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Genetic Interaction between MTMR2 and FIG4 Phospholipid Phosphatases Involved in Charcot-Marie-Tooth Neuropathies

Ilaria Vaccari1,2, Giorgia Dina3, Hélène Tronchère4, Emily Kaufman5, Gaëtan Chicanne4, Federica Cerri3, Lawrence Wrabetz5, Bernard Payrastre4, Angelo Quattrini3, Lois S. Weisman6, Miriam H. Meisler7, Alessandra Bolino1,2,*

1 Human Inherited Neuropathies Unit, INSPE-Institute for Experimental Neurology, Division of Neuroscience, San Raffaele Scientific Institute, Milan, Italy, 2 Dulbecco Telethon Institute, San Raffaele Scientific Institute, Milan, Italy, 3 Neuropathology Unit, INSPE-Institute for Experimental Neurology, Division of Neuroscience, San Raffaele Scientific Institute, Milan, Italy, 4 INSERM U1048 and Université Toulouse 3, I2MC, CHU Toulouse, Toulouse, France, 5 Biology of Myelin Unit, Division of Genetics and Cell Biology, San Raffaele Scientific Institute, Milan, Italy, 6 Life Science Institute, University of Michigan, Ann Arbor, Michigan, United States of America, 7 Department of Human Genetics, University of Michigan, Ann Arbor, Michigan, United States of America

Abstract

We previously reported that autosomal recessive demyelinating Charcot-Marie-Tooth (CMT) type 4B1 neuropathy with myelin outfoldings is caused by loss of MTMR2 (Myotubularin-related 2) in humans, and we created a faithful mouse model of the disease. MTMR2 dephosphorylates both PtdIns3P and PtdIns(3,5)P2, thereby regulating membrane trafficking. However, the function of MTMR2 and the role of the MTMR2 phospholipid phosphatase activity in vivo in the nerve still remain to be assessed. Mutations in FIG4 are associated with CMT4J neuropathy characterized by both axonal and myelin damage in peripheral nerve. Loss of Fig4 function in the plt (pale tremor) mouse produces spongiform degeneration of the brain and peripheral neuropathy. Since Fig4 has a role in generation of PtdIns(3,5)P2 and MTMR2 catalyzes its dephosphorylation, these two phosphatases might be expected to have opposite effects in the control of PtdIns(3,5)P2 homeostasis and their mutations might have compensatory effects in vivo. To explore the role of the MTMR2 phospholipid phosphatase activity in vivo, we generated and characterized the Mtmr2/Fig4 double null mutant mice. Here we provide strong evidence that Mtmr2 and Fig4 functionally interact in both Schwann cells and neurons, and we reveal for the first time a role of Mtmr2 in neurons in vivo. Our results also suggest that imbalance of PtdIns(3,5)P2 is at the basis of altered longitudinal myelin growth and of myelin outfolding formation. Reduction of Fig4 by null heterozygosity and downregulation of PIKfyve both rescue Mtmr2-null myelin outfoldings in vivo and in vitro.


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* E-mail: bolino.alessandra@hsr.it

Introduction

Phosphoinositides (PIs) constitute potent signaling molecules with a specific and restricted distribution at intracellular membranes that is strictly controlled by the concerted action of kinases and phosphatases [1,2]. PI-phosphatases are key regulators of membrane trafficking as they contribute to assembly of molecular machineries that promote and control membrane dynamics and vesicle movement, tethering and fusion. In the nervous system, both neurons and glia rely on efficient membrane trafficking for many functions, such as axonal transport or myelination.

Charcot-Marie-Tooth (CMT) neuropathies are very heterogeneous disorders from both the clinical and genetic point of view [3–6]. Several CMT genes encode proteins that regulate or are connected with PI metabolism, including FRABIN/FGD4, FIG4, DNM2, RAB7, SIMPLE, LRSAM1, SH3TC2, MTMR2, and MTMR13, supporting the idea that regulation of intracellular trafficking is a key process in peripheral nervous system biology [7] (http://www.molgen.ua.ac.be/CMTMutations/default.cfm).

We first demonstrated that loss of function mutations in the MTMR2 (Myotubularin-related 2) gene cause autosomal recessive demyelinating Charcot-Marie-Tooth type 4B1 (CMT4B1, OMIM #601362) neuropathy with myelin outfoldings [8]. MTMR2 is a phospholipid phosphatase that dephosphorylates both PtdIns3P and PtdIns(3,5)P2 phosphoinositides at the D3 position of the inositol ring, thus generating PtdIns5P [9–16]. We have generated a Mtmr2-null mouse which models the CMT4B1 neuropathy and we reported that loss of Mtmr2 specifically in Schwann cells is both sufficient and necessary to provoke myelin outfoldings [17,18]. We recently identified a potential mechanism using vivo and in vitro models of CMT4B1 and proposed that Mtmr2 belongs to a molecular machinery that titrates membrane trafficking during myelination. According to this model, myelin...
outfoldings arise as a consequence of the loss of negative control on the amount of membrane produced during myelination [19]. Despite these findings, the function of MTMR2 and the role of the MTMR2 phospholipid phosphatase activity in the nerve still remain to be assessed.

Loss of the FIG4/SAC3 phospholipid phosphatase in human provokes another form of autosomal recessive demyelinating CMT, the CMT type 4J (OMIM #611228) neuropathy [20,21]. FIG4 is a 5-phosphatase involved in the dephosphorylation of PtdIns(3,5)P2, a predicted substrate of MTMR2. Loss of Fig4 in the mouse causes the plt (pale tremor) phenotype, characterized by extensive neuronal vacuolization and degeneration and by a peripheral neuropathy [20,22]. Yeast Fig4 is localized at the vacuolar membrane-the yeast lysosomal compartment- and is required for both the generation and turnover of PtdIns(3,5)P2 [23,24]. In addition to the 5-phosphatase activity, yeast Fig4 appears to activate Fab1, the kinase that produces PtdIns(3,5)P2 from PtdIns3P [23,24]. Deletion of yeast Fig4 reduces rather than increases PtdIns(3,5)P2 leading to defects in vacuole homeostasis and function. A significant decrease of PtdIns(3,5)P2 was found also in plt (Fig4-null) fibroblasts, suggesting conserved enzymatic and cellular functions of Fig4 from yeast to mouse [20]. Moreover, the most common human mutation of FIG4 acts by reducing its affinity for the PtdIns(3,5)P2 biosynthetic complex [25].

Since Fig4 has a role in generation of PtdIns(3,5)P2 and MTMR2 catalyzes its dephosphorylation, these two phosphatases might have opposite effects in the control of PtdIns(3,5)P2 homeostasis and their mutations might have compensatory effects in vivo. To explore the role of the MTMR2 phospholipid phosphatase activity in vivo, we took advantage of the Fig4 and Mtmr2-null mice and generated and characterized the Mtmr2/−Fig4/− double null mutant. Here we provide strong evidence that Mtmr2 and Fig4 interact in both Schwann cells and neurons, and reveal for the first time a role of Mtmr2 in neurons in vivo. We also report that the imbalance of PtdIns(3,5)P2 might be at the basis of myelin outfullding in the nerve. Reduction of Fig4 by null heterozygosity and downregulation of PIKfyve both rescue Mtmr2-null myelin outfoldings in vivo and in vitro.

### Results

#### Generation of Mtmr2/−Fig4/−null mice

The generation and characterization of Mtmr2-null and Fig4-null mice have been reported [17,20]. Mtmr2/−Fig4/− double null mice and controls were analyzed in the F2 generation. At postnatal day three (P3) Mtmr2/−/−Fig4/−/− mice had reduced body size and dilute pigmentation of the coat similar to the Mtmr2/−/−Fig4/−/− mice in the same litter, and as reported for the plt mouse [20]. Tremor and abnormal gait developed in the second week after birth. Mtmr2/−/−Fig4/−/− mice show juvenile lethality and die around 1 month of age. The viability of Mtmr2/−/−Fig4/−/− mice was lower than for Mtmr2/−/+Fig4/−/− littermates. A reduced number of both Mtmr2/−/−Fig4/−/− and Mtmr2/−/−Fig4/−/− mice were present at P8, compared to the expected Mendelian ratio (Table 1 and Table 2). The longest survival of the double mutant was 20 days. The data indicate that loss of Mtmr2 reduces viability of Mtmr2/−/+Fig4/−/−. We therefore hypothesized that loss of Mtmr2 might provoke a worsening of the Mtmr2/−/+Fig4/−/− neurodegeneration.

Mtmr2 loss exacerbates Fig4/−null neurodegeneration

To explore this possibility, we performed semithin section analysis of DRG ganglia, brain and spinal cord from Mtmr2/−/−Fig4/−/− and Mtmr2/−/−Fig4/−/− mice. DRG ganglia from both Mtmr2/−/−Fig4/−/− and Mtmr2/−/−Fig4/−/− mice at P3 were severely affected, exhibiting neuronal loss and massive vacuolization (Figure 1A–1C). In the cerebellum of both Mtmr2/−/−Fig4/−/− and Mtmr2/−/−Fig4/−/− mice at P8 and at P20 we observed a thickening of the molecular layer as compared to wild-type, and cells with cytoplasmic vacuoles were present in the granular layer. At P20, a consistent loss of Purkinje and basket cells was observed in both genotypes (Figures S1 and S2). These cerebellar findings have not been previously reported in the plt mouse.

In the cortex and brainstem of Mtmr2/−/−Fig4/−/− mice at P3 we noted more cells with vacuoles and inclusions than in Mtmr2/−/−Fig4/−/− mice, which were never observed in wild-type animals (Figure 1D–1F). In particular, in the brainstem of Mtmr2/−/−Fig4−/− mice at P8 the number of neurons carrying pathological abnormalities was significantly increased as compared to Mtmr2/−/−Fig4−/− mice (Figure 1G–1I). We also analyzed the spinal cord of Mtmr2/−/−Fig4−/− and Mtmr2/−/−Fig4−/− mice at P3 and P8 (Figure 2A–2F). Vacuolated cells and cells with inclusions were observed, as previously described for the plt phenotype, which were not present in wild-type spinal cords [20,26]. At P8, we observed a significant decrease in the number of mice with Mtmr2/−/−Fig4−/−, Mtmr2/−/−Fig4−/− and Mtmr2/−/−Fig4−/− genotypes scored at P8 on a total of 608 mice.

#### Table 1. Number of mice with Mtmr2+/−Fig4−/−, Mtmr2+/−Fig4−/−, and Mtmr2/−/−Fig4−/− genotypes scored at P8 on a total of 608 mice

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Observed</th>
<th>Predicted</th>
<th>P values</th>
</tr>
</thead>
<tbody>
<tr>
<td>+/+</td>
<td>--/−</td>
<td>38</td>
<td>34</td>
</tr>
<tr>
<td>+/−</td>
<td>--/−</td>
<td>76</td>
<td>56</td>
</tr>
<tr>
<td>−/+</td>
<td>--/−</td>
<td>38</td>
<td>10</td>
</tr>
</tbody>
</table>

*“Predicted” refers to the number of expected mice among 608 mice born, based on Mendelian ratios. “Observed” refers to the mice identified by genotyping at P8. The number of mice decreases with loss of Mtmr2 on a Fig4-null background. P values were calculated for a binomial distribution, using the chi-square test.*

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of motor neurons and cells in Mtmr2−/−Fig4−/− mice as compared to Mtmr2+/+Fig4−/− mice (Figure 2H, 2I). These findings demonstrate that loss of Mtmr2 exacerbates the Mtmr2+/+Fig4−/− neurodegeneration.

This effect could be the consequence of loss of Mtmr2 in neurons or in other cells, such as in astrocytes. For example, in the plt mouse, a block of autophagy in astrocytes has been reported. In plt mice at 1 week of age, the p62 autophagy marker was increased in GFAP-positive astrocytes from brain regions severely affected at later stages, suggesting that autophagy impairment contributes to the pathogenesis [26]. Elevated p62 co-localized with LAMP2-positive late endosomes/lysosomes (LE/LY) in astrocytes, showing

Table 2. The relative risk (RR) of lethality for the Mtmr2+/+Fig4−/−, Mtmr2+/−Fig4−/−, and Mtmr2−/−Fig4−/− genotypes.

<table>
<thead>
<tr>
<th>Genotype 1</th>
<th>Genotype 2</th>
<th>RR (95%CI)</th>
<th>Chi-square</th>
<th>P values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mtmr2+/+Fig4−/−</td>
<td>Mtmr2+/−Fig4−/−</td>
<td>2.5 (0.92, 6.80)</td>
<td>2.91</td>
<td>0.09</td>
</tr>
<tr>
<td>Mtmr2+/+Fig4−/−</td>
<td>Mtmr2−/−Fig4−/−</td>
<td>7 (2.72, 18.03)</td>
<td>28.55</td>
<td>9×10−4</td>
</tr>
<tr>
<td>Mtmr2+/−Fig4−/−</td>
<td>Mtmr2−/−Fig4−/−</td>
<td>2.8 (1.84, 4.27)</td>
<td>21.42</td>
<td>4×10−5</td>
</tr>
</tbody>
</table>

The relative risk analysis was performed on the 608 animals scored at P8, as also reported in Table 1.

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Figure 1. Neurodegeneration in Mtmr2+/+Fig4−/− and Mtmr2−/−Fig4−/− mice. Semithin section analysis of DRG ganglia, cortex and brainstem of wild-type (A, D, G) and mutant mice (B, C, E, F, H, I). (B, C) DRG sensory neurons from Fig4-null and Mtmr2/Fig4 double null mice at P3 with massive vacuolization. (F, I) More cells carrying inclusions and vacuoles were observed in Mtmr2−/−Fig4−/− cortex (F) and brainstem (I) at P3 as compared to Mtmr2+/+Fig4−/− mice (E, H). (K, L) Brainstem analysis at P8 showed increased number of pathological neurons in Mtmr2−/−Fig4−/− (L) as compared to Mtmr2+/+Fig4−/− mice (K) and quantification in (L), P = 2.75×10−14. A total of 10030, 10290, and 9620 neurons were counted and analyzed in wild-type Mtmr2+/+Fig4−/−, and Mtmr2−/−Fig4−/− mice, respectively. All neurons scored were normal in wild-type, whereas 48.3% and 81.1% of neurons displayed pathological features in Mtmr2+/+Fig4−/− and Mtmr2−/−Fig4−/−, respectively. Arrows indicate inclusions; arrowheads vacuoles. Three mice per genotype were analyzed. Bar is 35 μm in (A–C), 10 μm in (D–L).

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that the block of autophagy occurred subsequent to the fusion of autophagosomes with LE/LY [26]. To determine whether loss of Mtmr2 in astrocytes might further impair autophagy, we evaluated p62 levels in total brain extracts from Mtmr2+/-Fig4-/- as compared to Mtmr2-/-Fig4-/- mice. Increased p62, LAMP1 and GFAP expression levels were confirmed in Mtmr2-/- as compared to wild-type but no differences were detected between Mtmr2+/+ and Mtmr2-/- double null mice (Figure 3A). This finding indicates that loss of Mtmr2 does not further impair the block in the autophagic process in astrocytes of Fig4-null mice.

To further investigate the cell autonomy of the Mtmr2/FIG4 interaction, we established dissociated Schwann cell/DRG neuron co-cultures from Mtmr2+/+ and Mtmr2-/- mice, in which mutant Schwann cells were replaced with exogenous wild-type rat Schwann cells. Mtmr2-/- DRG neurons cultured with wild-type Schwann cells were significantly more severely vacuolated (57.8%) as compared to Mtmr2+/+ cultures (37.4%) (Figure 3B-3D’ and 3H). This finding provides strong evidence that the loss of Mtmr2 in neurons leads to the worsening of the Fig4-null neurodegeneration.

Like neurons, mouse primary fibroblasts (MFs) from plt mutants display enlargement and vacuolization of the LAMP2-positive LE/LY compartment [20,22]. To provide further evidence for functional interaction between MTMR2 and FIG4, we established MF cultures from Mtmr2+/+ and Mtmr2-/- mice. By LAMP1 staining and confocal microscopy, we observed that the number of fibroblasts carrying enlarged LE/LY was significantly increased in Mtmr2-/- as compared to wild-type but no differences were detected between Mtmr2+/+ and Mtmr2-/- double null mice (Figure 4). This finding indicates that Mtmr2 loss exacerbates Fig4-null vacuolar phenotype by further impairment of the endo/lysosomal trafficking pathway.

**Mtmr2 exacerbates Fig4-null hypomyelination in sciatic nerve**

The plt mouse phenotype is characterized by a peripheral neuropathy. Loss of large diameter myelinated axons, hypomyelination (reduced myelin thickness), reduced amplitude of compound motor action potential (cMAP) and slowing of the nerve conduction velocity (NCV) have been reported in plt mice and the presence of demyelinating features in CMT4J patient biopsies such as onion bulb suggested that FIG4 has also a cell autonomous role in Schwann cells [22].

We investigated sciatic nerves from Mtmr2+/+ and Mtmr2-/- mice. At P3 and P8, mutant sciatic nerves showed a normal development. In both genotypes at P8, Schwann cells often contained cytoplasmic inclusions and occasionally
Figure 3. Mtmr2 loss in neurons exacerbates Fig4-null neurodegeneration. (A) Western blot analysis of brain extracts from wild-type, Mtmr2+/-Fig4-/-, and Mtmr2-/-Fig4-/- mice at P20. P62, GFAP, and LAMP1 are increased in Mtmr2+/-Fig4-/- as compared to wild-type as already reported for the plt mouse. No differences were detected between Mtmr2+/-Fig4-/- and Mtmr2-/-Fig4-/-, suggesting that loss of...
Mtmr2 does not further impair astrocytosis and the block in the autophagic process in astrocytes. (B–G) Dissociated DRG explants from wild-type (B, B', E, Mtmr2+/+Fig4+/−/− (C, C', F), and Mtmr2−/−Fig4−/−/− (D, D', G) mice, in which mutant Schwann cells were replaced by exogenous wild-type rat Schwann cells. (D, D') More Mtmr2−/−Fig4−/−/− neurons (n = 303 neurons on 523 total neurons) are vacuolated as compared to Mtmr2+/+Fig4−/−/− neurons (C, C'; n = 236 on 635 total neurons; P = 7.56271 × 10−8, 16 covers per genotype, and quantification in H). (G) Mtmr2−/−Fig4−/−/− co-cultures produced less myelinated MBP-positive segments after 7 days of ascorbic acid treatment (n = 52 segments on 16 coverslips) as compared to Mtmr2+/+Fig4−/−/− (F) (n = 234 segments on 12 coverslips; P = 1.9218 × 10−5) and quantification in (I). Both Mtmr2+/+Fig4+/−/− and Mtmr2−/−Fig4−/−/− explants are severely hypomyelinated as compared to wild-type (E) (n = 6560 segments on 16 coverslips). Green is neurofilament (NF-L); red is myelin basic protein (MBP), staining compact myelin. Arrowheads in (C, D, C', D') indicate vacuoles within sensory neurons. Bar in (B–G) is 50 μm. doi:10.1371/journal.pgen.1002319.g003

contained vacuoles, which were never observed in wild-type nerves. At P20, the latest time point of survival of Mtmr2+/+Fig4−/− double null mice, Mtmr2+/+Fig4−/−/− sciatric nerves were hypomyelinated (thin myelin sheath) with an increased g-ratio (diameter of the axon divided by the diameter of the fiber) as compared to wild-type nerves (Figure 5E). At this stage, sciatric nerves from Mtmr2−/−Fig4−/−/− double null mice were more severely hypomyelinated than Mtmr2+/+Fig4−/−/− mice with a higher g-ratio, demonstrating that Mtmr2 loss exacerbates the hypomyelination (thin myelin sheath) with an increased g-ratio, as compared to wild-type nerves (Figure 5E). The total number of fibers and the axonal diameter distribution at P20 were not significantly altered in mouse nerves of either genotype (Figure 5F). These observations indicate that the hypomyelination is not a developmental defect related to delayed axonal growth. Hypomyelination may result from a defective axonal/Schwann cell interaction due to the severe neuronal degeneration and/or from the loss of FIG4 in Schwann cells. We thus cultured dissociated DRG neurons from Mtmr2−/−Fig4−/−/− and Mtmr2+/+Fig4−/−/− mice, seeded with exogenous wild type rat Schwann cells. Following induction of myelination by ascorbic acid treatment, vacuolated DRG neurons from both Mtmr2−/−Fig4−/−/− and Mtmr2+/+Fig4−/−/− mouse embryos were able to produce myelinated segments, although significantly fewer than wild-type cultures. Moreover, DRG neurons from Mtmr2−/−Fig4−/−/− mice cultured with wild-type Schwann cell produced significantly fewer myelinated segments than Mtmr2+/+Fig4−/−/− neurons seeded with wild-type Schwann cells (Figure 5E, H and quantification in panel I). This observation suggests that the hypomyelination of Mtmr2−/−Fig4−/−/− represents at least in part the consequence of impaired Schwann cell-axonal interaction.

Figure 4. Enlarged late endosomal compartment in primary fibroblasts from Mtmr2+/+Fig4+/−/− and Mtmr2−/−Fig4−/−/− mutant mice. (A–C) The number of fibroblasts carrying enlarged late endosomal lysosomal compartment (LAMP1 positive, green) is significantly increased in Mtmr2−/−Fig4−/−/− (B) as compared to (A) Mtmr2+/+Fig4+/−/−. In red is phallolidin which stains actin filaments. Blue, DAPI. The quantification in (D) was made by counting cells established from three animals per genotype. n = 526 on 17 coverslips for Mtmr2+/+Fig4+/−/−; n = 583 cells on 20 coverslips for Mtmr2−/−Fig4−/−/−; P = 0.000079; confocal microscopy analysis. Asterisks mark cells carrying enlarged late endosomal compartments. Bar in (A–C) is 25 μm. doi:10.1371/journal.pgen.1002319.g004

Fig4 heterozygosity rescues Mtmr2-null myelin outfoldings

To further investigate Mtmr2 and Fig4 interaction in the nerve, we evaluated whether loss of Fig4 modifies the myelin outfolding phenotype. Myelin outfoldings in Mtmr2-null mice arise around the third to fourth week after birth, and the number of fibers containing myelin outfoldings and loops progressively increases with age (up to 6 months or even later). Since Mtmr2−/−/−Fig4+/−/− double mutants die before 1 month of age, we compared sciatric and peroneal nerves at 6 months of age from Mtmr2−/−/−Fig4+/−/− and Mtmr2−/−/−Fig4+/−/− (Fig4 heterozygous) mice. Using semithin section analysis, we measured the number of fibers carrying myelin outfoldings in mutant sciatric and peroneal nerves normalized to the total number of fibers. In Mtmr2−/−/−Fig4+/−/− nerves myelin outfoldings were significantly reduced as compared to Mtmr2−/−/−Fig4+/−/− mice (Figure 6A–6D). Since loss of Mtmr2 in Schwann cells is both sufficient and necessary to provoke myelin outfoldings, loss of Fig4 in Schwann cells (rather than axons) is likely to account for the rescue of the disease phenotype.

To further evaluate this finding, we established myelin-forming Schwann cell/DRG neuron co-cultures from Mtmr2−/−/−Fig4+/−/+ and Mtmr2−/−/−Fig4+/−/+ mouse embryos at E13.5 (Figure 7A, 7B). By measuring the number of MBP positive fibers carrying myelin outfoldings in the cultures, we confirmed that Mtmr2-null myelin outfoldings were rescued by Fig4 heterozygosity (Figure 7C).

Loss of Fig4 in plt fibroblasts leads to a significant decrease in PtdIns(3,5)P2, whereas Mtmr2 loss should lead to an increase in both PtdIns3P and PtdIns(3,5)P2 in vivo in the nerve [20]. Indeed, by performing a sensitive in vitro assay on Mtmr2-null Schwann cell/DRG neuron co-cultures, we found that in null cells PtdIns3P is significantly reduced (up to 70%) as expected by the loss of MTMR2 3-phosphatase activity on PtdIns(3,5)P2 (Figure 8A). We hypothesized that the observed rescue by Fig4 heterozygosity might be the consequence of a restored level of PtdIns(3,5)P2 in Schwann cells. Heterozygosity of Fig4 might decrease PIKfyve activity and therefore partially restore PtdIns(3,5)P2 levels in Mtmr2-null cells.

To test this hypothesis, we downregulated either the activity or expression of PIKfyve in Mtmr2-null co-cultures to rescue Fig4.
outfoldings. We transduced Mtmr2-null co-cultures with lentiviral vectors (LV) carrying PIKfyve shRNA and scored the number of myelinated MBP-positive fibers with myelin outfoldings. Titration of the PIKfyve shRNA LV was previously performed to determine the highest amount of virus which does not significantly affect myelination (Figure S3A, S3B). We found that myelin outfoldings were significantly rescued by downregulating PIKfyve expression (Figure 7D–7F).

On the other hand, we found that loss of Fig4 in Mtmr2-null fibroblasts leads to a significant increase in PtdIns(3,5)P2 levels due to loss of 50% of phosphatase activity (Figure 8C). Moreover, PtdIns(3,5)P2 was equally reduced in Fig4−/− and in Mtmr2−/− Fig4−/− cells (Figure 8B), possibly because the PtdIns(3,5)P2 substrate is already severely affected by loss of Fig4, and Mtmr2 acts downstream of Fig4 in the control of this lipid level.

To support the hypothesis that myelin outfoldings in Mtmr2−/− Fig4−/− co-cultures were rescued by restored PtdIns(3,5)P2 levels, we also measured PtdIns(3,5)P2 in Mtmr2−/− Fig4+/+ and Mtmr2−/− Fig4+/− fibroblasts. However, PtdIns(3,5)P2 did not differ in Mtmr2−/− Fig4+/+ and Mtmr2−/− Fig4+/− fibroblasts (Figure 8C). Small changes in PtdIns(3,5)P2 levels due to loss of 50% of phosphatase expression may be below the level of detection of this method.

Overall, these findings indicate that Mtmr2 and Fig4 control PtdIns(3,5)P2 with opposite effects. If Fig4 is totally absent and PtdIns(3,5)P2 is low, the absence of Mtmr2 which dephosphorylates PtdIns(3,5)P2 has no influence. On the other hand, when PtdIns(3,5)P2 is high due to loss of Mtmr2, a partial reduction in PIKfyve activity due to heterozygosity of Fig4 might lead to PtdIns(3,5)P2 rebalance and rescue of myelin outfoldings.

Finally, we tested for interaction between phosphatases using a pull-down assay. GST-MTM2R was not able to pull-down Fig4 enzyme such as myotubularin does not necessarily result in an imbalance of PtdIns3P, as already reported [9,30,31].

Figure 6. Fig4 heterozygosity rescues Mtmr2-null myelin outfoldings. (A, B) Asterisks indicate myelinated fibers from sciatic nerve carrying myelin outfoldings. (C) Quantification of fibers carrying myelin outfoldings in sciatic nerves from Mtmr2−/− Fig4−/− and Mtmr2−/− Fig4+/− mice and (D) in peroneal nerves, using semithin section analysis. Four animals per genotype were analyzed. The entire nerve section was reconstructed and the number of fibers carrying myelin outfoldings divided for the total number of fibers was counted. P = 0.0097 in (C) and P = 0.0086 in (D). Bar is 10 μm in (A, B).

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from brain or isolated rat Schwann cell lysates, suggesting that the functional interaction between MTMR2 and FIG4 demonstrated here is not mediated by physical interaction between the two proteins (Figure S3C–S3E).

Mammalian MTMR2 converts PtdIns(3,5)P$_2$ and PtdIns3P in yeast

The mutant yeast strain fig4A displays enlarged vacuoles caused by reduced PtdIns(3,5)P$_2$, which in yeast controls the homeostasis of the vacuole (the lysosomal compartment). To further test MTMR2 function, and further test functional interactions between Mtmr2 and Fig4, we transformed FLAG-MTMR2 in the mutant yeast strain fig4A. Overexpression of wild-type MTMR2 in fig4A caused a further enlargement of the vacuolar compartment and defects in vacuole fission whereas the catalytically inactive mutant FLAG-MTMR2C417S did not cause these changes (Figure 9). To determine the substrates and products of mammalian MTMR2 in yeast, we measured phosphorylated phosphoinositide lipid levels from cells expressing FLAG-MTMR2 as compared to the vector alone. To enhance the sensitivity of the assay, we subjected the

Figure 7. A rebalance of PtdIns(3,5)P$_2$ rescues Mtmr2-null myelin outfoldings in vitro. (A, B) Heterozygosity of Fig4 in Mtmr2-null explants rescues myelin outfoldings. (C) The number of MBP positive segments carrying myelin outfoldings on the total of MBP segments on random fields was assessed. Mtmr2$^{+/−}$/Fig4$^{++/+}$ n = 383 fibers; Mtmr2$^{+/−}$/Fig4$^{++/−}$ n = 405 fibers, on at least 5 covers per genotype; P = 1.9754 × 10$^{-2}$. (D) Decrease of PIKfyve expression on lysates from myelin-forming Schwann cells/DRG explants transduced with PIKfyve shRNA lentiviral vector (LV), as compared to a scramble sequence. (E–F$''$ and I) Myelin outfoldings are rescued in Mtmr2-null transduced by PIKfyve shRNA LV as compared to cultures transduced with a scramble sequence (PIKfyve, n = 428 and scramble, n = 484; P = 6 × 10$^{-6}$). (G, H, and J) Myelin outfoldings are rescued in Mtmr2-null treated with 70 nM of YM201636 PIKfyve inhibitor as compared to cultures treated with DMSO (YM201636 n = 913 and scramble, n = 857; P = 6.5 × 10$^{-6}$). Bar is 20 μm in (A, B, E–F$''$, and G, H). Green is neurofilament (NF-L); red is myelin basic protein (MBP), staining compact myelin. Arrowheads indicate myelin outfoldings.

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yarn to hyperosmotic shock. In wild-type yeast, this results in a transient increase in PtdIns(3,5)/P2 levels (green line) and concomitant decrease in PtdIns(3,5)/P2 (blue line) (Figure 10, 5 min time point). If MTMR2 acts on PtdIns(3,5)/P2, then there should be a decrease in PtdIns(3,5)/P2 and a corresponding increase in PtdIns3/P2. Moreover, if MTMR2 acts on PtdIns3/P2 there will be a decrease in that lipid as well. Each of these changes was observed (Figure 10, solid lines and Table S1). These observations demonstrate that MTMR2 and FIG4 coordinately regulate the PtdIns3/P2 pathway in vivo.

**Discussion**

The MTMR2 3-phosphatase activity toward PtdIns3/P and PtdIns(3,5)/P2 has been demonstrated by a number of studies using recombinant MTMR2 in vitro as well as conventional cell lines overexpressing MTMR2 [10–16]. Overexpressed MTMR2 has been co-localized with Rab7 in A431 cells at the level of late endosome/lysosomes, where PtdIns(3,5)/P2 is generated [16]. Interestingly, another phospholipid phosphatase, FIG4/SAC3, is involved in both the dephosphorylation and the production of PtdIns(3,5)/P2 in yeast and is mutated in autosomal recessive demyelinating CMT4J neuropathy [20]. Loss of Fig4 in mouse provokes the plt phenotype characterized by massive neurodegeneration and peripheral neuropathy. In Fig4-null fibroblasts a decrease in PtdIns(3,5)/P2 levels were demonstrated between the wild-type and Fig4−/− cells as compared to wild-type, thus also confirming previous findings [20], whereas Mtmr2+/-Fig4−/− and Mtmr2−/-Fig4−/− cells showed similar levels of this lipid (P = 0.0071 between wild-type and Fig4−/−; P = 0.0044, between wild-type and double null cells). Mcas of Mtmr2 in Mtmr2−/-Fig4+/- cells provoked a significant increase in PtdIns(3,5)/P2 (P = 0.0096), whereas no differences were detected between Mtmr2−/- and Mtmr2−/-Fig4+/- cells, possibly because heterozygous deletion of Fig4 leads to small PI changes hardly detectable using this method. (D) Comparison between the two independent experiments showed in (B) and (C), where each value was proportioned to the wild-type belonging to the same experiment and wild-type values were arbitrarily converted to 1 (P = 0.0055 when comparing Mtmr2−/-Fig4−/− and Mtmr2−/-Fig4+/- cells).

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in neurons was not excluded since axonopathies are length dependent and not easily reproduced in mice [18]. Interestingly, a role for MTMR2 in neurons in vitro has been recently reported suggesting that Mtmr2 is localized to excitatory synapses of central neurons via direct interaction with the PSD-95 scaffolding protein [33]. Knockdown of Mtmr2 in cultured neurons markedly

Figure 9. MTMR2 and FIG4 interaction in yeast. Expression of human FLAG-MTMR2 in the mutant yeast strain fig4Δ, caused enlarged vacuoles and a defect in vacuole fission. Wild-type and fig4Δ expressing pVT102U (vector), pVT102U-MTMR2 or pVT102U-MTMR2-C417S, were grown to mid-log phase and the vacuole membrane was labeled with SynaptoRed C2. Arrow indicates an example of a bud vacuole that has not properly separated from the mother vacuole. Scale bar, 6 μm.

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Figure 10. MTMR2 3-phosphatase activity in yeast. Mouse MTMR2 has 3-phosphatase activity in vivo and converts PtdIns(3,5)P2 and PtdIns3P to PtdIns5P and phosphatidylinositol, respectively. Expression of MTMR2 in yeast causes a decrease in PtdIns3P and PtdIns(3,5)P2 as well as an increase in PtdIns5P. Mouse MTMR2 cDNA was subcloned into a yeast expression vector, pVT102U (vector) to express MTMR2 from the ADH promoter, pVT102U-MTMR2. Yeast expressing MTMR2 or with vector alone were labeled with [3H]inositol for 18 h. Cells were exposed to 0.9 M NaCl for the times indicated, Lipids were extracted, deacylated, and the corresponding glycerol-inositol phosphates were quantitated by high-pressure liquid chromatography, reported as the percent of total [3H]phosphatidylinositol extracted. Two independent experiments are shown.

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reduced excitatory synapse density and function and it was proposed that the MTMR2/PSD95 complex contributes to the maintenance of excitatory synapses by inhibiting excessive endosome formation and destructive endosomal traffic to lysosomes.

Here, we assessed MTMR2 and FIG4 interaction in yeast and found that overexpression of MTMR2 reduces both PtdIns3P and PtdIns(3,5)P2 leading to an increase in vacuole size in the fig4Δ mutant. These findings support the in vivo role of MTMR2 as a 3-phosphatase that acts on both PtdIns3P and PtdIns(3,5)P2.

Fig4 heterozygosity rescues myelin outfoldings due to Mtmr2 deficiency both in vivo and in vitro, thus providing evidence of the Fig4 and Mtmr2 interaction in Schwann cells as well as neurons. Loss of Mtmr2 specifically in Schwann cells provokes myelin outfoldings. The presence of cytoplasmic inclusions in Schwann cells and the reduced NCV in the Fig4-null mouse, and the typical demyelinating features (onion bulbs) of CMT1J patients, all strongly support a Schwann cell autonomous role for Fig4. But how does loss of Fig4 in Schwann cells rescue Mtmr2-null myelin outfoldings? We hypothesized that a 50% reduction of Fig4 might be sufficient to rebalance the PtdIns(3,5)P2 elevation in Mtmr2-null cells (Figure 11), thus reducing myelin outfoldings. MTMR2 loss should lead to an increase of both PtdIns3P and PtdIns(3,5)P2, whereas FIG4 loss reduces PtdIns(3,5)P2 levels. In agreement with this model, we observed that downregulation of PIKfyve expression or inhibition of its activity in Mtmr2-null co-cultures reduced myelin outfoldings, as also observed with Fig4 heterozygosity (Figure 11). Our results therefore suggest that imbalance of PtdIns(3,5)P2 is at the basis of altered longitudinal myelin growth and formation of myelin outfoldings. The observed rescue of myelin outfoldings is likely mediated by restored PtdIns(3,5)P2 rather than PtdIns3P. PtdIns5P may be produced via dephosphorylation of PtdIns(3,5)P2 by MTMRs, and can also be generated, at least in vitro, by PIKfyve acting on phosphatidylinositol [34]. Therefore, Fig4 heterozygosity in Mtmr2-null cells would lead to a further reduction in PtdIns5P rather than restoration, as for PtdIns(3,5)P2.

PtdIns(3,5)P2 is thought to be localized to EE and the limiting membranes of LE/LY, although it cannot be excluded that this lipid might also be generated at other membranes. The lack of specific probes to detect PtdIns(3,5)P2 prevents the definition of other membrane localization [35,36]. Our studies raise the question of how dys-regulation of PtdIns(3,5)P2 leads to aberrant longitudinal myelin growth. Excessive longitudinal myelin growth and myelin outfoldings might arise as a consequence of reduced endocytosis/recycling and degradation or as a consequence of increased exocytosis. One can speculate that increased PtdIns(3,5)P2 due to loss of MTMR2 might favor exocytosis from...

Figure 11. Model of MTMR2 and FIG4 phosphatase activities in the control of PtdIns3P-PtdIns(3,5)P2 metabolism. In red, altered levels of PIs due to loss of MTMR2 and/or FIG4 are depicted. In blue, rebalanced levels of PtdIns(3,5)P2 in Mtmr2-null cells upon Fig4 heterozygosity and downregulation of either PIKfyve expression or inhibition of its activity are shown.

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the LE/LY compartment during myelin biogenesis. However, this mechanism, which has been recently suggested to occur in oligodendrocytes [37], accounts for the assembly of myelin components during the active phase of myelination. In myelin outfoldings, myelin thickness is normal, so a more subtle mechanism of regulation would be involved. Increased PtdIns(3,5)P_2 might alter autophagy dynamics. However, we did not observe any change on LC3II/I ratio and/or p62 levels in Mtmr2-null nerves or in myelin-forming DRG co-cultures (unpublished results).

Alternatively, MTMR2 may favor endocytosis and counteract ectocytosis during later stages of myelin biogenesis. The myelin outfoldings may thus arise as a consequence of the loss of negative control on the amount of membrane produced during myelination. Another alternative is that MTMR2 might control endocytosis of specific transmembrane proteins linking Schwann cell plasma membrane to the axonal plasma membrane, which then act as signaling molecules to control longitudinal myelin growth. Note that myelin outfoldings often contain axoplasm and axons branches at paranodal regions thus following myelin elongation. Other members of the MTMR family seem to possess similar biological functions. MTMR4 was recently demonstrated to regulate the sorting of MTMR family seem to possess similar biological functions. Note that myelin outfoldings may thus arise as a consequence of the loss of negative control on the amount of membrane produced during myelination.

Materials and Methods

Ethics statement

All experiments involving animals were performed in accordance with Italian national regulations and covered by experimental protocols reviewed by local Institutional Animal Care and Use Committees.

Mice

Mtmr2-null mice were backcrossed for at least 5 generations to strain C57BL/6N.

Fig4+/−/ heterozygous mice were maintained on the recombinant inbred line CB:plt derived predominantly from strains CAST/Ei and C57BL/6j [25%] [25]. Heterozygous Fig4+/− males were crossed with Mtmr2-null females to obtain Mtmr2−/−/−Fig4+/−/−/− double heterozygous mice. Double heterozygotes were crossed to generate Mtmr2−/−/−Fig4+/−/−/− null mice as well as Mtmr2−/−/−Fig4+/−/−/− mice for analysis. Genotyping was performed as described [17,20].

Morphological analysis

Semithin morphological analysis was performed as described previously [41]. For morphometric analysis in brainstem at P8, neuronal damage was evaluated in the facial nucleus at the level of the upper medulla oblongata (Bregma = −5.88). For each experimental sample, microscopic images [130 μm×90 um, nine images per slide, three slides for each brain] were taken with a digital camera and processed by Adobe Photoshop 7.0 software. To be counted, a cell (diameter >20 μm) had to be located in the facial nucleus and 100–150 cells were scored per section. Cells with abnormal cytoplasm vacuolization were scored as pathological. The average percentage of normal and damaged neurons for each sample was considered for each experimental group to represent the neuronal density. Counts were performed in double blind by 2 investigators on slides with a number-code system, and results were analyzed.

The number of motorneurons and of total cells in spinal cord was assessed by performing at least 15 sections for each spinal cord from three animals per genotype as before and by counting the number of cells per area-cell density (mm²).

The proportion of fibers carrying myelin outfoldings in Mtmr2-null nerves as compared to Mtmr2-null mice with Fig4+/−/− heterozygosity was determined by measuring the number of fibers carrying myelin outfoldings normalized to the total number of axons per section (the entire nerve section was reconstructed).

Ultrathin morphological analysis was conducted as reported previously [41]. For morphological analysis, three to five animals were evaluated at each time point in most cases.

Primary mouse fibroblast (MF) culture

MFs were established at P3 from tails and legs chopped in pieces and incubated after PBS washing with RPMI medium and 1 mL Collagenase Type II (Stock = 2000 U/mL in 1× PBS, Worthington, LS004204) overnight at 37°C. The next day, cells were plated in RPMI-1640 with 15% FBS/1× L-Glutamine/1× Pen/Strep. Cells were subjected to only two-three passages to allow maximum efficiency of metabolic labelling for PI measurement.

Phospholipid analysis

Fibroblasts were labeled for 16 h in phosphate free DMEM (Invitrogen) containing 200 μCi/ml [32P]orthophosphate (Perkin Elmer). Lipids were extracted, separated on Silica gel G60 plates and analyzed by HPLC as described previously [42].

PtdIns5P was quantified by mass assay as described [43]. Briefly total lipids were extracted from duplicate or triplicate plates of DRG co-cultures from Mtmr2+/− or Mtmr2−/− knock-out mice and separated on Silica gel G60 plate. Monophosphorylated Pls were scraped, eluted from silica and assessed for PtdIns(4,5)P_2 formation in vitro using the recombinant specific PIP4KIIalpha and [gamma-32P] ATP.

Antibodies

For western blot analysis and immunohistochemistry the following antibodies were used: rat anti-LAMP1 (Iowa Hybridoma bank), Guinea pig anti-P62 (Progen), rabbit anti-GFAP (Sigma), rabbit anti-MAG (kindly provided by Dr. J. Salzer), rabbit anti-AMP (kindly provided by Dr. V. Lee), mouse anti-MBP (Covance), rabbit anti-NF-L (Chemicon), mouse anti-tubulin (Sigma), and mouse anti-FG4 (NeuroMab).

Schwann cell/DRG neuron co-cultures

Myelin-forming Schwann cell/DRG neuron co-cultures were established from E13.5 mouse embryos as previously described [19,44]. Myelination was induced by treatment for 15 days with ascorbic acid (final concentration, 50 μg/ml (Sigma-Aldrich). Dissociated Schwann cell/DRG neuron co-cultures were established as described but DRGs were first incubated with trypsin (0:25%) for 45 min at 37°C. Cells were also mechanically dissociated and then plated at a concentration of one to two DRGs per glass coverslip. Isolated rat Schwann cells were prepared as reported previously and cultured using DMEM with 10% of fetal calf serum, 2 ng/ml recombinant human nerve growth factor, and 2 mM forskolin (Calbiochem).
YM201636 was provided by Symansis. A titration of the compound starting from 800 nM until 30 nM was performed on co-cultures to select the maximum amount of compound which did not affect myelination. As already described [27,28], 100 or 800 nM of compound provoked extensive cell vacuolization after overnight incubation. YM201636 was provided to co-cultures at 70 nM every other day together with ascorbic acid for 13 days to achieve full myelination.

Immunohistochemistry

Schwann cell/DRG neuron co-cultures were fixed for 15 min in 4% paraformaldehyde, permeabilized for 5 min in ice-cold methanol at −20°C, blocked for 20 min with 10% normal goat serum (Dako), 1% bovine serum albumin (BSA) (Sigma-Aldrich), and then incubated with primary antibody for 1 h. After extensive washing, the coverslips where incubated with the secondary antibody for 30 min, washed, and mounted. For double immunostaining with anti-NF-L and anti-MBP antibody, the coverslips were blocked with 1% BSA, 10% NGS for 20 min on ice, and primary antibodies were incubated overnight at 4°C. For LAMP1 staining, fibroblasts were permeabilized using 0.1% saponin after fixation. For immunolabeling, secondary antibodies included fluorescein-conjugated (FITC) and rhodamine (tetramethylrhodamine isothiocyanate) (Jackson ImmunoResearch). Coverslips were analyzed using TCS SP3 laser-scanning confocal (Leica) or Olympus BX (Olympus Optical) fluorescent microscope, and Zeiss Axiovert S100 TV2 with Hamamatsu OrcaII-ER.

Analysis of myelination

To quantify the amount of myelination, the number of MBP positive segments in each explant/cover-slip was assessed. As myelination is also a function of the amount of neurites/axons and of the Schwann cell number in the culture, the network of NF-L-positive filaments and the number of Schwann cells (DAPI) were also evaluated in each explant. To quantify MBP-positive fibers displaying myelin outfoldings, at least 200 MBP-positive myelinated fibers per explant/cover-slip were evaluated, in at least ten different explants/cover-slip. The percentage of MBP-positive fibers showing myelin outfoldings among the total number of MBP-positive fibers was counted.

Analysis of fibroblasts with enlarged late endosome/lysosomes

Fibroblasts were stained using LAMP1 antibody and images were acquired using a confocal microscope. Images were then processed using the Image J software and those cells displaying almost all LAMP1 positive endosomes bigger than 1.67 μm (only occasionally observed in wild-type cells) were considered as carrying enlarged late endosome/lysosomes.

Imaging and statistical analysis

Micrographs were acquired using a digital camera (Leica F300), and figures were prepared using Adobe Photoshop, version 7.0 and 8.0 (Adobe Systems). Statistical analysis was performed using the Student t test; two tails, unequal variants, and alpha = 0.005 were used. Error bars in the graphs represent SEM.

Lentiviral vector (LV) preparation

To downregulate PIKfyve expression, a shRNA cloned into pLKO.1 LV (human U6 promoter) without a GFP reporter was used (clone ID TRCN0000150081). Non-concentrated LVs were used for RNA interference. The transfer constructs were transfected into 293FT cells together with packaging plasmids Δ6.9 and pCMV-VSGV using Lipofectamine 2000 (Invitrogen). As control, a vector encoding a shRNA to a nonspecific sequence (helicase) was used. Viral supernatants were collected 48 h after transfection, centrifuged at 3000 rpm for 15 min, and frozen at −80°C.

To check for PIKfyve depletion, freshly plated rat Schwann cells (10⁶ cells per 100-mm plate) were incubated with the LV’s in DMEM, 10% FBS, and 2 mM L-glutamine plus forskolin and rhNRG-1 (EGF domain, R&D). Cells were expanded for an additional week and maintained in MEM, 10% FBS, 2 mM L-glutamine and 2 μM forskolin before use. A western blot using an anti-PIKfyve antibody (Santa Cruz) was performed. Using non-concentrated LV, transduction of Schwann cell/DRG neuron co-cultures was performed 4–5 days after dissection by incubating the cells with LVs overnight. Cells were then supplemented with C-media, and myelination was induced after 2 days.

Glutathione S-transferase–binding assays

Glutathione S-transferase (GST) fusion proteins were expressed in *Escherichia coli* BL21 cells and purified directly from bacterial extract on glutathione-Sepharose 4 Fast Flow beads. Rat isolated Schwann cells and mouse brains were homogenated, and protein lysates were prepared using a binding buffer with 1%NP-40, 50 mM Tris buffer, pH 7.4, 10% glycerol, 100 mM NaCl, 10 mM NaF, 1 mM Na-vanadate. Equal amounts of protein lysates were incubated for 4 h at 4°C with immobilized GST fusion proteins and GST as control. After three washes with a buffer containing 0.5% NP-40, the pellets were dissolved in SDS sample buffer and analyzed by SDS-PAGE and immunoblotting. To show the relative amount of GST fusion proteins used, beads were dissolved in SDS sample buffer and analyzed by SDS-PAGE, and the gel was stained with Coomassie.

Yeast analysis and phospholipid measurement in yeast

Yeast cells were labeled with SynaptoRed C2 (Biotium, Inc., CA). 0.1 units of cells (at 600 nm) were collected and resuspended in 250 μl fresh media. 6 μl of SynaptoRed C2 (10 μg/μl dissolved in dimethyl sulfoxide) was added to the cells and incubated at 24°C for 1 hour. Cells were then washed 2 times with fresh media and chased for 2.5 hours. Fluorescence and differential interference contrast (DIC) images were generated using a DeltaVision RT Microscope System (Applied Precision, WA). Images were processed using Softworx and Adobe Photoshop.

Measurement of phosphoinositide levels were performed as described previously [45]. Cells were grown in selective media to mid-log phase, harvested, washed, and resuspended in synthetic media lacking inositol. 1–4×10⁶ cells were inoculated into 5 ml of media lacking inositol containing 5 μCi of myo-[2-3H]-inositol. Cells were labeled for 18 h at 24°C, harvested by centrifugation, washed, and resuspended in 100 μl of inositol-free media. For hyperosmotic shock, an equal volume of 1.8 M NaCl was added to cells (for a final concentration of 0.9 M NaCl) and the resulting suspension was incubated at 24°C for the times indicated. 800 μl of ice cold 4.5% perchloric acid [46] was added to the cells. Cells were lysed in the presence of 0.5-mm zirconia beads (Biospec, Bartlesville, OK) on a Beadbeater (Biospec) for three cycles of 2 min at room temperature followed by 2 min on ice. Cell extracts were centrifuged at 14,000 rpm for 10 min at 4°C. Precipitates were washed with 1 ml of 100 mM EDTA, centrifuged 14,000 rpm for 10 min at 4°C, and resuspended in 30 μl of sterile distilled deionized water.

Lipids were decylated by treatment with methylamine [47,1 ml methylamine reagent (10.7% methylamine, 45.7% methanol, 11.4% n-butanol) was added to each sample and
incubated at 55°C for 1 h. Samples were dried in a SpeedVac and the pellets were resuspended in 300 µl of sterile water, centrifuged at 14,000 rpm for 2 min and the supernatants were transferred to new Eppendorf tubes. 300 µl of butanol/ethyl ether/formic acid ethyl ester (20:4:1) was added to each. The samples were vortexed and centrifuged at 14,000 rpm for 2 min. The aqueous phase (bottom layer) was transferred to new tubes and the extraction was repeated. At the end of the second extraction the aqueous phase was dried in a SpeedVac. Samples were resuspended in 20 µl of sterile water and 15 µl of each was analyzed by HPLC using an anion exchange, Partispher SAX (Whatman), column. The column was developed with a gradient of 1 M (NH₄)₂HPO₄, pH 3.8 (pH adjusted with phosphoric acid); 0% for 5 min, 1–2% over 15 min, 2% for 80 min, 2–10% over 20 min, 10% for 65 min, 10–80% over 40 min, 80% for 20 min and finally 80-0%; flow rate, 1.0 ml/min [48]. The value of each glycerol-inositol corresponding to PtdIns3P, PtdIns4P, PtdIns5P, PtdIns(3,5),P2, and PtdIns(4,5),P2 is reported as percent of total phosphoinositol, to normalize number of cells and incorporation of [³H] inositol.

Supporting Information

**Figure S1** Semithin section analysis of the cerebellum from Mtmr2+/+Fig4−/− and Mtmr2−/−Fig4−/− mice at P8. A disorganization of both the molecular and granular layer was observed in the cerebellum of Mtmr2+/+Fig4−/− and Mtmr2−/−Fig4−/− mice. Middle panels show loss of Purkinje and basket cells which are not aligned at the border of the granular layer. Inset is showing a cell carrying vacuoles in the granular layer. Bar is 80 µm in (A–C); 50 µm in (A’–C’); 80 µm in (A’’) and 50 µm in (B’, C’). (TIF)

**Figure S2** Semithin section analysis of the cerebellum from Mtmr2+/+Fig4−/− and Mtmr2−/−Fig4−/− mice at P20. The loss of Purkinje and basket cells is even more evident at P20 at the border of the granular layer. More vacuolated cells are present in the granular layer. Arrows indicate vacuolated cells. Bar is 80 µm in (A–C) and 50 µm in (A’–C’). (TIF)

References


**Table S1** PI measurements from wild-type yeast strains transformed with FLAG-MTMR2. Values listed are the percent of total phosphatidylinositol. Values for two independent experiments, as well as the averages (Avg) for each time point are presented. (DOC)

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**Author Contributions**
Conceived and designed the experiments: AB HT BP LSW MHM. Performed the experiments: IV GD HT GC EK FC. Analyzed the data: IV HT ER BPLW AQ LSW MHM AB. Contributed reagents/materials/analysis tools: MHM. Wrote the paper: AB LSW MHM. Provided pli mice: MHM. Planned experiments: AB BP AQ LSW MHM.


