S-methionine (Promega) following the manufacturing instructions. The *in vitro* translated proteins were mixed and allowed to interact for 1 h at 4°C. IPs were carried out using either anti-myc antibody (9E10) or anti-FLAG M2 beads (Sigma). After binding, the beads were washed five times with RSB-100 buffer (20 mM Tris–HCl, pH 7.5, 1200 mM NaCl, 2.5 mM MgCl2, 0.5% Triton X100 and complete EDTA-free protease inhibitors) and boiled in SDS–PAGE loading buffer. The IPs were analyzed by NuPAGE and exposed to a Biomax XAR film (Kodak).

**Affinity purification of IGHMBP2 protein complex**

FLAG-IGHMBP2 and control cell lines were grown in the presence of G418 (100 μg/ml). Total cell extracts were prepared by resuspending cell pellets in RSB-100 (10 mM Tris–HCl, pH 7.4, 2.5 mM MgCl2, 100 mM NaCl) buffer containing 0.1% Nonidet P-40 and protease inhibitors. Cells were sonicated briefly. Following centrifugation at 10 000 rpm for 15 min, supernatants were added to anti-FLAG beads (Sigma) pre-washed with the same buffer. Extracts were incubated with anti-FLAG beads for 2 h at 4°C. Supernatants were discarded, and beads were extensively washed with RSB-100 containing 0.02% Nonidet P-40. Three high-salt washes were performed with 10 bed volumes of RSB-300 (300 mM NaCl) buffer containing 0.02% Nonidet P-40 for 15 min at 4°C. Following three washes with RSB-100 containing 0.02% Nonidet P-40, bound complexes were eluted with 10 bed volumes of the same buffer containing 0.25–0.5 mg/ml 3X-FLAG peptide (Sigma) for 1 h at 4°C.

**Mass spectrometry**

Bands were excised from Coomassie-stained NuPAGE gel, *in gel*-digested with trypsin and peptides were identified by tandem MS-MS. The peptide sequences obtained from mass spectrometry for selected proteins that co-immunoprecipitate with FLAG-IGHMBP2 (Fig. 2) were RuB-like 2 (Reptin/Tip49α, Tip48; Q9Y230): AVLIAGQPGTGK, VYSLFLDESR, AVFVDLEPTVIDEVR; HSP70 (P08107): EIDLVLDR, AVFDLEPTVIDEVR; HSP70 (P08107): SAVEDEGLK, LLQDDFFNGR, VEILANDQGNR, TTPSY-VAFDTDTER; Zfp-161 (O43829): HENNMHSER.

**Reverse transcriptase–PCR**

Total RNA was extracted from forebrain or spinal cord of wild-type, nmd or nmd-Tg27K3 mice using TRIzol reagent (Invitrogen). RNA was treated with DNase I (Roche) and reverse-transcribed with oligo-dT using Advantage RT-for-PCR kit (Clontech). The cDNA was used as a template for PCR with primers E4F (5′-CGCGTGAAGAGCAGCTGATGAC) and E5R (5′-TGTTGT TAGAAGAGAGTGGGG), and the PCR products were analyzed on 2% Agarose electrophoresis gel.

**Immunofluorescence**

NT2 cells plated on glass coverslips were briefly washed with PBS, fixed in 2% formaldehyde-PBS for 30 min at room temperature and permeabilized in 0.5% Triton X-100-PBS for 5 min at room temperature. Cells were blocked in 3% bovine serum albumin in PBS for 30 min at room temperature. Double-label immunofluorescence experiments were performed by separate sequential incubations of each primary antibody diluted in PBS containing 3% bovine serum albumin followed by the specific secondary antibody coupled to fluorescein isothiocyanate or TRITC. All incubations were at room temperature for 1 h.

**SUPPLEMENTARY MATERIAL**

Supplementary Material is available at *HMG* online.

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