ANIMAL MODELS

Motor Deficit in a Tauopathy Model Is Induced by Disturbances of Axonal Transport Leading to Dying-Back Degeneration and Denervation of Neuromuscular Junctions

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Several neurodegenerative diseases are characterized by both cognitive and motor deficits associated with accumulation of tau aggregates in brain, brainstem, and spinal cord. The Tg30 murine tauopathy model expresses a human tau protein bearing two frontotemporal dementia with Parkinsonism linked to chromosome 17 pathogenic mutations and develops a severe motor deficit and tau aggregates in brain and spinal cord. To investigate the origin of this motor deficit, we analyzed the age-dependent innervation status of the neuromuscular junctions and mutant tau expression in Tg30 mice. The human transgenic tau was detected from postnatal day 7 onward in motoneurons, axons in the sciatic nerve, and axon terminals of the neuromuscular junctions. The development and maturation of neuromuscular junctions were not disrupted in Tg30 mice, but their maintenance was disturbed in adult Tg30 mice, resulting in a progressive and severe muscle denervation. This muscle denervation was associated with early electrophysiological signs of muscle spontaneous activities and histological signs of muscle degeneration. Early loss of synaptic vesicles in axon terminals preceding motor deficits, accumulation of Gallyas-positive aggregates, and cathepsin-positive vesicular clusters in axons in the sciatic nerve suggest that this denervation results from disturbances of axonal transport. This physiopathological mechanism might be responsible for motor signs observed in some human tauopathies, and for synaptic dysfunction resulting from alterations at the presynaptic level in these diseases. (Am J Pathol 2015, 185: 2685–2697; http://dx.doi.org/10.1016/j.ajpath.2015.06.011)

Tauopathies are a class of neurodegenerative diseases characterized by the presence of aggregates of abnormally hyperphosphorylated forms of the microtubule-associated protein tau (MAPT) in nerve cells and/or glial cells.1 Tauopathies include Alzheimer disease, some forms of frontotemporal lobar degeneration, Pick’s disease, progressive supranuclear palsy, and corticobasal degeneration, among others. Several of these tauopathies are characterized by both cognitive and motor deficits and tau pathology in brain, brainstem, and spinal cord. Frontotemporal lobar degeneration patients can develop both dementia and a motoneuron disease with loss of motoneurons in spinal cord.2 The Guam amyotrophic lateral sclerosis/Parkinsonism-dementia complex is characterized by a motor neuron disease with widespread neurofibrillary tangles (including some in the spinal cord), Parkinsonism, and dementia.3 Frontotemporal dementia with Parkinsonism linked to chromosome 17 (FTDP-17) is caused by mutations in the MAPT gene4 and presents with behavioral, language, and motor abnormalities.5

Tau is mainly expressed in neuron, and its primary function, by binding to microtubules, is to regulate or...
modulate the stability of axonal microtubules. Other functions of tau include a role in signal transduction, the regulation of axoplasmic transport, and interaction with actin cytoskeleton. FTDP-17 tau mutations reduce the ability of tau to interact with microtubules, promote the assembly of tau into filaments, or perturb the normal ratio of tau isoforms, leading to abnormal aggregation of tau.

Expression of FTDP-17–associated mutant tau proteins in transgenic mice induces development of tau pathology and neurofibrillary tangles, and several of these models develop motor deficits. To better understand the development of motor impairment in tauopathies, we have analyzed the age-dependent expression of mutant tau and the age-dependent innervation status of the neuromuscular junction in the Tg30tau transgenic model. Tg30tau transgenic mice express a human tau protein bearing two FTDP-17 pathogenic mutations (P301S and G272V) in the forebrain and in the spinal cord and develop tau pathology and neurofibrillary tangles. Tg30tau mice develop a severe motor deficit beginning at 8 months, an axonopathy and severe muscle atrophy. We observed that the expression of human transgenic tau impaired the maintenance of neuromuscular junctions (NMJ), leading to muscle denervation in Tg30 mice. Loss of synaptic vesicles at the NMJ and accumulation of tau inclusions and cathepsin-positive vesicles suggest the existence of disturbance of axonal transport in the sciatic nerve of Tg30 mice. Similar alterations might explain the development of a motoneuron phenotype in some human tauopathies.

Materials and Methods

Animals

Tg30tau transgenic mice were generated and characterized as described. Briefly, these mice express a 1N4R human tau isoform mutated at positions G272V and P301S, under the control of a Thy1 promoter. Only heterozygous animals were used here, and nontransgenic littermates were used as wild-type (WT) controls. Mice were genotyped by PCR as reported. All mice were maintained on a 12-hour light–dark cycle, with food and water ad libitum. All studies on animals were performed in compliance and following approval of the ethical committee for the care and use of laboratory animals of the medical School of the Free University of Brussels.

Tissue Preparation

Newborn and adult mice were sacrificed by decapitation or cervical dislocation, respectively. The brain, the lumbar portion of the spinal cord, sciatic nerve, and left hindlimb muscles were dissected and then formalin-fixed overnight at room temperature. Samples were embedded in paraffin and cut at 7 μm thickness using a microtome (Leica Leitz 1400; Leica Microsystems, Diegem, Belgium). Right hindlimb muscles were fixed in 4% paraformaldehyde in phosphate-buffered saline (PBS), pH 7.4, at 4°C for 20 to 60 minutes according to their developmental stage. After fixation, the tissues were rinsed three times in PBS and then cryoprotected in 20% sucrose/PBS for 48 hours. Tissues were embedded in Tissue Tek (Prolabo; VWR International, Leuven, Belgium) and cut at 20 μm thickness using a cryostat (Leica CM3050 S). Cryosections were dried at room temperature and stored at −20°C.

Immunofluorescence Labeling of Neuromuscular Junctions

Muscle cryosections (tibialis anterior, gastrocnemius, and quadriceps) were treated with 0.1 mol/L glycine in PBS for 30 minutes before processing. After washes with PBS, sections were permeabilized with PBS/0.3% TritonX-100 for 10 minutes and treated with PBS/10% horse serum/0.1% Triton (buffer A) for 1 hour at room temperature. The primary antibodies were diluted in the buffer A and incubated overnight at 4°C on tissue sections. After washes in PBS/Triton 0.1%, the secondary antibodies and z-bungarotoxin (BTX) were diluted in buffer A and incubated on tissue sections for 1 hour at room temperature. After washes in PBS/Triton 0.1%, the slides were mounted with Gelvatol containing DABCO 0.1 g/mL (Sigma-Aldrich, Diegem, Belgium).

Primary antibodies for immunofluorescence were mouse monoclonal anti-neurofilament (NF; clone SM132, NE1023, dilution 1:1000; Millipore, Overijse, Belgium), rabbit monoclonal anti-synaptophysin (#01-1019, dilution 1:500, Millipore) and rabbit polyclonal anti-human tau (BR21, dilution 1:500). Secondary antibodies were goat anti-mouse antibody conjugated to fluorescein isothiocyanate (F-1010, dilution 1:250; Sigma-Aldrich) or a goat anti-rabbit antibody conjugated to biotin (BA-1000, 1:300; Vector Laboratories, Burlingame, CA) followed by streptavidin conjugated to AlexaFluor 488 (S-11223, dilution 1:250; Life Technologies, Merelbeke, Belgium). BTX conjugated to Alexa594 (B13423, dilution 1:1000; Life Technologies) was incubated with the secondary antibodies. Pictures were taken with an Axiovert 200M fluorescence microscope (Carl Zeiss, Zaventem, Belgium). For all images, brightness and contrast were adjusted with Adobe Photoshop CS3 extended version 10.0 (Adobe Systems, San Jose, CA) after acquisition to match with the observation.

Analysis of the Maturation and the Innervation Status of Motor Endplates

Motor endplate maturation was evaluated in postnatal day (P)14 newborn mice based on criteria previously described. A minimum of 100 junctions was analyzed for each animal (n = 3). The innervation status of motor endplates was assessed after labeling with the synaptophysin antibody and BTX by classifying the synaptophysin distribution into three categories: i) perfect superposition of
the BTX and synaptophysin labeling; ii) partial superposition of synaptophysin and BTX labeling; and iii) absence of synaptophysin labeling. The innervation status of NMJ was also assessed after labeling with NF and BTX by categorizing them as innervated if there was any overlap between the two labels or denervated if NF was completely absent at the synaptic junction. A minimum of 300 junctions was analyzed for each mouse (n = 3).

Immunocytochemistry

The immunohistochemical labeling was performed using the ABC method. Briefly, tissue sections were treated with peroxide for 30 minutes to inhibit endogenous peroxidase. After washes in Tris buffer saline (TBS), sections were incubated with the blocking solution (10% horse serum or goat serum in TBS/0.13% sodium azide). The primary antibodies were diluted in 1% horse serum or goat serum/TBS/0.13% NaN3 and incubated on tissue sections overnight at room temperature. After washes in TBS, sections were sequentially incubated with either horse anti-mouse or goat anti-rabbit antibodies conjugated to biotin (Vector Laboratories) for 30 minutes at room temperature, followed by the ABC complex (Vector Laboratories). After washes in TBS, the peroxidase activity was detected using diaminobenzidine as chromogene (Vector Laboratories). In some case, the slides were counterstained with hematoxylin and eosin (H&E). The slides were mounted with Depex (VWR International).

The primary antibodies for immunocytochemistry were mouse monoclonal anti-NeuN (NEUN-A60, dilution 1:10,000; Millipore), rabbit polyclonal anti-human tau (BR21, dilution 1:5000),12 and rabbit polyclonal anti-cathepsin (BR6, dilution 1:5000).17

Gallyas Staining

Gallyas staining was used for the detection of tau aggregates. Formalin-fixed, paraffin-embedded sections of the sciatic nerve (7 μm thick) from 7- and 10-month-old WT and Tg30 mice were stained with a modified Gallyas silver-staining method10,18 and counterstained with 0.1% Nuclear Fast Red.

Motor Neuron Counts

The number of motoneurons in lumbar spinal cord sections was quantified after staining with cresyl violet stain and after immunohistochemistry for NeuN. Counting of motoneurons was performed on the left and right ventral sides of three spinal cord sections for each mouse (n = 3). Motoneurons were identified on the basis of the large size of their cell body and the cytoplasmic abundance of Nissl bodies.

Fiber Diameter Analysis

A fragment of hindlimb muscle was dissected at 1 month and the tibialis anterior, gastrocnemius, and quadriceps muscles were dissected at 3 and 10 months. They were formalin-fixed, paraffin-embedded, cut (7 μm thick), and stained with H&E. Pictures were acquired with a DMS5000 microscope with LAS EZ software version 3.0 (Leica Microsystems). The muscle fiber cross-sectional area (CSA) was measured on 200 fibers for quadriceps and gastrocnemius muscles and on 100 fibers for tibialis and hindlimb muscles of mice, age 1 month, in each animal (n = 3) using the ImageJ software version 1.49s (NIH, Bethesda, MD; http://imagej.nih.gov/ij).

Electromyography

Electromyography recordings were performed in control and Tg30 mice ages 2 months to 11 months; measurements were conducted at 2 (n = 4), 3 (n = 4), 6 (n = 5), 8 (n = 3) and 11 (n = 4) months of age. Repeated measurements were obtained at 1-month intervals for four mice and at 5-month intervals for four other mice. The animals were first sedated by intraperitoneal injection of 1 mg/g body weight of medetomidine (Sedator; Eurovet Animal Health, Bladel, the Netherlands). Anesthesia was induced and maintained using isoflurane (Isoflo, 1.5% in oxygen; Abbott Laboratories, Wavre, Belgium). During the anesthesia, the
Figure 2 Early muscle hypertrophy and late myopathy in mutant tau Tg30 mice. A and B: H&E staining of transverse sections of hindlimb muscles from 1-month-old WT (A) or Tg30 (B), showing a similar aspect in both groups. C: Distribution of muscle fiber cross-sectional areas in hindlimb muscles of 1-month-old WT (black bars) or Tg30 (gray bars) mice. The distribution of fiber areas in posterior muscle is similar in WT and in Tg30 mice at 1 month. D and E: H&E staining of transverse sections of quadriceps muscle from 3-month-old WT (D) or Tg30 mice (E). Some signs of muscle degeneration are observed in Tg30 mice as shown by the presence of nuclei in the center of muscle fibers (arrowhead) and of clusters of fibers with reduced size. Numerous fibers in Tg30 mice have a large size. F: Distribution of muscle fiber cross-sectional areas in quadriceps muscle of 3-month-old WT or Tg30 mice. The distribution of fiber areas in quadriceps muscle indicates that there is a significantly increased proportion of the largest muscle fibers in Tg30 mice at 3 months. G and H: H&E staining of transverse sections of quadriceps muscle from 10-month-old WT (G) or Tg30 (H) mice. There is a reduction in cross-sectional areas of many muscle fibers and frequent nuclei are located in the center of muscle fibers (arrowhead), indicative of muscle degeneration. I: Distribution of muscle fiber areas in quadriceps muscle of 10-month-old WT or Tg30 mice. There is a significantly increased proportion of the smallest muscle fibers in Tg30 mice at 10 months. J–M: Electromyogram recordings of gastrocnemius muscle in WT (J) or Tg30 (K–M) mice aged 2 (J and K), 9 (L), and 11 (M) months. The number of spontaneous muscle activities has a net tendency to increase with aging in Tg30 mice. n = 3 (F and I), P < 0.001 by χ² Pearson’s test [comparison of 1 month (C) with 3 months (F), and 10 months (I) with 1 month (C)]. Scale bars: 50 μm (A and D). H&E, hematoxylin and eosin; WT, wild type.
animals were placed on a heated pad (38°C). A Medelec Sierra II apparatus (Medelec Instruments, New Delhi, India) connected to a personal computer running the program Sierra XP version 3.3.3 was used for both stimulation and recordings. The stimulation of the sciatic nerve was performed using percutaneous needle electrodes placed caudal to the greater trochanter, close to the sciatic nerve. Recordings were documented using subcutaneous needle electrodes inserted at the level of the gastrocnemius muscle in a belly-tendon configuration; a subcutaneous ground electrode was positioned midway between the stimulating cathode and the recording electrode. The maximum amplitude of the muscle compound action potential was recorded, and its decrement during a train of 10 stimulations at 3 per second was measured (calculated as the difference between the amplitude of the tenth and first response divided by the amplitude of the first response). Detection electromyography was performed with concentric needle electrode (25 × 0.3 mm; Technomed, Maastricht, the Netherlands) inserted in the leg muscles at least at three locations, and spontaneous muscle activities were recorded. The amplitudes of the muscle potential evoked by the sciatic nerve stimulation were compared using a t-test for paired samples in the 2 × 4 mice groups.

Figure 3 The initial formation of NMJ and their innervation is not disrupted in Tg30 mice. A–D: Double labelling on muscle fiber sections of acetylcholine receptors in muscle motor endplate by α-bungarotoxin (A and C) and immunofluorescence detection of synaptophysin in motor nerve terminal (B and D) in WT (A and B) or Tg30 (C and D) mice at P14. Insets in B and D show overlapping BTX/SYN labelling illustrated in A and B, and C and D, respectively. The progressively increased apposition of the motor nerve terminals to motor endplates observed during the maturation of the motor endplates happens normally in Tg30 mice. E: Quantification of NMJ maturation in WT or Tg30 mice at P14 based on four morphological stages of AChR patterns of distribution (M1: oval plaque; M2: single perforation; M3: multiple perforations; M4: mature junction). The proportion of each stage is similar in WT and Tg30 mice, indicating that the maturation of the NMJ junctions is normal in Tg30 mice. Error bars indicated means ± SEM. F and G: Labeling of acetylcholine receptors by α-bungarotoxin and immunofluorescence detection of neurofilament on muscle fiber sections of WT (F) and Tg30 (G) mice at P14. The innervation of motor endplates is normal in Tg30 mice. H: Quantification of muscle mono- or polyinnervation at P14 using NF/BTX overlap as criterion. The proportion of mono- or polyinnervated endplates is similar in WT and Tg30 mice. The t-test showed nonsignificant results. n = 3 (E and H, each group). Scale bar = 5 μm. AChR, acetylcholine receptor; BTX, α-bungarotoxin; NF, neurofilament; NMJ, neuromuscular junctions; P, postnatal day; SYN, synaptophysin; WT, wild type.
Muscle Denervation in a Tauopathy Model

Statistics

Data are presented as means ± SEM. Unpaired t-test and χ² Pearson’s test were used to determine statistical significance using GraphPad Prism version 4 software (San Diego, CA). Differences at \( P < 0.05 \) were considered significant.

Results

Human Tau Transgenic Protein Is Expressed in the Spinal Cord of Tg30 Mice from P7

To determine the timing of postnatal expression of human tau transgenic protein, an immunocytochemical labeling with the BR21 antibody specific for human tau was performed on traverse sections of the lumbar spinal cord of Tg30 and WT mice at P1, P7, and P14. At P1, the human tau transgenic protein was not detected in the spinal cord of Tg30 newborn mice (Figure 1A) but was detected at P7 in the cell bodies and in some axonal processes in motoneurons of the ventral horn of Tg30 mice (Figure 1B). The expression of human tau was increased in Tg30 mice at P14, and the protein was clearly detected in neuronal processes (Figure 1C). Human tau protein was also detected from P7 in the brain of Tg30 mice (data not shown). Thus, human tau transgenic protein was expressed early in motoneurons of the ventral horn in the spinal cord and in the brain from P7 in Tg30 newborn mice.

Tg30 Mice Show an Early Hypertrophy of Muscle Fibers and a Late Hypotrophy of Muscle Fibers

Tg30 mice develop a motor deficit of hindlimbs from 8 months of age that is severely worsened at 12 months. To analyze the evolution of muscular atrophy, the mean CSA of muscle fibers in hindlimb muscles was measured at 1, 3, and 10 months in Tg30 mice. The mean CSA and the distribution of muscle fibers areas were similar in WT and Tg30 mice at 1 month (Figure 2, A–C). By contrast, at 3 months, the CSA in quadriceps muscle (Figure 2, D–F) and gastrocnemius muscle (data not shown) increased in Tg30 mice and was associated with a significantly increased proportion of muscle fibers with large sizes (Figure 2F). At 3 months, the CSA in tibialis muscle was similar in WT and Tg30 mice (data not shown). Furthermore, nuclei in the center of muscle fibers were frequently observed in Tg30 mice at 3 months, suggesting that Tg30 muscle fibers undergo degeneration (Figure 2, D–F). At 10 months, the CSA in gastrocnemius, quadriceps, and tibialis muscles was decreased in Tg30 mice and was associated with a significantly increased proportion of muscle fibers with small sizes (Figure 2I). In addition, muscle fibers with central nuclei were frequently observed in Tg30 mice at 10 months (Figure 2, G–I).

Thus, at 3 months, Tg30 mice showed an increased proportion of hypertrophic muscle fibers in gastrocnemius and quadriceps muscles associated with the beginning of muscle fiber degeneration. Whereas at 10 months, Tg30 mice showed an increased proportion of hypotrophic muscle fibers with robust signs of muscle fiber degeneration.

Absence of Motoneuron Loss in the Spinal Cord of Tg30 Mice

To rule out the possibility that muscle atrophy in Tg30 mice could be the consequence of a denervation due to neuronal loss in the spinal cord, the number of motoneurons was evaluated in the lumbar spinal cord of WT and Tg30 mice at different ages (P1, P7, P14, 1, 3, and 10 months). The density of lumbar motoneurons was similar in WT and Tg30 mice at all investigated ages (Figure 1D). These results

Figure 4 Expression of human tau transgenic protein leads to NMJ denervation in adult Tg30 mice. A–F: Labeling of acetylcholine receptors by α-bungarotoxin (red, A–C) and immunofluorescence labeling of synaptophysin (green, D–F) on muscle fiber sections of 1-month-old WT (A and D) or Tg30 (B, C, E, and F) mice. Insets in D, E, and F show overlapping BTX/SYN labeling illustrated in A and D, B and E, and C and F, respectively. A and D: In WT mice, NMJ display the expected pretzel-like shape and show perfect apposition of the synaptophysin labeling with the motor endplate. B and E: In Tg30 mice, a majority of NMJ are very similar to those observed in WT animals. C and F: In 1-month-old Tg30 mice, a minority of NMJ junctions show a defective localization of synaptophysin, which only partially overlaps with AChR labeling (the asterisk in F shows an area of a motor endplate devoid of overlapping synaptophysin labeling). G: Quantification of patterns of synaptophysin distribution in NMJ of WT or Tg30 mice at 1 month. Both in WT and in Tg30 mice, the vast majority of NMJ shows a perfect superposition of synaptophysin and α-bungarotoxin labelling (black), and a minority shows a partial overlap (gray). H and I: Labeling of acetylcholine receptors by α-bungarotoxin (red) and immunofluorescence detection of neurofilament (green) on muscle fiber sections of 1-month-old WT (H) or Tg30 (I) mice. All of the junctions were innervated in Tg30 mice. J: Quantification of muscle innervation at 1 month in WT and Tg30 mice using NF/BTX overlap as criterion. The vast majority of junctions are innervated in both WT and Tg30 mice. K–R: Labeling of acetylcholine receptors by α-bungarotoxin (red, K–N) and immuno-fluorescence labeling of synaptophysin (green, O–R) on muscle fibers section of 10-month-old WT (K and O) or Tg30 (L–R) mice. Insets in O, P, and Q show overlapping BTX/SYN labeling illustrated in K and O, L and P, and M and Q, respectively. K and O: In WT mice, NMJ display the expected pretzel-like shape and show perfect apposition of the synaptophysin labeling to the motor endplate. L and P: In Tg30 mice, a minority of NMJ are very similar to those observed in WT animals. M, N, and R: A majority of junctions in Tg30 mice show defective localization of synaptophysin, which only partially overlaps with AChR labeling (M and Q, arrowhead) or is absent (N and R, double asterisks). S: Quantification of synaptophysin distribution in NM2 of WT or Tg30 at 10 months. Synaptophysin labeling is properly superposed to the α-bungarotoxin labeling in a vast majority of junctions in WT mice (black), but only in a minority of NMJ in Tg30 mice, and shows most often a partial overlap (gray), or is absent from the terminal nerve (white). T–V: Labeling of acetylcholine receptors by α-bungarotoxin (red) and immunofluorescence labeling of neurofilament (green) on muscle fiber sections of 10-month-old WT (T) or Tg30 (U and V) mice. T: All of the junctions are innervated in WT mice. U: Some junctions are innervated in Tg30 mice, but others are denervated (V). Arrows point on the junctions (T–V). W: Quantification of muscle innervation at 10 months in WT and Tg30 mice using NF/BTX overlap as criterion. The vast majority of junctions are innervated in WT mice (black), but a significant proportion are denervated (gray) in Tg30 mice. * * P < 0.05 (G), * * * P < 0.01 (S), and ** * * P < 0.001 (S and W) by t-test. n = 3 (J, S, and W). Scale bars: 5 μm (A and K). BTX, α-bungarotoxin; NF, neurofilament; NMJ, neuromuscular junctions; SYN, synaptophysin; WT, wild type.
confirm our previous observation of absence of neuronal loss in 12-month-old Tg30 mice and indicated that the muscular atrophy and the motor deficit in Tg30 mice cannot be simply due to neuronal loss.

The Formation and the Maturation of Neuromuscular Junctions Are Not Affected in Tg30 Mice

We next investigated the formation and the maturation of NMJ in WT and Tg30 mice. The development of NMJ was studied between birth and P14, using BTX as a marker of acetylcholine receptors (AChR) in muscle motor endplates and immunolabelling for synaptophysin to detect synaptic vesicles in motor nerve terminals. The development of NMJ was normal in Tg30 hindlimb muscles. Indeed, at birth, AChR appeared as plaques overlaid with sparse dots of synaptophysin-positive clusters likely corresponding to presynaptic vesicles (data not shown). At P7, motor endplates reorganized and started to display classical perforations whereas morphological matching between motor nerve terminals and motor endplates increased (data not shown). At P14, the most mature junctions displayed a pretzel-like shape, whereas synaptophysin labeling was superposed to the motor endplates, indicative of a precise apposition of the motor nerve terminals to the motor endplates (Figure 3, A–D). To confirm this result, the maturation of NMJ was evaluated at P14 based on four morphological stages of maturation of motor endplates (based on AChR patterns) (Figure 3E) and their innervation status (based on neurofilament immunoreactivity) (Figure 3, F and G). A similar distribution of NMJ maturation stages (Figure 3E) and a similar proportion of monoinnervated motor endplates (Figure 3, F and G) was observed in WT and Tg30 mice (Figure 3, F and G). Taken together, these observations suggested that the formation and initial maturation of NMJ were not altered in hindlimb muscles of Tg30 mice.

Tg30 Mice Develop a Progressive Denervation of Neuromuscular Junctions with Aging and Show an Abnormal Muscle Electrical Activity

To determine whether more tardive consequences on the stabilization of NMJ might be present in Tg30 mice, the morphology of NMJ was studied at 1 and 10 months in WT and Tg30 mice. Labeling of AChR by α-BTX unveiled that NMJ displayed a pretzel-like shape in WT and Tg30 mice at 1 month, indicative of a normal morphology of motor endplates on muscle fibers (Figure 4, A, B, D, and E). However, the localization of synaptophysin was defective in a proportion of NMJ in 1-month-old Tg30 mice (Figure 4, C and F). In WT mice, 95.58% ± 0.49% of the NMJ showed extensive superposition of BTX and of synaptophysin labeling (Figure 4, D and G), indicative of precise apposition between motor nerve terminals and motor endplates. By contrast, in Tg30 mice, 90.36% ± 0.65% of NMJ exhibited precise superposition of BTX and of synaptophysin labeling (Figure 4, E and G), and in 9.63 ± 0.65% cases, synaptophysin was partly superposed to the BTX labeling (Figure 4, F and G). Despite this defect of synaptophysin distribution at 1 month, the motor endplates were correctly innervated.
by motor nerve terminals in Tg30 mice as judged by NF/BTX labeling (Figure 4, H–J).

In 10-month-old WT mice, 96.81% ± 0.41% of the NMJ showed extensive superposition of BTX and of synaptophysin labeling (Figure 4, K, O, and S), indicative of perfect apposition of motor nerve terminals to the motor endplates. By contrast, in Tg30 mice, only 39.64% ± 3.11% of NMJ exhibited perfect superposition of motor nerve terminals to the motor endplates (Figure 4, L, P, and S). In the other NMJ, the synaptophysin labeling was either partly superposed to the BTX labeling (Figure 4, M, Q, and S) or absent (Figure 4, N, R, and S). Furthermore, in WT mice and in a proportion of Tg30 mice, NMJ were correctly innervated (Figure 4, T, U, and W). But in 41.36% ± 2.28% of Tg30 NMJ, motor endplates were denervated without apposition of motor nerve terminals (Figure 4, V and W). Taken together, these data demonstrate a disturbance of NMJ stabilization in Tg30 mice leading to a progressive denervation of motor endplates with aging.

Furthermore, the denervation of motor endplates observed in Tg30 mice was confirmed by electromyographic recordings (Figure 2, J–M). Positive sharp waves and/or

**Figure 6**  Human tau expression leads to accumulation of Gallyas-positive inclusions and cathepsin B—positive vesicular clusters in sciatic nerve. A and B: Gallyas silver staining of longitudinal sections of sciatic nerve of Tg30 mice at 7 (A) and 10 (B) months counterstained with Nuclear Fast Red. Insets show the same staining on longitudinal sections of sciatic nerve of WT mice. Gallyas-positive aggregates are observed in Tg30 mice at 7 months and 10 months. C–G: Immunocytochemical labeling with the BR6 antibody to cathepsin B on longitudinal sections of the sciatic nerve of WT (C and D) or Tg30 (E–G) mice at 7 months (C and E) and 10 months (D, F, and G) counterstained with hematoxylin and eosin. The distribution of cathepsin B—positive vesicular structures is similar at 7 months in WT and Tg30 mice. Local accumulations of clusters of cathepsin B—positive vesicular structures (arrowheads) are observed in the sciatic nerves of Tg30 mice at 10 months. Scale bars: 25 μm (A and B); 30 μm (C–F); 10 μm (G). WT, wild type.
fibrillation were detected in some Tg30 mice at 2 months of age that became more abundant in older animals. At 10 months of age, abundant spontaneous activities could be recorded in Tg30 mice (Figure 2M). Spontaneous activities were not observed in 2-month-old WT mice (Figure 2J). To test whether this abnormal electrical activity reflected an altered neurotransmission at the NMJ, we recorded the muscle evoked response to repetitive nerve stimulation. If neuromuscular transmission was affected, a sharp decrease (>10%) in the amplitude of the electrical response should be observed after repeated stimulations. But muscle compound action potential amplitude decrement ranged from −0.4% to −9.9% in Tg30 mice, indicating that neuromuscular transmission did not seem compromised using this test.

Human Tau Transgenic Protein Is Expressed in Sciatic Nerve and in Motor Nerve Terminals But Not in Muscle in Tg30 Mice

We further analyzed the expression and the localization of the human tau protein in the NMJ, sciatic nerve, and muscle of Tg30 mice. To analyze tau expression in NMJ, we performed a double labeling of AChR by α-BTX combined with an immunofluorescent labeling of human tau transgenic protein on hindlimb tissue sections in Tg30 mice at P7, P14, and 1 and 10 months. At P7, human tau protein was not detected in the NMJ (data not shown). At P14, the human tau protein was expressed in axons and in motor nerve terminals (Figure 5, A and B). A superposition of α-BTX and of human tau protein was observed to some extent in NMJ. The human tau protein was also expressed in axons and in motor nerve terminals at 1 and 10 months in Tg30 mice (Figure 5, C–F). We also immunolabelled sections of muscle tissue and of sciatic nerve with the BR21 anti-human tau antibody. The human tau protein was expressed in axons of sciatic nerves of 3- and 10-month-old Tg30 mice (Figure 5, G and H). Human tau was not detected in muscle (data not shown).

Accumulation of Gallyas-Positive Inclusions in Sciatic Nerve in Tg30 Mice

Gallyas-positive tau aggregates develop in the brain and the spinal cord of Tg30 mice with age. To assess whether aggregates were also present in sciatic nerves in Tg30 mice, we stained sciatic nerves with the Gallyas silver-staining method. A few elongated Gallyas-positive inclusions and clusters of dot-like inclusions were detected along axons in sciatic nerves of 7-month-old Tg30 mice (Figure 6A). Gallyas-positive aggregates were much more numerous in 10-month-old Tg30 mice and of larger size. Gallyas-positive aggregates were also encountered in focal areas of axonal degeneration (Figure 6B). These Gallyas-positive aggregates were not encountered in WT mice (Figure 6, A and B). Thus, Gallyas-positive aggregates develop also in sciatic nerves and increase with age in Tg30 mice.

Accumulation of Cathepsin B—Positive Vesicular Clusters in Sciatic Nerve in Tg30 Mice

 Tau proteins regulate the stability of axonal microtubules and the axonal transport, and abnormal expression of tau could disrupt axonal transports and lead to the observed NMJ denervation in Tg30 mice. Disturbances in the axoplastastic transport leads to accumulation of cathepsin-positive autophagic vacuoles. To document potential disturbances in the axoplasmic transport in axons in Tg30 axons, we compared in WT and Tg30 mice the distribution of cathepsin B immunoreactivity in the sciatic nerve at 3, 7, and 11 months (Figure 6, C–G). At 3 months of age, the cathepsin B immunoreactivity was similar in WT and Tg30 mice (data not shown). Occasional small clusters of cathepsin B—positive vesicular structures were observed along axons at 7 months in sciatic nerve of Tg30 mice, but not in WT mice (Figure 6, C and E). Numerous clusters of such cathepsin B—positive vesicular structures were observed at 10 months in sciatic nerve of Tg30 mice (Figure 6, F and G), but not in WT mice (Figure 6D).

Discussion

We show in this study that the Tg30 tauopathy murine model develops a muscular degeneration due to a progressive NMJ denervation, leading to a severe motor deficit. This motor deficit is preceded by an early defective localization of synaptophysin in NMJ, spontaneous muscle electrical activity,
Muscle fibers degeneration, and accumulation of tau aggregates in sciatic nerve axons. The sequence of appearance of these pathogenic events is schematically represented in Figure 7.

The Motor Phenotype and NMJ Denervation in Tg30 Mice Is Not Due to Motoneuron Loss

We did not observe a loss of spinal motoneurons in Tg30 mice from P0 to 10 months, extending our previous results showing no neuronal loss in 12-month-old Tg30 mice. This absence of neuronal loss is striking in view of the severe motor phenotype developed by Tg30 mice from 8 months and onward, and demonstrates that the muscular atrophy and NMJ denervation in Tg30 mice is not secondary to death of spinal motor neurons. A denervation of the neuromuscular junctions despite an absence of death of spinal motor neurons was also observed in a TDP-43 knockout mice model developing a motor impairment around the age of 60 weeks, a degeneration of large motor axons, grouped atrophy of the skeletal muscle. These observations indicate that a severe motoneuron phenotype can develop in the absence of significant motoneuron death.

Early Expression of Human Tau Transgenic Protein up to the Nerve Terminals of Motoneurons Is Not Associated with Denervation

We observed that the human tau protein was expressed in cell body and dendrites of motoneurons in the spinal cord of Tg30 mice from P7. This is consistent with the observation that the Thy1 promoter starts to drive expression in neurons at P6. From P14, the human tau protein was also expressed in motor axons and in motor nerve terminals. We previously reported that the human tau protein was detected at P18 in brain and spinal cord of Tg30 mice. Despite this early expression of human mutant tau up to the nerve terminals of motoneurons, the initial maturation of NMJ up to P14 was not affected in Tg30 mice. The first abnormality that we detected was a defect of synaptophysin distribution in a moderate proportion of NMJ at 1 month that was not associated with morphological signs of denervation and without evidence of motor deficits. These observations suggest that the toxic effect of human tau is a relatively slow process that manifests itself only in mature NMJ and leads to a progressive denervation only in adulthood.

Tg30 Mice Develop a Progressive Muscle Denervation

We observed a two-phase evolution of muscle fibers changes in Tg30 mice. Although the size of muscle fibers was similar at 1 month in WT and Tg30 mice, a higher proportion of hypertrophic muscle fibers (and some signs of muscle fibers degeneration) was seen at 3 months in Tg30 mice, followed by a higher proportion of hypertrophic muscle fibers (with robust signs of muscle fiber degeneration) at 10 months. This two-phase evolution most probably corresponds to a slow process of muscular denervation: the onset of muscular degeneration concerns a selected population of muscle fibers but initially this is associated to a compensation mechanism with nearby/neighbor muscle fibers increasing their size. The ongoing denervation then concerns an increasing population of muscle fibers showing hypotrophy. The presence of both hypotrophic and hypertrophic muscle fibers has been observed in other models of progressive neurodegenerative diseases. For example, in aged SOD1G93A mice, the gastrocnemius muscles show severe changes, including a small group of atrophic muscle fibers, centrally placed nuclei, and hypertrophic fibers.

This early compensation mechanism might explain why Tg30 mice do not develop a clinical motor deficit before 8 months of age, i.e., starting when the ongoing muscle degeneration leads to the observed hindlimb muscle weakness at this age. This compensation mechanism might also explain the absence of difference in the amplitude of the muscle compound action potential at an early age in Tg30 mice, because collateral re-innervation may mask the real progress of neuromuscular junction loss. Muscle spontaneous activities were, however, recorded in the majority of mice, already at early ages, indicative of the presence of denervated muscle fibers, as supported also by histological findings in the NMJ. Three-Hertz repetitive stimulations may detect postsynaptic impairment at the neuromuscular junction level, whereas 30-Hz repetitive stimulations, not attempted here, might have disclosed a presynaptic problem similar to the one reported in this study.

Disturbance of Axonal Transports as a Mechanism Leading to NMJ Denervation in Tg30 Mice

We previously observed in Tg30 mice the occurrence of axonal spheroids and axonal accumulation of cytoplasmic organelles and neurofilaments in the spinal cord. Accumulations of degraded organelles were also observed in axons in the sciatic nerve. These ultrastructural abnormalities were indicative of impairment in axonal transports. We further document these disturbances of axoplasmic transport by showing that clusters of cathepsin B-positive vesicles accumulated in sciatic nerve axons in Tg30 mice. Cathepsin B is a marker of lysosomes/autophagosomes, and disturbances in axoplasmic transport lead to accumulation of cathepsin-positive autophagic vacuoles in experimental models. Accumulations of autophagic vacuoles in distended, dystrophic neurites is frequently encountered in Alzheimer disease and other neurodegenerative diseases. Interestingly, Gallyas-positive aggregates accumulated also in sciatic nerve axons in Tg30 mice, and might also play a role in impairments in axonal transports. We suggest that the progressive reduction of synaptic vesicles observed in the motor nerve terminals in Tg30 mice also results from impairments in their axonal transports, and participates in an alteration of synaptic transmission and the functional denervation of NMJ. In the Thy22 mutant tau transgenic model expressing the same mutant tau
protein as the present Tg30 tau model, defects in retrograde transport in the septohippocampal pathway were also previously demonstrated.24 In drosophila overexpressing a human tau protein, a disruption of synaptic transmission at the neuromuscular junction was observed as a consequence of disturbance of axonal transports and reduction of the number of mitochondria in axons.28

Disturbances of anterograde and retrograde transports have been implicated as a pathogenic mechanism in several neurodegenerative diseases.29–33 In models of amyotrophic lateral sclerosis, expression of mutant SOD1 leads to impaired fast axonal transport at an early asymptomatic stage preceding loss of spinal motor neurons.22 Mutations in dynnein (a motor protein complex that moves along microtubules in the minus-end direction) result in progressive motor neuron degeneration34 and the formation of Lewy-like inclusions leading to defects in retrograde transport.35 In Alzheimer disease, a reduction of axonal transport is an early pathogenic mechanism,36–38 affecting particularly retrograde transport.33 Tau proteins play a role in modulation of axonal transport, and tau pathology plays a crucial role in disturbances of these axonal transports in Alzheimer disease.39–43 The role of pathological tau at the dendritic level has been pointed out in several studies on tauopathy models.44,45 Much evidence points, however, to dying-back axonal degeneration due to axoplasmic transport defects induced by pathological tau as a physiopathological mechanism in Alzheimer disease.46 Consistently, the alterations at the level of the NMJ in Tg30tau mice illustrate a toxic effect of abnormal tau at the presynaptic level. A recent study also pointed to disturbances at the presynaptic level in mossy fibers in the hippocampus in a tauopathy model.47

To conclude, we show in this study that expression of a human mutant tau protein in the motoneurons and their processes leads to dying-back axonopathy and an early alteration of synaptophysin distribution in motor nerve terminals from 1 month in Tg30 mice. An axonopathy develops during the following months, characterized by accumulation of aggregates and organelles. These alterations progressively result in a defect in synaptic transmission leading to denervation of NMJ, muscular atrophy, and motor deficit.

We suggest that abnormalities developing at the NMJ in Tg30 mice will be a useful model to unravel mechanisms of synaptic dysfunction resulting from disturbances of axoplasmic transports in neurodegenerative diseases.

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References