Dido mutations trigger perinatal death and generate brain abnormalities and behavioral alterations in surviving adult mice

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Nearly all vertebrate cells have a single cilium protruding from their surface. This threadlike organelle, once considered vestigial, is now seen as a pivotal element for detection of extracellular signals that trigger crucial morphogenetic pathways. We recently proposed a role for Dido3, the main product of the death inducer-obliterator (*dido*) gene, in histone deacetylase 6 delivery to the primary cilium [Sánchez de Diego A, et al. (2014) *Nat Commun* 5:3500]. Here we used mice that express truncated forms of Dido proteins to determine the link with cilium-associated disorders. We describe *dido* mutant mice with high incidence of perinatal lethality and distinct neurodevelopmental, morphogenetic, and metabolic alterations. The anatomical abnormalities were related to brain and orofacial development, consistent with the known roles of primary cilia in brain patterning, hydrocephalus incidence, and cleft palate. Mutant mice that reached adulthood showed reduced life expectancy, brain malformations including hippocampus hypoplasia and agenesis of corpus callosum, as well as neuromuscular and behavioral alterations. These mice can be considered a model for the study of ciliopathies and provide information for assessing diagnosis and therapy of genetic disorders linked to the deregulation of primary cilia.

ciliopathies | brain patterning | perinatal lethality

The primary cilium is a unique, mainly nonmotile, microtubule-based organelle found in nearly all noncycling cells in vertebrates. Its main function is to transduce extracellular signals that regulate a wide range of functions from fluid flow to cell proliferation, differentiation, and migration; hence, it alters cell polarity and tissue development. Defects in cilia growth, resorption, or stability lead to deregulation of these pathways, which results in a number of functional defects (ciliopathies) that affect diverse organs, giving rise to complex pleiotropic phenotypes such as those of Joubert, Meckel-Gruber, or Bardet-Biel syndromes (1). In many cases, these pathologies are associated with embryonic or perinatal lethality (2). Mechanically or chemically stimulated signaling pathways are associated with the cilium; hedgehog (Hh), canonical and noncanonical Wnt, Notch, fibroblast growth factor, and platelet-derived growth factor are some of the pathways involved in development and are altered in organisms with dysfunctional cilia (3, 4).

The cognitive impairment found in human ciliopathies is manifested as behavioral changes in the mouse. Anatomical changes in the brain are usual in both species, reflecting alterations in central nervous system (CNS) development (2) that include defects in neural tube patterning and closure (5, 6) and in hippocampal neurogenesis, leading to neuropsychiatric phenotypes (7, 8). All these processes are closely linked to Hh signaling, and most CNS-related ciliopathies appear to be associated with altered Hh signaling (9).

Morphogenetic activity of sonic hedgehog (Shh) in the neural tube requires coordinated antagonistic expression of bone morphogenetic protein (BMP) (10, 11) and involves a physical interaction between Smad and Gli proteins (12). Death inducer-obliterator (*dido*) is a BMP4-specific Smad-regulated target gene (13), and we recently demonstrated a role for Dido3, the main *dido* product, in actin-dependent histone deacetylase 6 (HDAC6) delivery to the primary cilium (14). HDAC6 counteracts the activity of α-tubulin acetyl transferase (Atat1) (15), which acetylates tubulin and thus stabilizes primary cilium structure. Recent studies also established a relationship between HDAC and tumors as well as neurological and immunological diseases. HDAC6 in particular appears to be involved in mood control, in tau-driven neurological disorders, in the progression of Alzheimer's and Huntington's diseases, in immune synapse formation, and in regulatory T-cell homeostasis (16).

In humans and mice, the *dido* locus encodes three isoproteins, Dido1, Dido2, and Dido3, by alternative splicing (17). We reported generation of a mouse mutant that expresses an N-terminal–truncated Dido protein that lacks the initial 422 amino acids (Fig. S1). Studies in homozygous $dido^{\Delta Nt/\Delta Nt}$ mice on a mixed genetic background showed an essential role for Dido3 in the regulation of the spindle assembly checkpoint (SAC), control of centrosome number, chromosome stability, and cytokinesis. Pups were nonetheless born at Mendelian frequencies and reached weaning with no obvious anatomical or behavioral abnormalities,

Significance

The primary cilium is an organelle protruding from most postmitotic vertebrate cells. A growing body of data supports the crucial role of primary cilia in developmental signaling pathways. Recent studies describe the main stages in ciliogenesis at the morphological level and components of some of the mechanisms involved, including the selective acetylation of tubulin. How this acetylation is modulated in cilia nonetheless remains poorly understood. Here we show that the death inducer-obliterator (*dido*) gene product, which regulates histone deacetylase 6 deacetylase activity, is necessary for orofacial development in the mouse embryo and influences brain patterning and neuromuscular activity. Mice deficient in dido function present neonatal mortality and various ciliopathies including cleft palate and hydrocephalus, as well as hippocampal and commissural dysplasia.

Author contributions: R.V. and C.M.-A. designed research; R.V., J.G., A.F., V.T., and F.G.d.B. performed research; R.V. and J.G. analyzed data; and R.V. and C.M.-A. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission.

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This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1419300112/-/DCSupplemental.

NEUROSCIENCE

although they developed a myelodysplasia/myeloproliferative-like syndrome (MDS/MPD) (17).

We also generated a *dido* mouse mutant that lacks exon 16, which encodes the Dido3 isoform-specific 1080 C-terminal amino acids. This $dido3^{\Delta Ct}$ allele is lethal in homozygosis at Theiler embryonic stage ts10–11 and is associated with DNA damage, apoptosis, and growth arrest during embryonic stem (ES) cell differentiation in vitro (18). The $dido^{\Delta Nt}$ allele rescued embryonic lethality of the $dido3^{\Delta Ct}$ mutation as well as the ability of $dido3^{\Delta Ct}$ mutant mouse ES cells to differentiate (18). During these studies, we identified a synthetic phenotype in the $dido^{\Delta Nt/\Delta Ct}$ double heterozygotes that drives severe perinatal lethality. Here we describe this phenotype, which identifies an important role for the *dido* gene in craniofacial development. The few $dido^{\Delta Nt/\Delta Ct}$ mice that reached adulthood had metabolic and neuromuscular alterations that highlight *dido* involvement in many physiological functions, including behavioral abnormalities.

Results

Dido Function Is Necessary in Vivo for Primary Cilium Control. We previously reported the ability of the $dido^{\Delta Nt}$ allele to rescue the embryonic lethality of the $dido3^{\Delta Ct}$ mutation (18). In addition, we described dissociation between Dido function in stem cell differentiation and its known role in the SAC, centrosome amplification, and cytokinesis (17, 19).

We also previously identified the role of Dido3 in the control of cilium size. To analyze the phenotype of the $dido^{\Delta Nt}$ allele-rescued embryonic lethality of the $dido3^{\Delta Ct}$ mutation in vivo, we crossed $dido^{\Delta Nt/+} \times dido^{\Delta Ct/+}$ mice. To test the relevance of normal Dido activity in cilia development and maintenance, we quantified primary cilia in the hippocampal granular cell layer of adult mouse brain. Adenylyl cyclase III (ACIII) is a prominent ciliary marker (20); to compare brain sections from 3-mo-old wild type (WT) and $dido^{\Delta Nt/\Delta Ct}$ mice, we used anti-ACIII staining and confocal microscopy (Fig. 1 *A* and *B*). The percentage of ciliated cells was reduced in $dido^{\Delta Nt/\Delta Ct}$ mice (Fig. 1*C*), although mean cilium length did not differ significantly (Fig. 1*D*). Distribution of cilium length is altered in $dido^{\Delta Nt/\Delta Ct}$ brains, with increased frequency of longer and shorter cilia (Fig. 1*E* and Fig. S2).

Perinatal Lethality in $dido^{\Delta Nt/\Delta Ct}$ Mice. To analyze the phenotype of the $dido^{\Delta Nt}$ allele-rescued embryonic lethality of the $dido3^{\Delta Ct}$ mutation in vivo, we crossed $dido^{\Delta Nt/+} \times dido^{\Delta Ct/+}$ mice. After a 1-y follow-up of the colony, $dido^{\Delta Nt/\Delta Ct}$, $dido^{\Delta Nt/+}$, $dido^{\Delta Ct/+}$, and WT genotypes appeared at frequencies of 2.0, 37.2, 30.7, and 30.2% at weaning, respectively (Fig. 2*A*), indicating selective loss of most double-heterozygous mice ($P < 10^{-5}$). Genotype analysis at various days postcoitum (dpc) of embryos in a $dido^{\Delta Nt/+} \times dido^{\Delta Ct/+}$ cross showed expected Mendelian frequencies up to birth (Fig. 2*B*), but most $dido^{\Delta Nt/\Delta Ct}$ mice died shortly thereafter.

Neonatal death after 24 h postpartum is usually associated with inability to suckle rather than with respiratory failure or homeostatic deficiencies (21). We found no gross morphological alterations or hemorrhage in $dido^{\Delta Nt/\Delta Ct}$ pups, nor were there obvious breathing defects in most $dido^{\Delta Nt/+} \times dido^{\Delta Ct/+}$ litters during the first 24 h after birth, although some $dido^{\Delta Nt/\Delta Ct}$ mice showed cyanosis by days 1 and 2 after birth.

Another cause of perinatal lethality is linked to macroautophagy defects (22). As a cilium-associated protein, Dido3 is potentially linked to the autophagosome (23); knockout mice for another cilium-associated protein, autophagy-related 5 (Atg5), show shortened primary cilia (24). We tested the Dido/autophagosome relationship by monitoring microtubule-associated protein 1 light chain 3B (LC3B). Cytosolic LC3B is processed for autophagosome formation; Apg7p and Apg3p activate cytosolic LC3B-I, which is phospholipid conjugated and forms membrane-bound, autophagosome-associated LC3B-II (25). We studied WT and $dido^{\Delta Nt/\Delta Ct}$ embryos obtained by cesarean section at 19 dpc (full term). At time 0 and after 6 h in starvation conditions, heart protein was isolated and analyzed in Western blot for LC3BI/II expression. We found no differences between WT and mutant embryos at any time (Fig. 3*A*). Adult primary fibroblasts and EBV-immortalized murine embryonic fibroblasts (MEFs) cultured in starvation medium showed no LC3B processing defect, either in primary lung (Fig. 3*B*) or in immortalized fibroblasts (Fig. 3*C*).

$dido^{\Delta Nt/\Delta Ct}$ pups showed no size differences compared with WT or heterozygous littermates at birth (Fig. 4*A*), whereas those that survived >24 h showed reduced body size, suggesting a nutrition defect (Fig. 4*B*). Mice were inspected visually through the semitransparent skin; from ∼12 h postpartum onwards all $dido^{\Delta Nt/\Delta Ct}$ mice showed little or no milk in the stomach (Fig. 4 *A* and *B*), which was dilated and air filled (Fig. 4*C*). A possible glucose mobilization deficiency was ruled out by glycemia measurement and by estimation of glycogenolysis activity by RT-PCR of hepatic glucose-6-phosphatase (Fig. S3).

The few $dido^{\Delta Nt/\Delta Ct}$ mice that reached adulthood showed no external morphological abnormalities, except that size and weight were approximately one-half to two-thirds that of WT or heterozygous littermates (Fig. 4*D*), and life expectancy at birth was <1 y (Fig. 4*E*). Adults were infertile and showed limited cachexia. Hemograms, blood biochemistry profiles, and proteinograms of $dido^{\Delta Nt/\Delta Ct}$ mice showed normal values, with only slight hypoproteinemia compared with those of littermates (Fig. S4). Histochemical studies showed anemia in some individuals as assessed by the Turnbull blue reaction (Fig. 4*F*), and visual examination showed differences in body fat accumulation (Fig. 4*G*), implying metabolic problems. Various nonrecurrent tumors were occasionally found in necropsies, which suggests nonspecific activity that facilitates tumor growth.

Delayed Closure of the Palate Primordium During $dido^{\Delta Nt/\Delta Ct}$ Embryo Development. The snout of some $dido^{\Delta Nt/\Delta Ct}$ fetuses was shorter

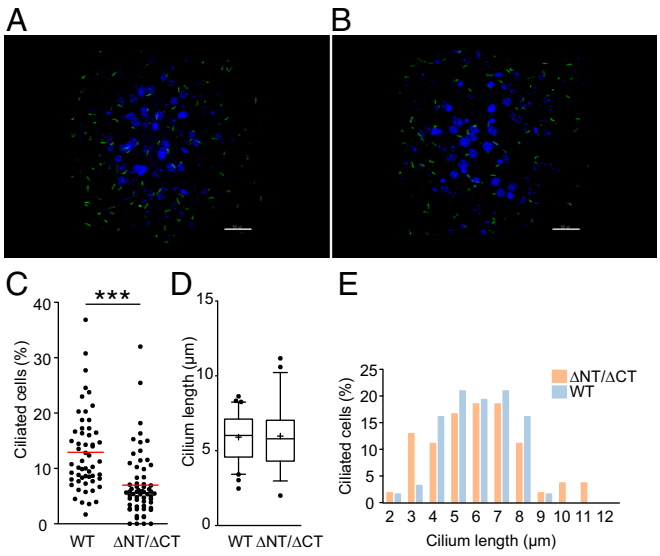

Fig. 1. Quantification of primary cilia in the adult hippocampal granular cell layer. (*A* and *B*) Immunofluorescent visualization of ACIII-positive primary cilia (green) of WT (*A*) and $dido^{\Delta NT/\Delta CT}$ (*B*) mice (z-projection of confocal stack; DAPI-stained nuclei in blue) (Scale bar, 30 μm.) (*C*) $dido^{\Delta NT/\Delta CT}$ genotype is associated with a lower percentage of ciliated cells. Cilia were counted manually in 30 microscopy fields of 5-μm-thick brain sections from two WT and two $dido^{\Delta NT/\Delta CT}$ mice. Individual values are shown; means are calculated for each genotype. Unpaired Student's *t* test with Welch's correction, ***$P < 0.001$. (*D* and *E*) Length of individual cilia for WT ($n = 62$) and $dido^{\Delta NT/\Delta CT}$ mice ($n = 54$) were determined (*Materials and Methods* and Fig. S2). (*D*) The median value and the 50th (boxes) and 10–90th percentile range (whiskers) are shown. Student's *t* test indicates no significant differences. (*E*) Distribution of cilia length in $dido^{\Delta NT/\Delta CT}$ brain is different from WT. Snedecor's F distribution, $P = 0.01$.

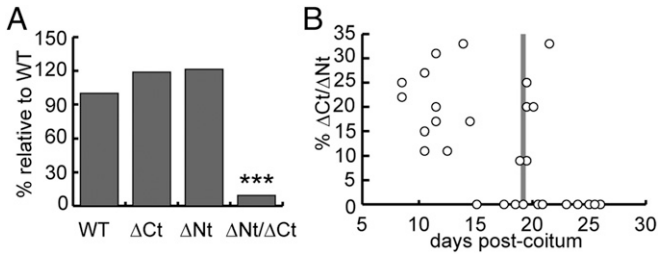

Fig. 2. Viability of *dido*$^{\Delta NT/\Delta CT}$ mutant mice. (*A*) The N- and C-terminal Dido3 deletions studied have no effect on viability when each allele is in heterozygosis, and only a few *dido*$^{\Delta NT/\Delta CT}$ mutant mice reached weaning. Relative frequency of each genotype is shown ($n = 409$); the χ^2 test was used to compare frequency for each group frequency with that of WT. ***$P < 0.001$. (*B*) Double heterozygotes did not show imbalanced proportions during the embryonic or perinatal periods, but dropped abruptly by 24–48 h postpartum (pp). Genotype results are shown for 15 litters between 9 and 19 dpc, 3 litters at ~24 h pp, and 7 litters between 2 and 7 d (d)pp. Mean litter size ± SD was 10.47 ± 2.0.

than in WT mice (Fig. 5*A*). We dissected the oral cavity of *dido*$^{\Delta Nt}$ × *dido*$^{\Delta Ct}$ F1 pups at 19.5 dpc (full-term pregnancy) to expose tongue, throat, and palate. The secondary palate was 15% shorter in *dido*$^{\Delta Nt/\Delta Ct}$ than in WT mice (Fig. 5 *B* and *C*), in agreement with the consistent lack of ruga 7b (the last-formed ruga in normal development), indicative of abnormal palate growth. The few double-heterozygous *dido*$^{\Delta Nt/\Delta Ct}$ mice that survived to adulthood showed normal palatal rugae number and palate size (Fig. S5), again suggesting defective palate development as a cause of perinatal death.

To determine the origin of the craniofacial and palate defects in newborn *dido*$^{\Delta Nt/\Delta Ct}$ mice, we analyzed palate features in *dido*$^{\Delta Nt}$ × *dido*$^{\Delta Ct}$ F1 embryos. At 15.5 dpc, the palatal shelves had already fused in WT embryos, whereas the mouth roof in *dido*$^{\Delta Nt/\Delta Ct}$ embryos remained open, with some degree of variability at this age (Fig. 5*D*). No palatal clefts were observed in *dido*$^{\Delta Nt/\Delta Ct}$ neonates, however, suggesting that the mutation causes delayed growth that gives rise to a shorter, defectively ossified palate.

Brain and Neurobehavioral Abnormalities in Surviving Adult dido$^{\Delta Nt/\Delta Ct}$ Mice. Previous reports of *dido* expression in the CNS, in embryo brain (www.emouseatlas.org/emage/, EMAGE:2631) and adult cerebellum (Allen Mouse Brain Atlas, mouse.brain-map.org/, experiment 69262318), and in gray matter of the spinal cord (Allen Mouse Spinal Cord Atlas, mouse.brain-map.org/, experiment 100014937), prompted us to analyze the effect of *dido* mutations on CNS development and neuron function. Histological examination of the brain showed enlarged ventricles with proportional reduction of caudoputamen and lateral striatum, agenesis of the corpus callosum, and hippocampus dysplasia (Fig. 6*A*).

Whereas WT mice displayed a normal grasping reflex, *dido*$^{\Delta Nt/\Delta Ct}$ mice normally stretched their forelimbs toward the wires and gripped them effectively, but tucked in their hind limbs and maintained them clasped.

Spontaneous locomotor activity of the mice was measured in a chamber equipped with infrared sensors. There was no notable difference in horizontal activity, defined as exploratory movements around and across the base of the chamber (Fig. 6*B*); in contrast, *dido*$^{\Delta Nt/\Delta Ct}$ mice showed reduced vertical activity (Fig. 6*C*), which assesses exploratory movements while standing upright, usually on the walls of the chamber. Pawprint tests along a covered straight path showed no marked gait or stride differences between *dido*$^{\Delta Nt/\Delta Ct}$ and WT mice. To further test sensorimotor performance, we used a thermal plantar test to assess adult *dido*$^{\Delta Nt/\Delta Ct}$ and WT mice. Latency to paw withdrawal following plantar heating with an infrared beam was similar in the two mouse groups (Fig. 6*D*); results were similar at different heat potencies, as well as after hand or foot stimulus. The results on the whole suggested mild distal motor neuropathy.

Differences in α-Tubulin Deacetylation of the Sciatic Nerve in dido$^{\Delta Nt/\Delta Ct}$ Mutant Mice. Sciatic nerves were dissected from thighs of *dido*$^{\Delta Nt/\Delta Ct}$ and WT mice of ages between 7 and 15 mo. A sample of each nerve was processed for histological study and another used to prepare whole protein extracts. Axon number and average myelin thickness were determined (Fig. S6); no differences were found in axon density or myelin integrity (Fig. 6*E*). Relative amounts of total and acetylated α-tubulin were evaluated by Western blot (Fig. S7). In age-matched mice, we found a significant difference in the amount of acetylated α-tubulin normalized to total amounts; each *dido*$^{\Delta Nt/\Delta Ct}$ mouse had lower levels of acetylated α-tubulin than an age-matched WT littermate (Fig. 6*F*). No significant differences were observed in total α- + β-tubulin levels between mutant and WT mice.

Discussion

Dido3 is a key determinant of cilium size (14), and Dido3 mutations cause chromosome segregation defects, meiosis prophase alterations, and stem cell differentiation blockade (17, 19, 26, 27). Misregulation of the *dido* gene is linked to increased incidence of genomic instability, myelodysplasia, and myeloproliferation, melanoma, and infertility (13, 17, 26, 28). Elucidating the mechanisms that govern the complex activities could help clarify the role of Dido in promoting stem cell differentiation, cilia size, tumorigenesis and, as shown here, life expectancy, brain development, and neural behavior alterations.

Homozygous *dido*$^{\Delta Nt/\Delta Nt}$, heterozygous *dido*$^{\Delta Nt/+}$, and *dido*$^{\Delta Ct/+}$ mice are viable, although mice of the first two genotypes occasionally develop MDS/MPD. We anticipated *dido*$^{\Delta Nt}$ complementation of the severe early embryonic lethality associated with the *dido*$^{\Delta Ct}$ allele in homozygosis (18), but low *dido*$^{\Delta Nt/\Delta Ct}$ frequency in weaned offspring suggested complementation deficiencies. This lack of complementation affects ciliogenesis in vivo, as noted by abnormal cilium size and numbers in the adult brain.

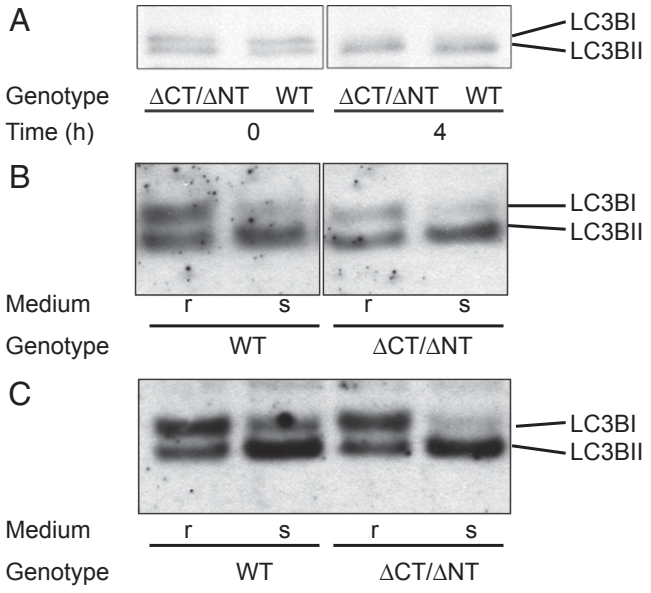

Fig. 3. Autophagy test in primary cells, cell lines, and fetuses. (*A*) Fetuses at 19.5 dpc were starved for 4 h and killed. Heart protein extracts were analyzed by Western blot and by LC3BI and LC3BII detection. The LC3BII/LC3BI ratio was quantified and showed no differences. (*B*) Adult lung primary fibroblasts and (*C*) immortalized embryonic fibroblasts were cultured in rich (r) or starvation (s) medium and analyzed as in *A*; no LC3B processing deficiency was found. All experiments were repeated twice.

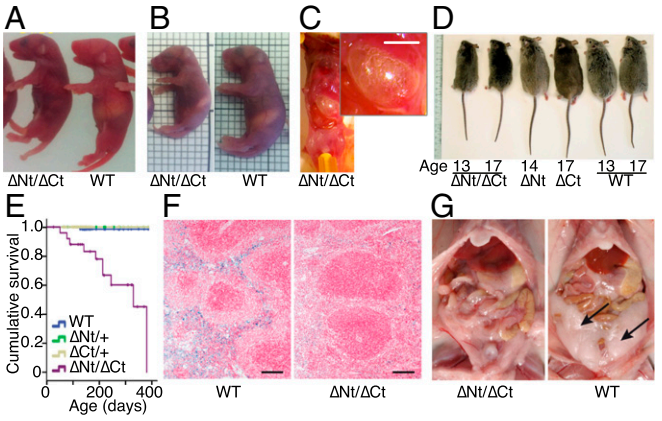

Fig. 4. Phenotypic alterations associated with $dido^{\Delta NT/\Delta CT}$. (*A*) At 0.5 dpp, mutant pups were similar in size to WT littermates, but their stomachs contained little or no milk. (*B*) At 1.5–2 dpp, the size of surviving mutant pups was approximately two-thirds that of WT littermates, with very little milk in the stomach. Grid lines = 2 mm. (*C*) $dido^{\Delta NT/\Delta CT}$ mutant pups showed accumulated air in the stomach. A peritoneal incision to expose the stomach shows inflation due to excess air pressure. (*Inset*; Scale bar, 5 mm.) (*D*) Reduced body size was maintained throughout life. At 13–17 mo of age, weight of $dido^{\Delta NT/\Delta CT}$ mice was about one-half that of littermates, whereas $dido^{\Delta Ct/+}$ and $dido^{\Delta Nt/+}$ showed no size differences compared with WT mice. (*E*) Life expectancy was greatly reduced for $dido^{\Delta NT/\Delta CT}$ mice, again with no effect for $dido^{\Delta Ct/+}$ and $dido^{\Delta Nt/+}$ genotypes. (*F*) Turnbull's blue stain shows depletion of ferrous iron (II) storage in spleen of $dido^{\Delta NT/\Delta CT}$ compared with WT mice. (Scale bar, 100 μm.) Original magnification, 6×. (*G*) Abdominal fat tissue (arrows) is virtually absent in 9-mo-old female $dido^{\Delta NT/\Delta CT}$.

Normal mouse pups gain 50–70% body weight from birth to day 2 (29), which highlights the importance of efficient nutrition in the first 48 h. We thus infer that the primary cause of death of $dido^{\Delta Nt/\Delta Ct}$ pups is an inability to feed adequately, although the metabolic anomalies detected in adult mutants could contribute to lethality. Inadequate feeding could be due to the mechanics of suckling itself, but also to deficient recognition or processing of pheromonal and/or complex odor signals (30). Although primary cilia in olfactory ganglion cells are reported to be necessary for pheromone detection (31), we suspect that the inability of $dido^{\Delta Nt/\Delta Ct}$ mice to feed is a consequence of their craniofacial abnormalities or/and neurological behavior defects.

Palatal shelves grow horizontally and fuse to each another by 15.5 dpc (32). At this age, $dido^{\Delta Nt/\Delta Ct}$ fetuses showed a marked delay in this process, albeit with considerable individual variation. Because of its frequency in man, the genetic basis of cleft palate has been studied extensively. Although the list of genes with a role in this condition is long (33), to our knowledge it does not include *dido*. Major alterations in Dido proteins might be too deleterious, as for the homozygous $dido^{\Delta Ct}$ condition in mice, and only specific allelic combinations on a given genetic background would result in this ciliopathy. It is also possible that *dido* does not primarily drive human cleft palate-like syndromes, but mediates the effects of other genes. Alterations in BMP are among those most often found in orofacial malformations (34); Braig and Bosserhoff (13) reported that *dido* is a notable mediator of BMP downstream effects. The *dido* gene is a target of the BMP-dependent Smad family of transcriptional activators, which are implicated directly in experimental cleft palate induced by all-*trans* retinoic acid.

Dido also regulates expression of integrin αV (13), which probably has a role in correct precursor cell migration from the neural crest of the developing embryo toward the zone that becomes the jaw primordium. Mouse *dido* expression correlates negatively with body weight and jaw length in several strains (35, 36). Further work is needed to determine how Dido mutations act to produce the defects in mouse palate development, as well as the mechanisms by which *dido* mediates or prompts action in palate development by molecules such as BMP or integrin αV. Specific focus on human *Dido* gene status in studies of cleft palate etiology will help to elucidate its clinical relevance.

The few surviving adult $dido^{\Delta Nt/\Delta Ct}$ mice showed hindlimb dyskinesia. As clasping behavior is common in mouse models of various human central and peripheral neuromuscular pathologies, we analyzed the possibility of a neuromuscular system disorder in $dido^{\Delta Nt/\Delta Ct}$ mice. Equilibrium and neuromuscular tests (tightrope walking, rotarod running, or grip strength) did not indicate consistent differences, arguably due to the disparity in body weight between mice of each genotype. It is unclear whether the reduced vertical activity observed in $dido^{\Delta Nt/\Delta Ct}$ mice is the result of peripheral neuropathy, a muscular defect, or both, but it coincides with clasping behavior to imply spinal and/or hindlimb neuromuscular disease. If so, it would be essentially a

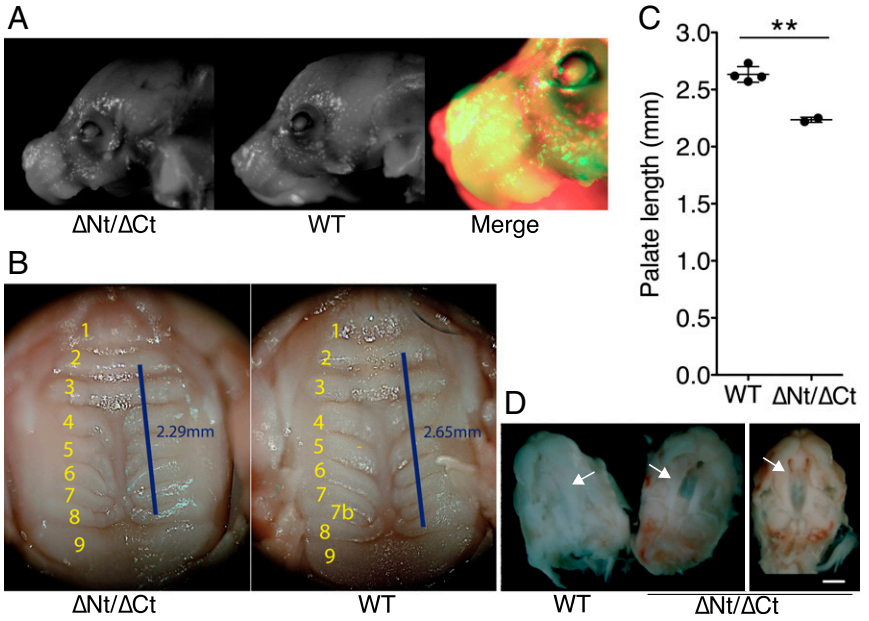

Fig. 5. Craniofacial defects in $dido^{\Delta NT/\Delta CT}$ neonates. (*A*) $dido^{\Delta NT/\Delta CT}$ mice showed altered craniofacial development at birth, with a shortened snout compared with WT littermates. Representative images of both genotypes are shown, with a false-color merged image for comparison. (*B* and *C*) Palate was fully closed at birth, but the distance between rugae 2 and 8 was 15% shorter in $dido^{\Delta NT/\Delta CT}$ than in WT pups (WT, $n = 4$; $dido^{\Delta NT/\Delta CT}$, $n = 2$; Student's t test $P = 0.0016$). (*D*) Delayed palate closure in $dido^{\Delta NT/\Delta CT}$ embryos. At 15.5 dpc, embryos were extracted and the lower jaw and tongue removed to expose the developing palate. The secondary palate was fully closed in WT embryos (*Left*), whereas $dido^{\Delta NT/\Delta CT}$ embryos showed varying degrees of delay in palate closure, with partial (*Center*) or almost no (*Right*) horizontal growth of palatal halves (arrows) toward the midline at the time analyzed ($n = 8$). (White scale bar, 5 mm.)

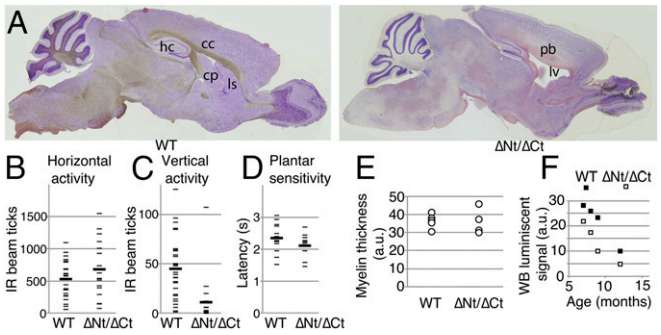

Fig. 6. Neural anomalies in *dido* mutant mice. (*A*) Sagittal sections of brain from a representative $dido^{\Delta NT/\Delta CT}$ mouse, showing ventricle enlargement and dysplasia of corpus callosum (cc) and hippocampus (hp). cp, caudoputamen; ls, lateral striatum; lv, lateral ventriculum; Pb, Probst bundles. (*B*) Horizontal and (*C*) vertical spontaneous locomotor activity in surviving adult $dido^{\Delta NT/\Delta CT}$ mice. Each mouse was tested three times over a 4-mo period. One-tailed Student's *t* test, $P = 0.04$ for vertical differences. (*D*) Plantar sensitivity to thermal stimulus in surviving adult $dido^{\Delta NT/\Delta CT}$ mice. (*B–D*) As one WT and two $dido^{\Delta NT/\Delta CT}$ mice died during this period, two values for *n* are given for each group; $n = 10$ (9) for WT, $n = 7$ (5) for $dido^{\Delta NT/\Delta CT}$ mice. (*E*) Toluidine-stained myelin was photographed in cross-sections of left and right sciatic nerves from adult $dido^{\Delta NT/\Delta CT}$ mice and WT littermates, and myelin sheet thickness was measured with Adobe Photoshop ($n = 6$ for WT, $n = 4$ for $dido^{\Delta NT/\Delta CT}$); 20 axons per nerve were evaluated in a blind manner. (*F*) Peripheral nerve α-tubulin deacetylation in $dido^{\Delta NT/\Delta CT}$ and WT mice ($n = 4 + 4$). Optical density of acetylated tubulin bands in Western blots of sciatic nerve extracts, normalized to total α- + β-tubulin levels. One-tailed paired Student's *t* test; $P = 0.02$.

motor pathology, because study of plantar thermal sensitivity by the Hargreaves method showed no defect in sensorial competence.

The normal axon count and myelin coating of sciatic nerves from $dido^{\Delta Nt/\Delta Ct}$ mice are consistent with the mild nature of the hindlimb motor defect. Peripheral neuromotor defects in mice can be associated with decreased axon tubulin content or an abnormal acetylated-to-total α-tubulin ratio (15). Tubulin is the main component of microtubules, which are crucial for efficient transport of organelles and effector molecules along axons. Microtubule stability is especially important for the function of long axons such as those of the sciatic nerve and depends on the acetylation of α-tubulin Lys40. As we found increased HDAC6 activity in embryo fibroblasts from $dido^{\Delta Nt/\Delta Ct}$ mice (14), we reasoned that increased deacetylation could destabilize axon microtubules along the $dido^{\Delta Nt/\Delta Ct}$ mouse sciatic nerve, leading to hindlimb motor defects of the type observed. Indeed, all $dido^{\Delta Nt/\Delta Ct}$ mice had less acetylated α-tubulin in the sciatic nerve. Deacetylation-driven microtubule destabilization would impair nervous transmission along the sciatic and other peripheral nerves, contributing to neuromotor defects that affect the hindlimbs.

Increased tubulin deacetylation has a major role in murine distal neuropathies related to human diseases. The symptoms of Charcot-Marie-Tooth disease (CMT), the most common inherited peripheral nervous system disorder in man, were reversed by HDAC6 inhibition in a mouse model of CMT (15). Correct control of cytoskeleton stability is crucial for balanced proliferation, neurogenesis, migration, differentiation, and connectivity in the brain, and mutations in genes such as *TUBA1a*, which encodes α-tubulin, are responsible for neurodevelopmental disorders, such as cortical digenesis (37), usually associated with mental retardation, epilepsy, or autism. A recent genome-wide association study (38) identified polymorphisms of genes such as *TUBA1a*, *TUBA4a*, and *KIF1B* (kinesin family member 1B, associated with CMT). An exome-wide rare variant analysis (39) also associated *TUBA4a* mutations with familial amyotrophic lateral sclerosis, which supports the destabilization of cytoskeleton as a cause of neurodegenerative diseases. Whether *dido* is involved in these or other distal neuropathies in humans requires further study, which will also identify factors other than increased tubulin deacetylation that affect the distal neuropathy in $dido^{\Delta Nt/\Delta Ct}$ mice. Whereas the limited survival of these mice to adulthood poses difficulties in their use, we are currently generating conditional *dido* mutants, which will be valuable for identifying the many roles of this gene.

It remains to be determined whether the phenotype associated with *dido* mutations is related not only to primary, nonmotile cilia regulation but is also modified by effects on motile cilia, which could account for phenotypic characteristics such as infertility or adult hydrocephalus.

The previously demonstrated role of Dido in the control of cilium length, based on its interaction with HDAC6, might be the basis for understanding the brain alterations observed in $dido^{\Delta Nt/\Delta Ct}$ mutant mice. Lack of Atat1, whose action is opposite that of HDAC6, leads to hydrocephalus and hippocampal dysplasia (40), brain alterations similar to those associated with the $dido^{\Delta Nt/\Delta Ct}$ genotype. Although Atat1 and HDAC6 are respectively the major α-tubulin acetyltransferase (41) and deacetylase (40) in mice, $atat1^{-/-}$ and $hdac6^{-/-}$ mice are viable and fertile (42). This is not the case for *dido* ($dido^{\Delta Ct/\Delta Ct}$ mice show early embryonic lethality), which suggests that Dido3 function is not restricted to the control of α-tubulin acetylation/deacetylation or that of other HDAC6 targets.

In a number of cases, *dido* mutation shows a distinct manifestation of cilia defects during brain development, namely agenesis of corpus callosum. It is tempting to speculate on a relationship of this gene with autism, as this aplasia is a distinctive characteristic of both the human condition (43) and its murine model (44). Of particular interest in human patients are subtelomeric microdeletions involving band 20q13.33, where the *Dido* gene is located. In a study of six subjects with this deletion (45), only one showed a thin corpus callosum and autistic disorders (46); this subject differed from others with normal MRI in the deletion of just 12 loci. These loci did not include *Dido*, but the breakpoint mapped only ∼60 kb downstream of the *Dido* transcription unit. This is compatible with neotelomere formation that could affect control of *Dido* expression, supporting the hypothesis of *Dido* involvement in human autistic behavior.

In summary, we describe various neurodevelopmental and morphogenetic phenotypes associated with murine *dido* mutations. Our model will help to understand the pathogenic mechanisms of some genetic or epigenetic disorders and could improve diagnosis and choice of treatment.

Materials and Methods

Mice. Heterozygous $dido^{\Delta Nt}$ and $dido^{\Delta Ct}$ mice were generated as described (17, 18) and maintained on a mixed genetic background (62.5% Sv129, 37.5% CD1). Double heterozygous $dido^{\Delta Nt/\Delta Ct}$ offspring were obtained from heterozygous parents, and sex-matched WT littermates were used as controls. Mice were handled according to national and European Union guidelines, and experiments were approved by the Comité Ético de Experimentación Animal, Centro Nacional de Biotecnología, Consejo Superior de Investigaciones Científicas.

Histology. Mouse brains were fixed in PBS/4% (wt/vol) paraformaldehyde (PFA; 24 h) and cryoprotected in PBS/30% (wt/vol) sucrose (>48 h). Trimmed brains (cut along the longitudinal axis) were embedded in OCT (Sakura). Floating sections (30 μm) were stained in 0.1% cresyl violet in 1% acetic acid (5 min).

For cilia analysis, brains were fixed in PBS/4% PFA, 24 h, and paraffin embedded. Sections (5 μm thick) were stained with rabbit anti-rat ACIII antibody (1:500; Santa Cruz) and goat anti-rabbit Alexa-488 (BD Systems); cell nuclei were DAPI stained. Confocal microscopy was performed using an IX81 microscope (Olympus). For 3D reconstruction and cilium measurement, we used Imaris 7 software.

For Turnbull staining, 4-μm-thick paraffin sections were hydrated, placed in 0.06 N potassium ferricyanide staining solution (1 h), washed in 1% acetic acid, and counterstained with nuclear-fast red (5 min). Sciatic nerve portions (7–8 mm long) were collected. Proximal (next to spinal segments L5–6) and distal segments (∼1 mm) were used for histochemistry; central segments were processed for protein analysis. Cryosections (10 μm) were stained with 0.5% toluidine blue.

Autophagy Assay. Progesterone (2.5 mg Depo-Provera; Upjohn) was administered to pregnant females at 17.5 and 18.5 dpc. Fetuses were extracted by cesarean section at 19.5 dpc and placed in a humidified chamber (30 °C). Heart samples were extracted at time 0 and 4 h.

Immortalized (19) or primary fibroblasts (obtained from 3-mo-old mice by mechanical disaggregation and trypsin digestion) were cultured in rich medium (DMEM + 10% FBS). At second passage, cells were plated at 75% confluence, cultured overnight, washed twice with HBSS, and incubated in starvation (HBSS) or rich medium (4 h).

Cell and tissue samples were lysed in cold PBS, 1% SDS with protease inhibitors. Equalized samples were fractionated in NuPAGE 12% Bis-Tris gels (Life Technologies) and transferred to Hybond-ECL nitrocellulose membranes (General Electric). LC3B I/II was detected with rabbit polyclonal antibody ab51520 (Abcam), anti-rabbit Ig-HRP (Dako), and Amersham-ECL (General Electric). Images were captured with X-ray film or a Proxima 2700 device (Isogen). Autograph density was analyzed with Adobe Photoshop CS5.

Behavioral Tests. Spontaneous locomotor activity was assessed in a flat 40 × 40 cm Perspex cage with two sets of IR emitter/sensor arrays for automatic monitoring of horizontal and vertical activity (Ugo Basile). Each mouse was tested for 5 min on 3 different days, and scores were averaged.

Latency to paw withdrawal after limb heating was assessed by the Hargreaves method using a plantar test instrument (Ugo Basile). All limbs were tested daily on each mouse for 4 d, and scores for fore- and hindlimbs were averaged separately.

Acetylated Tubulin Measurement. Left and right sciatic nerves from five WT and four mutant mice were dissected (see *Histology*). Protein extracts (8 μg) were loaded on 12% polyacrylamide gels. Acetylated α-tubulin was assessed by Western blot using mouse anti–α-acetylated tubulin mAb (clone 6-11B-1, Sigma) and normalized to total tubulin using DM1A + DM1B anti–α- and β-tubulin mAb (ab44928 Abcam) and GAPDH (anti-GAPDH (ab8245 Abcam). Luminescent signals were developed with HRP-conjugated rabbit anti-mouse IgG (Dako) and ECL reagents (Amersham/General Electric). Luminescence was detected and quantified with a Proxima 2700 device (Isogen).

ACKNOWLEDGMENTS. We thank Dr. J. R. Naranjo for advice and help with neurobehavioral tests, A. Alonso-Guerrero for technical assistance, R. Gutiérrez for animal handling, and C. Mark for editorial assistance. This work is financed by Grants SAF2010-21205, PIB2010BZ-00564, and SAF2013-42289-R from the Spanish Ministerio de Economía y Competitividad.

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