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TDP-43 accumulation in IBM muscle suggests a common pathogenic mechanism with Frontotemporal dementia

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Abstract

TDP-43 is found in ubiquitinated inclusions (UBIs) in some frontotemporal dementias (FTD-U). One form of FTD-U, due to mutations in VCP, occurs with an inclusion body myopathy (IBMPFD). Since IBMPFD brain has TDP-43 in UBIs, we looked for TDP-43 inclusions in IBMPFD muscle. In normal muscle TDP-43 is present in nuclei. In IBMPFD muscle TDP-43 is additionally present as large inclusions within UBIs in muscle cytoplasm. TDP-43 inclusions were also found in 78% of sIBM muscles. In IBMPFD and sIBM muscle TDP-43 migrated with an additional band on immunoblot similar to that reported in FTD-U brains. This study adds sIBM and hereditary inclusion body myopathies to the growing list of TDP-43 positive inclusion diseases.

Keywords

TDP-43; inclusion body myositis; ubiquitin; frontotemporal dementia

Introduction

TAR DNA binding protein-43 (TDP-43) is one component of the ubiquitinated inclusions in brains of patients with FTD-U and ALS¹. Moreover, a phosphorylated form of TDP-43 was identified to be more prevalent specifically in FTD-U tissue as well¹. Little is known about the function of TDP-43. It is a ubiquitously expressed, highly conserved nuclear protein that may be a transcription repressor or activator of exon skipping as well as a scaffold for nuclear bodies through interactions with survival motor neuron protein¹. The significance of TDP-43 accumulation and phosphorylation is currently unclear but may relate to ubiquitin-proteasome

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system (UPS) dysfunction. One rare form of FTD-U, inclusion body myopathy, Paget's disease of the bone and frontotemporal dementia (IBMPFD) has ubiquitinated and TDP-43 positive inclusions in affected neurons^{2, 3}. While IBMPFD muscle contains rimmed vacuoles and UBIs consistent with an inclusion body myopathy (IBM)⁴.

The pathogenesis of IBM and the more common sporadic inclusion body myositis (sIBM) is unknown but may also be due to UPS dysfunction⁵. Affected muscle has UBIs that contain proteins, such as β -amyloid and phosphorylated tau, known to aggregate in CNS degenerative disorders^{6, 7}. This has led to the suggestion that sIBM is related pathophysiologically to neurodegenerative diseases. It is not known whether TDP-43 is a component of the inclusions in IBMPFD and sIBM muscle tissue. We evaluated the localization of TDP-43 in normal, IBMPFD and sIBM skeletal muscle tissue.

Methods

Patients with sIBM had typical patterns of muscle weakness on physical examination, an abnormal EMG with myopathic motor units and spontaneous activity, and a muscle biopsy with myopathic changes, rimmed vacuoles within muscle fibers and endomysial inflammation with focal invasion of muscle fibers⁸. Five patients with IBMPFD and missense mutations in the p97/VCP gene (four with R155H and one with N387H)⁹ were participants in the IRB approved study. All muscle biopsies were processed and evaluated in the Washington University Neuromuscular Laboratory. Cryostat sections of rapidly frozen muscle were processed for muscle histochemistry and immunocytochemistry in our standard fashion^{4, 9}. The presence of vacuoles was evaluated via routine histochemical methods such as hematoxylin and eosin or modified gomoritricrome stains. Immunocytochemistry for each antibody was performed on tissue from patients and compared to normal tissue controls processed simultaneously. Primary antibodies used in this study were directed against CD8 (clone M7103), phosphorylated neurofilaments or other phosphorylated epitopes (SMI-31 Covance, Berkeley, CA), FK2 antibody to ubiquitinated proteins (PW8810-0500 BioMol, Plymouth Meeting, PA), TDP-43 antibodies: rabbit polyclonal antibody (ProteinTech Antibody Group, Chicago, IL) and mouse monoclonal antibody 2E2-D3 (Abnova, Taipei, Taiwan). Double-labeling immunofluorescence was performed as previously described using Alexa Fluor 488 and 594 conjugated secondary antibodies (Molecular Probes, Eugene, OR). Immunoperoxidase was performed as previously described using peroxidase conjugated secondary antibodies (Sigma; St. Louis, MO, USA). The specificity of TDP-43 immunostaining was confirmed with two different commercial antibodies and incubation with secondary antibody alone (either fluorescent or peroxidase conjugated). Immunoblots were performed as previously described¹⁰.

Results

In 4 normal muscle biopsies TDP-43 was localized within scattered myonuclei with no evidence of sarcoplasmic staining (Figure 1A). In contrast, all IBMPFD patients muscle tissue had large peripherally based TDP-43 positive sarcoplasmic inclusions that did not localize to myonuclei (Figure 1B–C). These inclusions consistently co-localized with FK2, an antibody that recognizes ubiquitinated proteins (Figure 1D) and in some cases with other proteins known to aggregate. SMI-31 binding was less prominent than TDP-43 in IBMPFD patient muscle tissue (Figure 1E).

A distinctively different pattern of TDP-43 immunostaining was seen in 21 of 27 sIBM patients muscle. TDP-43 immunostained multiple small sarcoplasmic aggregates, most commonly in small angular muscle fibers (Figure 2A). These inclusions did not co-localize with myonuclei. TDP-43 was also present in debris surrounding some rimmed vacuoles (Figure 2A). The

TDP-43 inclusions in sIBM were usually ubiquitin negative (Figure 2B), but occasionally co-localized with FK2. TDP-43 also co-localized with T-cells at sites of inflammatory infiltrates (Figure 2C). In contrast to sIBM, TDP-43 positive inclusions were found in only 1 of 12 steroid responsive polymyositis patient biopsies.

Immunoblots of normal, IBMPFD and sIBM patient tissue with an antibody to TDP-43 demonstrated an increase in TDP-43 immunoreactivity present at 43kDa as well as a higher migrating band similar to the phosphorylated form seen in FTD-U patient tissue (Figure 2D).

Discussion

Disruptions in the UPS may be associated with the pathogenesis of several degenerative disorders. In particular, IBM muscle and FTD-U brain have UBIs that contain aggregated proteins. However, the principal molecular constituents of the UBIs seen in these diseases have been incompletely defined. Recent studies have identified TDP-43 as a component of the UBIs in FTD-U, including IBMPFD, and ALS brain tissue^{1, 3}. Since IBMPFD muscle also has UBIs, we examined the localization of TDP-43 in normal, sIBM and IBMPFD patient skeletal muscle. We found that TDP-43 localized to myonuclei in normal muscle, but, in IBMPFD and sIBM patient muscle, TDP-43 was additionally present as sarcoplasmic inclusions. This is associated with an increase in TDP-43 protein levels as well as a higher molecular weight band seen in sIBM and IBMPFD patient muscle tissue via immunoblot when compared with normal patient tissue. A similar higher molecular weight band was identified as phosphorylated TDP-43 in patients with FTD-U and ALS¹. Whether the band seen in IBMPFD and sIBM muscle tissue is the same or another post-translationally modified form (i.e. ubiquitinated) is not known.

TDP-43 inclusions were present in 100% of IBMPFD and 78% of sIBM patient muscle biopsies, while 0% of normal muscle and 8% of steroid responsive polymyositis patient muscle biopsies had similar TDP-43 inclusions. This suggests that TDP-43 immunohistochemistry may be helpful in confirming the diagnosis of sIBM. At present, the most reliable antibody marker for the diagnosis of sIBM is SMI-31 (an antibody against phosphorylated tau epitopes) which is not present in all sIBM biopsies and found in other diseased muscle tissue¹¹. For example in our hands SMI-31-positive aggregates are present in 66% of sIBM patients compared with 17% of steroid responsive polymyositis patients (unpublished observations).

It is notable that the TDP-43 immunostaining pattern was different when comparing sIBM and IBMPFD muscle. sIBM had small punctuate TDP-43 positive inclusions throughout the sarcoplasm that occasionally co-localized with ubiquitin. IBMPFD muscle had large peripherally based TDP-43 positive inclusions that always co-localized with ubiquitin. Both sIBM and IBMPFD patient muscle had evidence of a higher molecular weight TDP-43 species on immunoblot.

It is unknown if TDP-43 accumulation in FTD-U, ALS or IBM is pathogenic or merely a pathologic feature of the diseased tissue¹². Our findings in skeletal muscle are similar to those found in CNS disease¹. TDP-43 inclusions in FTD-U and ALS are cytoplasmic, similar to the predominantly sarcoplasmic inclusions in IBM muscle. The presence of ubiquitinated TDP-43 in disparate tissues (i.e. IBMPFD muscle and brain) in which the only commonality is mutant VCP, suggests that VCP dysfunction may play a role in TDP-43 aggregation.

TDP-43 contains RNA binding motifs that may be associated with its function. Interestingly, another hereditary IBM, oculopharyngeal muscular dystrophy, is due to mutations in another RNA binding protein PABPN1¹³. Mutant PABPN1 forms intranuclear inclusions that contain ubiquitin and polyadenylated mRNA¹⁴. It is not known whether a similar process occurs with TDP-43 in sIBM and IBMPFD.

Diseases that develop TDP-43 inclusions include FTD-U, ALS, IBMPFD and sIBM^{1,3} suggesting that similar pathogenic mechanisms may be present. Further studies will be needed to further define the role of these proteins and their dysregulation in central nervous system and skeletal muscle tissue.

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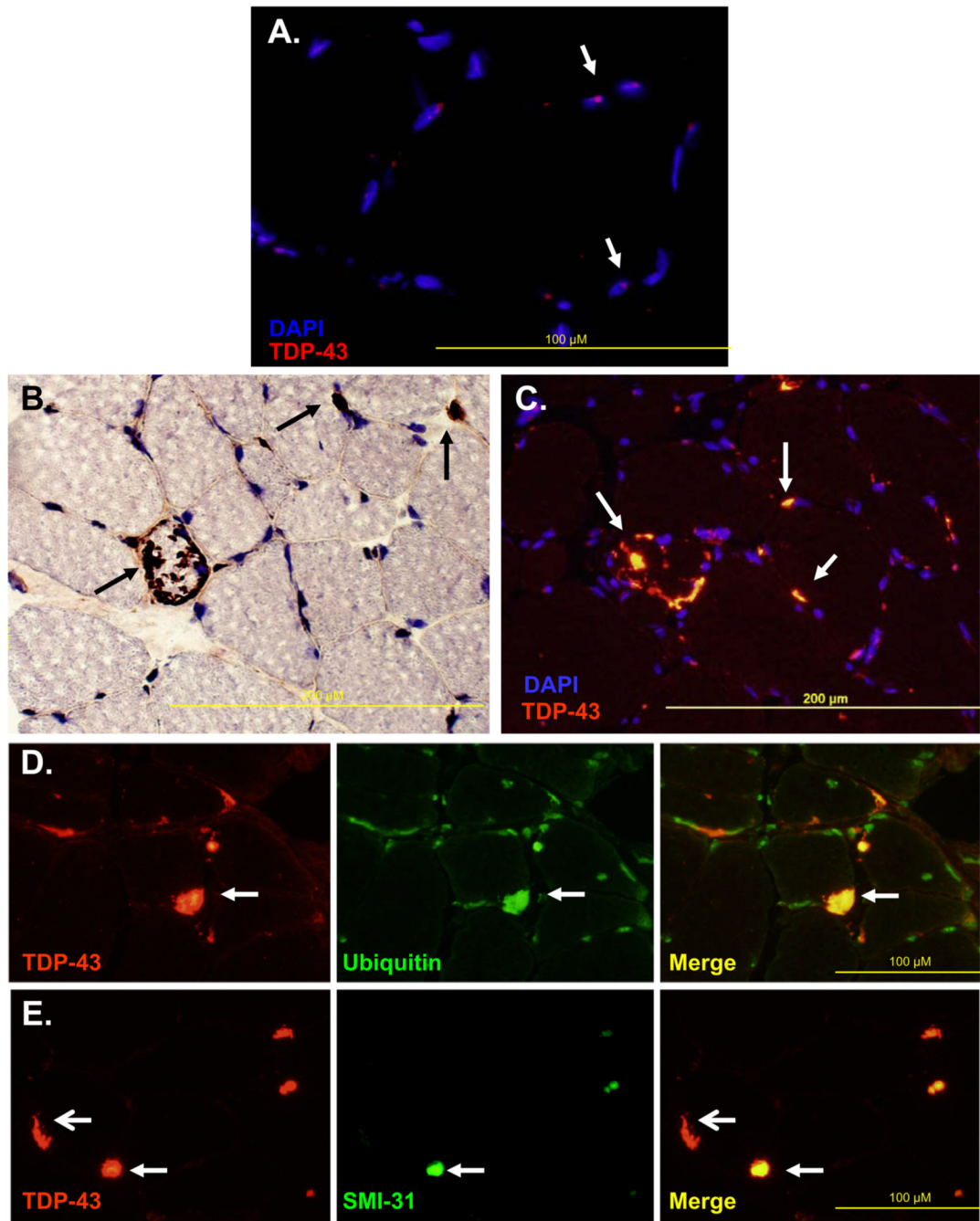


Figure 1.

A) Normal muscle immunostained with anti-TDP-43 antibody and counterstained with DAPI to allow visualization of nuclei. Figure is overlay of anti-TDP-43 and DAPI images. Arrows denote blue nuclei with TDP-43 (red dots) in scattered myonuclei. B) IBMPFD patient tissue immunostained with anti-TDP-43 (brown) and counterstained with Congo red to allow visualization of nuclei and myofibers. Arrows denote large inclusions, some of which are peripherally based. C) Overlay of anti-TDP-43 (orange) and DAPI (blue) of IBMPFD patient tissue. Note that large peripheral inclusions do not localize within nuclei (arrows). D) IBMPFD patient tissue co-immunostained with anti-TDP-43 (red) and FK2 (green). Note that TDP-43 inclusions co-localize with FK2 (ubiquitinated proteins) (arrows). E) IBMPFD patient tissue

co-immunostained with anti-TDP-43 (red) and SMI-31, an antibody against phosphorylated tau epitopes (green). Note that some TDP-43 inclusions co-localize with SMI31 (closed arrows) and others do not (open arrows).

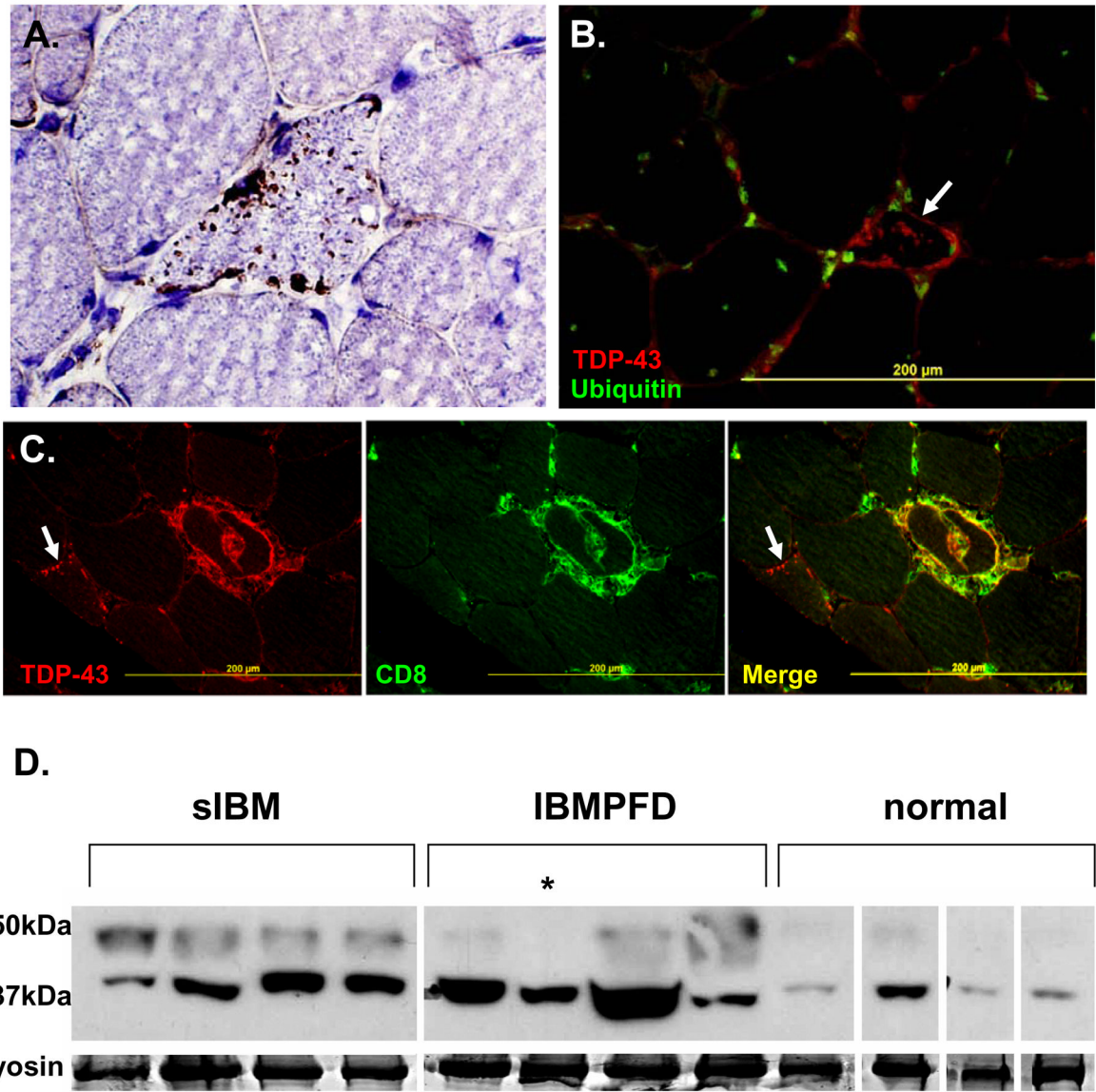


Figure 2.

A) sIBM patient tissue immunostained with anti-TDP-43 (brown) and counterstained with Congo red to allow visualization of nuclei and myofibers. TDP-43 inclusions are small, in angular fibers and occasionally surround rimmed vacuoles. B) Overlay of sIBM patient biopsy immunostained with TDP-43 (red) and FK2, for ubiquitinated proteins (green). TDP-43 inclusions do not co-localize with ubiquitin. C) sIBM patient tissue co-immunostained with anti-TDP-43 (red) and anti-CD8 (green at focal sites of inflammation). Note the co-localization of TDP-43 and CD8. This is in contrast to an adjacent fiber (arrow) with sarcoplasmic TDP-43 inclusions that does not co-localize with CD8. D) Muscle tissue from sIBM, IBMPFD and normal patient biopsies were homogenized and separated by SDS-PAGE. The subsequent gel was transferred to nitrocellulose and immunoblotted with an antibody to TDP-43. Normal tissue has a discrete band at 43kDa consistent with TDP-43, while sIBM and IBMPFD have a more prominent band and also a higher migrating band. Myosin is show as a loading control. All tissues are from flash frozen biopsies except the lane noted with an * which is from autopsy muscle. All bands are from the same autoradiograph and moved for presentation purposes.