

# MBNL1<sub>42</sub> and MBNL1<sub>43</sub> gene isoforms, overexpressed in DM1-patient muscle, encode for nuclear proteins interacting with Src family kinases

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Myotonic dystrophy type-1 (DM1) is the most prevalent form of muscular dystrophy in adults. This disorder is an RNA-dominant disease, caused by expansion of a CTG repeat in the *DMPK* gene that leads to a misregulation in the alternative splicing of pre-mRNAs. The longer muscleblind-like-1 (*MBNL1*) transcripts containing exon 5 and the respective protein isoforms (MBNL1<sub>42-43</sub>) were found to be overexpressed in DM1 muscle and localized exclusively in the nuclei. *In vitro* assays showed that MBNL1<sub>42-43</sub> bind the Src-homology 3 domain of Src family kinases (SFKs) via their proline-rich motifs, enhancing the SFK activity. Notably, this association was also confirmed in DM1 muscle and myotubes. The recovery, mediated by an siRNA target to *Ex5-MBNL1*<sub>42-43</sub>, succeeded in reducing the nuclear localization of both Lyn and MBNL1<sub>42-43</sub> proteins and in decreasing the level of tyrosine phosphorylated proteins. Our results suggest an additional molecular mechanism in the DM1 pathogenesis, based on an altered phosphotyrosine signalling pathway.

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Myotonic dystrophy type-1 (DM1, OMIM 160900) is an autosomal dominant multisystemic disorder with considerable phenotypic variation between individuals; clinical aspects include myotonia, progressive muscle weakness, cataracts, insulin resistance and cardiac conduction defects.<sup>1</sup> The genetic mutation responsible for DM1 is an expanded (CTG)*n*, in the 3'-untranslated region (UTR) of the dystrophin myotonia protein kinase (*DMPK*) gene (OMIM 605377), on chromosome 19q13.<sup>2-4</sup> DM1 pathogenesis is linked to the expression of expanded mutant RNA transcripts, which accumulate in the nuclei to form nuclear foci (ribonuclear inclusions).<sup>5</sup> This clue prompted the theory that a toxic gain-of-function effect, resulting from these inclusions, alters the proportion-activity of two RNA-binding proteins: muscleblind-like-1 (MBNL1) and CUG-BP1. The splicing of several MBNL1 and CUG-BP1 pre-mRNA targets is subsequently severely altered in DM1-affected tissues. The disruption of the *Mbnl1* gene in transgenic mouse models in fact recapitulates the splicing abnormalities of DM1 patients,<sup>6-8</sup> as well as the myotonic-dystrophy-like phenotype, with skeletal muscle myotonia and cataracts. Moreover, the myotonic-dystrophy-like phenotype of mice expressing (CUG)*n* can be reversed by

overproduction of MBNL1,<sup>9</sup> and long-term MBNL1 overexpression prevents CUG-induced myotonia, myopathy and alternative splicing abnormalities in DM1 mice.<sup>10</sup> MBNL1 is also sufficient to largely abolish DM1 foci *in vivo*.<sup>11</sup> In humans, the *MBNL1* gene contains 12 exons and the coding sequence is distributed over 10 exons (numbered 1 to 10 in this paper; Figure 1a). The inclusion/exclusion of exons 3, 5, 7 and 9 generates several mRNA isoforms<sup>12,13</sup> developmentally regulated and reportedly altered in DM tissues.<sup>7,9,14,15</sup> *MBNL1* exons encode for protein domains with different functions: exons 1, 2 and 4 are essential for RNA binding,<sup>16</sup> exons 5 and 6 for controlling the nuclear localization of MBNL1; exon 3 strongly enhances the affinity of MBNL1 for its pre-mRNA target sites; exons 3 and 6 encode for a splicing regulatory domain; and exon 7 enhances MBNL1 self-dimerization.<sup>17</sup> Although *MBNL1* isoforms have been partly characterized in human skeletal muscle, recent work on DM1-patient brain revealed the existence of a combination of foetal and adult isoforms, with the expression of additional, longer variants assuming different functional roles (still unexplored).<sup>9,15</sup> The MBNL1 protein contains three proline-rich motifs (PRMs), known to bind Src-homology 3 (SH3)

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**Abbreviations:** c-TNT, cardiac troponin T; DAPI, 4',6-diamidino-2-phenylindole; DM1, myotonic dystrophy type-1; GST, glutathione S-transferase; IR, insulin receptor; MBNL1, muscleblind-like-1; PRMs, proline-rich motifs; pTyr, phosphotyrosine; SFKs, Src family kinases; SH, Src homology domain; SPR, surface plasmon resonance  
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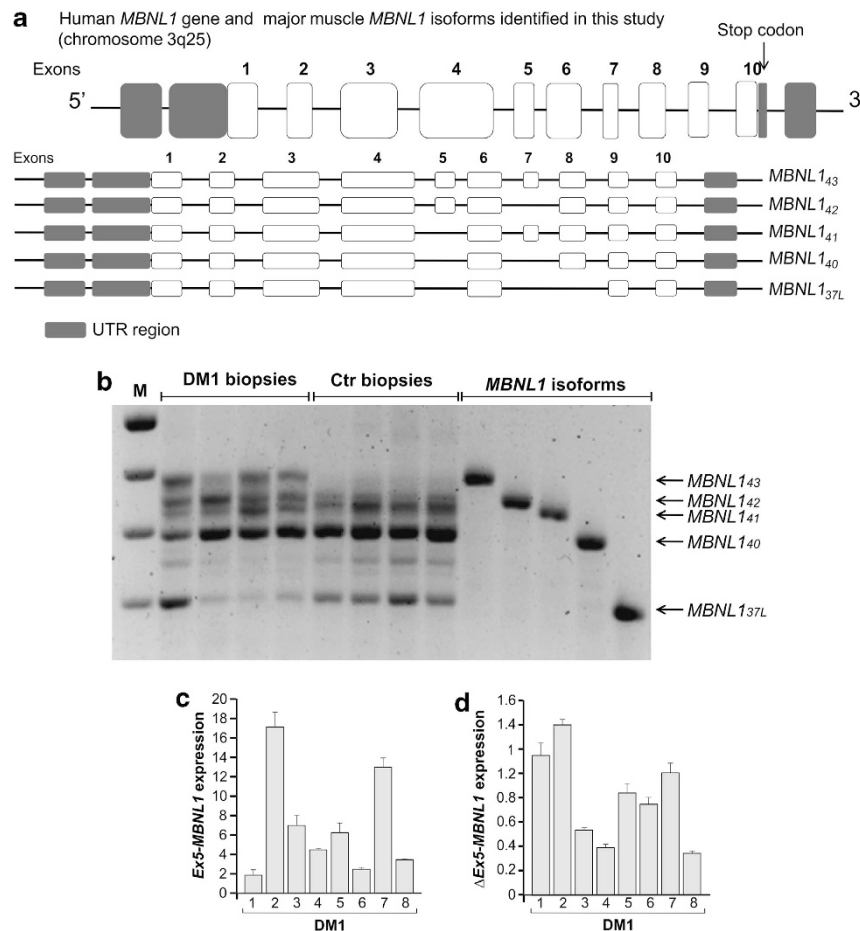
domains present in many signalling proteins (Supplementary Figure SF1). This observation suggests that the different MBNL1 isoforms might regulate the activity and/or localization of the tyrosine (Tyr) kinases of the Src family (SFKs), triggering a signalling cascade mediated by Tyr phosphorylation. Interestingly, CUG-BP1 was also recently found hyperphosphorylated in DM1 tissues and in a DM1 mouse model.<sup>18</sup>

Intracellular localization, regulation of the alternative splicing of model pre-mRNA, ability to interact with SFKs through SH2 and SH3 domains and susceptibility to Tyr phosphorylation of MBNL1 isoforms in DM1 muscle and myotubes were the aspects specifically addressed by our investigation.

## Results

**MBNL1<sub>42-43</sub> isoforms are overexpressed in the muscle from DM1 patients compared with controls, and localize in the nuclei.** The complete splicing pattern of the MBNL1

gene was first determined in muscle tissues from DM1 ( $n=8$ ) patients compared with controls ( $n=4$ ), by RT-PCR analysis. The number assigned to each MBNL1 exon varies in the literature; here, we use the exon numbering of Pascual and co-workers<sup>19</sup> and coding exons have been labelled from 1 to 10 (Figure 1a). Five major MBNL1 transcripts were identified; their relative expression in DM1 and control samples was found to be significantly different. The major transcripts expressed in the muscle from controls corresponded to MBNL1<sub>40</sub> (NM 207292.1) and MBNL1<sub>41</sub> (NM 021038.3) isoforms. We also identified and characterized a novel MBNL1 isoform lacking exons 5, 7 and 8 – named MBNL1<sub>37L</sub> – in view of the predicted molecular weight (MW) of the encoded protein (Figure 1a). In the DM1 muscle, in addition to these MBNL1 transcripts, we observed higher levels of the MBNL1<sub>42</sub> (NM 207293.1) and MBNL1<sub>43</sub> isoforms (Figure 1b), whereas the expression of MBNL1<sub>43</sub> and MBNL1<sub>42</sub> was absent and greatly reduced, respectively, in the control



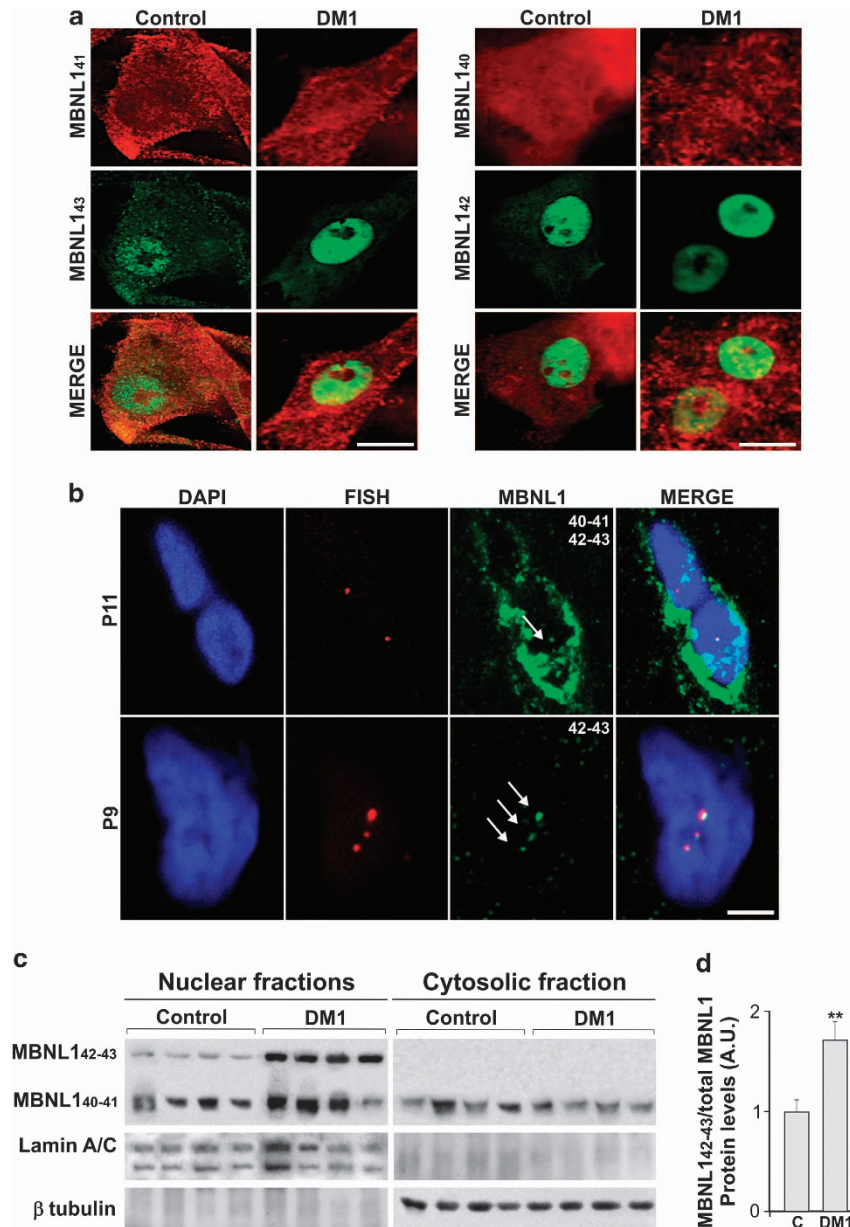
**Figure 1** Qualitative and quantitative expression analysis of the major MBNL1 isoforms expressed in the muscle from DM1 patients and controls. (a) Representation of the MBNL1 gene and major muscle transcripts identified in this study. The human MBNL1 gene contains 10 coding exons, represented as white boxes; untranslated regions (UTRs) are shown in grey, and introns as lines. The names of each isoform and the exon order are indicated on the right following Pascual *et al.*<sup>19</sup> (b) Representative gel electrophoresis of the RT-PCR products corresponding to the entire coding sequences of the MBNL1 gene. Five major muscular transcripts were identified (MBNL1<sub>37L-40-41-42-43</sub>) that showed relative expression differences between four DM1 (lanes 2–5) and four control (lanes 6–9) muscle samples. (c and d) Results from Syber Green Real Time PCR analysis using primers allowing detection of MBNL1 isoforms (c) including (Ex5-MBNL1<sub>42-43</sub>) and (d) not including (ΔEx5-MBNL1<sub>40-41</sub>) exon 5 in muscle from DM1 ( $n=8$ , grey bars). DM1 samples express significantly higher level of Ex5-MBNL1<sub>42-43</sub> isoforms compared with controls ( $P < 0.01$  by Mann–Whitney test), whereas no differences were found in ΔEx5-MBNL1<sub>37L-40-41</sub> transcript expression. All experiments were in triplicate; the median value of controls was set as one, and the level of  $\beta$ -actin transcript was used for sample normalization

muscle. The *MBNL1*<sub>43</sub> mRNA is expressed in human foetal brain and mouse skeletal muscle<sup>9,15</sup> and corresponds to the *MBNL1*<sub>42</sub> sequence of Pascual's classification with the addition of exon 7 (Figure 1a). RT-PCR analysis, using primers designed to discriminate between *MBNL1* isoforms containing (*Ex5-MBNL1*<sub>42-43</sub>) or not containing ( $\Delta$ *Ex5-MBNL1*<sub>37L-40-41</sub>) exon 5, confirmed that DM1 tissues express higher levels of *Ex5-MBNL1*<sub>42-43</sub> mRNAs than controls. *Ex5-MBNL1*<sub>42-43</sub> isoforms were overexpressed in DM1 muscle 1.9- to 17.3-fold compared with controls, whereas  $\Delta$ *Ex5-MBNL1*<sub>37L-40-41</sub> transcripts vary over the range from 0.38 to 1.40 (Figures 1c and d). These results are in accordance with the previous observation that MBNL1 autoregulates the splicing of its own pre-mRNA inducing the exclusion of exon 5 by binding to a response element located in intron 4 of the gene.<sup>20</sup> The functional depletion of the MBNL1 protein in DM1 tissues could therefore lead to the aberrant inclusion of exon 5 within the *MBNL1* transcripts. We then subcloned the coding sequences of the *MBNL1*<sub>40-41-42-43</sub> major muscular *MBNL1* isoforms into appropriate expression vectors to localize the encoded proteins. The exon 5-dependent nuclear localization of the *MBNL1*<sub>42-43</sub> isoforms has been reported in HeLa cells and control myoblasts.<sup>15,17</sup> Consistently, in DM1 myoblasts, the localization of *MBNL1*<sub>40-41-V5</sub> isoforms was diffuse in the cytosol and nucleus, whereas *MBNL1*<sub>42-43</sub>-myc isoforms were exclusively nuclear (Figure 2a), with no differences between DM1 and control primary myoblasts (Figure 2a). This observation suggests that in this *in vitro* model, the MBNL1 cell localization is unaffected by the expression of the pathological CUG expansions. Two novel anti-MBNL1-specific antibodies (anti-P9 and -P11) were synthesized to distinguish the MBNL1 isoforms. Anti-P9, targeted to the 18 amino acids encoded by exon 5, was designed to recognize only the *MBNL1*<sub>42-43</sub> isoforms, whereas anti-P11, designed against the constitutive exon 6, binds to all isoforms (see Supplementary Figures S1 and S2). Immunofluorescence analysis showed that the antibody anti-P9 (*MBNL1*<sub>42-43</sub>) colocalizes with the ribonuclear foci of DM1 muscle, as well as the anti-P11 antibody (*MBNL1*<sub>40-41-42-43</sub>) (Figure 2b). However, the immunofluorescence signal of anti-P11 antibody revealed a nuclear and cytosolic expression, providing further evidence of a ubiquitous cell distribution of *MBNL1*<sub>40-41</sub> (Figure 2b). Consistent with the results of mRNA expression, western blotting (WB) analysis showed significant increase ( $P < 0.01$ ) of the signal recognized by the anti-P9 antibody (target to exon 5) in DM1 muscle nuclei, as compared with controls, but was absent in the cytoplasmic fraction of all tested samples (Figures 2c and d). Tests were performed to establish the extent to which the various MBNL1 isoforms differed in function, properties and interactions. Minigene assays on the cardiac troponin T (*c-TNT*) and insulin receptor (*IR*) pre-mRNAs splicing<sup>21,22</sup> revealed no differences in the *in trans* regulatory splicing properties of the characterized MBNL1 isoforms (Supplementary Figure S3).

**MBNL1<sub>42-43</sub> proteins interact with SFKs, enhance cellular Tyr phosphorylation and induce the nuclear translocation of Lyn Tyr kinase.** The four above MBNL1 isoform sequences exhibit three PRMs, encoded by MBNL1 exon 2

(<sup>111</sup>PLQVPV<sup>116</sup>), exon 3 (<sup>161</sup>PGVPVP<sup>166</sup>) and exon 6 (<sup>295</sup>PPLPKRP<sup>301</sup> of *MBNL1*<sub>42-43</sub> proteins, corresponding to amino acids 277–283 in *MBNL1*<sub>40-41</sub> proteins), respectively (Supplementary Material and Supplementary Figure S1); these sequences display the consensus motif for interaction with the SH3 domain within the SFKs. As the PRM binding to the SH3 domain of SFKs is described to bring about activation,<sup>23</sup> turnover<sup>24</sup> and subcellular localization<sup>25,26</sup> of SFKs themselves, we wondered whether the *MBNL1* isoforms could affect these SFK properties, and/or whether Lyn could be implicated in the pathogenesis of DM1. SFKs are non-receptor Tyr kinases, classifiable into two groups, based on the phylogenetic tree,<sup>27</sup> namely Src-related (Src, Yes, Fyn and Fgr) and Lyn-related SFKs (Lyn, Hck, Lck and Blk).<sup>28</sup> Their common modular structure consist of (i) an N-terminal SH4 domain for acylation; (ii) the aforementioned SH3 domain; (iii) an SH2 domain, which recognizes phospho-Tyr (pTyr) motifs; and (iv) a kinase domain (SH1), followed by a C-terminal tail having a negative regulatory role when phosphorylated by the inhibitory kinases.<sup>23</sup> SFKs are normally maintained in a closed, inactive conformation through two major intramolecular inhibitory interactions: the binding of the phosphorylated C-terminal tail to the SH2 domain, and the interaction of a polyproline type-II helical motif containing the core sequence P–X–X–P, where X is any amino acid, within the SH2 kinase linked with the SH3 domain.<sup>29</sup> Conformational transition of SFKs from inactive to active form is induced by an array of factors causing dephosphorylation of the C-terminal tail by several Tyr phosphatases and/or displacement of the SH2 or SH3 domains by specific ligands, ultimately leading to disruption of the inhibitory intramolecular interactions.<sup>30-34</sup>

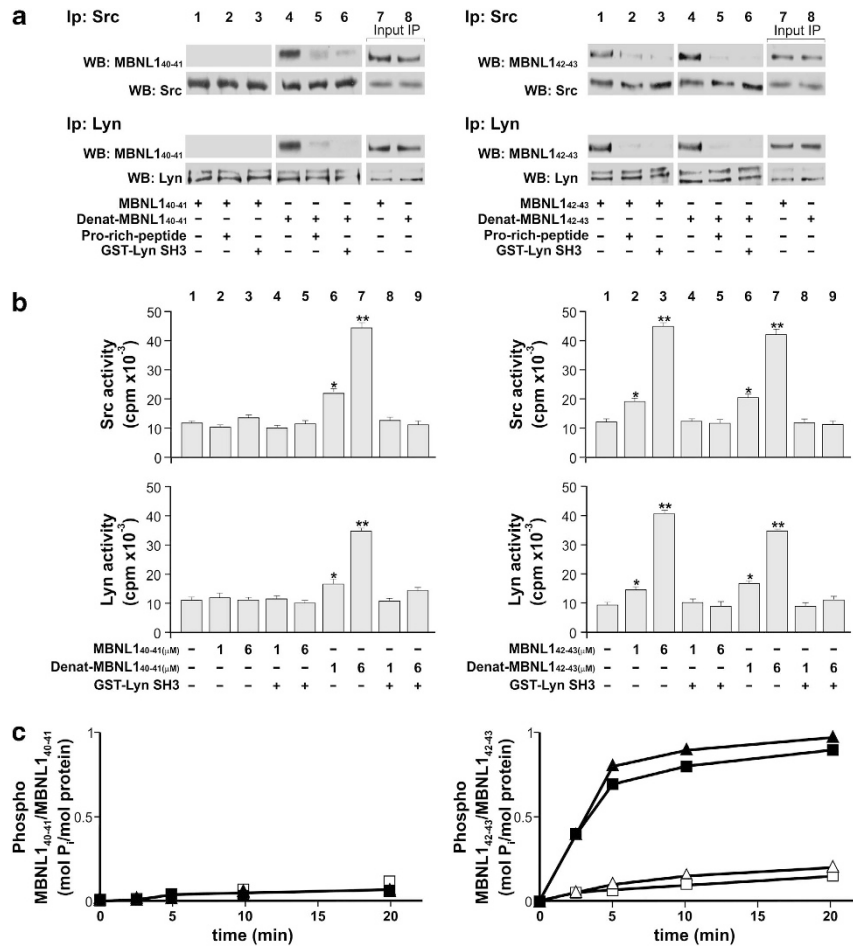
To assess whether *MBNL1*<sub>40-41</sub> and *MBNL1*<sub>42-43</sub> interacted with the SFK SH3 domain through their PRMs, we performed competition assays by incubating the *MBNL1* isoforms with Lyn or Src (the prototypical members of the two SFK groups) in the absence or presence of a synthetic proline-rich peptide (KGGRSRLPLPLPPPG), which is known to interact with Lyn SH3 domain, or the recombinant glutathione S-transferase (GST)-Lyn SH3 domain. The following immunoprecipitation with anti-Lyn and anti-Src antibodies showed that only the *MBNL1*<sub>42-43</sub> associated with Src and Lyn via interaction between the MBNL1 PRM and the SFK SH3 domain (Figure 3a, left panel). This interaction also determined stimulation of SFK activity (Figure 3b, right panel) and phosphorylation of *MBNL1*<sub>42-43</sub> itself (Figure 3c, right panel). Yet, denaturation by heating at 55 °C allowed *MBNL1*<sub>40-41</sub> to acquire the same ability to bind to and activate Src and Lyn as *MBNL1*<sub>42-43</sub> (Figures 3a and b, left panel), suggesting that the binding motif in the latter isoforms is accessible owing to a structural conformation conferred by the insert encoded by exon 5. As to phosphorylation, denaturation prevented recognition of *MBNL1*<sub>42-43</sub> as a substrate by Src and Lyn (data not shown), suggesting the significance of not only the interaction due to exposure of specific motifs but also of an appropriate three-dimensional structure. Surface plasmon resonance (SPR) studies confirmed the specific interaction of the Lyn kinase SH3 domain with only *MBNL1*<sub>43</sub> isoform (Supplementary Material and Supplementary Figure S4).



**Figure 2** Nuclear localization and amount of MBNL1<sub>42-43</sub> isoforms in DM1 myoblasts and muscle tissue. **(a)** Representative confocal images of DM1 and control primary myoblasts, overexpressing MBNL1<sub>40-41-42-43</sub> isoforms. Immunofluorescence with specific anti-V5 (MBNL1<sub>40-41</sub>) or -myc (MBNL1<sub>42-43</sub>) tags was performed after 48 h. Scale bar, 10  $\mu$ m. **(b)** Representative images of nuclear foci (red) from fixed DM1 muscle, obtained by fluorescence *in situ* hybridization with a Texas Red labelled (CAG)<sub>10</sub> probe. The MBNL1 antibodies (green) anti-P9 and anti-P11 co-localized with foci. Anti-P9 was localized only in the nucleus, in contrast with anti-P11, which was also present in the cytosol. Scale bar, 2  $\mu$ m. **(c)** Representative WB analysis of MBNL1<sub>40-41-42-43</sub> in nuclear and cytosolic fractions from four DM1 patients and four control muscle samples. Proteins, separated in 7.5–17.5% T30C4 SDS-PAGE gels, were blotted onto nitrocellulose and probed with anti-P9 (specific for MBNL1<sub>42-43</sub> isoforms) and anti-P11 antibodies. Lamin A/C and  $\beta$ -tubulin protein levels were used to normalize the samples from nuclear and cytosolic fractions, respectively. **(d)** The graph represents the mean  $\pm$  S.D. of at least three experiments. In each lane the ratio between the signal density of MBNL1<sub>42-43</sub> (upper band) and the sum of the signal density of the two MBNL1 bands (upper and lower) was calculated. The values were expressed as percentage of the value obtained in the control muscle, considered as reference unit. \*\*Significant difference for  $P < 0.01$  by Mann–Whitney test

Our *in vitro* data provide novel evidence of a strong functional discrimination between the long and short isoforms of MBNL1 in terms of interaction with SFKs and regulation of their activity. These novel functions of the MBNL1 proteins were also analyzed in 293T cells co-overexpressing Lyn and one MBNL1<sub>40-41-42-43</sub> isoform. As 293T cells express a low basal level of SFKs, the amount of Lyn-encoding plasmid was

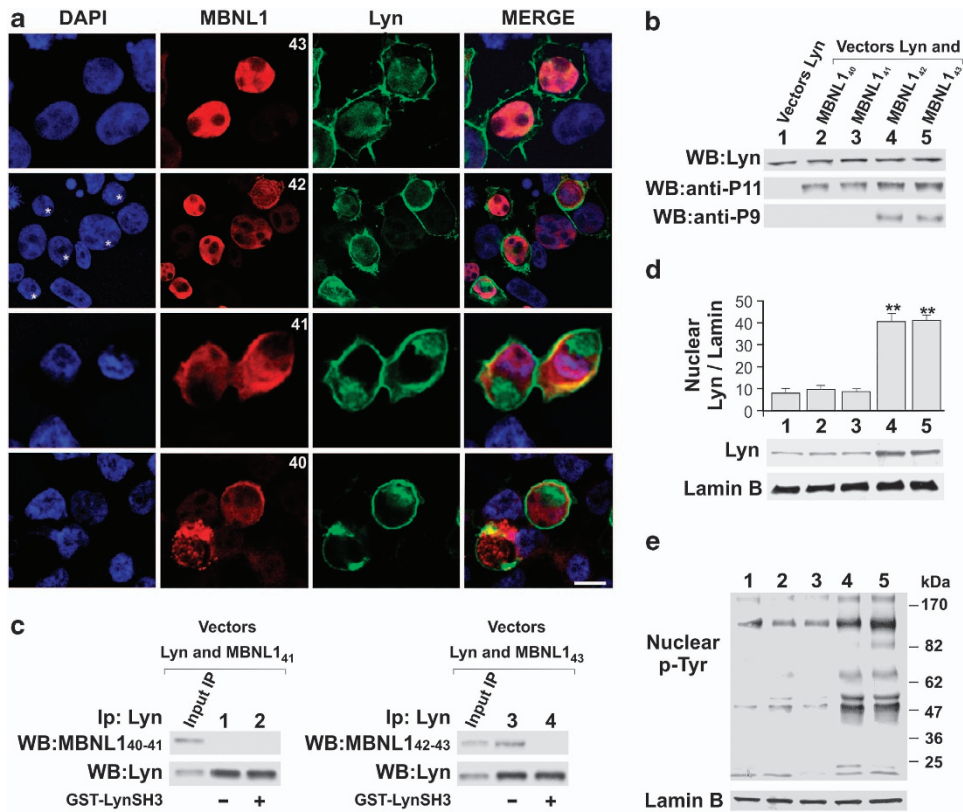
standardized to the expression of Lyn within HeLa cells (Supplementary Figure S5). Figure 4a shows colocalization of MBNL1<sub>42-43</sub> and Lyn in the nucleus, where Lyn translocated after the overexpression of MBNL1<sub>42-43</sub> proteins, and with Lyn being equally expressed under all the experimental conditions and regardless of the cotransfection with MBNL1 isoforms (Figure 4b). To assess the existence of a MBNL1/Lyn complex



**Figure 3** Analysis of the interaction between the SH3 domain of the SFKs Src and Lyn and the proline-rich motif of recombinant MBNL1<sub>40-41</sub> and MBNL1<sub>42-43</sub> isoforms. (a) The interaction between SH3 domain of two SFK (Src and Lyn) and the proline-rich motifs of recombinant MBNL1<sub>40-41</sub> and MBNL1<sub>42-43</sub> isoforms. The 0.2 μg of recombinant MBNL1<sub>40-41</sub> or MBNL1<sub>42-43</sub> (as such or heated at 55 °C for 10 min, denat. MBNL1) was incubated with 0.1 μg of Src or Lyn in the absence or presence, respectively, of a proline-rich peptide or the GST-Lyn SH3 domain, for 10 min at 30 °C. Three-fourths of the sample underwent immunoprecipitation with anti-Src or anti-Lyn antibodies, respectively. Immunocomplexes were then probed with anti-P11 or anti-P9 antibodies, and subsequently with anti-Src or anti-Lyn antibodies. The presence of MBNL<sub>42</sub> or MBNL<sub>43</sub> (right panel), but not of MBNL<sub>40</sub> or MBNL<sub>41</sub> (left), indicates its association with the two SFKs and suggests that the stretch encoded by exon 5 might have a role in this interaction, possibly through a conformational change that renders the PRMs available to binding. Moreover, competition assays carried out in the absence or presence of a synthetic proline-rich peptide (5'-KGGRSRLPLPLPPG-3') known to interact with the Lyn SH3 domain, or the GST-Lyn/SH3 domain, confirmed that this interaction was mediated by PRMs and the SH3 domain. The remainder of the sample underwent WB analysis (lanes 7 and 8) (b) As this interaction was also a prerequisite for enhancement of SFK activity relative to the basal state, we tested Tyr kinase activity of Src (top panel) and Lyn (bottom) on the Src-specific peptide substrate cdc2 in the absence (lane 1) or presence of increasing concentrations of MBNL<sub>41</sub> (left panels, lanes 2–5) and denat. MBNL<sub>41</sub> (left panels, lanes 6–9) or MBNL<sub>43</sub> (right panels, lanes 2–5) and denat. MBNL<sub>43</sub> (right panels, lanes 6–9), alternatively supplemented with the GST-Lyn SH3 domain (lanes 4 and 5, and 8–9) as described in the Materials and Methods section. (c) As this interaction was also a prerequisite for recognition of MBNL<sub>43</sub> as a substrate by SFKs, we tested whether MBNL<sub>41</sub> or MBNL<sub>43</sub> (each 1 μM) were phosphorylated by Lyn and Src in the absence (solid triangles and solid squares, respectively) or presence (open triangles and open squares, respectively) of the GST-Lyn SH3 domain. At various time points, reactions were stopped and the phosphate incorporated was analyzed after SDS-PAGE on the Cyclone Plus. The amount is expressed as mol P incorporated/mol prot. (values represent the mean of three separate experiments ± S.D.). As expected, only MBNL<sub>43</sub> is phosphorylated by Src. The same set of experiments conducted with the MBNL<sub>140</sub> and MBNL<sub>142</sub> isoforms demonstrated that these two proteins behave similarly to MBNL<sub>141</sub> and MBNL<sub>143</sub>, respectively (data not shown). Only MBNL<sub>142-43</sub> isoforms were able to significantly increase Src and Lyn activity, suggesting a novel function of these long isoforms in kinase enzyme regulation. Data are expressed as mean ± S.D. from three separate experiments. \*P < 0.05; \*\*\*P < 0.001 by Kruskal–Wallis analysis

*in vivo*, immunoprecipitation assays with anti-Lyn antibody on 293T cells coexpressing Lyn and each of the MBNL1 isoforms were carried out. This experiment revealed that MBNL1<sub>42-43</sub> (but not MBNL1<sub>40-41</sub>) co-immunoprecipitated with Lyn, this event occurring as a result of PRM/SH3 domain interaction, as confirmed by the competition assay with GST-Lyn-SH3 (Figure 4c). Moreover, the WB analysis of the nuclear fraction of these cells highlighted that the nuclear levels of Lyn rose up fourfold (Figure 4d, lanes 4 and 5 *versus* lane 1) when the

MBNL1<sub>42-43</sub> was coexpressed, whereas the localization of Lyn remained unchanged in the presence of MBNL1<sub>40-41</sub> (Figure 4d, lanes 2 and 3 *versus* lane 1). To explore whether this interaction determined SFK activation in the nuclei, as a result of SH3 engagement by PRMs,<sup>35</sup> WB analysis with anti-pTyr antibody revealed a dramatic increase in Tyr phosphorylation in the nuclear fraction of 293T cells when MBNL1<sub>42-43</sub>, and not MBNL1<sub>40-41</sub>, were coexpressed with Lyn (Figure 4e).



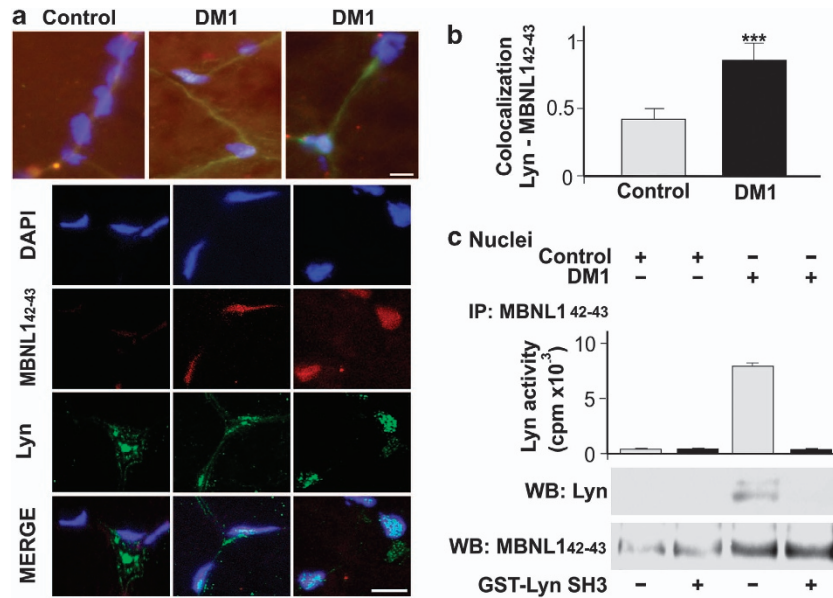
**Figure 4** Translocation of Lyn to the nuclei of 293T cells occurs only upon co-expression with MBNL1<sub>42-43</sub>. (a) Representative confocal images of 293T cells co-transfected with plasmids carrying different MBNL1 isoforms (MBNL1<sub>40-41-42-43</sub> in pEF1-Myc/HisA vector) and Lyn (pCMV6-XL4/Lyn). Nuclei were labeled with DAPI (blue), MBNL1 with anti-P9 (MBNL1<sub>42-43</sub>) or anti-P11 (MBNL1<sub>40-41</sub>) antibodies (red) coupled to immunodetection of Lyn (green). Scale bar 10  $\mu$ m. (b) Representative WB analysis of whole-cell lysates from 293T cells transfected with expression vectors for Lyn, MBNL1<sub>40</sub>, MBNL1<sub>41</sub>, MBNL1<sub>42</sub> and MBNL1<sub>43</sub> probed with anti-Lyn antibody (top strip), anti-P11 antibody (middle strip) and anti-P9 antibody (bottom strip). (c) Representative co-immunoprecipitation of four-fifths of whole-cell lysates from 293T cells, overexpressing MBNL1<sub>41</sub> (pEF1/MBNL1<sub>41</sub> (left panel), lanes 1 and 2), or MBNL1<sub>43</sub> (pEF1/MBNL1<sub>43</sub> (right panel), lanes 3 and 4) and Lyn (pCMV6-XL4/Lyn, lanes 1–4). Thirty-six hours after transfection, whole-cell lysates were immunoprecipitated with anti-Lyn in the absence (lanes 1 and 3) or presence (lanes 2 and 4) of the GST-Lyn SH3 domain. The immunocomplexes and one-fifth of the whole-cell lysates underwent WB analysis with anti-P11 or anti-P9 antibodies, respectively, and subsequently with anti-Lyn antibodies. The figure represents one of three independent experiments. The same set of experiments conducted with MBNL1<sub>40</sub> and MBNL1<sub>42</sub> isoforms demonstrated that these two proteins behave similarly to MBNL1<sub>41</sub> and MBNL1<sub>43</sub>, respectively, again supporting the hypothesis that it is exon 5 that enables MBNL1s to interact with SFKs (data not shown). (d) Representative WB analysis of nuclear lysates from 293T cells transfected with expression vectors for Lyn (lanes 1–5), MBNL1<sub>40</sub> (lane 2), MBNL1<sub>41</sub> (lane 3), MBNL1<sub>42</sub> (lane 4) and MBNL1<sub>43</sub> (lane 5). The membranes were probed with anti-Lyn and with anti-Lamin B antibody as loading control. The resulting bands underwent densitometric analysis. The values (arbitrary units) represent the mean  $\pm$  S.D. of at least three experiments. Reference unit: lane 1 sample (only transfected with Lyn). \*\*Significant difference for  $P < 0.01$  by Kruskal–Wallis analysis. (e) Representative WB analysis of nuclear lysates from 293T cells, transfected with expression vectors for Lyn (lanes 1–5), MBNL1<sub>40</sub> (lane 2), MBNL1<sub>41</sub> (lane 3), MBNL1<sub>42</sub> (lane 4) and MBNL1<sub>43</sub> (lane 5) and probed with anti-p-Tyr antibody. The membranes were re-probed with anti-Lamin B antibody as loading control.

### Muscle and myotubes from DM1 patients show increased nuclear localization of Lyn and Tyr hyperphosphorylation.

These data strongly suggest that MBNL1<sub>42-43</sub> protein isoforms, whose abundance is elevated in DM1 muscle, might induce an inappropriate activation of Lyn in DM1 muscle cell nuclei: as a direct test, we first examined muscle sections from DM1 patients ( $n=4$ ) and controls ( $n=4$ ). Confocal immunofluorescence images of the MBNL1<sub>42-43</sub> and Lyn proteins showed a nuclear localization of both proteins only in DM1 muscle (Figure 5a), confirmed by Manders overlap coefficient analysis (Figure 5b). Moreover, to assess whether the existence of an MBNL1/Lyn complex was also present in the nuclei of muscle tissue, co-immunoprecipitation analysis with anti-P9 (against MBNL1<sub>42-43</sub>) antibody was performed on DM1 and control muscle samples. Consistently with the results obtained with 293T cells and in line with the increased amount of

MBNL1<sub>42-43</sub> in DM1 muscle tissue (Figures 2c–d), we found that MBNL1<sub>42-43</sub> co-immunoprecipitated with Lyn only in the nuclear fraction of DM1 muscles, in parallel with an increased nuclear activity of Lyn (Figure 5c). This event occurred as a result of PRM/SH3 domain interaction, as confirmed by the competition assay with GST-Lyn-SH3 (Figure 5c).

The same parameters were analyzed in 15-day-differentiated myotubes from two DM1 patients and three controls. As expected and reported elsewhere,<sup>36</sup> RT-PCR-based analysis of MBNL1 splicing showed that DM1 myotubes had a statistically significant (2.8-fold) increase in the expression of *Ex5-MBNL1* transcripts relative to controls (Figures 6a and b). Consistently, WB analysis demonstrated a significantly increased amount of Lyn (570%,  $P < 0.02$ ) and MBNL1<sub>42-43</sub> (680%,  $P < 0.005$ ) (Figures 6c, e and f) in the nuclei of the same DM1 myotubes, as compared with controls. The increased nuclear localization of Lyn in DM1 myotubes



**Figure 5** The nuclear colocalization of Lyn and MBNL<sub>42-43</sub> was detected in DM1 muscle tissues. (a) Representative images at epifluorescent (low magnification) and confocal (high magnification) microscope of nuclei from muscle samples (four controls and four DM1 patients). The muscle sections were probed with anti-P9 (red) and anti-Lyn (green), nuclei were marked with DAPI. Scale bar, 10  $\mu$ m. (b) Quantification of Lyn and MBNL<sub>42-43</sub> signal colocalization in control and DM1 muscle samples. Values represent Manders' overlap coefficient (the fraction of total fluorescence signal of Lyn overlapping with MBNL<sub>42-43</sub> signal), obtained from confocal images of nuclear and perinuclear regions as in (a) and as described in Materials and Methods section. Statistical significance was evaluated by two-tailed Student's *t*-test ( $n = 50$  nuclei,  $***P < 0.001$ ). (c) MBNL<sub>42-43</sub> immunoprecipitation from nuclear fractions, obtained from two controls and two DM1 muscle samples. Immunoprecipitate was used for Lyn Tyr kinase activity assay (upper panel) and for WB analysis (below), in the absence or presence of binding competitor domain GST-Lyn/SH3, as described in Materials and Methods section. Lyn co-immunoprecipitates with MBNL<sub>42-43</sub> and displays Tyr kinase activity only in DM1 muscle samples. Both the interaction and the activity are abolished in the presence of competing GST-Lyn/SH3 domain

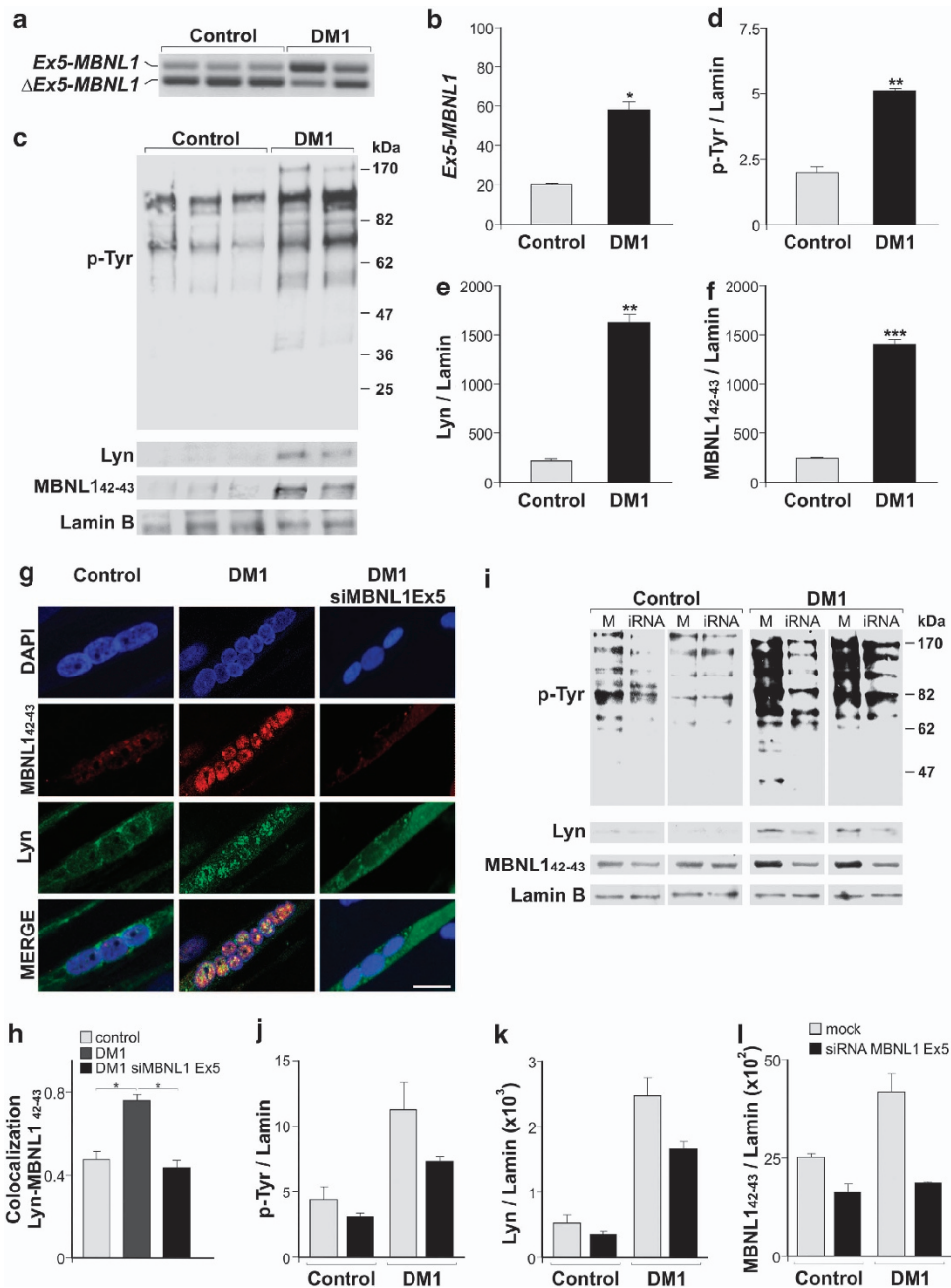
caused a Tyr hyperphosphorylation (250%,  $P < 0.02$ ), as revealed by WB analysis (Figures 6c and d). Immunofluorescence analysis with anti-Lyn and anti-P9 (against MBNL<sub>42-43</sub>) antibodies in 15-day-differentiated myotubes gave further confirmation of the MBNL<sub>42-43</sub> and Lyn cellular distribution in DM1 and control cells (Figure 6g), validated by Manders' overlap coefficient analysis (Figure 6h).

**siRNA-mediated downregulation of MBNL<sub>42-43</sub> isoforms in myotubes from DM1 patients restores cytoplasmic localization of Lyn.** We used an siRNA-mediated approach to test whether both cytoplasmic localization of Lyn and increased nuclear Tyr phosphorylation could be restored in myotubes from DM1 patients. An siRNA specific to target *MBNL1-exon 5* (siMBNL1-Ex5) was designed and tested on DM1 myotubes overexpressing *Ex5-MBNL1<sub>42-43</sub>* transcripts (Figure 6a). Splicing analysis, using primers designed to discriminate between *Ex5-MBNL1<sub>42-43</sub>* and  $\Delta$ *Ex5-MBNL<sub>37L-40-41</sub>* isoforms, revealed a marked reduction of *MBNL1<sub>42-43</sub>* (from 70 to 85%), but not of the *MBNL1<sub>40-41</sub>* transcripts (from 97 to 115%) (data not shown). Immunofluorescence, on differentiated myotubes from DM1 patients 48 h after siRNA transfection, revealed that the signal corresponding to MBNL<sub>42-43</sub> isoforms decreased in the nuclei along with that of Lyn, which instead increased almost throughout the myotube cytoplasm (Figure 6g). These data have been further validated by colocalization analysis, showing the significant decrease of Manders' overlap coefficient in DM1 siRNA-transfected myotubes (Figure 6h)

to the control value. WB analysis confirmed the decrease in the nuclear amount of Lyn and MBNL<sub>42-43</sub> to 67% and 44%, respectively, relative to the cells transfected with empty vector (mock) (Figures 6i, k and l), in parallel with a 35% decrease of Tyr phosphorylation (Figures 6i and j). These results strengthen the case for a functional link between the Ex5-containing *MBNL1* isoforms and the Tyr kinase Lyn.

## Discussion

MBNL1 expression is crucial in the DM pathophysiology, and the loss of MBNL1 protein leads to the spliceopathy in the muscle tissues of DM patients and mouse models.<sup>8,37</sup> The MBNL1 pre-mRNA itself undergoes tissue-specific alternative splicing.<sup>17</sup> In this paper, we showed that muscle tissue from DM1 patients overexpresses *MBNL1<sub>42</sub>* and *MBNL1<sub>43</sub>* isoforms containing exon 5 (*Ex5-MBNL1<sub>42-43</sub>*). In contrast, control muscle showed a prevalent expression of the *MBNL1<sub>40</sub>* and *MBNL1<sub>41</sub>* isoforms, without exon 5 ( $\Delta$ *Ex5-MBNL<sub>37L-40-41</sub>*). The altered splicing of the *MBNL1* gene has also been reported in DM1 brain, whereas  $\Delta$ *Ex5-MBNL<sub>40-41</sub>* isoforms are expressed in adult tissues.<sup>15</sup> These data support the hypothesis that the DM1 phenotype is in part due to a change in the splicing pattern linked to the *trans*-dominant effect of the CUG and CCUG repeats, resulting in the mixed expression of adult and foetal isoforms of various proteins, including MBNL1 itself. The overexpression of *Ex5-MBNL1<sub>42-43</sub>* isoforms in the muscle from DM1 patients, ensuring their restrictive nuclear



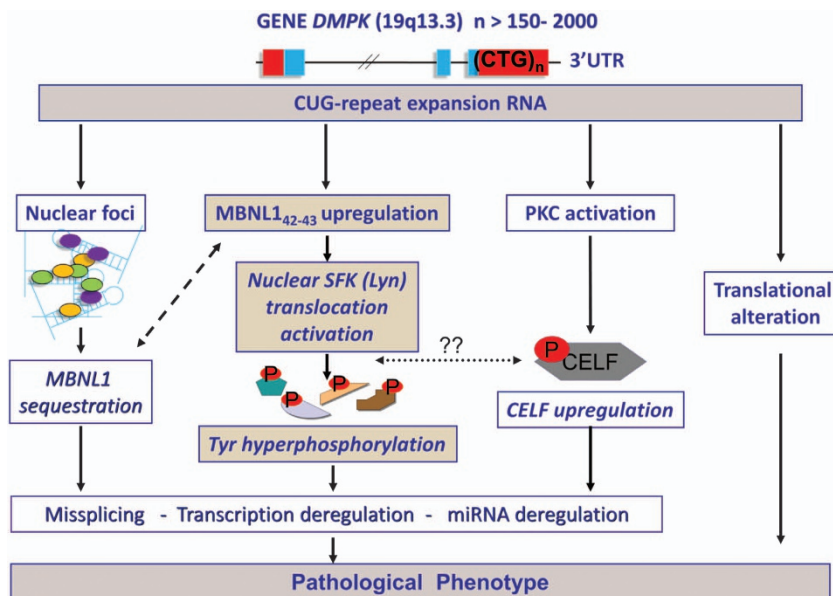
**Figure 6** In 15-day-differentiated DM1 myotubes, nuclear MBNL1<sub>42-43</sub> isoforms are associated with increased presence and activity of Lyn in the nuclei. **(a)** RT-PCR analysis of *MBNL1*<sub>Ex5</sub> and *MBNL1*<sub>ΔEx5</sub> transcripts in 15-day-differentiated myotubes from three controls and two DM1 patients. **(b)** Quantification of *MBNL1*<sub>Ex5</sub> transcript expressed as the percentage of the total *MBNL1* transcripts. **(c)** Representative WB analysis with anti-p-Tyr, anti-Lyn, anti-P9 (against MBNL1<sub>42-43</sub>) of nuclear lysates from 15-day-differentiated myotubes of three controls and two DM1 patients. Equal loading was monitored by anti-Lamin B antibody. **(d-f)** Quantification of at least three independent WB analyses as in **(c)**. Values, normalized to the loading control, were expressed as mean  $\pm$  S.D. and compared with controls for statistical evaluation. \* $P < 0.05$ ; \*\* $P < 0.02$ ; \*\*\* $P < 0.005$  by Mann-Whitney test. **(g)** Representative confocal images of 15-day-differentiated myotubes, from two controls and two DM1 patients, showing different localization of MBNL1<sub>42-43</sub> and Lyn (obtained using identical exposure time and acquisition setting). Myotubes from DM1 patients were also transfected with siMBNL1-Ex5, designed to target specifically the MBNL1<sub>42-43</sub> isoforms. Nuclei were labelled with DAPI (blue), MBNL1 with anti-P9 (red) coupled to immunodetection of Lyn (green). Scale bar 10  $\mu$ m. **(h)** Quantification of Lyn and MBNL1<sub>42-43</sub> colocalization in control and DM1 myotubes. Values represent Manders' overlap coefficient (the fraction of total fluorescence signal of Lyn overlapping with MBNL1<sub>42-43</sub> signal), obtained from confocal images of nuclear and perinuclear regions as in **(g)** and as described in Materials and Methods section. Statistical significance was evaluated by two-tailed Student's *t*-test ( $n = 30$  nuclei, \* $P < 0.05$ ). **(i)** Representative WB analysis with anti-p-Tyr, anti-Lyn and anti-P9 (against MBNL1<sub>42-43</sub>) antibodies of nuclear lysates from 15-day-differentiated myotubes of two controls and two DM1 patients, transfected with empty vector (M) and siMBNL1-Ex5 (iRNA). Equal loading was monitored by anti-Lamin B antibody. **(j-l)** Quantification of two WB analyses as in **(i)**. Values, normalized to the loading control, were expressed as mean  $\pm$  S.D.



localization, indicates an important feature for the sequestration within the foci. To address this issue, we developed novel anti-MBNL1 antibodies that allow us to discriminate different MBNL1 isoforms on the basis of the inclusion/exclusion of sequences encoded by exon 5. We showed that all four MBNL1 isoforms analyzed can bind to the nuclear foci of CUG-expanded repeats in DM1 muscle tissue; however, MBNL1<sub>42-43</sub> protein levels are increased in the nuclei of DM1 muscle compared with controls.

Our data demonstrate for the first time that MBNL1<sub>42-43</sub> isoforms are able to interact with SFKs, in particular with Lyn, which is one of the most ubiquitous members of this class of Tyr kinases. As MBNL1<sub>40-41</sub> isoforms cannot form this complex, it is possible that the amino-acid stretch encoded by exon 5 enables MBNL1 protein to interact with SFKs, possibly through a conformational change whereby the PRMs become available for binding. The MBNL1<sub>42-43</sub>-SFK interaction is also a prerequisite for enhancement of the SFK activity relative to the basal state, evidencing a strong functional discrimination between the long and short isoforms of MBNL1 in terms of interaction with SFKs and regulation of their activity. We also show that MBNL1<sub>42-43</sub> overexpression induces the nuclear translocation of Lyn and enhancement of its catalytic activity, resulting in increased levels of Tyr phosphorylation of nuclear proteins, which also occurs in DM1-patient myotubes. These findings are consistent with the accumulating evidence that SFKs, Lyn in particular, localize to a variety of intracellular organelles,<sup>38-41</sup> in addition to the plasma membrane<sup>42-44</sup> Importantly, the acylation state—especially the combination of myristate and palmitoylate—of the N-terminal SH4 domain of SFKs, has been shown to be critical in directing SFKs to their final location.<sup>45,46</sup> However, recent studies have noted that an additional mechanism involved in targeting SFKs to cell compartments is the

engagement of the SH3 domain by PRM-bearing proteins. Examples include p13, a human T-cell leukaemia virus-1 accessory protein with both a C-terminal PRM and a mitochondrial localization signal; this has been shown to bind to and carry SFKs into the mitochondria in a cell culture model.<sup>47</sup> Another striking effect is the increase in SFK activity, with an elevation of Tyr phosphorylation within the specific compartment in which SFKs operate. We therefore hypothesize a mechanism by which the transfer of SFKs to various cell districts might exert a more general action on cell fate by modifying the phosphorylation state and the function of important substrates in the targeted cell compartments. The nucleus is another organelle to which Lyn has been shown to localize, with a role in the DNA damage response and cell cycle control by associating with and/or phosphorylating critical signalling molecules involved in cell survival and apoptosis.<sup>48-51</sup> Although far from fully elucidated, nuclear translocation of Lyn seems to depend on the level of its catalytic activity.<sup>52</sup> In our study, MBNL1<sub>42-43</sub> seem to act as carriers for Lyn into the nucleus and they boost its catalytic activity; the effects of phosphorylation on the target proteins possibly representing an additional pathogenetic pathway in DM1, as proposed in Figure 7. Of the Tyr-phosphorylated proteins localized in the nuclei, the ‘RNA-processing’ group consists predominantly of players involved in transcription and post-transcriptional events.<sup>53</sup> Of the proteins found to be ‘RNA-related’, six were hnRNP (heterogeneous nuclear ribonucleoproteins), known to complex with hnRNA and to influence pre-mRNA transcription, processing and export, ultimately regulating gene expression and RNA processing, in particular splicing and alternative splicing. In this scenario, phosphorylation of pre-mRNA splicing factors may either cause their removal from pre-mRNA processing events or result in changing the selection of a splice site.<sup>54,55</sup>



**Figure 7** Additional molecular pathogenetic mechanisms in DM1. Results suggest an additional altered mechanism in DM1 nuclei, triggered by CUG-containing *DMPK* transcripts and mediated by the longer Ex5-MBNL1<sub>42-43</sub> protein isoforms: Tyr hyperphosphorylation of nuclear proteins may influence and mediate changes in gene expression and RNA processing, expanding the known mechanisms of RNA toxicity in DM1 cells

The relationship between members of the Src family and splicing regulators has already been demonstrated and may coordinate and regulate apoptosis, thus promoting cell proliferation and formation of metastasis.<sup>56</sup> Possibly, the phosphorylation of splicing regulators, mediated by MBNL1<sub>42-43</sub> overexpression and SFK activation, allows the cells to lower temporarily their active concentration, further aggravating the DM spliceopathy, resulting from the MBNL1 sequestration into the nuclear foci. Therefore, the altered pattern of MBNL1 expression may further contribute to the multisystemic manifestation of DM not only through the mere loss-of-function due to the titration of the RNA-binding proteins by the CUG expansions. The importance of these data is underscored by the ongoing research for functional differences between the MBNL1 isoforms, which, when better delineated, would open up the possibility that the MBNL1 proteins may have different, still unexplored functional roles. A recent paper by Wang *et al.*<sup>57</sup> suggests that MBNL proteins, besides regulating the splicing, also contribute to mRNA translation, protein secretion and localization of alternative 3'-UTR isoforms.<sup>57</sup> Our results broaden this scenario showing for the first time that MBNL1<sub>42-43</sub> proteins, overexpressed in the DM muscle, are Lyn-interacting partners that have an important role in regulating the nuclear-cytoplasmic shuttling and catalytic activity of Lyn itself, and suggest Lyn as a novel factor implicated in the onset and progression of DM1. In this respect, additional functional studies should help clarifying how the altered Lyn-dependent signalling pathways are affected by the dysregulation of Lyn itself, thereby paving the way for discovery of potential novel targets of therapeutic interest in DM1 muscle.

## Materials and Methods

**Skeletal muscle biopsies.** After written consent, DM1 ( $n=8$ ) muscle samples were obtained from vastus lateralis by diagnostic open biopsies. Control samples (also from vastus lateralis,  $n=4$ ) were obtained from subjects deemed free of neuromuscular disorders. Molecular diagnosis of DM1, using a combination of long PCR and Southern blot analysis in the peripheral blood, showed that DM1 patients considered in this study had a CTG expansion between 800 and 1500 repetitions.<sup>58</sup>

**RNA extraction, RT-PCR and QRT-PCR analysis.** Total RNA was extracted from frozen muscle biopsies and 1  $\mu$ g of total RNA was reverse-transcribed according to the protocol of the High Capacity cDNA Archive kit (Applied Biosystem, Foster City, CA, USA). To assess the complete MBNL1 splicing pattern of muscle samples, PCR was performed using primers corresponding to sequences in exons 2 and 10 of the gene (numbered according to Pascual *et al.*,<sup>19</sup> for primer sequences see Supplementary Table 1). The MBNL1 isoform expression levels were measured by SYBR Green semiquantitative QRT-PCR (Applied Biosystems) using primer pairs designed to amplify MBNL1 exons 4, 5 and 7 to distinguish MBNL1<sub>40-41</sub> from MBNL1<sub>42-43</sub> (Supplementary Table 1). The splicing pattern of the MBNL1 gene across exon 5 in DM1 myoblasts and myotubes has been determined as described previously.<sup>14</sup> Transcripts from the exogenous *IR* and *c-TNT* minigenes have been measured using primers representing unique sequences in the cloning vectors (Supplementary Table 1).

**Cloning of MBNL1 isoforms, molecular constructs and MBNL1 recombinant proteins.** To clone the sequences of the MBNL1<sub>40-41-42-43</sub> isoforms, the entire coding sequence of the MBNL1 gene was amplified using the primers MBNL1-F and MBNL1-R listed in Supplementary Table 1. After sequencing, PCR products were cloned into pEF1-Myc/HisA and pEF4-V5/HisA (Invitrogen Carlsbad, CA, USA). Minigene B and minigene RTB300 have been described previously.<sup>21,22</sup> MBNL1 recombinant proteins have been produced in *Escherichia coli* and purified as described in the Supplementary Experimental Procedures.

**Cell culture and plasmid transfection.** Primary myoblasts were obtained and differentiated as described previously.<sup>36</sup> Cells, plated 24 h before transfection at 80–90% confluence, were transiently transfected with 4  $\mu$ g of plasmid DNA and Lipofectamine 2000 (Invitrogen). For siRNA experiments, myotubes from DM1 patients were transfected with 50 nM of siMBNL1-Ex5 and analyzed after 48 h. Rhabdomyosarcoma clones with permanent overexpression of MBNL1<sub>41</sub> and MBNL1<sub>43</sub> isoforms, named RD41 and RD43, respectively, were selected with G418.

**Immunofluorescence analysis and RNA fluorescent *in situ* hybridization.** Lyn and different MBNL1 isoforms were localized as detailed in the Supplementary Experimental Procedures. CUG-containing foci were detected in muscle sections and myoblasts as described.<sup>36</sup> Colocalization was evaluated with JACoP plugin<sup>59</sup> for the ImageJ software by calculating Manders' overlap coefficients  $M1$  (fraction of A overlapping B) and  $M2$  (fraction of B overlapping A). Threshold was arbitrarily set to non-saturating levels for Lyn and MBNL, whereas it was left saturated for 4',6-diamidino-2-phenylindole (DAPI) images. Sets of seven  $M1$  and  $M2$  coefficients per pairwise analysis were collected by increasing the threshold. At least three different images were used for each condition. Data are shown as mean  $\pm$  S.E.M.

**Anti-P9 and anti-P11 antibodies: generation, immunoblotting and immunoprecipitation.** MBNL1 antibodies were raised in rabbits against amino acids 270–288 (TQSAVKSLKRPLEATFDL; this epitope is encoded by exon 5 of MBNL1 gene), hereafter defined as P9, and 294–308 (LPPLPKRPALEKTNG, encoded by exon 6 of MBNL1 gene), hereafter defined as P11, of the human MBNL1<sub>43</sub> protein (Supplementary Material and Supplementary Figure S1A) with an extra cysteine at the C terminus. Peptide synthesis, rabbit immunization and purification of anti-MBNL1 antisera and use of antibodies in immunoblotting and immunoprecipitation experiments are described in detail in the Supplementary Experimental Procedures.

**Surface plasmon resonance.** The interaction between MBNL1 isoforms and Lyn was performed as described below. Recombinant MBNL1 isoforms were immobilized on channels 1 and 3 of a His Cap research-grade sensor chip (Nomadics, ICx Nomadics Bioinstrumentation, Group, Oklahoma City, OK, USA) by nickel-affinity interaction using the manufacturer's protocols. Channel 2 was used as reference for all experiments. The Lyn kinase SH3 domain was the analyte and it was injected at 30  $\mu$ l/min. SPR measurements were carried out in HBS buffer with 0.005% P20 surfactant at 20 °C using a SensiQ Pioneer Instrument (ICx Technologies, Stillwater, OK, USA). The data were analyzed with the Qdat evaluation analysis software (ICx Nomadics Bioinstrumentation, Group).

**Phosphorylation and interaction assays.** Tyr kinase assays were performed in 40  $\mu$ l reaction mixture containing 50 mM Tris/HCl, pH 7.5, 10 mM MnCl<sub>2</sub>, 30  $\mu$ M ATP/[ $\gamma$ -<sup>33</sup>P]ATP (specific activity 1000 c.p.m./pmol) (Perkin-Elmer, Waltham, MA, USA), 100  $\mu$ M sodium orthovanadate, 200  $\mu$ M Src-specific peptide substrate cdc2(6–20) as substrate and 20 ng of Src, or Lyn. Following incubation for 5 min at 30 °C, the reaction was blocked by adding 5  $\times$  SDS buffer and the samples were subjected to SDS-PAGE. Peptide phosphorylation was evaluated using the Cyclone Plus Storage Phosphor System (Perkin-Elmer). In time-course experiments, MBNL1<sub>40-41</sub> and MBNL1<sub>42-43</sub> (each 1  $\mu$ M) were phosphorylated in the reaction mixture and the reactions were stopped at different time points as indicated above. The degree of protein phosphorylation was evaluated by analysis on the Cyclone Plus Storage Phosphor System (Perkin-Elmer). Interaction assay (between MBNL1 isoforms and Lyn) followed by immunoprecipitation were performed as described in the Supplementary Experimental Procedures.

**Statistical analysis.** Statistical analysis was performed only where three or more experimental values were available. Quantitative data were presented as mean  $\pm$  S.D. The statistical comparisons were performed using Mann–Whitney and Kruskal–Wallis tests, for comparison between two or more groups, respectively. In every analysis, values of  $P < 0.05$  were considered significant.

## Conflict of Interest

The authors declare no conflict of interest.

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