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How the discovery of ISS-N1 led to the first medical therapy for spinal muscular atrophy

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Abstract

Spinal muscular atrophy (SMA), a prominent genetic disease of infant mortality, is caused by low levels of survival motor neuron (SMN) protein owing to deletions or mutations of the *SMN1* gene. *SMN2*, a nearly identical copy of *SMN1* present in humans, cannot compensate for the loss of *SMN1* due to predominant skipping of exon 7 during pre-mRNA splicing. With the recent FDA approval of nusinersen (SpinrazaTM), the potential for correction of *SMN2* exon 7 splicing as a SMA therapy has been affirmed. Nusinersen is an antisense oligonucleotide that targets intronic splicing silencer N1 (ISS-N1) discovered in 2004 at the University of Massachusetts Medical School. ISS-N1 has emerged as the model target for testing the therapeutic efficacy of antisense oligonucleotides using different chemistries as well as different mouse models of SMA. Here we provide a historical account of events that led to the discovery of ISS-N1 and describe the impact of independent validations that raised the profile of ISS-N1 as one of the most potent antisense targets for the treatment of a genetic disease. Recent approval of nusinersen provides a much-needed boost for antisense technology that is just beginning to realize its potential. Beyond treating SMA, the ISS-N1 target offers myriad potentials for perfecting various aspects of the nucleic-acid-based technology for the amelioration of the countless number of pathological conditions.

Introduction

Spinal muscular atrophy (SMA) is a prominent and devastating genetic cause of infant mortality with an incidence of ~1 in 10,000 live births.^{1,2} SMA results from low levels of survival motor neuron (SMN) protein due to deletions of or mutations in the *SMN1* gene.^{3,4} SMN is an essential protein involved in diverse functions including snRNP assembly, snoRNP assembly, DNA repair, transcription, telomerase biogenesis, translation, RNA trafficking, selenoprotein synthesis, stress granule formation and cell signaling.⁵ While

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The ISS-N1 target (US7838657) was discovered in the Singh laboratory at UMass Medical School (MA, USA). Inventors, including RN Singh, NN Singh, EJA and UMASS Medical School, are currently benefiting from licensing of the ISS-N1 target to Ionis Pharmaceuticals. Iowa State University holds intellectual property rights on GCRS and ISS-N2 targets. Therefore, inventors including RN Singh, NN Singh and Iowa State University could potentially benefit from any future commercial exploitation of GCRS and ISS-N2 targets.

motor neurons and neuromuscular junctions are the first to be affected in SMA,⁶ other tissues including the cardiovascular system, lung, bone, intestine, liver, pancreas, spleen and testis are also impacted by low levels of SMN.^{7–18} The disease spectrum of SMA, divided into four types (I through IV) is broad and ranges from infant mortality to adult onset.¹ In general, SMA severity correlates with *SMN2* copy number: the lower the *SMN2* copy number, the lower the SMN and the more severe the disease.¹ Lorson and Hahnen, working in the Androphy laboratory at Tufts Medical School, found that *SMN2*, a nearly identical copy of *SMN1*, fails to compensate for the loss of *SMN1* due to defective splicing.¹⁹ The skipping of *SMN2* exon 7 is attributed to a silent c.840C>T transition that corresponds to the 6th position of the 54-nucleotide (54-nt)-long exon 7.^{19,20} Since this silent mutation does not affect the protein sequence, correction of *SMN2* exon 7 splicing has the potential to restore the fully functional SMN protein.^{21,22} The *SMN2* exon 7-skipped mRNA itself leads to the production of SMN 7, a partially stable protein.^{23–26}

Pre-mRNA splicing is a complex process involving multiple RNA-protein, protein-protein and RNA-RNA interactions unique for the removal of each individual intron.²⁷ Several cis-elements and transacting factors have been implicated in regulation of *SMN* exon 7 splicing. Many recent reviews describe the mechanism of splicing regulation of *SMN* exon 7.^{28–30} The generation and nature of various splice variants of *SMN* have been described elsewhere.^{31–34} Here we provide background information pertaining to the discovery of the RNA regulatory element that produced nusinersen (SpinrazaTM), the first US Food and Drug Administration (FDA)-approved drug for SMA.³⁵ Several investigations had demonstrated that C6U weakens the 3'-splice site (3' ss) of exon 7 due to its close proximity to the 3' ss.^{36–38} However, a subsequent analysis using live cells expressing an exon 7 splicing cassette revealed that a weak 5' ss is also a limiting factor for inclusion of exon 7.³⁹ In vivo selection is an unbiased approach in which the significance of every nucleotide at each exonic position is tested employing a large pool of unique molecules (>10¹²) with random mutations.^{40,41} Surprisingly, the results of in vivo selection revealed that an adenosine residue at the last position (A54) of exon 7 has a more detrimental effect on *SMN2* exon 7 splicing than the natural C6U substitution known to cause *SMN2* exon 7 skipping.³⁹ In agreement with the results of in vivo selection, an A54G substitution restored exon 7 inclusion even in the absence of positive cis-elements considered to be important for inclusion of exon 7 in both *SMN1* and *SMN2*.³⁹ The strong stimulatory effect of A54G on *SMN2* exon 7 splicing was attributed to several factors, including the disruption of an inhibitory structure that sequesters the 5' ss of exon 7 and/or lengthening of the RNA:RNA duplex formed between U1 snRNA and the 5' ss of exon 7.^{39,40} Both of these mechanisms would result in enhanced recruitment of U1 snRNP at the 5' ss of exon 7. Indeed, a follow up study confirmed the presence of an inhibitory terminal stem-loop structure (TSL2) that facilitates *SMN2* exon 7 skipping by sequestering the 5' ss of this exon.⁴² It was also confirmed that increasing the size of the duplex formed between U1 snRNA and the 5' ss of exon 7 has a strong stimulatory effect on *SMN2* exon 7 inclusion.⁴²

These insights into the processing of the exon 7 *SMN2* mRNA brought immediate attention to the role of the 5' ss in regulation of *SMN* exon 7 splicing. Subsequent studies in the Singh and Androphy labs at the University of Massachusetts Medical School (UMMS) in Worcester, MA, USA discovered Intronic Splicing Silencer N1 (ISS-N1) as the major

inhibitory element in *SMN* exon 7 splicing regulation.^{35,43,44} Here we describe how the discovery of ISS-N1 transformed our understanding of *SMN* exon 7 splicing regulation and paved the way for an antisense-oligonucleotide (ASO)-based therapy for SMA. Nusinersen (Spinraza™), the recently FDA-approved drug for SMA, is an ISS-N1 targeting ASO (Figure 1). We will present what lessons can be learnt from the successes of Spinraza™ for future ASO-based therapies of SMA and other diseases.

Defining features of the ISS-N1 target

ISS-N1 is 15-nt long motif located immediately downstream of the 5' splice site of exon 7. Deletion of ISS-N1 fully restored inclusion of *SMN2* exon 7.⁴³ Interestingly, the effect of ISS-N1 deletion was similar to what was observed with A54G mutation, since positive cis-elements became dispensable for exon 7 inclusion when ISS-N1 was deleted.^{39,43} Hence, Baralle and colleagues termed ISS-N1 as the master checkpoint of *SMN2* exon 7 splicing.⁴⁵ Consistent with the strong inhibitory effect of ISS-N1, an ASO complementary to and thereby blocking ISS-N1 fully restored *SMN2* exon 7 inclusion in cultured fibroblasts derived from the SMA patient.⁴³ The ASOs employed in this study carried phosphorothioate backbone and the 2'-O-methyl (2'OMe) modifications in all sugar residues. The stimulatory effect of the ISS-N1-targeting ASO was robust even at the very low concentration of 5 nM, suggesting that the ISS-N1 target is structurally accessible.⁴³ Demonstrating the target specificity, two mismatch mutations within the ISS-N1 target as well as in an ASO that targeted it, completely abolished the stimulatory effect on splicing of exon 7.⁴³ Additional validation of specificity came from an antisense walk in which the effect of three control ASOs that annealed to sequences downstream of ISS-N1 were evaluated. As expected, none of these control ASOs had any appreciable effect on *SMN2* exon 7 splicing.⁴³ These observations presented a rare example of a highly specific ASO with desired high activity at a record low nanomolar concentration. Equally significant was the target location within an intron, since an ASO annealing to an intronic sequence cannot have an adverse effect on mRNA export and protein translation. Consistent with the splicing correction, the ISS-N1-targeting ASO increased SMN protein in human SMA derived fibroblasts at the low nanomolar concentration.⁴³ Overall, these findings were highly instructive in suggesting that the devastating consequences of exon 7 skipping could be fully overcome by abrogating the activity of an intronic inhibitory element. For commercial exploitation and drug development, UMMS filed for intellectual property rights with the US Patent Office in 2004 and subsequently secured a series of patents on the ISS-N1 target.^{35,44}

At the time the report on the ISS-N1 target was published in 2006, Ionis Pharmaceuticals (previously ISIS Pharmaceuticals, Carlsbad, CA, USA), in collaboration with the Krainer group (Cold Spring Harbor Laboratory, Long Island, NY, USA), was working to develop an ASO-based therapy for SMA. In 2007 Ionis Pharmaceuticals presented the results of screening of a large number of ASOs that targeted exon 7.⁴⁶ The ASOs used in these experiments carried phosphorothioate backbone and the 2'-O-methoxyethyl (MOE) modifications in all sugar residues. While the findings of this study validated the outcome of in vivo selection of exon 7, none of the ASOs identified emerged as a contender for therapeutic application. Hoping to find a better lead candidate, Ionis Pharmaceuticals continued with another MOE ASO library screen, this time targeting intronic sequences

upstream and downstream of exon 7 including the ISS-N1 region.⁴⁷ This study was particularly significant since it allowed a side-by-side comparison of an ISS-N1-targeting ASO with a likely novel lead ASO. The ISS-N1-targeting ASO (ASO 10–27) emerged as the best ASO in this study.⁴⁷ Thus, ISS-N1 became the first antisense target for SMA therapy to be independently validated by employing an ASO with different chemistry.

Encouraged by these findings, the Krainer laboratory in collaboration with Ionis Pharmaceuticals and others, performed a series of in vivo studies with the ISS-N1-targeting ASO (ASO 10–27).^{48–50} The results of these studies showed its unprecedented therapeutic efficacy, including the record lifespan extension benefits in the severe SMA models (described below). While in vivo studies on the ISS-N1 target were still underway, Ionis Pharmaceuticals achieved another milestone in 2010 by securing an exclusive license from UMMS to develop SMA drug based on the ISS-N1 target.³⁵ These advances put Ionis Pharmaceuticals in 2011 at the forefront of launching a clinical trial of an ISS-N1-targeting drug (ISIS-SMN_{RX}) for the treatment of SMA. Subsequent years also witnessed a series of in vivo studies employing phosphorodiamidate morpholino oligonucleotides (PMOs) targeting ISS-N1.^{13,15,51–54} These studies provided additional independent validations of the efficacy of ISS-N1-targeting ASOs.

Mechanism of Action of ISS-N1-targeting ASOs

The mechanism by which an ISS-N1-targeting ASO stimulates *SMN2* exon 7 inclusion appears to be complex. Sequences downstream of ISS-N1 harbor TIA1 binding sites.⁵⁵ Overexpression of TIA1 restores inclusion of *SMN2* exon 7, suggesting that factor(s) interacting with ISS-N1 interfere with the recruitment of this protein.⁵⁵ Owing to the presence of two hnRNP A1/A2 motifs spanning the last 14 positions of the 15-nt long ISS-N1, it has been proposed that the inhibitory effect of ISS-N1 is due to interaction of ISS-N1 with hnRNP A1/A2.⁴⁷ ISS-N1 overlaps with an 8-nt long GC-rich sequence (GCRS) spanning from the 7th to 14th positions of intron 7.⁵⁶ Sequestration of GCRS by an 8-mer 2'OMe ASO also restores *SMN2* exon 7 inclusion. The GCRS-targeting ASO partially sequestered only one of the two putative hnRNP A1/A2 motifs and yet this ASO produced the robust stimulatory effect on *SMN2* exon 7 splicing. These surveys suggest that ISS-N1 may be a composite landing site for several splicing factors.

The most compelling evidence against the role of hnRNP A1/A2 in the negative effect of ISS-N1 came from a systematic study employing two 14-mer 2'OMe ASOs called L14 and F14.⁵⁷ F14 and L14 bind to the first and the last fourteen nucleotides of the 15-nt long ISS-N1, respectively. As expected, F14 restored *SMN2* exon 7 inclusion. Surprisingly, L14 (which sequestered both hnRNP A1/A2 sites but not the 1st position of ISS-N1) strongly inhibited *SMN2* exon 7 splicing.⁵⁷ Similar results were observed with two 14-mer ASOs with different chemistry, namely locked nucleic acid, suggesting that chemistry is not the cause of the L14-induced skipping of *SMN2* exon 7.⁵⁷ These observations ruled out the role of hnRNP A1/A2 as the sole regulator(s) associated with the negative effect of ISS-N1 and called for a unique mechanism in which sequestration of the first position of ISS-N1 is necessary for the stimulatory effect of an ISS-N1 targeting ASO.

The opposite effect of F14 and L14 on exon 7 splicing was investigated in a recent study where the 1st position of ISS-N1 was shown to be involved in the formation of a unique long-distance interaction (LDI) with a deep intronic sequence termed ISS-N2 (Figure 2).⁵⁸ In particular, an internal stem formed by a LDI (ISTL1) sequesters the 1st position of ISS-N1 in addition to sequestering a portion of the 5' ss of exon 7.⁵⁹ An ISS-N1-targeting ASO was found to disrupt ISTL1 and increase the accessibility of the 5' ss of exon 7 for the recruitment of U1 snRNP.⁵⁹ Supporting the inhibitory role of ISTL1, an ISS-N2-targeting ASO that disrupted ISTL1, also produced a strong stimulatory effect on *SMN2* exon 7 splicing.⁵⁸ Hence, the stimulatory effect of an ISS-N1-targeting ASO is at least in part due to the breaking of the inhibitory structure, ISTL1. It is also likely that the sequestration of ISS-N1 promotes recruitment of TIA1 at sequences downstream of ISS-N1.⁵⁹ In turn, an enhanced binding of TIA1 is likely to promote recruitment of U1 snRNP to the 5' ss of exon 7 causing its inclusion similarly as recently described.^{59,60} Longer morpholino ASOs that target ISS-N1 as well as downstream sequences also stimulated *SMN2* exon 7 splicing.⁵² These results suggest that the recruitment of TIA1 downstream of ISS-N1 may not be critical once the ISS-N1 site is sequestered and ISTL1 is disrupted. However, longer ASOs may bring additional changes within the intron 7 structure and these changes may obviate the requirement for TIA1 in *SMN* exon 7 splicing regulation.

In vivo studies with ISS-N1-targeting ASOs

ISS-N1-targeting ASOs have shown remarkable efficacy in ameliorating the phenotype of several SMA mouse models when administered soon after birth (Table 1).^{13,15,48–54,61–72} Initially, the 2' OMe ISS-N1-targeting ASO⁴³ increased SMN protein in the central nervous system (CNS) and improved motor function in the α -7 mouse.⁶¹ This finding led to studies from Krainer and colleagues, and independently from the Burghes (The University of Ohio School of Medicine, Columbus, OH, USA) and Muntoni groups (University College London, London, UK), to further explore the efficacy of the ISS-N1 inhibitor, albeit they used contrasting delivery strategies and ASOs of different lengths and chemistries.^{48,50–53} The Krainer group first published results using an 18-mer MOE ASO (targeted to intron 7 nucleotides 10 to 27) in Taiwanese SMA mice, which can be bred to express only the human *SMN2* transcript.^{48,50} Remarkably, in these mice subcutaneous (SC) administration of the MOE ASO extended median lifespan 25-fold and improved neuromuscular junction development.⁵⁰ This survival benefit remains one of the greatest increases observed for all SMA treatments.²¹ Interestingly, MOE ASO delivery directly to the CNS provided relatively little survival benefit.⁵⁰ Comparatively, the Burghes group observed that a single 27, 54 or 81 μ g intracerebroventricular (ICV) dose of a 20-mer ISS-N1-targeting PMO (targeted to intron 7 nucleotides 10 to 29) increased median lifespan nearly 7-fold non-dose dependently in the relatively less severe α -7 mouse model, which is transgenic for human *SMN2* plus the *SMN* γ cDNA.⁵¹ It remains unclear which ASO chemistry, length and route of delivery would provide the most efficacious treatment strategy for SMA. ICV injection is relatively invasive, but provides targeted ASO delivery to the CNS. Since the blood brain barrier (BBB) in neonatal mice is leaky, ASO delivered systemically could enter the CNS. However, BBB leakiness is much less prominent in humans and systemic administration (SC or intravascular) would most likely require a high dose to effectively distribute the ASO in the

human CNS. High doses could potentially produce toxic effects especially in the liver and kidneys. The Krainer laboratory aimed to determine whether SMN restoration in the CNS is required to rescue the SMA phenotype in Taiwanese mice.⁶⁷ SC delivery of an 18-mer ISS-N1-targeting MOE ASO accompanied with an ICV administration of a decoy MOE ASO complementary to the ISS-N1 targeting ASO mitigated the SMA phenotype similarly to SC 18-mer ISS-N1-targeting MOE ASO administration alone.⁶⁷ Thus, a SMN increase in peripheral organs rather than strictly in the CNS appears important to the therapeutic outcome. This finding is not entirely surprising, since a host of recent studies have revealed peripheral organ defects in mouse models of SMA.⁵ Overall, regardless of their chemistry and length, ISS-N1-targeting ASOs consistently show the greatest efficacy in improving the phenotype of SMA mouse models.²¹ Collectively, these findings confirm the potency of the ISS-N1 target in stimulating *SMN2* exon 7 inclusion and for the treatment of SMA. Because of this potency, the ISS-N1 target can be useful in investigating and perfecting emerging antisense technologies, including chemical modifications that could target the ASO to specific organs. For instance, Wood and colleagues have recently reported promising CNS delivery of systemically-administered peptide-conjugated PMOs.^{70,71} However, the pharmacokinetics and tolerability of these conjugated ASOs remains to be further examined in non-human primates. Nevertheless, these modifications could be particularly informative for other genetic diseases in which nucleic-acid-based treatment may be beneficial.

ISS-N1-targeting ASOs also provide a powerful tool to further understand SMA disease mechanisms. Using ISS-N1-targeting ASOs allows researchers to examine the temporal impact of increasing SMN protein. For example, recent studies show impaired intestine and liver development in Taiwanese mice.^{13,15} SC delivery of a 25-mer PMO targeted to ISS-N1 normalized the development of these organs indicating that SMN is required for their development.^{12,14} ISS-N1-targeting ASOs can also validate biomarkers that could be examined in SMA patients in clinical trials. Namely, electrophysiological abnormalities and serum biomarkers in the *7* mouse are partially corrected with administration of a 20-mer ISS-N1-targeting PMO.^{63,65,66} Finally, these ISS-N1-targeting ASOs may also develop new models of the disease. The Wirth group generated an intermediate SMA mouse model through SC administration of suboptimal doses of the 18-mer ISS-N1-targeting MOE ASO in severe Taiwanese mice.⁷³ This new model allowed for examination of the benefit of the SMA genetic modifier *plastin3*.⁷³ Intermediate models of the disease would also allow one to capture potential defects in organ development that may not be observed in severe models with early postnatal lethality.

Therapeutic development

Based on the strong pre-clinical data on the efficacy of ISS-N1, Ionis Pharmaceuticals commenced clinical trials with nusinersen, an 18-mer ASO with their proprietary MOE chemistry. Given the neuromuscular nature of SMA, nusinersen was delivered intrathecally through a lumbar puncture for these clinical trials.^{74–76} Lumbar puncture effectively distributed nusinersen throughout the CNS and did not produce adverse effects beyond what had been previously reported for this procedure.⁷⁵ In the phase I study, nusinersen was well-tolerated and led to an increase in SMN protein in the cerebrospinal fluid at 9–14 months post treatment.⁷⁴ In addition, the highest dose (9 mg) increased motor function up to 9–14

months post treatment as assessed by the Hammersmith Functional Motor Score Expanded (HFMSE). The HFSME score increase was particularly promising since it constituted what would be considered a clinically meaningful outcome and led to commencement of larger phase 2 and 3 clinical trials.⁷⁴ A phase 2 clinical trial enrolled severe type I SMA infants with symptom onset between 3 weeks and 4 months.⁷⁶ Multiple doses of nusinersen were well-tolerated and the majority of the adverse effects were mild. Nusinersen was broadly distributed throughout the CNS, increased full-length *SMN2* transcript and SMN protein and most infants exhibited improved motor function as well as survival and electrophysiological function.⁷⁶ The promising results from these clinical trials led to a placebo-controlled double-blinded phase III clinical trials. While the results have yet to be published, the trials were terminated early because the primary endpoint at interim analysis was met;⁷⁷ FDA approved Spinraza™ on December 23, 2016. Hence, Spinraza™ became the first FDA-approved drug for SMA as well as the first antisense drug to treat the major population of a genetic disease through splicing correction.

Conclusions and Future directions

Correction of *SMN2* exon 7 splicing has long been considered as the most efficacious therapeutic option for most SMA patients who retain at least one copy of the *SMN2* allele. The specificity with which an oligonucleotide corrects splicing remains unmatched. The major obstacle in the way of an effective ASO-based therapy of SMA is the delivery of the oligonucleotide across the BBB. Other concerns include the stability and the tolerability of the oligonucleotide. With the recent FDA approval of Spinraza™, much of the skepticism of the oligonucleotide-based therapy of SMA has been put to the rest. Animal studies underscore that an early restoration of SMN is key to achieving the maximum therapeutic benefit. The availability of Spinraza™ provides SMA patients with their first opportunity to receive a drug that could increase SMN levels. This development puts SMA among a handful of rare diseases with at least one FDA-approved drug. This is also a great win for nucleic-acid therapeutics, which holds the promise for the treatment of rare and orphan diseases.

Despite the anticipated enthusiasm among the SMA community and caregivers, it is understandable that Spinraza™ may not act with equal efficacy in all SMA patients. This is due to a variety of reasons including a limited number of *SMN2* alleles, delayed age of drug administration and the presence or absence of other disease modifying factors. There may be some instances in which Spinraza™ may be completely unusable due to an acute immune response. These concerns call for a continued progress towards developing alternative therapies for SMA. Fortunately, all oligonucleotide chemistries incorporated into ISS-N1-targeting ASOs have shown promising results. These independent validations of target efficacy suggest bright prospects for developing advanced ASO-based SMA therapies. We now know that low levels of SMN affect most tissues. The launch of Spinraza™ has invigorated the field of nucleic acid-based therapeutics to further translate the potential use of ASOs targeting other regulatory elements in *SMN2*, and modifications that enable penetration into all tissues when delivered employing noninvasive procedures. Recent reports have also validated the efficacy of additional antisense targets, including Element 1 in *SMN2* intron 6⁷⁸ and ISS-N2 in *SMN2* intron 7⁷⁹ in SMA mouse models. In addition, the

potential for dual-masking ASOs that simultaneously sequester two targets could increase the repertoire of ASOs for SMA therapy.⁸⁰ Given the spectrum of the SMA phenotype, it will be useful to have multiple ASO targets so as to treat best treat patients who may not respond to specific ASOs.

Success of an ASO-based therapy for SMA has implications for other diseases in which nucleic acid-based therapy remains an option. Progress thus far exhibits a classic example of smooth transition from fundamental discovery of the ISS-N1 target to independent pre-clinical validations of ISS-N1-targeting ASOs to independent clinical trials of Spinraza™. The coming years will be critical for evaluating the long-term efficacy of Spinraza™ in large cohorts comprised of the appropriate age-matched patients. However, there remains a concern of drug affordability as the high cost of Spinraza™ might limit the number of patients who might otherwise benefit from this drug. Irrespective of these concerns, there is every reason to believe that tomorrow's SMA patients will have more options thanks in part to the new insights brought forth from the basic investigations into the regulatory elements that control *SMN2* exon 7 alternative splicing.

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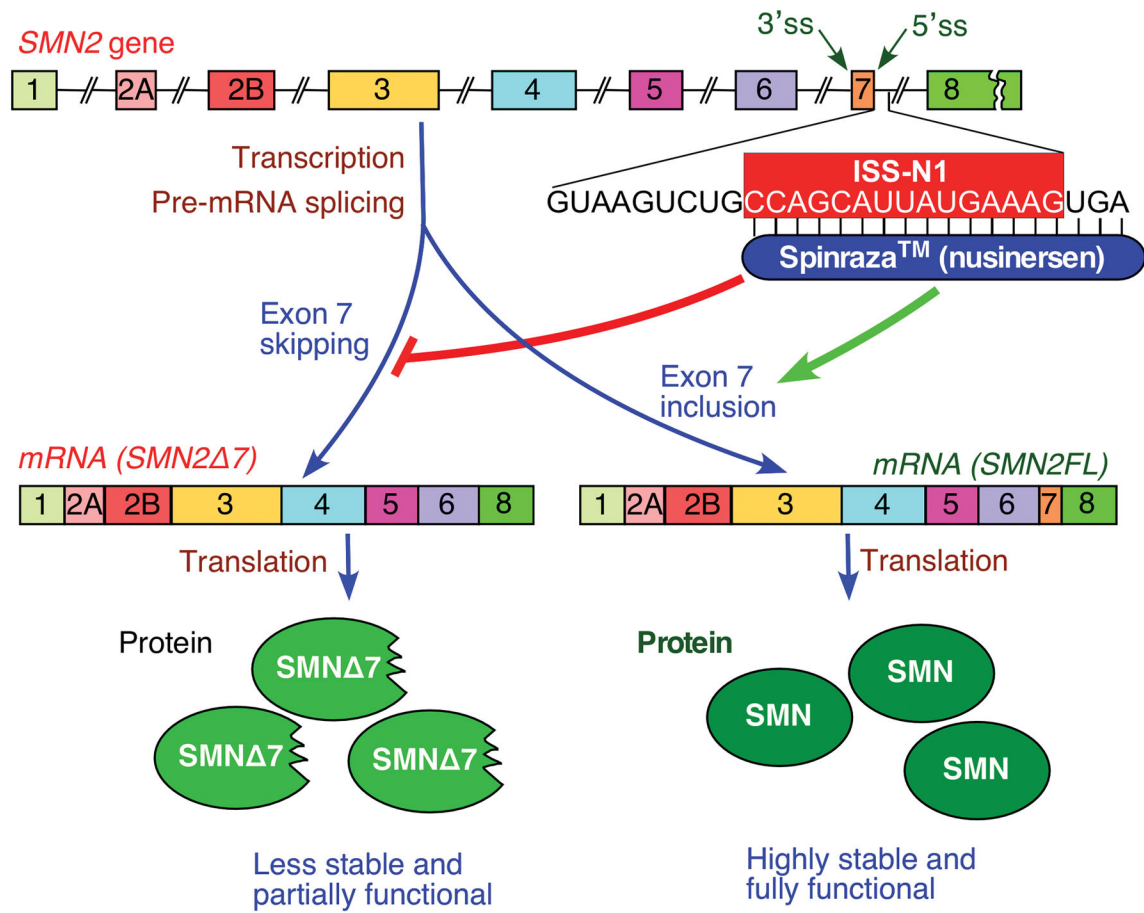


Figure 1.

Diagrammatic representation of *SMN2* gene and Spinraza mode of action. *SMN2* exons are represented by colored boxes, while introns are shown as broken lines. Intronic sequence immediately downstream of exon 7 is given. ISS-N1 region within this sequence is highlighted in pink box. Positions to which Spinraza anneals are indicated. *SMN2* pre-mRNA splicing results in exon 7-included (*SMN2**FL*) and exon 7-skipped (*SMN2* Δ 7) transcripts, translation of which leads to production of the full length functional SMN protein and a truncated less stable isoform, respectively. Targeting of ISS-N1 by Spinraza prevents exon 7 skipping and as a consequence increases levels of the full length SMN.

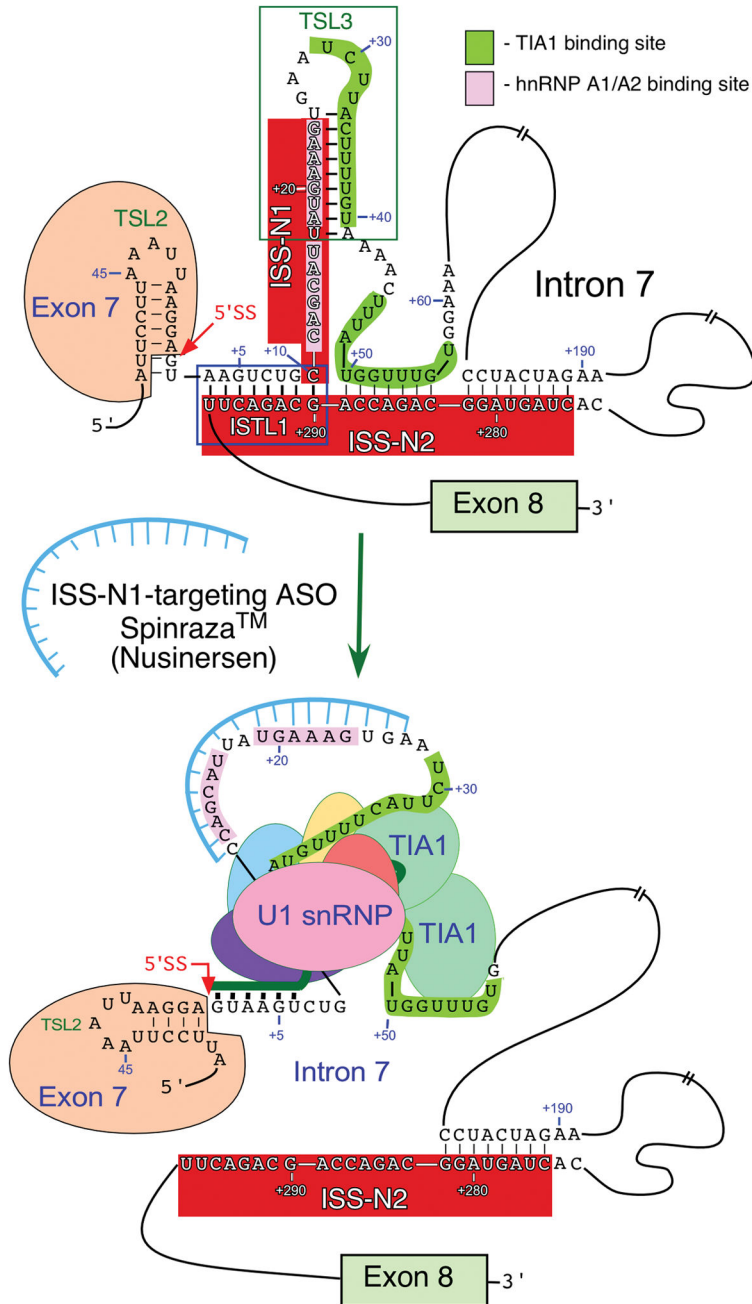


Figure 2.

Mechanism of exon 7 splicing correction by an ISS-N1 targeting ASO. Only relevant portions of exon 7 and intron 7 are shown. Exonic and intronic sequences are presented in the context of experimentally derived structures.^{42,58} Neutral and positive numbering start from the first position of exon 7 and the first position of intron 7, respectively. Splicing regulatory cis-elements and structures, such as the 5' ss of exon 7, ISS-N1, ISS-N2, TSL2 and ISTL1 are highlighted. Binding sites for hnRNPA1/A2 and TIA1 are indicated. Annealing positions of the ASO and U1 snRNA are shown. Targeting of ISS-N1 by ASO causes structural rearrangements, such as disruption of TSL3 and ISTL1 and blocks the

binding sites of hnRNP A1/A2. As the result TIA1 binding sites become accessible and the recruitment of U1 snRNP to the 5' ss of exon 7 is increased. Abbreviations: ASO, antisense oligonucleotide; ISS, intronic splicing silencer; ISTL, internal stem-loop structure; ss, splice site; TSL, terminal stem-loop structure.

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Table 1

In vivo studies using ISS-N1-targeting ASOs

Chemistry (Length)	Treatment Details	Outcome Measures	Ref
Studies in 7 mouse model			
PS 2'OMe (20)	ICV P1, P3, P5, P7, P10 (1 µg)	Increased SMN protein in the CNS; improved motor function; increased body weight	61
PS MOE (18)	ICV P0 (4 µg)	Median survival increased 16 d to 26 d; improved muscle and NMJ development	49
PS 2OMe (21 ^a)	ICV P1, P3, P5 (1 or 10 µg)	Median survival increased ~10 d to ~20 d; improved motor function	62
PMO (20)	ICV P0 (27, 45 or 81 µg)	Median survival increased 15 d to ~100 d; improved motor function; increased body weight	51
PMO (20, 22 and 25)	ICV P0 (0.5, 1, 2, 4 or 6 mM)	25-mer dose-dependently increased median survival 15 d to 37–126 d	53
PMO (20)	ICV P1 (40 µg)	Age-dependent normalization of CMAP and MUNE	63
PMO and Dendrimeric octoguanadine (Vivo)-PMO (25)	ICV P0, P3 (2 nM)	Median survival increased 17 d to 23 d	64
	SC P0, P3 (10, 12 or 24 nM)	Median survival increased 17 d to ~27 d	
	ICV P0 (2, 5, 10 or 12 nM) and SC P0, P3 (2, 5, 10 or 12 nM)	Median survival dose-dependently increased 17 d to between 40–120 d; combination of highest doses improved NMJ maturation and motor function	
PMO (20)	ICV P0–P2, P4 or P6 (40 µg)	Treatment at P2 improved CMAP, MUNE and EIM measures when examined at P12	65
PMO (20)	ICV P1 (40 µg)	Normalization of several SMA biomarkers	66
Studies in Taiwanese type I mouse model			
PS MOE (18)	ICV P1 (20 µg)	Median survival increased 10 d to 16 d	50
	SC P0–1, P2–3 (40, 50, 80 or 160 µg/g)	Median survival dose-dependently increased 10 d to between 84–248 d; improved NMJ maturation, motor function; motor neuron protection	
	ICV P1 (20 µg) and SC P0–1, P2–3 (50 µg/g)	Median survival increased 10 d to 173 d	
PMO (18 and 25)	ICV P1 (20 or 40 µg/g)	Median lifespan increased 9.5 d to 12 d or 32 d (18-mer) or 43 d or 85.5 d (25-mer)	52
PMO (25)	IV P0 (40 µg/g)	Median lifespan increased 9.5 d to >230 d	
PMO (25)	IV P0 (40 µg/g) and IP or SC P3 (40 µg/g)	Median survival increased 9.5 d to 93.5 d	
Vivo-PMO (25)	IV P0 10 µg/g and IP P3 10 µg/g	Median survival was 16 d	
PMO (25)	ICV P0 (10, 20 or 40 µg/g)	Median survival increased 9.5 d to 22 d (10 µg/g), 24 d (20 µg/g) or 212 d (40 µg/g)	54
	SC P0 (10, 20 or 40 µg/g)	Median survival increased 9.5 d to 22 d (10 µg/g), 58 d (20 µg/g) or 261 d (40 µg/g); improved NMJ and muscle maturation	
	SC P0 (10 µg/g), SC P5 and every 2 weeks (10 µg/g)	Survival increased 9.5 d to >150 d; improved muscle and NMJ maturation	
PS MOE (18)	SC P0, P2 (120 mg/kg per day)	Median survival increased 10 d to 237 d; improved NMJ maturation and motor function; motor neuron protection	67
PS MOE (18) with decoy ASO	SC P0, P2 (120 mg/kg per day); ICV decoy ASO P0 (10 µg), P20 (20 µg)	Median survival increased 10 d to 212 d; improved NMJ maturation and motor function and protected motor neurons	
PMO (20)	SC P0 (80 µg/g)	Mean lifespan increased from 7.7 d to 19.7 d; FDB-2 muscle protection	68

Chemistry (Length)	Treatment Details	Outcome Measures	Ref
PMO (25)	SC P0 (40 µg/g)	Normalized intestine development	13
PMO (25)	SC P0 (40 µg/g)	Normalized select miRNA levels in SC, muscle, serum	69
PMO and Pip6a-conjugated PMO (20)	IV P0 (10 µg/g)	Median survival increased 12 d to 167 d (Pip6a-PMO); no effect with unconjugated PMO	70
	IV P0, P2 (5 or 10 µg/g each day)	Median survival increased 12 d to 57 d (unconjugated) or 283 d (5 µg/g Pip6a-conjugated) or 457 (10 µg/g Pip6a-conjugated) days; improved weight gain, NMJ maturation, motor function	
PMO (25)	SC P0 (40 µg/g)	Normalized liver development	15
PMO and branched ApoE derivative-conjugated (20)	IV P0, P2 (10 mg/kg each day)	Median survival increased 10.5 d to 29 (unconjugated) or 78 d (ApoE-PMO)	71
Studies in Taiwanese type III mouse model			
PS MOE (18)	ICV infusion for 7 days in adult mice (10, 25, 50, 100 or 150 µg per day)	Increased <i>SMN2</i> exon 7 inclusion and SMN protein in the CNS	48
	ICV E15 (2.5, 5, 10 or 20 µg)	Dose-dependent increase in tail length and delayed tail necrosis	
PMO (25)	IV P0 (20 µg/g)	Tail necrosis onset delayed from 3 to 8 weeks	52
Vivo-PMO (25)	IV P0 (10 µg/g)	Tail necrosis onset delayed from 3 weeks to 9 months	
PS MOE (18)	IP P0, P2 (40, 80 or 120 mg/kg)	Prevented tail necrosis, but tails significant shorter than control mice	67
Study in Burgheron mouse model			
PS MOE (20)	IP P10, P12 (80 µg/g)	Significantly extended lifespan; prevented tail necrosis; improved NMJ maturation	72
	IP P25, P27 (80 µg/g)	Significantly extended lifespan	

^aExamined and ISS-N1-targeting ASO as well as ISS-N1-targeting ASO conjugated to sequence to recruit either SF2/ASF or hTra2β1

Abbreviations: 2'OMe, 2'-O-methyl; CMAP, compound muscle action potential; d, days; EIM, electrical impedance myography; ICV, intracerebroventricular; IP, intraperitoneal; IV, intravascular; MOE, 2'-O-methoxyethyl; MUNE, motor unit number estimation; NMJ, neuromuscular junction; P, postnatal day; PMO, phosphorodiamidate morpholino oligonucleotide; PS, phosphorothioate; SC, subcutaneous