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PITHD1 is a proteasome-interacting protein essential for male fertilization

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The proteasome is a protein-degrading molecular complex that is necessary for protein homeostasis and various biological functions, including cell cycle regulation, signal transduction, and immune response. Proteasome activity is finely regulated by a variety of proteasome-interacting molecules. PITHD1 is a recently described molecule that has a domain putatively capable of interacting with the proteasome. However, it is unknown whether PITHD1 can actually bind to proteasomes and what it does in vivo. Here we report that PITHD1 is detected specifically in the spermatids in the testis and the cortical thymic epithelium in the thymus. Interestingly, PITHD1 associates with immunoproteasomes in the testis, but not with thymoproteasomes in the thymus. Mice deficient in PITHD1 exhibit severe male infertility accompanied with morphological abnormalities and impaired motility of spermatozoa. Furthermore, PITHD1 deficiency reduces proteasome activity in the testis and alters the amount of proteins that are important for fertilization capability by the sperm. However, the PITHD1-deficient mice demonstrate no detectable defects in the thymus, including T cell development. Collectively, our results identify PITHD1 as a proteasome-interacting protein that plays a nonredundant role in the male reproductive system.

Protein degradation is a crucial process in cellular homeostasis. The proteasome is a large enzyme complex responsible for protein degradation in cells (1). In most cases, proteasome-dependent proteolysis targets ubiquitinated proteins (2, 3). Protein degradation by the proteasome is important for not only

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the maintenance of protein homeostasis, but also various biological functions, including cell cycle regulation, signal transduction, and immune response (3, 4). The proteasome consists of a 20S catalytic core proteasome and regulatory particles that bind to the core proteasome (5). The 20S proteasome is composed of 28 subunits, namely, 2 sets of 7 different α subunits $(\alpha 1 - \alpha 7)$ and 7 different β subunits $(\beta 1 - \beta 7)$. Among the 20S proteasome components, β 1, β 2, and β 5 have catalytic activities (6). The proteasome containing β 1, β 2, and β 5 is called the constitutive proteasome and expressed in most tissues. On the other hand, antigen-presenting immune cells, hematopoietic cells, and cells exposed to proinflammatory cytokines express the immunoproteasome in which β_{1i} , β_{2i} , and β_{5i} are incorporated in place of β 1, β 2, and β 5, respectively (7). Immunoproteasomes possess altered proteolytic activity and participate in the efficient processing of major histocompatibility complex class I-restricted antigens (8). In the thymus, another catalytic subunit β 5t is specifically expressed in cortical thymic epithelial cells (cTECs)³ and incorporated into the 20S proteasome together with β_{1i} and β_{2i} , thereby forming the thymoproteasome (9, 10). Thymoproteasomes in cTECs are essential for the production of self-antigens that induce the positive selection of CD8⁺ T cells (9, 11). Thymoproteasome-dependent positive selection in the thymus also fine-tunes antigen responsiveness of $CD8^+$ T cells in the periphery (12).

The function of the 20S proteasome is modulated by interactions with regulatory proteins. A wide variety of proteasomeinteracting proteins have been identified, the most well-known of which is the 19S regulatory particle that binds to both ends of the 20S proteasome to form the 26S proteasome complex (10, 13, 14). The 19S regulatory particle consists of 6 ATPase subunits and 13 non-ATPase subunits, and pre-processes sub-

³ The abbreviations used are: cTEC, cortical thymic epithelial cell; PITH, proteasome-interacting thioredoxin; mTEC, medullary thymic epithelial cell; tdTomato, tandem dimeric tomato fluorescent protein; DP, double positive; VAP, average path velocity; VCL, curvilinear velocity; VSL, straight-line velocity; TMT, tandem mass tag; PI, propidium iodide; FDR, false discovery rate; gRNA, guide RNA; DAPI, 4',6-diamidino-2-phenylindole; ACN, acetonitrile; GAPDH, glyceraldehyde-3-phosphate dehydrogenase.



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This article contains Figs. S1 and S2 and Table S1.

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Figure 1. Association capability of PITHD1 with proteasomes in thymus and testis. *A*, relative mRNA expression of *Pithd1* in the indicated cells and organs isolated from 2-week-old C57BL/6 mice. The expression levels (mean \pm S.E.) of *Pithd1* measured by quantitative RT-PCR were normalized to that of *Gapdh* and compared with the level measured in thymocytes. *, p < 0.05, *n.s.*, not significant. Plotted are the results of three independent experiments using samples obtained from three to nine mice. *B* and *C*, thymus lysates were immunoprecipitated with anti- β 5t antibody (*B*), anti- α 6 antibody (*C*), or control IgG followed by immunoblotting with anti-PITHD1, anti- β 5t, and anti- α 6 antibody (*D*), anti- α 6 antibody (*D*), anti- β 5t antibody (*E*), anti- α 6 antibody (*F*), or control IgG followed by immunoblotting with anti-PITHD1, anti- α 5s, and anti- α 6 antibodies. *G* and *H*, thymus lysates from β 5t-deficient (*G*) or β 5t-sufficient (*H*) mice were immunoprecipitated with anti- β 5i antibody or control IgG followed by immunoblotting with anti- β 5t, and anti- α 6 antibodies. *Red arrowheads* indicate each protein. *Black arrowheads* indicate IgG light chain. All images in *panels B--H* are representative results of three independent experiments.

strates for degradation by the 20S proteasome. Furthermore, ubiquitin chain-binding receptors, de-ubiquitinating enzymes, and E3 ubiquitin ligases associate with the 19S regulatory particle, and assist or modulate proteasomal functions (13, 15). In addition to the 19S regulatory particle, proteasome activators and inhibitors, such as PA200, PA28, and PI31, directly bind to the 20S proteasome (13–16).

In the course of our explorative study for thymoproteasome modulators in cTECs, we noticed that a functionally unidentified gene encoding PITHD1 was abundantly expressed in cTECs. PITHD1 has a putative proteasome-interacting thioredoxin (PITH) domain that potentially binds to the proteasome. Another PITH domain-containing protein, thioredoxin-like protein 1 (TXNL1), possesses redox activity (17). Unlike TXNL1, PITHD1 lacks redox activity due to the absence of the activesite motif of thioredoxin (18). It has been reported that the ectopic expression of PITHD1 in human leukemia cell lines promotes megakaryocyte development by regulating the expression of RUNX1, a transcription factor that is important for lineage determination between megakaryocytes and erythrocytes in hematopoiesis (18, 19). However, whether PITHD1 can actually bind to proteasomes and what it does in vivo are not known. We report herein that PITHD1 is highly expressed in cTECs and the testis. PITHD1 associates with immunoproteasomes in the testis, but not with thymoproteasomes in cTECs.

By producing mice deficient in PITHD1, we show that PITHD1 deficiency causes infertility accompanied with morphological abnormalities and impaired motility of spermatozoa, without detectable defects in the development and function of cTECs. We further show that PITHD1 deficiency reduces proteasome activity in the testis and alters the amount of several proteins that are important for fertilization capability by the sperm. Our study reveals a novel and nonredundant function of PITHD1 as a proteasome-interacting protein essential for the male reproductive system.

Results

Detection of PITHD1 in cTECs and testis

We have previously reported that the β 5t-containing thymoproteasome specifically expressed in cTECs is important for the positive selection of CD8⁺ T cells in the thymus (9–12). Accordingly, we explored proteasome-interacting proteins that could affect the function of thymoproteasomes in cTECs. We found that *Pithd1*, which encodes PITHD1 that contains a putative proteasome-interacting domain, was highly expressed in cTECs in the transcriptomic profiles of thymic epithelial cells (20). Quantitative RT-PCR analysis confirmed that the expression of *Pithd1* was higher in cTECs than other thymic cells, including medullary thymic epithe-

lial cells (mTECs), CD45⁺ thymocytes, and non-TEC stroma cells (Fig. 1*A*). Moreover, we detected a significantly higher expression of *Pithd1* in the testis than other organs (Fig. 1*A*). Thus, we found that *Pithd1* is specifically and abundantly expressed in cTECs and the testis.

The detected PCR products of *Pithd1* were 328 bp in size, derived from spliced RNA, and not 9558 bp from genomic DNA (Fig. S1A). PCR detection of *Hprt* reconfirmed that the amplified signals in our RT-PCR analysis were primarily 249 bp derived from spliced RNAs and not 1087 bp derived from genomic DNAs contaminated in the total RNA samples (Fig. S1A).

PITHD1 associates with immunoproteasomes in testis but not with thymoproteasomes in cTECs

We next examined the association capability of PITHD1 proteins with proteasomes in cTECs and the testis. Immunoprecipitation of mouse thymus lysates using antibody specific for the cTEC-specific thymoproteasome component β 5t showed that PITHD1 protein was absent from the precipitates (Fig. 1B). This was not due to the failure of immunoprecipitation by anti- β 5t antibody because in addition to β 5t protein, α 6 protein, which is a 20S proteasome component, was immunoprecipitated by this antibody (Fig. 1B). Similarly, PITHD1 protein in the thymus was not immunoprecipitated by anti- α 6 antibody, whereas β 5t protein was co-precipitated by the same anti- α 6 antibody (Fig. 1C and Fig. S1B). In the testis, the expression of immunoproteasomes and testis-specific proteasomes has been reported (21–23). In the testis-specific proteasome, the α 4s subunit is incorporated into the constitutive proteasome in place of the $\alpha 4$ subunit (21). Immunoprecipitation of testis lysates using antibody specific for α 4s showed the absence of PITHD1 protein in the precipitates (Fig. 1D). Again, this was not due to the failure of immunoprecipitation by anti- α 4s antibody, because in addition to α 4s protein, α 6 protein was immunoprecipitated by this antibody (Fig. 1D). On the contrary, immunoprecipitation by anti- β 5i antibody or anti- α 6 antibody revealed the presence of PITHD1 protein in the precipitates (Fig. 1, *E* and *F*). Therefore, these results indicate that PITHD1 associates with β 5i-containing immunoproteasomes in the testis either directly or indirectly, whereas it is incapable of binding to α 4s-containing testis-specific proteasomes in the testis or ß5t-containing cTEC-specific thymoproteasomes in the thymus.

In β 5t-deficient cTECs, β 5i is compensatively expressed, thereby forming immunoproteasomes instead of thymoproteasomes (11). Therefore, we examined whether PITHD1 could associate with immunoproteasomes in β 5t-deficient cTECs or β 5t-sufficient cTECs. We found that immunoprecipitation of β 5t-deficient thymus lysates using antibody specific for β 5i showed that PITHD1 protein was absent from the precipitates (Fig. 1, *G* and *H*). These results indicate that PITHD1 is incapable of binding to thymoproteasomes or immunoproteasomes expressed in cTECs.

Generation of PITHD1-deficient mice

To investigate the physiological role of PITHD1, we next generated PITHD1-deficient mice by introducing a targeting

ation site in exon 1 of the *Pithd1* gene along with neighboring genomic sequences into HK3i embryonic stem cells for homologous recombination (Fig. 2A). Thus, the tdTomato fluorescent protein would be detectable in the cells where Pithd1 was transcribed in this mouse. PCR analysis, Southern blot analysis, and sequencing analysis of genomic DNA isolated from the offspring mice indicated successful germline recombination at the *Pithd1* locus (Fig. 2, *B* and *C*). Quantitative RT-PCR analysis showed that Pithd1 mRNA expression in the thymus and the testis was lost in PITHD1 $^{-/-}$ mice (Fig. 2D). Immunoblot analysis also confirmed the loss of PITHD1 protein in the thymus and the testis in PITHD1^{-/-} mice (Fig. 2*E*). The tdTomato fluorescent protein was detectable in the Ly51-expressing cortical region of the PITHD1^{+/-} thymus (Fig. 2F). Flow cytometric analysis of thymic cells showed tdTomato expression in \sim 3% of cTECs (Fig. 2*G*), suggesting that PITHD1 is expressed in a minor subpopulation of cTECs. Intense tdTomato fluorescence was detected in the testicular regions surrounding the lumen of seminiferous tubules (Fig. 2H). Flow cytometric analysis showed that 37% of testicular cells highly expressed the tdTomato fluorescent protein (Fig. 21). Microscopic inspection of testicular cells revealed that the tdTomato fluorescence was detectable in the cytoplasm of elongated spermatids, whereas it was not detected in round spermatids or other testicular cells (Fig. 2J). The tdTomato fluorescence was also detectable in the cytoplasmic droplets of sperm where proteins are transiently stored and eventually removed in mature sperm (Fig. 21). We obtained two independent mouse lines exhibiting an identical phenotype, and will show the results from one PITHD1 $^{-/-}$ mouse line in this report.

vector that contained a gene encoding the tandem dimeric

tomato fluorescent protein (tdTomato) at the translation initi-

No abnormalities in development and function of cTECs in PITHD1-deficient mice

We then examined the impact of PITHD1 deficiency in the thymus. Hematoxylin and eosin staining and immunofluorescence analysis of the thymic sections showed unaffected corticomedullary architectures in PITHD1^{-/-} mice (Fig. 3, A and B). Similar to the WT thymus, the cortical regions of PITHD1^{-/-} mice were enriched with $\beta 5t^+$ cTECs and the PITHD1^{-/-} thymic medullas contained Aire⁺ mTECs essential for the generation of self-tolerant T cells (Fig. 3B). Flow cytometric analysis of liberase-digested thymic cells showed that the cellularity of CD45⁻EpCAM⁺UEA1⁻Ly51⁺ cTECs and CD45⁻EpCAM⁺UEA1⁺Ly51⁻ mTECs, and their proteasome activity were undisturbed in PITHD1^{-/-} mice (Fig. 3, C and D). Furthermore, neither the development nor the cellularity of thymocytes, including CD4/CD8 subpopulations and CD25/CD44 CD4⁻CD8⁻ double-negative subpopulations, was altered by PITHD1 deficiency (Fig. 3, E and F). The development of CD4⁺CD8⁺ double-positive (DP) thymocytes has been subdivided into three subpopulations: $CD69^{-}TCR\beta^{-/low}$ preselected DP cells (population I), CD69⁺TCR β^{low} post-selected DP cells (population II), and CD69⁺TCR β^{high} post-selected DP cells (population III) (24). The cellularity of DP thymocytes in each subpopulation was comparable between WT mice and PITHD1^{-/-} mice (Fig. 3G). Consistent with the normal thymo-





Figure 2. Generation of PITHD1-deficient mice. *A*, schematic diagram of Pithd1 locus, targeting vector, and targeted allele. *DT-A*, diphtheria toxin A. *pgk-Neo-pA*, phosphoglycerate kinase I promoter-driven neomycin resistance gene followed by polyadenylation signal sequence. The 3' probe is for Southern blot analysis. *Arrows* indicate primers for genotyping PCR. *B*, genotyping PCR analysis of genomic DNA from the indicated mice. Gel electrophoresis of amplified WT allele fragment (763 bp) and targeted allele fragment (1058 bp). Positions of the primers are shown in *A*. *WT*, PITHD1^{+/+} mice. *Het*, PITHD1^{+/-} mice. *KO*, PITHD1^{-/-} mice. *C*, Southern blot analysis of BamHI-digested genomic DNA from the indicated mice. Probe is shown in *A*. Signals at 5.4 and 9 kbp indicate *WT* and targeted alleles, respectively. *D*, relative mRNA expression of *Pithd1* in thymus and testis isolated from 4-week-old mice. The expression levels (mean \pm S.E.) of *Pithd1* measured by quantitative RT-PCR were normalized to that of *Gapdh*. ****, *p* < 0.001. Plotted are the results of three independent experiments using the samples obtained from three mice per group. *E*, immunoblot analysis of PITHD1 protein in thymus and testis isolated from 4-week-old mice. *B*-Actin was examined as loading control. *F*, immunofluorescence analysis of tdTomato (*red*), Ly51 (*green*), and UEA1 (*blue*) in thymic section of PITHD1^{+/-} (PITHD1^{+/tdTomato}) mice. *Bar*, 100 µm. *G*, detection of tdTomato fluorescence in cECs analyzed by flow cytometry. *H*, immunofluorescence analysis of tdTomato fluorescence in testicular cells analyzed by flow cytometry. *J*, immunofluorescence analysis of fur *A*-week-old floor soft doed by flow cytometry. *J*, mice at 8-week-old. *Scale bars*, 20 µm. All images (*B*, *C*, *E*, *F*, *H*, and J) and histograms (*G* and J) are representative results of at least three independent experiments.

cyte development in PITHD1^{-/-} thymus, the cellularity of splenic CD4⁺ T cells and CD8⁺ T cells was comparable with those in WT mice (Fig. 3*H*). Therefore, these results indicate that despite the physiological expression of PITHD1 in cTECs, PITHD1 deficiency has no detectable impact on the development and function of the thymus.

We also examined whether PITHD deficiency induces the phenotypic alteration of the thymus in $\beta 5t^{-/-}$ mice. The cellularity of cTECs and mTECs in $\beta 5t^{-/-}$ mice was not altered by PITHD1 deficiency (Fig. 4*A*). Furthermore, the cellularity of TCR β^{high} CD4⁻CD8⁺ thymocytes and splenic CD8⁺ T cells, which was reduced in $\beta 5t^{-/-}$ mice, was neither further decreased nor recovered by PITHD1 deficiency (Fig. 4, *B* and

C). Taken together with the finding that PITHD1 is incapable of binding to thymoproteasomes in β 5t-sufficient cTECs or immunoproteasomes in β 5t-deficient cTECs, these results indicate that PITHD1 plays no apparent roles in the thymus in the presence or absence of the thymoproteasome.

PITHD1-deficient mice exhibit male infertility

We then examined the functional significance of PITHD1 in the testis. The weights of the testis and the epididymis were comparable between PITHD1^{+/-} and PITHD1^{-/-} mice (Fig. 5*A*). The morphology of the testis and the epididymis was undisturbed by PITHD1 deficiency (Fig. 5*B*). However, when PITHD1^{-/-} male mice were mated with C57BL/6 female mice



Figure 3. Characterization of the thymus in PITHD1-deficient mice. A, hematoxylin and eosin staining of thymic sections from 4-week-old mice. Representative data from three independent mice analyzed in three independent experiments are shown. Scale bar, 500 μ m. B, immunofluorescence analysis of eta5t (green), Aire (red), and UEA-1-binding molecules (blue) in thymic sections from 4-week-old mice. Representative data from three independent mice analyzed in three independent experiments. Scale bars, 100 µm. C, flow cytometric analysis of liberase-digested thymic cells isolated from 4-week-old mice. Shown are dot plots of EpCAM and CD45 expression in total cells (*left*) and UEA1 reactivity and Ly51 expression in CD45⁻EpCAM⁺-gated epithelial cells (*right*). Plots on the right show cell number (mean ± S.E.) of CD45⁻EpCAM⁺UEA1⁻Ly51⁺ cTECs and CD45⁻EpCAM⁺UEA1⁺Ly51⁻ mTECs in individual mice, measured in six independent experiments. D, histograms show the detection of proteasome activity by cell-permeable triple-leucine substrate-based fluorescent probe in cTECs and mTECs of PITHD1^{+/+} mice (*blue line*) and PITHD1^{-/-} mice (*red line*). *Shaded area* and *black line* represent background fluorescence profiles without the addition of proteasome probe in PITHD1^{+/+} TECs and PITHD1^{-/-} TECs, respectively. Plots on the *right* show mean fluorescence intensity (mean \pm S.E.) of indicated cells in individual mice, measured in three independent experiments. E, flow cytometric analysis of total thymocytes isolated from 4-week-old mice. Shown are dot plots of CD4 and CD8 expression in propidium iodide⁻ (PI⁻) thymocytes. Plots on the right show cell number (mean ± S.E.) of indicated cell populations in individual mice, measured in five independent experiments. F, flow cytometric analysis of CD44 and CD25 expression in CD4⁻CD8⁻ thymocytes. Plots on the right show cell number (mean ± S.E.) of CD44⁺CD25⁻ cells (DN1), CD44⁺CD25⁺ cells (DN2), CD44⁻CD25⁺ cells (DN3), and CD44⁻CD25⁻ cells (DN4) in individual mice, measured in five independent experiments. G, flow cytometric analysis of CD69 and TCR β expression in CD4⁺CD8⁺ thymocytes. Plots on the *right* show cell number (mean ± S.E.) of CD69⁻TCR β^{-rlow} cells (population I), CD69⁺TCR β^{low} cells (population I), and CD69⁺TCR β^{high} cells (population III) in individual mice, measured in three independent experiments. H, flow cytometric analysis of spleen cells. Shown are dot plots of CD4 and CD8 expression in CD3⁺ splenic T cells. Plots on the *right* show cell number (mean ± S.E.) of indicated cell populations in individual mice, measured in five independent experiments. Numbers in dot plots indicate frequency of cells within indicated area. n.s., not significant.

for 2 months, no pregnancy was observed in female mice even though vaginal plug formation was detectable (Fig. 5*C*). On the other hand, PITHD1^{+/-} male mice were able to impregnate female mice and 7 pups were produced on average per pregnancy (Fig. 5*C*). An *in vitro* fertilization assay confirmed the

fertilization incapability of sperm isolated from PITHD1^{-/-} mice in both cumulus-intact and cumulus-free conditions (Fig. 5*D*). However, PITHD1^{-/-} sperm were able to fertilize an egg under the zona pellucida-free condition (Fig. 5*D*). Therefore, these results indicate that sperm from PITHD1^{-/-} mice are





Figure 4. T cell development in β **5t/PITHD1 double-deficient mice.** *A*, flow cytometric analysis of liberase-digested thymic cells from the indicated mice. Shown are dot plots of UEA1 reactivity and Ly51 expression in CD45⁻EpCAM⁺-gated epithelial cells. Plots on the *right* show cell number (mean \pm S.E.) of CD45⁻EpCAM⁺UEA1⁻Ly51⁺ cTECs and CD45⁻EpCAM⁺UEA1⁺Ly51⁻ mTECs in individual mice, measured in three independent experiments. *B*, flow cytometric analysis of total thymocytes. Shown are dot plots of CD4 and CD8 expression in Pl⁻TCR^{high} thymocytes. Plots on the *right* show cell number (mean \pm S.E.) of indicated cell populations in individual mice, measured in three independent experiments. *C*, flow cytometric analysis of spleen cells. Shown are dot plots of CD4 and CD8 expression in Pl⁻TCR^{high} thymocytes. Plots on the *right* show cell number (mean \pm S.E.) of CD4 and CD8 expression in CD3⁺ splenic T cells. Plots on the *right* show cell number (mean \pm S.E.) of the indicated cell populations in individual mice, measured in three independent experiments. *C*, flow cytometric analysis of spleen cells. Shown are dot plots of CD4 and CD8 expression in CD3⁺ splenic T cells. Plots on the *right* show cell number (mean \pm S.E.) of the indicated cell populations in individual mice, measured in three independent experiments. Numbers in dot plots indicate frequency of cells within indicated area. *, *p* < 0.05; *n.s.*, not significant.

defective in fertilization capability due to passage disability through egg layers such as cumulus and zona pellucida. We noticed no defects in the fertility of female $PITHD1^{-/-}$ mice (data not shown).

PITHD1-deletion mice exhibit similar phenotype to PITHD1deficient mice

The results so far indicated that PITHD1 is important for male reproductive function rather than cTEC function. However, it remained possible that the artificial expression of tdTomato in $PITHD1^{-/-}$ mice might have hindered the phenotypes that could have resulted simply from the loss of PITHD1. Therefore, we next generated a new set of PITHD1-deletion mice (Δ/Δ) in that the genomic region spanning the middle of exon 1 to exon 3 was deleted by CRISPR/Cas9 technology (Fig. 6A). PCR analysis of genomic DNA isolated from the offspring mice indicated successful deletion of the *Pithd1* genomic sequence (Fig. 6B). Quantitative RT-PCR analysis and immunoblot analysis confirmed the loss of PITHD1 at mRNA and protein levels (Fig. 6, C and *D*). Like PITHD1^{-/-} mice, the cellularity of Ly51⁺ cTECs and UEA1⁺ mTECs as well as those of thymocyte subpopulations and splenic T cells was undisturbed in PITHD1^{Δ/Δ} mice (Fig. 6, E-G). The morphology of the testis and the epididymis

was also undisturbed in PITHD1^{Δ/Δ} mice (Fig. 6*H*). However, an *in vitro* fertilization assay of sperm isolated from PITHD1^{Δ/Δ} mice revealed fertilization incapability in both cumulus-intact and cumulus-free conditions, like PITHD1^{-/-} mice and unlike PITHD1^{WT/ Δ} mice (Fig. 6*I*). Therefore, these results indicate that the aberrant expression of the tdTomato fluorescent protein has essentially no impact on the phenotype of PITHD1^{-/-} mice, and reinforce that the impaired fertilization capability of sperm is caused by PITHD1 deficiency.

PITHD1 deficiency results in abnormal morphology and impaired motility of sperm

To understand the mechanism behind male infertility caused by PITHD1 deficiency, we next examined the morphology of mature sperm isolated from the cauda epididymis of PITHD1^{-/-} mice (Fig. 7*A*). The number of sperm was comparable between PITHD1^{+/-} mice and PITHD1^{-/-} mice (data not shown). However, the frequency of sperm with normal morphology was significantly reduced in PITHD1^{-/-} mice compared with PITHD1^{+/-} mice (Fig. 7*B*). On the other hand, the frequency of sperm with multiheads was significantly increased in PITHD1^{-/-} mice (Fig. 7*B*). The reduction of morphologically normal sperm and the increase of sperm with



Figure 5. Male infertility in PITHD1-deficient mice. *A*, weights (mean \pm S. E., individually measured for four mice) of testis and epididymis in 8-week-old mice. *n.s.*, not significant. *B*, hematoxylin and eosin staining of testis and epididymis sections from 8-week-old mice. Representative data from three independent mice analyzed in three independent experiments are shown. *Scale bar*, 100 μ m. *C*, four PITHD1^{+/-} male mice and four PITHD1^{-/-} male mice were mated with C57BL/6 female mice, respectively. Plots on the *left* show pregnancy rate (pregnancy/vaginal plug formation) in female mice mated with indicated male mice. Plots on the *right* show the number of pups born in one pregnancy. Mean \pm S.E. are shown by *black bars*. *D*, *in vitro* fertilization assay. Sperm from indicated male mice were incubated with cumulus-intact, cumulus-free, or zona-free WT eggs, and the numbers of eggs carrying pronuclei (zona-free eggs; *n* = 176) and of eggs from three independent experiments are plotted.

abnormal midpiece and multiheads were also detected in PITHD1^{Δ/Δ} mice (Fig. 7*C*). We next examined the motility of sperm incubated in TYH medium for 10 min or 2 h by measuring the average path velocity (VAP), the curvilinear velocity (VCL), and the straight-line velocity (VSL). The values of these motility parameters for PITHD1 $^{-/-}$ sperm were comparable with those for PITHD1^{+/-} sperm after 10 min incubation (Fig. 7D). However, a significant reduction of these velocities was detected in PITHD1^{-/-} sperm after 2 h incubation (Fig. 7*D*). Similarly, motility reduction in PITHD1 $^{\Delta/\Delta}$ sperm was detected after 2 h incubation (Fig. 7E). Morphological analysis revealed that motile sperm from PITHD1^{+/-} and PITHD1^{WT/ Δ} mice were predominantly morphologically normal, whereas a majority of motile sperm from $\text{PITHD1}^{-/-}$ and $\text{PITHD1}^{\Delta/\Delta}$ mice exhibited morphological abnormalities (Fig. 7, F and G). Collectively, these results indicate that PITHD1 deficiency causes impaired sperm motility accompanied with morphological abnormalities.

PITHD1 regulates proteasome activity in testicular cells

So far, we have shown that PITHD1 associates with β 5i-containing immunoproteasomes in the testis and that its deficiency causes impaired sperm motility. We therefore speculated the possibility that PITHD1 deficiency might affect proteasome activity in the testis. In mice that carried the *Pithd1* knock-out tdTomato knock-in allele (Fig. 2A), tdTomato fluorescence expression enabled the detection of cells in that *Pithd1* was transcribed. In the testis of these mice, tdTomato fluorescence

was detected in the elongated spermatids and the cytoplasmic droplets of sperm (Fig. 2J). We examined proteasome activity in tdTomato⁺ and tdTomato⁻ cells in the testis by using a cellpermeable triple-leucine substrate-based fluorescent probe that shows high affinity binding to the chymotrypsin-like catalytic site of proteasomes (25). The fluorescence intensity detected with this probe could reflect the chymotrypsin-like activity and/or the abundance of proteasomes. We found that the fluorescence intensity in tdTomato⁺ testicular cells was significantly reduced in PITHD1^{-/-} mice compared with PITHD1^{+/-} mice (Fig. 8, A and B, and Fig. S2A). We also found that the fluorescence intensity was unaltered in tdTomato⁻ testicular cells, tdTomato⁺ sperm, and tdTomato⁻ sperm by PITHD1 deficiency (Fig. 8, A and B). It was possible that the reduced proteasome activity was due to the reduced abundance of proteasomes. Indeed, the amount of the immunoproteasome component β 5i in tdTomato⁺ testicular cells was significantly reduced in PITHD1 $^{-/-}$ mice compared with that in PITHD1^{+/-} mice (Fig. S2B). We also found that the proteasome activity measured by the fluorescent probe was higher in tdTomato⁺ testicular cells than tdTomato⁻ testicular cells, and subsequently reduced in tdTomato⁺ sperm and tdTomato⁻ sperm in both PITHD1^{+/-} mice and PITHD1^{-/-} mice (Fig. 8, A and B). Correlated with the higher proteasome activity in tdTomato⁺ testicular cells than sperm, immunoblot analysis using lysate from PITHD1^{+/+} cells showed that the amount of PITHD1 protein was high in the testis but reduced in caput





Figure 6. Generation of PITHD1-deletion mice by CRISPR/Cas9 technology. A, schematic diagram of Pithd1 locus and targeted allele. Bold arrows indicate the position of guide RNA. Primer 1, Primer 2, and Primer 3 are for genotyping PCR. B, genotyping PCR analysis of genomic DNA from indicated mice. Gel electrophoresis of amplified WT allele fragment (198 bp) and targeted allele fragment (167 bp). Positions of the primers are shown in A. Representative data from three independent mice analyzed in three independent experiments are shown. C, relative mRNA expression of Pithd1 in testis. The expression levels (mean \pm S.E.) of *Pithd1* measured by quantitative RT-PCR were normalized to that of *Gapdh*. *, *p* < 0.05. Plotted are the results of three independent experiments using three mice per group. *D*, immunoblot analysis of PITHD1 protein in testis isolated from 8-week-old mice. β -Actin was examined as loading control. Representative results of three independent experiments are shown. E, flow cytometric analysis of liberase-digested thymic cells isolated from 8-week-old mice. Shown are dot plots of EpCAM and CD45 expression in total cells (*left*) and UEA1 reactivity and Ly51 expression in CD45⁻EpCAM⁺ epithelial cells (right). Plots on the right show cell number (mean ± S.E.) of CD45⁻EpCAM⁺UEA1⁻Ly51⁺ cTECs and CD45⁻EpCAM⁺UEA1⁺Ly51⁻ mTECs in individual mice, measured in three independent experiments. *F*, flow cytometric analysis of total thymocytes isolated from 8-week-old mice. Shown are dot plots of CD4 and CD8 expression in propidium iodide⁻ (*PI*⁻) thymocytes. Plots on the *right* show cell number (mean ± S.E.) of indicated cell populations in individual mice, measured in three independent experiments. G, flow cytometric analysis of total spleen cells isolated from 8-week-old mice. Shown are dot plots of CD4 and CD8 expression in CD3⁺ splenic T cells. Plots on the right show cell number (mean ± S.E.) of indicated cell populations in individual mice, measured in three independent experiments. Numbers in dot plots indicate frequency of cells within indicated area. n.s., not significant. H, hematoxylin and eosin staining of testis and epididymis sections from 8-week-old mice. Representative data from three independent experiments are shown. Scale bar, 100 µm. I, in vitro fertilization assay. Sperm from indicated male mice were incubated with cumulus-intact, cumulus-free, or zona-free WT eggs, and the numbers of eggs carrying pronuclei (zona-free eggs; n = 177) and of eggs that had reached the 2-cell stage (cumulus-intact eggs; n = 101; cumulus-free eggs; n = 38) were counted. The frequencies (mean \pm S.E.) of fertilized eggs from four independent experiments are plotted.

epididymal sperm and cauda epididymal sperm (Fig. 8*C*). Taken together, these results indicate that PITHD1 regulates proteasome activity including its abundance in testicular cells including elongated spermatids.

We finally performed MS-based quantitative proteomic analysis of sperm isolated from PITHD1^{+/+} mice and PITHD1^{-/-} mice. Proteins extracted from sperm were digested with trypsin, and the peptides were labeled with iso-





Figure 7. Morphological abnormalities and impaired motility of PITHD1-deficient sperm. *A*, morphology of sperm isolated from PITHD1^{-/-} mice. All images are representative results of three independent mice in three independent experiments. *B* and *C*, frequency (mean \pm S.E.) of sperm exhibiting indicated morphologies, measured individually in PITHD1^{-/-} mice (*B*), PITHD1^{Δ/Δ} mice (*C*), and control mice (three mice per group). *, *p* < 0.05; **, *p* < 0.01; ***, *p* < 0.001, *n.s.*, not significant. *D* and *E*, sperm isolated from PITHD1^{-/-} mice (*D*), PITHD1^{Δ/Δ} mice (*E*), and control mice were allowed to swim out from the dissected cauda epididymis in TYH medium for 10 min (noncapacitating) or 2 h (capacitating). VAP, VCL, and VSL were measured using a computer-assisted sperm analysis system. Shown are graphs of each parameter (mean \pm S.E.) measured in sperm isolated individually from three to four mice per group. *, *p* < 0.05; **, *p* < 0.01; ***, *p* < 0.001; *n.s.*, not significant. *F* and *G*, frequencies of indicated morphologies observed in motile sperm isolated from PITHD1^{-/-} mice (*F*), PITHD1^{Δ/Δ} mice (*C*), and control mice *n*, sperm number measured in individual mice (three mice per group).

baric tandem mass tag (TMT) reagents. A high confidence analysis at 1% false discovery rate (FDR) of LC-tandem mass spectrometric (LC-MS/MS) profiles resulted in the identification and quantification of 3,727 protein species (Table S1). Although few proteins showed significant alteration in abundance (Q < 0.05) in the comparison between PITHD1^{+/+} mice and PITHD1^{-/-} mice, modest differences (Q < 0.15) in the amounts of several proteins were detected (Fig. 8D). Only one protein MROH7 showed significantly (Q < 0.15) higher expression in sperm from PITHD1^{-/-} mice than PITHD1^{+/+} mice, but its role in sperm is unknown (Fig. 8D). On the contrary, seven protein species were detected at significantly (Q <0.15) lower amounts in sperm from $PITHD1^{-/-}$ mice than PITHD1^{+/+} mice (Fig. 8D). Among these seven proteins, ACE, PRKAR1A, and TEX101 are known to be important for the fertilization capability of sperm (26-29). The other four proteins (RPS27A, PRSS39, and the 19S proteasomal regulatory subunits PSMC2 and PSMD1) were also implicated in sperm function (30-34). Therefore, these results suggest that the PITHD1-mediated regulation of proteasomes in testicular cells

including elongated spermatids is required for the quantitative control of proteins important for the fertilization capability of sperm.

Discussion

The present results demonstrate that PITHD1, which is highly expressed in cTECs and the testis, associates with the immunoproteasomes in the testis, but not with α 4s-containing testis-specific proteasomes or cTEC-specific thymoproteasomes. The present results also demonstrate that mice specifically deficient in PITHD1 show male infertility accompanied with morphological abnormalities and impaired motility of sperm. PITHD1 deficiency results in the reduction of proteasome activity and/or abundance in testicular cells including elongated spermatids and the alteration in the amount of proteins important for fertilization capability in sperm. Thus, this study reveals that PITHD1 is a proteasome-interacting protein that plays a nonredundant role in the male reproductive system.

Our proteomic analysis of sperm shows that the amounts of 7 proteins (ACE, PRKAR1A, RPS27A, TEX101, PRSS39, PSMC2,



Figure 8. Alteration of proteasome activity in testicular cells by PITHD1 deficiency. *A*, histograms show the detection of proteasome activity by cellpermeable triple-leucine substrate-based fluorescent probe in testicular cells and sperm of PITHD1^{+/-} mice (*blue line*) and PITHD1^{-/-} mice (*red line*). *Shaded area* and *black line* represent background fluorescence profiles without the addition of proteasome probe in PITHD1^{+/-} mice and PITHD1^{-/-} mice, respectively. Representative data from four to five independent mice measured in four independent experiments are shown. *B*, mean fluorescence intensity (mean ± S.E.) of a triple-leucine substrate-based fluorescent probe in indicated cells in PITHD1^{+/-} mice (*blue symbols*) and PITHD1^{-/-} mice (*red symbols*) from four independent experiments. *, *p* < 0.05. *C*, immunoblot analysis of PITHD1 protein in testis and sperm isolated from cauda epididymis and caput epididymis of PITHD1^{+/+} mice. *β*-Actin was examined as loading control. Representative results of four independent experiments are shown. Graph shows the intensity (mean ± S.E.) of PITHD1 signal detected by immunoblotting relative to *β*-Actin signal. *, *p* < 0.05; **, *p* < 0.01. *D*, volcano plot analysis of proteomes for sperm isolated from PITHD1^{+/+} mice and PITHD1^{-/-} mice (three mice per group) at 8-week-old. Detected proteins are plotted as log₂-fold changes *versus* -log₁₀ Q-values.

and PSMD1), which have all been implicated in male fertilization capability, are reduced by PITHD1 deficiency. It was previously reported that mice deficient in ACE and its specific substrate TEX101 exhibit male infertility due to the impaired ability of sperm to migrate into the oviduct (26-28). The haploinsufficiency of PRKAR1A was reported to cause the fertility defects accompanied with morphological abnormalities of sperm in both mouse and human (29). The association between male infertility and the reduced expression of RPS27A and the 19S regulatory subunit PSMC2 was shown in transcriptomic and proteomic analyses of human sperm (30, 31, 33). It was also suggested that PRSS39 and the 19S regulatory subunit PSMD1 play a role in zona penetration by sperm (32, 34). In this study, we demonstrated that the reduction of proteasome activity and/or abundance by PITHD1 deficiency is detectable in testicular cells including elongated spermatids, rather than sperm. It is therefore conceivable that the regulation of proteasomes by PITHD1 in testicular cells is important for the optimal expression of these 7 proteins in mature sperm.

Our results show that the reduction of proteasome activity and/or abundance in testicular cells is modest in PITHD1-deficient mice. It was recently reported that mice deficient in α 4s, a component of testis-specific proteasome, exhibit infertility in male, and proteins that are normally degraded during spermatogenesis are accumulated in testicular cells in α 4s-deficient mice, although the reduction of proteasome activity is also mild in the testis of these mice (22, 23). It was also reported that proteasome activity *per se* is not reduced in β 5t-deficient cTECs, although β 5t deficiency markedly impairs the positive selection of CD8⁺ T cells in the thymus (9–11, 20). Therefore, the modest reduction of proteasome activity and/or abundance in testicular cells by PITHD1 deficiency may sufficiently affect the function of key proteins that are important for fertilization capability.

Our immunoprecipitation experiments reveal that PITHD1 associates with immunoproteasomes in the testis but not with immunoproteasomes or thymoproteasomes in the thymus. It was previously demonstrated that the proteasome activator PA200 is highly expressed in the testis, and ~90% of proteasomes in the testis contain PA200 at either one or both ends of the 20S proteasomes (35, 36). On the contrary, in our recently published proteomic profiles, PA200 was undetectable in cTECs (20). Therefore, PA200 expression in the testis may support the association of PITHD1 with immunoproteasomes in the testis but not in cTECs. Our results also show that PITHD1 does not associate with α 4s-containing testis-specific protea-

somes. It was previously reported that the α 4s subunit is highly expressed in spermatocytes and spermatids, and the α 4s-containing testis-specific proteasomes contain PA200 (21–23, 36). However, α 4s is incorporated into the constitutive proteasomes and not the immunoproteasomes (21). It is possible that the type of 20S proteasomes, in addition to the association with other proteasome-interacting proteins, is important for the association of PITHD1 with proteasomes.

The role of proteasome-interacting proteins, including PA200 and PA28 γ , in the male reproductive system was studied previously. PA200 is essential for the acetylation-associated degradation of core histones to replace histones with protamines in elongated spermatids, and that is a necessary chromatin remodeling that gives rise to healthy sperm (36). Indeed, mice deficient in PA200 exhibit male subfertility accompanied with the reduced production of healthy sperm (37). It was also reported that PA28 γ -deficient male mice are subfertile due to the reduced production and motility of sperm (38, 39). Furthermore, PA200/PA28 γ double deficiency causes infertility with defect in sperm motility (40). Therefore, it is possible that PITHD1 functions in the male reproductive system coordinately with other proteasome-interacting proteins, including PA200 and PA28 γ .

We further show that the impaired motility and the morphological abnormalities of sperm are caused by PITHD1 deficiency. The morphological abnormalities of sperm from PITHD1-deficient mice could be the cause of the defect in sperm motility. However, \sim 50% of sperm showed normal morphology in PITHD1-deficient mice under macroscopic observation. The motility of sperm is generated by axoneme in flagellum, which consists of two central microtubules surrounded by nine pairs of microtubules (41). Therefore, morphologically normal sperm in PITHD1-deficient mice may indeed have a defect in the ultrastructure of flagellum, including microtubules, which is undetectable under macroscopic observation.

Finally, our results show that PITHD1-tdTomato protein is detected in \sim 3% of cTECs. In the thymus, both cTECs and mTECs consist of diverse subpopulations, and cTECs are heterogeneous in the expression of functionally important molecules, including interleukin-7, delta-like-4, and major histocompatibility complex class II (42). The heterogeneity of cTECs in PITHD1-tdTomato expression newly detected in this study offers an additional diversity in cTECs. Our results also show that PITHD1-deficient mice exhibit no apparent defects in the thymus, including the development of TECs and thymocytes. Our results further denote that the impaired generation of TCR β^{high} CD4⁻CD8⁺ thymocytes in β 5t-deficient thymus is neither further decreased nor recovered by PITHD1 deficiency. These results indicate that PITHD1 plays no apparent roles in the thymus in the presence or absence of the thymoproteasome. However, it is still possible that PITHD1 in the thymus plays a presently unknown role mediated by a minor subpopulation of cTECs.

In conclusion, the present study reveals the nonredundant function of the testis-specific proteasome-interacting protein PITHD1 in the male reproductive system in mouse. PITHD1 protein has been detected in the human testis (The Human Protein Atlas public database). Although the association between PITHD1 and infertility has not been reported in genome-wide association studies, PITHD1 could actually be an additional molecule important in human male fertility. Our findings may provide clues to better understand the molecular mechanisms in the male reproductive system common to mouse and human.

Experimental procedures

Mouse

C57BL/6 mice, B6D2F1 mice, and ICR mice were obtained from SLC. β 5t-deficient mice were described previously (9). All mouse experiments were performed with consent from the Animal Experimentation Committee of Tokushima University (T28-58), from the Animal Care and Use Committee of the Research Institute for Microbial Diseases, Osaka University (H28-08, H30-01), and from the Institutional Animal Care and Use Committee (IACUC) of RIKEN Kobe Branch (A2001-03).

Generation of PITHD1-deficient mice

The targeting vector shown in Fig. 2 was prepared by subcloning *PITHD1*-containing mouse genomic DNA and tdTomatoencoding cDNA (Clontech) into a plasmid containing a pgk-neo cassette. The linearized targeting vector was introduced into HK3i ES cells from C57BL/6N embryos (43). Targeted alleles were screened by genomic PCR analysis and Southern blot analysis. PITHD1⁻ mice (PITHD1^{tm1(tdTomato)}, accession no. CDB1274K) are available to the scientific community. The primers used for genotyping PCR were as follows: WT-F, 5'-ACGCTTGAACGGTACCACTG-3'; WT-R, 5'-GCTGCGA-ACTCAAAAGACCA-3'; tdTomato-R, 5'-GGTCTTGAACT-CCACCAGGT-3'.

Generation of PITHD1-deletion mice

As shown in Fig. 6A, guide RNAs, which recognize sequences in the exon 1 (5'-GCTGTACCTGCGCATCGACC-3) or exon 3 (5'-ACTCACAGTCTCATCTCCGA-3'), were designed using CRISPRdirect web tool. Their cleavage activity was validated by single-strand annealing assay (44). WT zygotes were obtained by mating superovulated B6D2F1 females with B6D2F1 males. The two gRNA/Cas9 complexes were electroporated into two pronuclear zygotes using a super electroporator NEPA21 (NEPA GENE, Chiba, Japan) (45). The zygotes were cultivated in KSOM medium overnight, and the developed two-cell stage embryos were transferred into the oviducts of pseudopregnant ICR females. The F0 pups obtained were genotyped by PCR using primers 1 and 2 for detecting WT allele or primers 1 and 3 for detecting knockout allele. The DNA sequence of the mutant alleles (PITHD1^{Δ} or PITHD1^{em1}) was further confirmed by Sanger sequencing. The crRNA target sequences and primers for genotyping were as follows: Primer-1, 5'-CGCCATGTCGCACGGCCACAG-3'; Primer-2, 5'-GAGCGGTCGGTCCGCTCCTC-3'; Primer-3, 5'-TAGGTG-GGAGTCTCCCGAGG-3'.

Southern blot analysis

Genomic DNA extracted from the liver was digested with BamHI, electrophoresed in 1% agarose, and transferred to

nylon membrane (Hybond-N⁺, GE Healthcare). Probes were labeled with a PCR DIG Probe Synthesis Kit (Roche), and hybridization was detected using anti-DIG-AP Fab fragment (Roche Applied Science), CDP-STAR (Roche Applied Science), and Light Capture II (Atto). The primers used for Southern blot analysis were as follows: forward, 5'-AGCTTGTCTGTCT-CTGTGGA-3'; reverse, 5'-CAGTGTCTGTAAGTCATG-GAGC-3'.

Quantitative RT-PCR analysis

Total cellular RNA was reverse-transcribed (RT) with Prime-Script Reverse Transcriptase (TaKaRa). Quantitative real-time PCR was performed using SYBR Premix Ex Taq (TaKaRa) and a StepOnePlus Real-Time PCR System or a 7900HT Sequence Detection System (Applied Biosystems). The primers used were as follows: *Pithd1*, 5'-CCGTTTACGGGCAATGT-CAA-3' and 5'-CATAGTTGCAGATGGTCACC-3'; *Gapdh*, 5'-CCGGTGCTGAGTATGTCGTG-3' and 5'-CAGTCTTC-TGGGTGGCAGTG-3'; *Hprt*, 5'-GCTGGTGAAAAGGACC-TCT-3' and 5'-CACAGGACTAGAACACCTGC-3'. Amplified products were confirmed to be single bands in gel electrophoresis and normalized to the amount of *Gapdh* amplification products.

Immunoprecipitation

Thymic cells were isolated from newborn mice. Testicular cells were isolated from 6-week-old mice. The cells were lysed in ice-cold lysis buffer containing 0.2% Nonidet P-40, 25 mM Tris-HCl, pH 7.5, 1 mM DTT, 2 mM ATP, and 5 mM MgCl₂, and clarified by centrifugation at 20,000 × g for 15 min at 4 °C. For immunoprecipitation, anti- β 5t antibody (MBL International) (20), anti- β 5i antibody (Proteintech) (46), anti- α 4s antibody (Proteintech) (22), or anti- α 6 antibody (Bethyl Laboratories) was bound to Protein G-Sepharose beads (GE Healthcare) and mixed with the cell lysates at 4 °C for 2 h. The beads were washed with lysis buffer, and immunoprecipitated molecules were eluted with 2× SDS sample buffer containing 125 mM Tris-HCl, pH 6.5, 4% SDS, 20% glycerol, 0.2 M DTT, and 0.01% bromphenol blue.

Immunoblot analysis

The immunoprecipitated molecules or the cell lysates were boiled with SDS sample buffer, separated by SDS-PAGE, and transferred to a polyvinylidene difluoride membrane. Membranes were probed with anti-PITHD1 antibody (Sigma-Aldrich), anti- β 5t antibody (MBL International) (20), anti- β 5i antibody (Proteintech) (46), anti- α 4s antibody (Proteintech) (22), anti- α 6 antibody (Bethyl Laboratories), or anti- β -actin antibody (Cell Signaling Technology) (47), and then with secondary antibodies conjugated to horseradish peroxidase (Santa Cruz). Signals were detected with ECL reagent (GE Healthcare) and Image analysis system (Atto).

Flow cytometric analysis

For the analysis of TECs, minced thymus were digested with 0.5 unit/ml of Liberase (Roche Applied Science) in the presence of 0.02% DNase I (Roche). Single-cell suspensions were stained with antibodies specific for EpCAM (CD326, BioLegend, clone

G8.8), CD45 (BioLegend, clone 30-F11), and Ly51 (CD249, Bio-Legend, clone 6C3), and for the reactivity with UEA1 (Vector Laboratories). For the analysis of thymocytes and spleen cells, cells were stained with antibodies specific for CD3 (BioLegend, clone 145-2C11), CD4 (BioLegend, clone RM4-5), CD8 α (Invitrogen, clone 5H10), TCR β (BioLegend, clone H57), CD25 (Bio-Legend, clone PC61), CD44 (eBioscience, clone IM7), and CD69 (eBioscience, clone H1.2F3). Multicolor flow cytometry was performed on FACSVerse.

Proteasome activity assay

Surface-stained thymus cells, testicular cells, and sperm were incubated with 1 μ M cell-permeable triple-leucine substratebased fluorescent probe (Me₄BodipyFL-Ahx₃Leu₃ VS, Boston Biochem), which shows high affinity binding to the chymotrypsin-like catalytic site of proteasomes, at 37 °C for 60 min, and then washed with FACS buffer. Where indicated, the cells were incubated at 37 °C for 1 h in the presence of the proteasome inhibitor MG132 (AdooQ Bioscience) prior to the incubation with the proteasome activity probe. Proteasome activity was measured by flow cytometry.

Tissue analysis

Frozen thymus, testis, and epididymis embedded in OCT compound (Sakura Finetek) were sliced into $10-\mu$ m-thick sections. Tissue sections stained with hematoxylin and eosin were examined under a light microscope. For immunofluorescence analysis, the thymus and the testis were fixed in 4% (g/volumn) paraformaldehyde and embedded in OCT compound. Frozen tissues were sliced into $10-\mu$ m-thick sections. The thymic sections were stained with anti- β 5t antibody, anti-Ly51 antibody, anti-Aire antibody, and UEA-1, followed by Alexa Fluor-conjugated secondary reagents. The testis sections were stained with DAPI. Images were analyzed under a TCS SP8 confocal laser-scanning microscope (Leica).

In vitro fertilization assay

Sperm were placed in 1 drop of TYH medium, covered with paraffin oil (Nacalai Tesque), and incubated at 37 °C for 2 h. Eight-week-old C57BL/6 female mice were injected with 5 units of pregnant mare serum gonadotropin 62 h before *in vitro* fertilization and with 5 units of human chorionic gonadotropin 14 h before *in vitro* fertilization. Eggs were collected from the oviduct. In some cases, eggs were treated with 330 g/ml of hyaluronidase (Sigma-Aldrich) or 1 mg/ml of collagenase (Sigma-Aldrich) at 37 °C for 10 min to prepare cumulus-free or zona-free eggs, respectively. The eggs were placed in 1 drop of TYH medium. The eggs were incubated with 2.0×10^5 sperm for 6 h at 37 °C, washed with TYH medium, further incubated for 18 h, and observed under a Hoffman modulation contrast microscope (IX71; Olympus).

Sperm motility assay

To assess sperm motility, sperm in the TYH medium at 37 °C were allowed to swim out from the cauda epididymis for 10 or 120 min and were placed on standard count 2-chamber slides (Leja) (depth 100 μ m) that had been preheated to 37 °C. Sperm motility parameters (VCL, VSL, and VAP) were evaluated by a

computer-assisted system (CEROS II sperm analysis system, software version 1.5.2; Hamilton Thorne Biosciences).

Proteomics analysis

Sperm isolated from $PITHD1^{+/+}$ mice and $PITHD1^{-/-}$ mice in triplicate were lysed in 150 μ l of 6 M guanidine HCl containing 100 mM Tris-HCl, pH 8.0, and 2 mM DTT. The lysates were dissolved by heating and sonication, followed by centrifugation at 20,000 \times g for 15 min at 4 °C. The supernatants containing 100 μ g of protein each were reduced in 5 mM DTT at room temperature for 30 min and alkylated in 27.5 mM iodoacetamide at room temperature for 30 min in the dark. Proteins were purified by methanol/chloroform precipitation and solubilized with 25 μ l of 0.1% RapiGest SF (Waters) in 50 mM triethylammonium bicarbonate buffer. After repeated sonication and vortexing, the proteins were digested with 1 μ g of trypsin/Lys-C mix (Promega) for 16 h at 37 °C. Peptide concentration was determined using Pierce Quantitative Colorimetric Peptide Assay (Thermo Fisher). Approximately 25 μ g of peptides for each sample was labeled with 0.2 mg of TMT6plex reagents (Thermo Fisher) for 1 h at room temperature. After the reaction was quenched with hydroxylamine, all the TMT-labeled samples were pooled, acidified with trifluoroacetic acid (TFA), and fractionated using a Pierce High pH Reversed Phase Peptide Fractionation Kit (Thermo Fisher). Nine fractions were collected using 5, 10, 12.5, 15, 17.5, 20, 22.5, 25, and 50% acetonitrile (ACN) in 0.1% triethylamine. Each fraction was evaporated in a SpeedVac concentrator and dissolved in 0.1% TFA.

LC-MS/MS analysis of the resultant peptides (1 μ g each) was performed on an EASY-nLC 1200 UHPLC connected to a Q-Exactive Plus mass spectrometer through a nanoelectrospray ion source (Thermo Fisher). The peptides were separated on a 75- μ m inner diameter \times 150-mm C18 reversedphase column (Nikkyo Technos) with a linear gradient of 4-20% ACN for 0-180 min and 20-32% ACN for 180-220 min, followed by an increase to 80% ACN for 220-230 min. The mass spectrometer was operated in the data-dependent acquisition mode with a top 15 MS/MS method. MS1 spectra were measured at the resolution of 70,000, the automatic gain control target of 3×10^6 , and a mass range of 375 to 1,400 m/z. HCD MS/MS spectra were acquired at the resolution of 35,000, the automatic gain control target of 1×10^5 , the isolation window of 0.4 m/z, the maximum injection time of 100 ms, and the normalized collision energy of 32. Dynamic exclusion was set to 30 s. Raw data were directly analyzed against Swiss-Prot database version 2017-10-25 restricted to Mus musculus (25,097 sequences) using Proteome Discoverer version 2.2 (Thermo Fisher) with Mascot search engine version 2.5 (Matrix Science) for identification and TMT quantification. The search parameters were (a)trypsin as an enzyme with up to two missed cleavages, (b) precursor mass tolerance of 10 ppm, (c) fragment mass tolerance of 0.02 Da, (d) TMT of lysine and peptide N terminus and carbamidomethylation of cysteine as fixed modifications, and (e) acetylation of protein N terminus and oxidation of methionine as variable modifications. Quantification was performed by using the Reporter Ion Quantification node of the Proteome Discoverer software. Peptides and

proteins were filtered at the FDR of 1% using the percolator node and the protein FDR validator node, respectively.

Statistical analysis

Statistical significance was assessed using the two-tailed unpaired Student's *t* test with Welch's correction for unequal variances. Q value was calculated using the multiple unpaired *t* test with the false discovery rate approach (48). The data are presented as mean \pm S.E. of at least three independent experiments.

Data availability

MS proteomics data have been deposited to the Proteome-Xchange Consortium via jPOST partner repository with the dataset identifier PXD015347 (49).

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