The Nucleoside Diphosphate Kinase Gene Nme3 Acts as Quantitative Trait Locus Promoting Non-Mendelian Inheritance

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Abstract

The t-haplotype, a variant form of the t-complex region on mouse chromosome 17, acts as selfish genetic element and is transmitted at high frequencies (>95%) from heterozygous (+/t) males to their offspring. This phenotype is termed transmission ratio distortion (TRD) and is caused by the interaction of the t-complex responder (Tcr) with several quantitative trait loci (QTL), the t-complex distorters (Tcd1 to Tcd4), all located within the t-haplotype region. Current data suggest that the distorters collectively impair motility of all sperm derived from t/+ males; t-sperm is rescued by the responder, whereas + sperm remains partially dysfunctional. Recently we have identified two distorters as regulators of RHO small G proteins. Here we show that the nucleoside diphosphate kinase gene Nme3 acts as a QTL on TRD. Reduction of the Nme3 dosage by gene targeting of the wild-type allele enhanced the transmission rate of the t-haplotype and phenocopied distorter function. Genetic and biochemical analysis showed that the t-allele of Nme3 harbors a mutation (P89S) that compromises enzymatic activity of the protein and genetically acts as a hypomorph. Transgenic overexpression of the Nme3 t-allele reduced t-haplotype transmission, proving it to be a distorer. We propose that the NME3 protein interacts with RHO signaling cascades to impair sperm motility through hyperactivation of SMOK, the wild-type form of the responder. This deleterious effect of the distorters is counter-balanced by the responder, SMOKTcr, a dominant-negative protein kinase specifically expressed in t-sperm, thus permitting selfish behaviour and preferential transmission of the t-haplotype. In addition, the previously reported association of NME family members with RHO signaling in somatic cell motility and metastasis, in conjunction with our data involving RHO signaling in sperm motility, suggests a functional conservation between mechanisms for motility control in somatic cells and spermatozoa.

Introduction

In general, diploid organisms transmit homologous chromosomes at the Mendelian (equal) ratio to their offspring. However, several types of non-Mendelian inheritance have been described, and in mammals a prominent example is transmission ratio distortion (TRD) in the mouse, which is caused by the t-haplotype. The t-haplotype is a variant form of the t-complex, which maps to the centromere-proximal third of chromosome 17. According to evolutionary studies, this haplotype originated more than one million years ago and, due to four large inversions, has since evolved devoid of meiotic exchange with the wild-type t-complex [1–3]. The t-haplotype is transmitted at an abnormally high ratio from heterozygous (+/t) males to their offspring [4]. This selective advantage is due to superior swimming behaviour of t-haplotype sperm as compared to t-sperm derived from the same male [5,6]. t-sperm does not, however, function superiorly to sperm derived from wild-type (+/+ ) males [7]. The t-haplotype rather encodes several t-complex-distorters (Tcd), which cumulatively affect sperm motility. This deleterious effect is rescued by the t-complex-responder (Tcr), but exclusively in t-haplotype carrying sperm. Thus, only sperm carrying the wild-type t-complex are affected.

The first evidence for how TRD is caused molecularly was obtained following the isolation of Tcr, which was found to encode a mutant, dominant-negative form of Sperm motility kinase 1 (SmoK1), termed Smok1Tcr [3]. Expression of wild-type Smok1 and of Smok1Tcr commences in haploid spermatids and, in contrast to other haploid expressed genes, neither their RNA nor their protein products are shared between haploid sperm cells, which are connected in a syncytium [9]. This exceptional behaviour provided a molecular explanation for the exclusive rescue of t-sperm from the deleterious effect of the distorters. The molecular nature of SMOK1Tcr revealed that TRD is caused by alterations in a signaling pathway involved in sperm motility, and led to the identification of Tcd genes, which were postulated to act upstream of SMOK1 in this signaling pathway [8]. The first Tcd isolated was Tcd1a, which was identified as a hypermorph of Tagap1, a GTPase activating protein (GAP) and
**Author Summary**

Selfish genetic elements, which promote their own propagation and thereby violate Mendel’s laws, have attracted much attention within the scientific community and by the public. The molecular principles underlying their exceptional behaviour are, in general, not well understood. A notable exception is the t-haplotype of the mouse, which was discovered in 1936 and has since been a paradigm for non-Mendelian inheritance in mammals. Recently we have revealed the molecular nature of several elements within this genetic region, including the t-complex responder and two t-complex distorters that interact to promote the high transmission rate of the t-haplotype from males (which carry this variant genetic region) to their offspring. Here we show that the nucleoside diphosphokinase gene, Nme3, acts as distorer and thus contributes to the high transmission rate of the t-haplotype. We show that Nme3 acts as a quantitative trait locus in sperm motility control. Since human NME family members have been associated with somatic cell motility (which is also controlled by RHO signalling) and cancer cell metastasis, we propose a functional conservation between motility control in somatic and sperm cells.

**Results/Discussion**

**Nme3 Is Expressed in Testis and Is Altered in the t-Haplotype**

The identification of the Rho small G protein regulators Tagap1 and Fgd2 as t-complex-distorters within the t-haplotype suggested that more genes involved in G protein signalling which may have a quantitative effect on t-haplotype transmission might be located in this chromosome segment. Therefore, we initiated a search for genes related to Rho signalling within this region of chromosome 17. We identified the gene Nme3, encoding a member of the nucleoside diphosphokinase (NDK) family, at position 25 Mb from the centromere (Figure 1A). Nme3 belongs to the group I Nme (Nme1–4) which are all catalytically active and share significant sequence homology [13,14].

In order to qualify as a Tsd candidate, a gene must be expressed in the testis and show variability between t- and wild-type alleles. Northern blot analysis showed expression of Nme3 in testes from the earliest stage after birth tested (7 days) to the adult (Figure 1B). Using in situ hybridization on sections of adult testes, Nme3 transcripts are detectable predominantly in early stages of spermatogenesis, while expression appears to be down-regulated in haploid cells (Figure 1C). However, the expression analysis of round spermatids using microarrays have shown that Nme3 transcripts are also present in spermatids ([15], and EMBL-EBI: Gene Expression Atlas). Thus, Nme3 transcripts apparently persist during spermiogenesis and allow translation of NME3 protein acting later in spermatooza. In conclusion, Nme3 was found to fulfill the first important criterion for a distorer.

Since the RNA expression level can be a good indicator of a QTL, as shown for the t-alleles of Fgd2 and Tagap1 [10,11], we analyzed the expression of Nme3 from the t-haplotype allele and compared it to the wild-type alleles from several mouse strains. The expression levels were found to be nearly identical between RNA derived from testes of C57BL/6 or other wild-type strains, and t-haplotype carrying strains (Figure 1D). c+/+ male testes showed marginally higher expression.

Since the analysis of RNA expression level revealed no significant variability between t- and wild-type alleles we examined sequence variation. We isolated cDNA clones by RT-PCR from testicular RNA from several wild-type and t-haplotype carrying strains and from a testis cDNA library prepared from c+/+ males. In addition, we analyzed genomic fragments derived from several t-haplotypes. All sequence analyses detected a t-specific C to T transition in the coding sequence of Nme3, a missense mutation resulting in the change of proline to serine at position 89 (P98S; Figure 1E). This mutation was found in all t-alleles tested, which carry the t-form of the inversion In(17)4 (p25, p32, p12, p1), but in none of the wild-type strains analyzed (C57BL/6, DBA/2, 129Sv, NMRI) (Figure 1E, and data not shown). Therefore, this P98S mutation in Nme3 distinguishes the t-allele from the wild-type allele. All group I NDK enzymes possess almost identical 3-D structures, and the mutation affects a highly conserved amino acid located between alpha-helix a2 and beta-sheet b3 of NME3 [13,14], which may alter the function of the protein. A similar proline to serine exchange was described in the Killer-of-prune mutation (awd6) of the abnormal-wing-disc (awd) gene of D. melanogaster (Figure 1E, [16]). The awd6 mutation was shown to decrease the nucleoside diphosphokinase kinase activity substantially with respect to the wild-type awd gene product [17].

The combined data identified the t-allele of Nme3 as a distorer candidate.

According to its position on chromosome 17 it was not clear whether the Nme3 gene is located within the Tsd2 or Tsd3 region. The proximal partial t-haplotype t6/+, which extends into inversion In(17)4, carries Tsd2, but not Tsd3, which maps more distally [18,19]. Therefore, analysis of t6/+ allows the assignment of Nme3 to either the Tsd3 or Tsd2 region. Genomic Southern blot analysis complemented by cDNA sequencing demonstrated that t6/+ carries the wild-type allele of Nme3 and thus, Nme3 is a Tsd2 candidate (Figure S1 and data not shown).

**A Knock-Out Allele of Nme3 Phenocopies a Distorter**

Distorter genes act as QTLs in the sense that up- or down-regulation of gene expression and/or activity has a qualitative effect on the phenotype; observed here as TRD. A proven method
Figure 1. *Nme3* is a *Tcd2* candidate. (A) The position of *Nme3* in the wild-type *t*-complex (t) and genetic mapping on *t*^18^ localized it to the *Tcd2* region of the *t*-haplotype (t) (see Figure S1). Symbols of genes verified to be involved in TRD are boxed, and map positions relative to the wild-type chromosome 17 (in Mb) are indicated. Molecularly unknown *Tcd* loci are also listed along with Chr17 inversions and their relative orientations in both, the wild-type and *t*-haplotype chromosome, as well as crucial genetic markers (*T*, *tf*, *H-2*). (B) Northern blot analysis of *Nme3* transcripts in testes from consecutive post partum stages reflecting the first round of spermatogenesis (P7–P24), and in testes of adult mice (ad). (C) *In situ* hybridization analysis of *Nme3* on testis cryosections from an adult male. Expression is predominant in cells near the basal lamina (dotted line) representing diploid cell types. Schematic view of a seminiferous tubule. (Se) Sertoli cells; (SG) spermatogonia; (SC) spermatocytes; (RS) round spermatids; (SZ) spermatozoa. Scale bar: 50 μm. (D) Northern blot analysis of *Nme3* expression in different wild-type strains and *t*-haplotypes. Quantification of the signals with respect to C57Bl/6 did not reveal significant differences. (E) Amino acid sequence comparison of several NME proteins encoded by *Mus musculus* (mmNME1 to mmNME3), the *t*-haplotype (mmNME3-t), *Homo sapiens* (hsNME3), and *Drosophila melanogaster* (dmAWD); a conserved proline at position 89 is altered in mmNME3-t (P89S), and a nearby proline residue was described as killer-of-prune mutation (*K-pn*) in the abnormal-wing-disc (*awd*) gene of *D. melanogaster* when mutated to serine (red boxed). doi:10.1371/journal.pgen.1002567.g001

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**Caption:**

*Figure 1. Nme3 is a *Tcd2* candidate.* (A) The position of Nme3 in the wild-type t-complex (t) and genetic mapping on *t*^18^ localized it to the *Tcd2* region of the *t*-haplotype (t) (see Figure S1). Symbols of genes verified to be involved in TRD are boxed, and map positions relative to the wild-type chromosome 17 (in Mb) are indicated. Molecularly unknown *Tcd* loci are also listed along with Chr17 inversions and their relative orientations in both, the wild-type and *t*-haplotype chromosome, as well as crucial genetic markers (*T*, *tf*, *H-2*). (B) Northern blot analysis of *Nme3* transcripts in testes from consecutive post partum stages reflecting the first round of spermatogenesis (P7–P24), and in testes of adult mice (ad). (C) *In situ* hybridization analysis of *Nme3* on testis cryosections from an adult male. Expression is predominant in cells near the basal lamina (dotted line) representing diploid cell types. Schematic view of a seminiferous tubule. (Se) Sertoli cells; (SG) spermatogonia; (SC) spermatocytes; (RS) round spermatids; (SZ) spermatozoa. Scale bar: 50 μm. (D) Northern blot analysis of *Nme3* expression in different wild-type strains and *t*-haplotypes. Quantification of the signals with respect to C57Bl/6 did not reveal significant differences. (E) Amino acid sequence comparison of several NME proteins encoded by *Mus musculus* (mmNME1 to mmNME3), the *t*-haplotype (mmNME3-t), *Homo sapiens* (hsNME3), and *Drosophila melanogaster* (dmAWD); a conserved proline at position 89 is altered in mmNME3-t (P89S), and a nearby proline residue was described as killer-of-prune mutation (*K-pn*) in the abnormal-wing-disc (*awd*) gene of *D. melanogaster* when mutated to serine (red boxed). doi:10.1371/journal.pgen.1002567.g001
for testing a possible effect of gene dosage on TRD is to inactivate the wild-type allele by gene targeting, assay the transmission of a t-haplotype from males carrying the knock-out allele on the homologous chromosome, and compare it to control males which carry the wild-type allele. We targeted the Nme3 gene in ES cells by replacing exon 1 and part of exon 2 with a Pgk-Neoycin resistance cassette, generating a null allele (Figure 2A). Successful integration of the targeting construct was verified by Southern blot analysis (Figure 2B). We introduced the targeted allele into the germ line and confirmed by RT-PCR that Nme3 transcripts are lacking in the testes of homozygous-null males (Figure 2C). We then generated males carrying the targeted allele on the wild-type chromosome and the wild-type allele on either of the partial t-haplotypes t6/p or th49. Littermates carrying the wild-type allele on both chromosomes 17, in conjunction with t6/p or th49, served as controls. For each male we determined the number of offspring that inherited the t-haplotype. In both tests the t-haplotype was transmitted at a significantly higher rate (15% and 8% respectively) from males carrying one targeted allele compared to littermates homozygous for the wild-type allele (Table 1). Therefore, a reduction of the gene dosage by half significantly increased the transmission rate of the t-haplotype. These data demonstrate that Nme3 acts as a QTL on t-haplotype inheritance.

The t-Allele of Nme3 Encodes a Hypomorph and Acts as Distorter

Although the genetic inheritance test proved the nature of Nme3 as a QTL, it did not verify that the Nme3 t allele acts as distorter gene. The latter requires that the t-allele itself alters the overall Nme3 activity in sperm, and causes a statistically significant change in t-haplotype transmission. To determine the mechanisms through which this may happen we first assessed the enzymatic activity of NME3-P89S relative to wild-type NME3.

We produced the enzyme in vivo using a coupled transcription/translation system in rabbit reticulocyte lysate, purified the complexes, and measured the activity in an enzymatic assay. NME3-P89S showed strongly reduced enzymatic activity compared to wild-type (1295 vs) NME3 protein (18% of wt activity; Figure 3A).

It is important to note that nucleoside diphosphate kinases function as hexamers. Therefore, in vivo the t-encoded NME3 monomers may form mixed hexamers with the wild-type protein, and in mixed complexes NME3-P89S might function as dominant-negative protein interfering with the function of the wild-type protein. Alternatively, it might form semi-functional complexes with wild-type monomers or have no effect. In order to discriminate between these possible effects, we assayed the enzymatic activity of a 1:1 mixture of NME3-P89S with wild-type protein, which would reflect the situation in a t/+ male. We combined plasmids encoding the wild-type and the t-allele at an equal ratio prior to in vitro transcription and translation, purified the complexes, and measured their enzymatic activity. The activity dropped to 58% of that of the wild-type protein, suggesting that both the wild-type and the mutant protein contribute to the total enzyme activity (Figure 3A).

In order to assess whether the Nme3 t-allele acts as an antimorph (dominant-negative) or hypomorphic (semi-functional) in vivo we took a transgenic approach. If the former were the case, a transgenic construct expressing the t-allele should enhance the transmission rate of a t-haplotype from a t/+ male, since a dosage increase of the t-allele should further reduce endogenous NME3 activity. In contrast, if the t-allele acts as hypomorph the transgene would provide extra NME3 activity to the endogenous gene products and thus the t-haplotype transmission should drop.

We created a transgene construct (Tg[Nme3]H17hgh, abbreviated Tg) expressing the Nme3 allele in haploid sperm cells using the testis-specific Ace promoter (Figure 3B) [20]. Northern blot analysis and quantification showed that the transgene construct was expressed at approximately 80% of the level of the two endogenous alleles from a t/+ male (Figure 3C). We generated t/+ mice carrying one wild-type and one t-allele, along with t/+ males additionally expressing the transgene. The former should maintain a 1:1 ratio of NME3 to NME3-P89S, while the latter should produce around 2.6-fold more NME3-P89S than NME3 (one wild-type allele and one t-allele plus approximately 1.6-fold over-expression of the t-allele from the Tg construct). We determined the transmission rate of t from t/+Tg/0 and compared it to t transmission from control littermates of the genotypic t/+.

Males (t/+)+ expressing one wild-type and one t-allele transmitted the t-haplotype to 92.3% of their offspring. An increase of the t-allele dosage in hemizygous transgenic males (t/+ +Tg/0) reduced the t-transmission to 89% of the offspring (p = 0.02; Table 1), indicating that the t-allele does not act as an antimorph, but as a hypomorph.

Biochemical testing confirmed this conclusion. Increasing the amount of NME3-P89S to 2.6-fold of the wild-type protein (1 part NME3; 2.6 parts NME3-P89S), which reflects the relative expression of wild-type and t-allele-derived Nme3 mRNA in t/+ +Tg/0 males, significantly increased the enzymatic activity in comparison to the 1:1 mixture (Figure 3A). Thus, the NME3- P89S protein contributes to the overall enzymatic activity, rather than interfering with activity of the wild-type protein. These biochemical data are consistent with and support the genetic data.

In summary, both data sets identify the t-allele of Nme3 as a hypomorphic allele acting as distorter of t-haplotype transmission.

The ability of the mouse t-haplotype to promote its transmission from t/+ males to a high proportion of their offspring is due to the unusual properties of the responder, SmokTcr, a dominant negative protein kinase which is retained in the haploid sperm cells expressing the gene and able to rescue the impairment of sperm motility caused by the distorters [9]. These latter act as QTLs, which additively contribute to the high transmission rate of the responder. We have previously identified two distorters, which act as hypermorphs, the Rho-GAP Tagap1 and the Rho-GEF Fgd2 [10,11]. Although the two proteins have antagonistic effects on Rho activity, excess activity of either gene enhances the transmission rate of the t-haplotype from t/+ males. Thus, we proposed that Tagap1 controls a negative regulator and Fgd2 an activator of SMOK, the wild-type form of the responder SMOKTcr [11]. Increased down-regulation of the negative regulator by Tagap1 and increased up-regulation of the activator by Fgd2 both contribute to hyperactivation of SMOK, leading to impairment of sperm motility. This deleterious effect of the distorters is counterbalanced by SMOKTcr, which exclusively rescues t-sperm resulting in TRD (Figure 4).

It is not yet clear how Nme3 interacts with the Rho signaling cascades involved in TRD. Recent reports have revealed negative interactions between NME1 and Rho small G protein signaling [21–23]. Nme1 can act as a negative regulator of CDC42 by binding to the PH domain of the CDC42-GEF DBL. It can also inhibit CDC42 by direct interaction [21,22]. Similarly, NME1 has been demonstrated to inhibit RAC1 activity by interacting with the RAC1 activator TIA1, an effect which is independent of its nucleoside diphosphate kinase activity [23]. It is unknown whether NME3 may function similarly. However, since genetic reduction of Nme3 promotes TRD, it is likely that the NME3

![Image](Image-58x24.png)

![Image](Image-76x41.png)
Figure 2. Targeted inactivation of the *Nme3* gene. (A) Gene targeting strategy: a neomycin selection cassette was used to replace the entire first and part of the second exon of *Nme3*. (B) Confirmation of correct homologous recombination by Southern blot analysis of *XbaI* digested genomic DNA derived from ES-cells or mice. (C) Loss of *Nme3* transcripts in homozygous knock-out mice (−/−) as determined by expression analysis using RT-PCR confirms that the targeted allele represents a null mutation. Genomic regions in (A) are not drawn to scale. Abbr.: LP, RP: left or right external probes; DT, dipheria toxin cassette for negative selection; wt, wild-type; +, wild-type allele; −, knock-out allele.

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protein is able to activate the inhibitory pathways controlling SMOK activity. Alternatively, it cannot be excluded that NME3 may inhibit the activating pathway or exert both, activating and inhibiting functions. The former effect might be caused by local increase of the GTP concentration promoting activation of Rho, inhibiting functions. The former effect might be caused by local increase of the GTP concentration promoting activation of Rho.

| Genotype of male | Offspring | Number of males | t | + | total % | t² | P |
|-----------------|-----------|-----------------|---|---|---------|----|---|
| Nme3<sup>3Amp</sup>/+ | 7 | 167 | 308 | 475 | 35.2 | 6.721 | 0.0095 |
| Nme3<sup>3Amp</sup>/+ | 7 | 117 | 317 | 434 | 27.0 | 11.701 | 0.0006 |
| Nme3<sup>3Amp</sup>/+ | 4 | 150 | 104 | 254 | 59.0 | 11.701 | 0.0006 |
| Nme3<sup>3Amp</sup>/+ | 5 | 139 | 175 | 314 | 44.3 | 5.337 | 0.0209 |
| T<sup>+/</sup> Tg/0 | 8 | 587 | 72 | 659 | 89.0 | 5.337 | 0.0209 |
| T<sup>+/</sup> | 8 | 646 | 50 | 696 | 92.9 |

Abb.: +, wild type; Tg(Nme3)H17bg/h, doi:10.1371/journal.pgen.1002567.t001

Materials and Methods

Ethics Statement

Animal experiments were approved by the ethics committee of the Regierungspräsidium Freiburg (registration number T-00/28) and the LAGeSo Berlin (registration numbers ZH120 and Reg 0248/03).

Transcript Analysis

We amplified Nme3 transcripts by RT-PCR of the Nme3 coding region from mouse testis RNA isolated from different wild-type strains and t-haplotypes (primers Nme3-s and Nme3-as). We isolated the amplicons (622 bp), cloned them into pBS-SK (Stratagene), and sequenced several independent clones for each genotype. We isolated Nme3 clones from a d<sup>3</sup>P<sup>3</sup> testes cDNA library by PCR-screening of subpools and colony hybridization [11] using the Nme3 cDNA as a probe and sequenced the library clones as above. We performed Northern blot analysis using the NorthernMax-Gly Kit (Ambion) according to the manufacturer’s instructions. For in situ hybridization analysis, DIG-labelled in vitro transcribed antisense RNA corresponding to the coding region of Nme3 (primers Nme3-s and Nme3-as) was hybridized to 10 µm frozen sections as described [30].

Gene Targeting and Transgene Constructs

We targeted the Nme3 locus in CJ7 ES-cells [31] by replacing exon 1 and part of exon 2 (bp 33550 to 33847 in BAC 126c8, accession number AF220941.1) with a Pgk1-neo-polyA (Pgk1-neo/flox3xPAG) - cassette [11] as depicted in Figure 2A. We included a diphtheria toxin cassette for negative selection. To construct the targeting vector, we isolated the left- and right homology arms (4695 bp and 2180 bp, respectively) of the targeting construct by PCR and cloned both arms on either side of the selection cassette, introducing a Sall site for linearization at the end of the left arm. We electroporated CJ7 ES-cells with the linearized targeting construct and selected, isolated, and analyzed clones according to standard procedures [32]. Correctly targeted ES-clones were identified by Southern hybridization of Xhol digested genomic DNA with the left (5’) probe (LP) and right (3’) probe (RP) (Primers LP-s and LP-as, 1052 bp probe fragment; RP-s and RP-as, 814 bp probe fragment). Both probes detect a 11.206 kb Xhol fragment in wild-type. Upon successful targeting, the LP detects a 7.228 kb fragment and the RP a 5.354 kb fragment. The Nme3<sup>+</sup> transgenic construct consists of the angiotensin converting enzyme (Ace) sperrimidogenesis-specific promoter including the transcriptional start site (position −91 to +17) [20] followed by the cDNA of Nme3 exon 1 to 4, and genomic sequence comprising intron 4/5, exon 5 and 141 bp from the 3’- untranslated region, ending 12 bp upstream of the endogenous polyadenylation (pA) signal, which we replaced by the SV40 pA sequence from pCS2+. We flanked this expression cassette with 2 copies of the chicken beta-globin insulator on both sides [33] (Figure 3B).

Biochemical Assays

We cloned the Nme3 alleles into the pET30c vector (Novagen) in-frame with a 6xHis and S-tag. Since expression of full-length recombinant proteins in E. coli was unsuccessful, we used the constructs for in vitro transcription/translation reactions (IVT) with the TNT T7 Quick Coupled Transcription/Translation System (Promega) according to the manufacturer’s instructions. Plasmid concentrations were kept constant relative to the reaction volumes. We purified recombinant NME3 using Ni-NTA agarose (Qiagen) (12.5 µl bed volume/50 µl IVT reaction). We added 0.6 ml Ni-

Table 1. The reduction of the wild-type Nme3 gene dosage and over-expression of the Nme3 t-allele have opposite effects on t-haplotype transmission.
Figure 3. The $t$-allele of Nme3 encodes a hypomorph. (A) Nucleoside diphosphate kinase activities of wild-type (+) and $t$-allele ($t$)-derived proteins or mixtures of both expressed in vitro. The 1:1 mixture reflects the relative contribution of both alleles to NME3 activity in $t^+/+$ heterozygotes, the 1:2.6 ratio of + and $t$ alleles corresponds to the expression of + and $t$ alleles measured in testes from $t^6+/+;Tg^t/0$ males (B, C). The columns show the mean of three experiments with the standard deviation. (B) Schematic representation of the transgene construct Tg(Nme3)H17bgh (Tg) generated for expression of the Nme3 $t$-allele during spermatogenesis. (C) Expression of the Nme3 $t$-allele from the transgene construct determined by Northern blot analysis of testis RNA derived from a $t^6+/+;Tg^t/0$ male. Quantification of endogenous ($t^6$, +) and transgene (Tg)-derived transcripts revealed a ratio of 1:0.8. Assuming that the $t^6$ allele contributes 50% of the endogenous Nme3 transcripts, the overall ratio of + to $t$ derived Nme3 transcripts expressed in a $t^6+/+;Tg^t/0$ male is therefore 1:2.6. Abbr.: Ace-prom, angiotensin converting enzyme promoter; SV40pA, Simian virus 40 polyadenylation signal fragment; ins, chicken beta globin insulator [33]; Ex, exon; Ndk, nucleoside diphosphate kinase.

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NTA buffer for binding (50 mM NaH$_2$PO$_4$, 30 mM NaCl, 20 mM Imidazole, 1× complete protease inhibitors EDTA-free (Roche), pH 8) and incubated at 4°C for 1.5 h with agitation. Samples were then washed 3× in Ni-NTA-buffer (0.8 ml) and NME3-protein was eluted with 45 µl elution buffer (Ni-NTA buffer containing 250 mM imidazole). For quantification of protein, we loaded 20 µl of eluted protein on a NuPAGE 4–12% Bis-Tris Gel (Invitrogen) and blotted with the I-blot system (Invitrogen) on a PVDF membrane (Millipore). After blocking 1–2 h at RT using 5% skim milk powder in TBS-T/0.2% Tween, we probed the blot with an anti-S-tag antibody (1:500 dilution, Delta Biolabs) overnight at 4°C, followed by incubation with donkey anti-rabbit HRP-coupled secondary antibody (1:10000 diluted, Jackson Immunoresearch), and detected the signal with the ECL Advance Western Blotting Detection Kit from GE Healthcare. Densitometry quantification of Western blots was performed using ImageJ software.

We analyzed NME3 protein preparations for nucleoside diphosphate kinase activity using a transphosphorylation assay followed by thin layer chromatography (TLC) essentially as described [34]. We used 20 µl NME3 protein eluate in a reaction mixture containing 10 mM HEPES, 20 mM NaCl and 2 mM MgCl$_2$, 2 mM ATP, 1 mM TDP and 2 mM [γ-32P] ATP. We took 6 µl samples after each 15, 30, and 45 minutes incubation at room temperature and stopped the reaction with 1 µl of 50 mM EDTA (pH 8.0). As a positive control we used 0.1 U of nucleoside 5'-diphosphate kinase from bakers yeast (Sigma Aldrich N0379). Reactions were analyzed by TLC on PEI-cellulose plates (Macherey-Nagel) using 0.75 M KH$_2$PO$_4$ (pH 3.65) as running buffer. We exposed dried TLC plates to phosphorimager screens and quantified with ImageJ. Wild-type NME3 and NME3-P89S protein activities were normalized to total protein levels.

**Supporting Information**

Figure S1  Nme3 maps to the Tcd2 region of the t-haplotype. (A) Genetic maps of the wild-type t-complex (+) and the t-haplotype (t). The proximal partial t-haplotype t$^{+\text{cd2b}}$ arose by recombination between a t-haplotype and a wild-type chromosome within inversion 4 [In(17)4] resulting in loss of the distorter Tcd2. (B) Genomic Southern blot analysis using the coding sequence of Nme3 as probe demonstrates the presence of a non-polymorphic Nme3 fragment in t$^{+\text{cd2b}}$, as shown by the presence of a 2.19 kb BglI fragment (+, 9) and of a 1.43 kb knock-out allele fragment in t$^{+\text{cd2b}}$/+; Nme3$^\text{tm5Bgh}$ genomic DNA. Since Nme3 is located distal to Fgd2 in the t-haplotype and Fgd2 carries the wild-type allele of Fgd2 (Figure 1a in [11]) we conclude that the 2.19 kb fragment in t$^{+\text{cd2b}}$ represents the wild-type Nme3 allele. In addition, we did not detect t-allele Nme3 transcripts in Nme3$^\text{tm5Bgh}$/+; t$^{+\text{cd2b}}$/+ testes-derived
Table S1 | Primer sequences for transcript and genomic analysis, generation of transgenic and gene targeting constructs, and genotyping.

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Author Contributions

Conceived and designed the experiments: BGH HB. Performed the experiments: HB JW SS BK YC. Analyzed the data: BGH HB SS YC. Wrote the paper: BGH HB.

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