

A Maternal-Zygotic Effect Gene, *Zfp57*, Maintains Both Maternal and Paternal Imprints

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SUMMARY

The mechanisms responsible for maintaining genomic methylation imprints in mouse embryos are not understood. We generated a knockout mouse in the *Zfp57* locus encoding a KRAB zinc finger protein. Loss of just the zygotic function of *Zfp57* causes partial neonatal lethality, whereas eliminating both the maternal and zygotic functions of *Zfp57* results in a highly penetrant embryonic lethality. In oocytes, absence of *Zfp57* results in failure to establish maternal methylation imprints at the *Snrpn* imprinted region. Intriguingly, methylation imprints are reacquired specifically at the maternally derived *Snrpn* imprinted region when the zygotic *Zfp57* is present in embryos. This suggests that there may be DNA methylation-independent memory for genomic imprints. *Zfp57* is also required for the postfertilization maintenance of maternal and paternal methylation imprints at multiple imprinted domains. The effects on genomic imprinting are consistent with the maternal-zygotic lethality of *Zfp57* mutants.

INTRODUCTION

Parental genetic influence on the development of offspring is evident in both the invertebrate and vertebrate animal kingdoms. Maternal effects are commonly mediated through deposition of the cytoplasmic transcripts or protein products in oocytes during oogenesis in the female germline. These then exert their effects on the fertilized zygotes and affect early embryonic development (De Robertis et al., 2000; Gosden, 2002; Melendez and Greenwald, 2000; Newman-Smith and Rothman, 1998; Nusslein-Volhard, 1991; Priess et al., 1987; Schier, 2001).

Genomic imprinting, the process that causes genes to be expressed according to their parental origin, occurs in plants and mammals and is mediated by epigenetic modifications that differ on the two parental chromosomes (Ferguson-Smith and Surani, 2001; McGrath and Solter, 1984; Reik and Walter, 2001; Surani et al., 1984; Tilghman, 1999; Verona et al., 2003). Over 80 imprinted genes have been discovered in mouse (<http://www.mgu.har.mrc.ac.uk/research/imprinting/>). Many are known to

be localized in clusters regulated by a *cis*-acting imprinting control region (ICR) that acquires heritable parental origin-specific differential DNA methylation in the male or female germline (Ben-Porath and Cedar, 2000; Lewis and Reik, 2006). The cycle of methylation programming at an ICR begins with erasure of methylation during the development of primordial germ cells. Subsequently, during oogenesis and spermatogenesis, de novo methylation is differentially established on the maternal and paternal chromosomes in the germline; these are the so-called germline imprints. After fertilization, the preimplantation embryonic genome loses much of its methylation through both active and passive demethylation events commencing in the zygote, and, subsequently, de novo methylation is acquired around the time of implantation (Reik et al., 2001). Germline imprints, however, appear to be resistant to this postfertilization methylation reprogramming, allowing them to be stably inherited from germline to offspring (Morgan et al., 2005).

Studies in which the de novo and maintenance DNA methyltransferase machinery has been mutated have proven that germline establishment and postfertilization maintenance of differential methylation is essential for the monoallelic activity of imprinted genes (Bourc'his et al., 2001; Kaneda et al., 2004; Li et al., 1993). In contrast, the mechanisms specifically rendering germline methylation imprints resistant to preimplantation, genome-wide demethylation are not understood. Oocyte-derived *PGC7/Stella* was shown to confer partial protection from demethylation at one class of repetitive sequence and at some imprinted domains (Nakamura et al., 2007). More recently, maternal and zygotic functions of *Dnmt1*, the maintenance methyltransferase, were found to maintain DNA methylation imprints in preimplantation embryos (Hirasawa et al., 2008).

The majority of germline methylation marks identified to date are established on maternally inherited chromosomes during oogenesis. These include the *Snrpn* imprinted region implicated in human Prader-Willi and Angelman syndromes. In general, these maternal methylation imprints are located at promoters that, in some cases, regulate large antisense noncoding RNAs expressed from the paternal chromosome that have been implicated in the repression of protein-coding genes in *cis* (Pauler et al., 2007). Only three imprinted domains are known to be regulated by a controlling element that is methylated in the paternal germline. Of these, the *Igf2-H19* and *Dlk1-Dio3* imprinting control elements regulate imprinted genes that are essential for normal prenatal growth and development (Edwards and

Ferguson-Smith, 2007; Kawahara et al., 2007; Thorvaldsen and Bartolomei, 2007).

KRAB zinc finger proteins form one of the largest transcription factor families in the mouse and human genomes (Looman et al., 2002). They act as potent transcriptional repressors through KRAB box-mediated interaction with KAP-1/TIF1 β corepressor complexes (Abrink et al., 2001; Friedman et al., 1996; Schultz et al., 2001, 2002). Mediated by the DNA-binding capacity of KRAB zinc finger proteins, KAP-1 functions to recruit factors associated with DNA methylation (Wiznerowicz et al., 2007) and the formation of repressive chromatin, including histone deacetylases and histone methyltransferases (Ayyanathan et al., 2003; Schultz et al., 2001, 2002). Despite this, there are very few known target genes of the KRAB zinc finger proteins. There is even more limited information as to the function of KRAB zinc finger proteins in vivo, except for the information regarding the mouse variant *rs1*, which affects sexually dimorphic gene expression in the liver (Krebs et al., 2003). We identified *Zfp57* during a gene trap-based screen for factors downregulated upon embryonic stem (ES) cell differentiation (Li and Leder, 2007). We generated a knockout mouse in the *Zfp57* locus that encodes a putative KRAB zinc finger protein. Our data indicate that *Zfp57* is an essential maternal-zygotic effect gene and is required for the establishment and reacquisition of the maternal methylation imprint at the *Snrpn* domain. It also maintains both paternal and maternal methylation imprints after fertilization at multiple imprinted regions.

RESULTS

ZFP57 Is a KRAB Zinc Finger Protein

Consistent with previous findings (Alonso et al., 2004), we found that mouse ZFP57 is a putative KRAB zinc finger protein (Figure 1A). A human homolog was identified, indicating conservation of this protein (Figure 1B). To determine whether ZFP57 contains a functional KRAB box, we performed a coimmunoprecipitation (co-IP) interaction assay in which both myc epitope-tagged mouse ZFP57 and KAP-1/TIF1 β , the obligate corepressor for KRAB zinc finger proteins, were overexpressed transiently in COS cells. Antibodies against the myc epitope were used to pull down ZFP57-associated proteins, and two different antibodies against two nonoverlapping regions of KAP-1/TIF1 β were used to probe the immunoprecipitated material (Figure 1C). KAP-1/TIF1 β was detected when it was coexpressed with ZFP57 (lane 5 in Figure 1C). These data prove that ZFP57 contains a conserved functional KRAB box that binds KAP-1/TIF1 β . Binding between endogenous ZFP57 and KAP-1/TIF1 β was confirmed in ES cells (see Figure S1A available online).

Zfp57 Is Expressed in Oocytes and in a Subset of Adult Tissues

As expected from our original screen, the transcription of *Zfp57* is downregulated when ES cells differentiate (Ahn et al., 2004; Akagi et al., 2005; Li and Leder, 2007). ES cells contain two prominent *Zfp57* transcript isoforms (Akagi et al., 2005; Okazaki et al., 1994). *Zfp57* also displays a restricted expression pattern in adult mouse organs, with the highest level of expression in testis and ovary and the lowest levels in lung and brain (Figure 1D). The longer transcript appears to be the main product

in lung and brain, and the short transcript is the predominant form in ovary. Testis has a third, intermediate-sized transcript isoform. *Zfp57* transcripts appear to be present at all postimplantation stages (Figure S1B).

RNA in situ hybridization demonstrates specific expression of *Zfp57* in oocytes within adult wild-type mouse ovaries (Figure 1E). Interestingly, no other cell types in the ovary, including the follicle cells surrounding the oocytes, express any detectable level of *Zfp57* transcripts.

Embryonic and Neonatal Lethality of *Zfp57* Mutants Indicates Distinct Zygotic and Maternal-Zygotic Effects

We generated a deleted null allele in the *Zfp57* locus (see Supplemental Data and Figure S2). From crosses between female and male heterozygous mice, only 7 out of 65 progeny mice (10.8%) at the time of weaning were homozygous mutants, which is less than half of the expected 25%. Likewise, from the crosses between female heterozygous mice and male homozygous mice, only half of the expected homozygotes were obtained (Table S1). This loss of homozygous offspring deficient in zygotic, but not maternal, *Zfp57* mainly occurred at the perinatal or neonatal stage (Table 1). These data indicate that loss of the zygotic function of *Zfp57* results in partial lethality (Table S1).

In contrast to the crosses generating zygotic mutants for *Zfp57*, mating homozygous females to heterozygous or homozygous males resulted in no homozygous animals surviving to weaning (Table S1). Analysis of prenatal lethality in these crosses indicated that lethality of homozygous embryos commenced around midgestation and progressively increased such that by E14.5–E16.5 ~80% of null embryos derived from null female mice were dead (Table 1). This earlier lethality of homozygotes from null female mice compared to those from heterozygous female mice indicates a maternal effect of *Zfp57* on the survival of animals (Figure S3). Crosses with oocytes derived from conditionally ablating *Zfp57* in female mice with ZP3-driven Cre-recombinase resulted in the same outcome, proving the maternal effect of the deletion (Table S2). Interestingly, loss of only the maternal or the paternal function of *Zfp57* did not cause either embryonic or neonatal lethality, indicating zygotic rescue (Table 1).

Defective Imprinting in *Zfp57* Maternal-Zygotic Mutants

Microarray analysis comparing transcript expression patterns between the live homozygous E11.5 embryos and heterozygous littermates in a 129 Sv/Ev background derived from the same homozygous (null) female mouse crossed to a heterozygous male was conducted. One of the genes most strongly affected was the imprinted *Dlk1* gene. Data indicate that expression of *Dlk1*, a paternally expressed mRNA gene, is dramatically reduced (six-fold decrease) in the homozygous mutant embryos. By contrast, the adjacent *Gtl2* gene, a noncoding RNA gene expressed from the maternally inherited chromosome, is expressed at a two-fold increased rate in homozygous mutant embryos. Downregulation of *Dlk1* transcripts was confirmed by northern blot analysis (Figure 2A) and quantitative RT-PCR (X.L. and P.L., unpublished data). *Dlk1* and *Gtl2* are members of a cluster of imprinted genes that also include *Rtl1* and *Dio3* expressed from the paternally inherited chromosome (Lin et al., 2003). Both *Rtl1* and *Dio3* (which is over 800 kb away from *Dlk1*) are downregulated in the homozygous mutant embryo compared

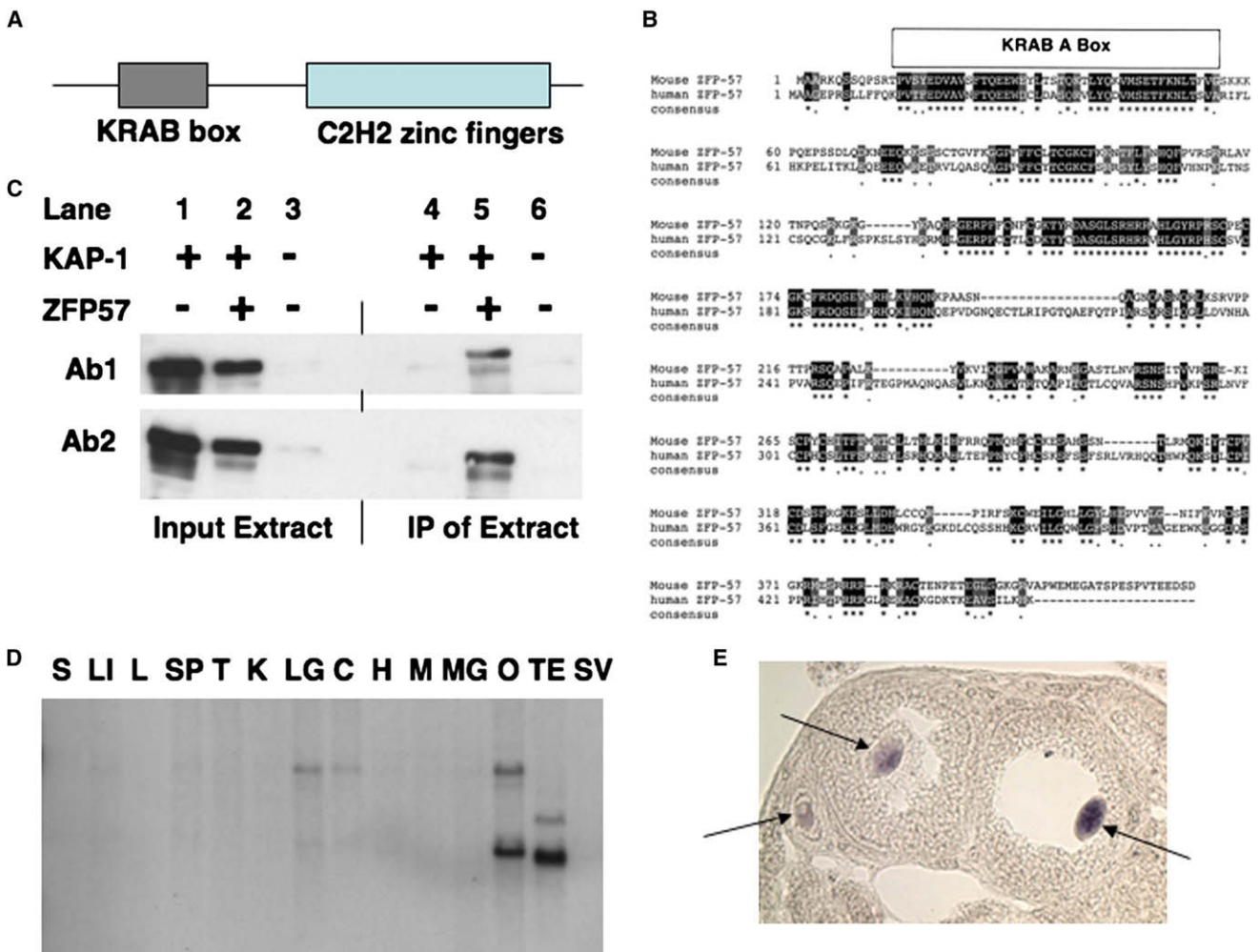


Figure 1. ZFP57 Is a Putative KRAB Zinc Finger Protein

(A) Schematic diagram of the ZFP57 protein.

(B) Sequence alignment between the mouse ZFP57 and the human ZFP57 proteins. The identical amino acids are boxed in black, and the similar residues are shaded.

(C) Coimmunoprecipitation assays were carried out for KAP-1 and myc epitope-tagged ZFP57. The three left lanes are the western blot of the total cell lysate ("Input") from the KAP-1-transfected (lane 1), the KAP-1 and myc epitope-tagged ZFP57 cotransfected (lane 2), and the untransfected cos cells (lane 3). Lanes 4–6 are the western blot of the immunoprecipitate (IP) derived from these three samples when the antibodies against myc epitope were used to pull down ZFP57-associated proteins. The rabbit polyclonal antibodies used for the western blot are Ab1 (anti-KAP-1 RBCC) and Ab2 (anti-KAP-1 CT) (Schultz et al., 2001).

(D) Adult mouse organ northern blot of *Zfp57*. A 1.2 kb cDNA fragment encompassing the entire open reading frame of *Zfp57* was used to probe a northern blot. An equal amount of polyA RNA was loaded on each well (Chester et al., 1998) and was prepared from the various organs as follows: S, stomach; LI, large intestine; L, liver; SP, spleen; T, thymus; K, kidney; LG, lung; C, cerebrum; H, heart; M, muscle; MG, mammary gland; O, ovary; TE, testis; SV, seminal vesicle.

(E) RNA in situ hybridization reveals that *Zfp57* is expressed specifically in the maturing oocytes (purple stain). Frozen sections of the wild-type ovary were probed with an antisense riboprobe derived from a 0.5 kb fragment of the 5' portion of the *Zfp57* cDNA. Arrows point to the labeled oocytes inside the cavity of the follicles.

with control heterozygous littermates (Figures 2B and 2C). Consistent with the microarray data, the amount of *Gtl2* transcripts in homozygous mutant embryos is about twice that in the heterozygous littermate control embryos (Figure 2D). Because all of the active alleles of all of the protein-coding genes become repressed on the paternally inherited chromosome, concurrent with an increase in the expression of the normally paternally repressed *Gtl2*, we deduced that genomic imprinting across the whole *Dik1-Dio3* imprinted domain may be perturbed in homozygous mutant embryos derived from a null female

mouse. Like *Dik1-Dio3*, the *Igf2-H19* domain is regulated by an ICR that acquires methylation during spermatogenesis. Both microarray and northern blot analysis indicated that expression at *Igf2-H19* was not affected in *Zfp57* mutants (data not shown).

Loss of *Zfp57* Affects the Maintenance of DNA Methylation Imprints

Germline-derived differentially methylated regions (DMRs) at imprinting control centers are required for genomic imprinting (Kobayashi et al., 2006; Spahn and Barlow, 2003). At the

Table 1. Embryonic and Neonatal Lethality of *Zfp57* Mutant Progeny Indicates Distinct Zygotic and Maternal-Zygotic Effects

Cross	Stage	Percentage of Dead Progeny/Total	Percentage of Dead $-/-$ Mutant/Total $-/-$	Type of Mutant
$+/-$ (f) \times $+/-$ (m)	E18.5	4.7% (n = 85)	16.7% (n = 18)	zygotic
	P1 pup	9.8% (n = 92) ^a	45% (n = 20) ^a	zygotic
$+/-$ (f) \times $-/-$ (m)	E18.5	0% (n = 32)	0% (n = 12)	zygotic
	P1 pup	20.2% (n = 84) ^a	39.5% (n = 43) ^a	zygotic
$-/-$ (f) \times $+/(m)$ ^b	E18.5	0% (n = 16)	NA	NA
	P1 pup	0% (n = 22)	NA	NA
$+/(f)$ \times $-/-$ (m) ^c	E18.5	3% (n = 33)	NA	NA
	P1 pup	0% (n = 12)	NA	NA
$-/-$ (f) \times $+/-$ (m)	E17.5-18.5	34.8% (n = 23)	80% (n = 10)	
	E11.5-E13.5	9.8% (n = 34)	38.1% (n = 21)	Mat-Zyg
$-/-$ (f) \times $-/-$ (m)	E17.5-19.5	87.5% (n = 8)	87.5% (n = 8)	Mat-Zyg
	E14.5-E16.5	81.4% (n = 59)	81.4% (n = 59)	Mat-Zyg
	E11.5-E13.5	43.4% (n = 83)	43.4% (n = 83)	Mat-Zyg
	E9.5-E10.5	7.9% (n = 38)	7.9% (n = 38)	Mat-Zyg

Embryos of the different embryonic stages were dissected out from the pregnant female mice in the mixed genetic background. Similar results were obtained with the mice in the 129 Sv/Ev background (Li and Leder, data not shown). Live embryos were confirmed under the dissection microscope by the heartbeat and body movement of the embryos. $+/-$ (f), heterozygous female mouse; $+/-$ (m), heterozygous male mouse; $-/-$ (f), homozygous mutant female mouse; $-/-$ (m), homozygous mutant male mouse; $+/(f)$, wild-type female mouse; $+/(m)$, wild-type male mouse; NA, not applicable; Mat-Zyg, maternal-zygotic mutant.

^aTotal dead P0 or P1 pups.

^bAll progeny from this cross are heterozygous, and they lost just the maternal function of *Zfp57*.

^cAll progeny from this cross are heterozygous, and they lost just the paternal function of *Zfp57*.

Dlk1-Dio3 imprinted domain, the germline DMR resides in the intergenic region between *Dlk1* and *Gtl2* (IG-DMR). In the absence of the IG-DMR, imprinting is lost on the maternally inherited chromosome (Lin et al., 2003). Methylation of the IG-DMR on the paternal chromosome has been shown to be required for *Dlk1* expression because, in the absence of the maintenance methyltransferase *Dnmt1*, *Dlk1* becomes downregulated and *Gtl2* levels increase (Schmidt et al., 2000). The methylation status of the IG-DMR in *Zfp57* mutants was therefore assessed. In the *Zfp57* homozygous mutant embryos, DNA methylation at the IG-DMR is indeed affected when genomic DNA samples of different classes of embryos were analyzed (left panel of Figure 3A; Figures S4A and S6A). Figure 3A illustrates relative differential DNA methylation levels quantified from Southern blots, an example of which is shown. Consistent with its requirement for *Dlk1* activity, differential DNA methylation is completely absent at the IG-DMR in the homozygous mutant embryos derived from a null mother when both the maternal as well as the zygotic *Zfp57* is eliminated (lane 5 of Figure 3A, 0%). This indicates that methylation of the paternal chromosome cannot be maintained in the absence of both maternal and zygotic *Zfp57* (maternal-zygotic effect). By contrast, loss of only the maternal function (maternal effect) in the heterozygous embryos derived from a null mother does not appear to cause any significant loss of differential DNA methylation at the IG-DMR (lane 4 of Figure 3A, 90.9%), indicating that depletion of maternal *Zfp57* alone is not sufficient to affect methylation maintenance. We also observed partial loss of differential DNA methylation in the homozygous mutant embryos derived from a heterozygous mother when only the zygotic function of *Zfp57* is missing (zygotic effect, lane 3 of Figure 3A, 67.5%). Differential methylation at the pro-

motor of *Gtl2* (*Gtl2*-DMR), which is regulated by methylation at the IG-DMR after fertilization, was similarly affected (Figure S4B). These results suggest that paternally derived germline methylation cannot be maintained in embryos lacking both maternal and zygotic *Zfp57*. However, providing this factor to a null zygote via the oocyte is sufficient to maintain at least some methylation at the domain.

Other imprinted domains were surveyed to see if differential DNA methylation at ICRs was affected in the *Zfp57* homozygous mutant embryos. Differential DNA methylation in the *H19* DMR (Tremblay et al., 1997) is normal in the *Zfp57* homozygous mutant embryos (Figure S4C). Methylation at two families of repeat sequences (*Line1* and *IAP*) was also unaffected (Figure S5).

DMRs that acquire their methylation in the maternal germline were assessed. These include *Snrpn*, *Peg1*, *Peg3*, *Igf2r*, and *Nnat/Peg5* (Lucifero et al., 2002). Similar to the finding in the *Dlk1-Dio3* imprinted region, differential DNA methylation was lost at the *Snrpn*, *Peg1*, *Peg3*, and *Peg5/Nnat* DMRs in all homozygous mutant embryos (maternal-zygotic) derived from a null mother (Figure 3B; Figure S6; and data not shown). Partial loss of DNA methylation at the DMRs was observed when only zygotic *Zfp57* is missing, and the extent of this varied between individuals (Figure S6). Differential DNA methylation is intact in the heterozygous embryos derived from a heterozygous mother. Methylation imprints at the *Igf2r* ICR were generally not affected in the homozygous embryos, except in one embryo lacking both maternal and zygotic *Zfp57* (Figure S6D).

These findings indicate that, in addition to failing to maintain imprints at *Dlk1-Dio3*, loss of both maternal and zygotic *Zfp57* results in failure to maintain methylation imprints at multiple maternally methylated DMRs. Zygotic *Zfp57* is able to rescue

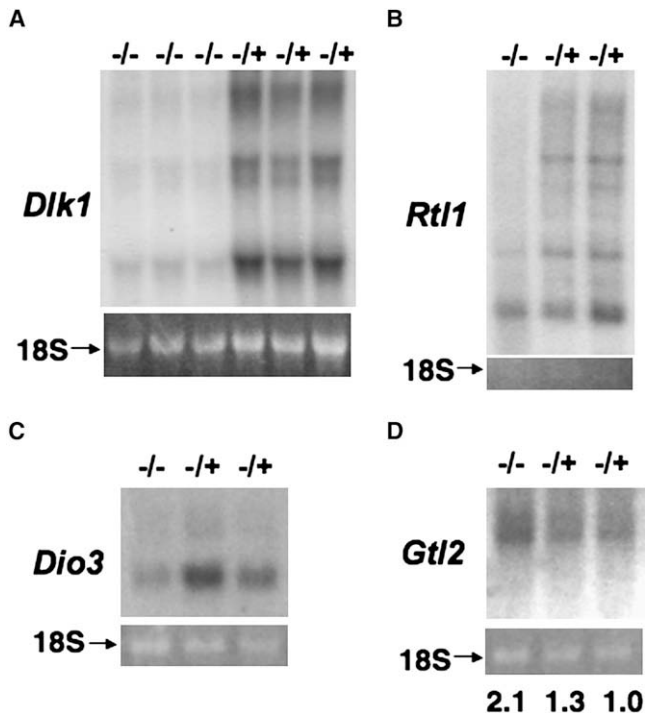


Figure 2. Loss of *Zfp57* Affects Expression of the Genes in the *Dlk1-Dio3* Imprinted Region

(A) Total RNA samples from three homozygous ($-/-$) and three littermate heterozygous ($-/+$) embryos derived from the same null mother were probed with *Dlk1* cDNA in northern blot.

(B–D) Equal amounts of polyA RNA samples from a homozygous ($-/-$) and two littermate heterozygous ($-/+$) embryos derived from the same null mother were hybridized with a probe derived from *Rtl1*, *Dio3*, or *Gtl2*, respectively. Numbers in (D) indicate the band intensity of *Gtl2* transcripts in each lane based on phosphor image analysis. Similar band intensity of 18S ribosomal RNA shown in the bottom panels of all figures indicates equal loading of the RNA samples.

All embryos are in the pure 129 Sv/Ev background.

loss of maternal *Zfp57* to allow imprint maintenance. Absence of zygotic *Zfp57* can sometimes compromise the maintenance of methylation conferred by maternal *Zfp57*. Therefore, for effective maintenance of methylation, both maternal and zygotic *Zfp57* are required.

Zfp57* Is Required for the Establishment of Maternal Methylation Imprints at *Snrpn

Given the absence of methylation at multiple maternally methylated ICRs, it was important to determine whether maternal germline methylation marks were appropriately established. Using both COBRA analysis and bisulphite sequencing, we determined that the *Snrpn* DMR is not methylated in oocytes derived from homozygous female mice (Figure 4B; Figures S7 and S8). In contrast, the *Snrpn* DMR showed normal methylation in control oocytes from heterozygous females (Figure 4A; Figures S7 and S8). Thus, *Zfp57* is required for the establishment of the maternal methylation imprint at *Snrpn* in the female germline.

Germline methylation marks were assessed at four other maternally methylated ICRs (Figure S7). Unlike *Snrpn*, germline methylation was found to occur normally at the DMR regions

of *Peg1*, *Peg3*, and *Nnat/Peg5* in unfertilized oocytes derived from either homozygous or heterozygous female mice. Methylation imprints appear to be partially affected at *Igf2r* in the oocytes produced by homozygous or heterozygous female mice; however, the significance of this is not clear.

Assessment of paternal germline methylation at the IG-DMR and *Igf2-H19* ICRs was conducted in DNA isolated from sperm purified from heterozygous and homozygous mutant males. All samples exhibited normal methylation marks for both regions (Figures S9), indicating that *Zfp57* does not affect the establishment of these paternal germline methylation imprints. Together, these data suggest that maternal *Zfp57* is required for the establishment of the maternal methylation imprint at the *Snrpn* imprinted region, but is dispensable for *Peg1*, *Peg3*, and *Nnat/Peg5* imprinting in the female germline. Consistent with the genetic outcome indicating a maternal-zygotic effect for *Zfp57*, *Zfp57* is not required for the establishment of paternal germline methylation. The data also indicate that *Zfp57* is not a general maternal germline imprinting factor.

Postimplantation Acquisition of Differential Methylation at the *Snrpn* DMR

Homozygous embryos generated from homozygous females are not methylated at the *Snrpn* DMR. This absence of methylation is heritable at preimplantation stages, as both heterozygous ($-/+$) and homozygous ($-/-$) E3.5 embryos derived from null females are unmethylated at the *Snrpn* DMR (Figure 4C; Figure S8C). As expected, the *Snrpn* DMR was methylated in heterozygous ($+/-$) E3.5 embryos derived from wild-type female mice (Figure 4C). Surprisingly, however, methylation was evident at *Snrpn* in approximately half of the heterozygous midgestational embryos derived from homozygous female mice (Figures 3B, 4D, and 4E). These findings suggest that the acquisition of methylation at the *Snrpn* DMR in heterozygous midgestational embryos derived from null female mice requires zygotic *Zfp57* and occurs after E3.5.

Interspecies crosses were carried out to determine whether postfertilization DNA methylation at *Snrpn* was acquired specifically on the maternally inherited chromosomes or at random on both maternal and paternal chromosomes. Homozygous female mice (129 Sv/Ev) were mated with wild-type male mice (DBA/2). Polymorphisms exist between these two strains at the *Snrpn* DMR (Hiura et al., 2006) (Figure S10). Based on the bisulphite sequencing results of eight heterozygous embryos from this cross (Figure 4E), methylation was only observed at the 129 Sv/Ev *Snrpn* DMR, i.e., the maternally derived DMR, but not at the paternally derived DBA/2-specific region (see Figure S11 for detailed information). As expected, differential methylation was maintained on the maternally derived *Snrpn* DMR in the control cross between 129 Sv/Ev female and DBA/2 male wild-type mice (Figure S12).

Endogenous ZFP57 Binds to the *Snrpn* DMR

Affinity-purified rabbit polyclonal antibodies against the N-terminal half of ZFP57 protein were generated. These purified antibodies specifically detect exogenous ZFP57 expressed in COS cells and endogenous ZFP57 present in wild-type ES cells. Immunoreactivity is absent in null ES cells (Figure 5A).

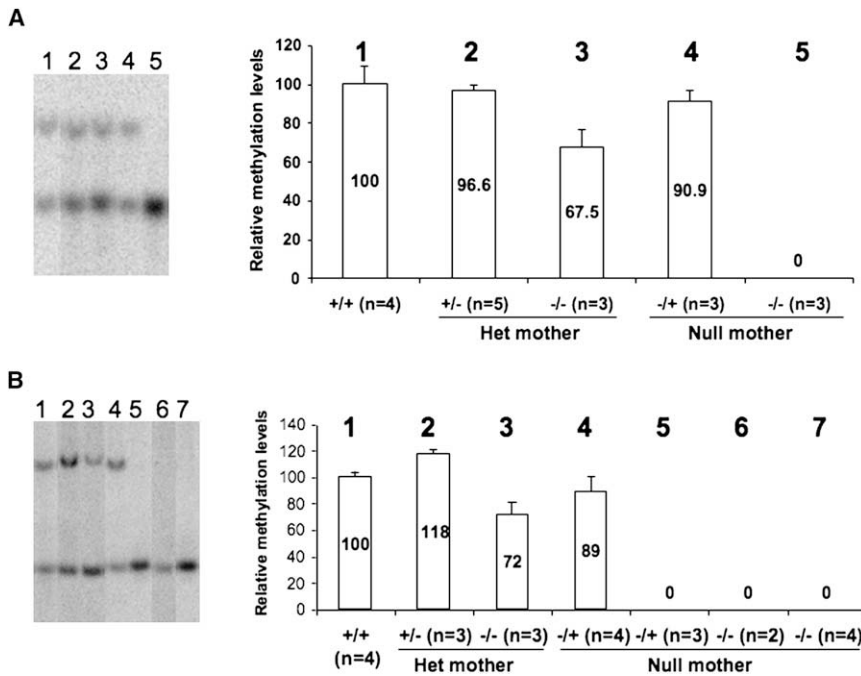


Figure 3. Loss of *Zfp57* Affects Differential DNA Methylation at the *Dlk1-Dio3* and the *Snrpn* Imprinted Regions

(A and B) Differential DNA methylation was assessed with the methylation-sensitive restriction enzyme HhaI. (A) StuI-digested and (B) PstI-digested genomic DNA samples from embryos were HhaI digested and hybridized with a probe derived from the differentially methylated regions by using Southern blot analysis (left panel). Histograms represent relative differential methylation levels based on quantitative phosphor image analysis of the band intensity of the Southern blots shown on the left. The numbers above each bar indicate the representative lane on the Southern blot. Numbers within the bars indicate the percentage of differential DNA methylation relative to that of wild-type embryos. Zero indicates an absence of methylation. Error bars are standard error. Genotypes and numbers analyzed for each class of embryo are indicated along the horizontal axis. All embryos are between E11.5 and E13 on a 129 Sv/Ev background. (A) The IG-DMR from the *Dlk1-Dio3* region. Lane 1, wild-type embryos; lane 2, heterozygous (+/-) embryos from a heterozygous (het) mother; lane 3, homozygous (-/-) embryos from a het mother; lane 4, heterozygous (-/+) embryos from a homozygous (null) mother;

lane 5, -/- embryos from a null mother. (B) The *Snrpn* DMR. Lane 1, wild-type embryos; lane 2, +/- embryos from a het mother; lane 3, -/- embryos from a het mother; lanes 4 and 5, -/+ embryos from a null mother; lane 6, -/- embryos from a null mother and a het father; lane 7, -/- embryos from a null mother and a null father. These *Snrpn* Southern data were further confirmed by COBRA analysis on samples including a subset analyzed by Southern blotting. A total of 25 midgestation heterozygous embryos from null mothers were assessed; 14 exhibited methylation acquisition, and 11 remained unmethylated (data not shown).

To determine whether ZFP57 directly interacts with the *Snrpn* DMR, chromatin immunoprecipitation (ChIP) was performed to determine whether ZFP57 binds this DMR. Mouse ES cells were used as a model system because they display high de novo methylation activities (Humpherys et al., 2001; Lei et al., 1996; Ooi et al., 2007). Furthermore, *trans*-acting factors that bind to the control elements within the *Snrpn* DMR are present in undifferentiated ES cells (Kantor et al., 2004). Analysis of DNA purified after ChIP revealed binding of endogenous ZFP57 to the *Snrpn* DMR in wild-type ES cells and in the ES cells containing one floxed allele and one targeted allele at the *Zfp57* locus (lanes 1 and 2 of Figure 5C), but not to the *Snrpn* DMR of ES cell clones containing two deleted alleles of *Zfp57* (lanes 3 and 4 of Figure 5C). These experiments, which were reproducible for three independent immunoprecipitations, indicate that ZFP57 can directly bind to the *Snrpn* DMR, whereas the control ChIP experiments involving the unaffected *H19* DMR or a distant upstream region of the *Snrpn* DMR did not show any binding activity (Figures 5D and 5E).

Maternal ZFP57 Is Present in Preimplantation Embryos

Embryos isolated from several different crosses were subjected to immunofluorescence staining. As shown in Figures 6A and 6D, there is no detectable ZFP57 present in the homozygous E1.5 or E3.5 embryos derived from a null mother. However, in the embryos derived from the cross between heterozygous female mice and homozygous male mice, ZFP57 was detected in the homozygous E1.5 and E3.5 embryos (Figures 6B and 6G) as well as in the littermate heterozygous E1.5 and E3.5 embryos (Figures 6C and 6F). These data suggest that the maternal

gene product of *Zfp57* is carried over from oocytes and is deposited in early embryos, as illustrated in Figure S3. Zygotic ZFP57 was detectable in the heterozygous E3.5 embryos from the cross between homozygous female mice and wild-type male mice (Figure 6H).

DISCUSSION

Mechanisms of establishment and maintenance of germline-derived methylation imprints are not fully understood. In a screen for genes downregulated upon ES cell differentiation, we identified a maternal-zygotic effect gene, *Zfp57*, that contributes to the stable maintenance of methylation imprints during development.

Consistent with its expression in the female germline, *Zfp57* is required for the establishment of the germline methylation imprint at the *Snrpn* imprinted region. More detailed biochemical analysis needs to be undertaken to determine whether ZFP57 is directly involved in the acquisition or maintenance of this methylation in the female germline. In contrast, maternal *Zfp57* is not essential for the establishment of the genomic imprints at other examined maternally methylated regions (*Peg1*, *Peg3*, and *Nnat/Peg5*) in oocytes, or in the male germline. Consistent with the maternal-zygotic effect evident from the genetic data, a role for *Zfp57* in imprinting establishment appears to be specific to the female germline.

In contrast, maternal *Zfp57* plays a much broader role in the maintenance of both paternal and maternal methylation imprints in embryos. Genomic imprints are not maintained at the paternally methylated IG-DMR of the *Dlk1-Dio3* domain and at maternally methylated *Peg1*, *Peg3*, and *Nnat/Peg5* in the absence of

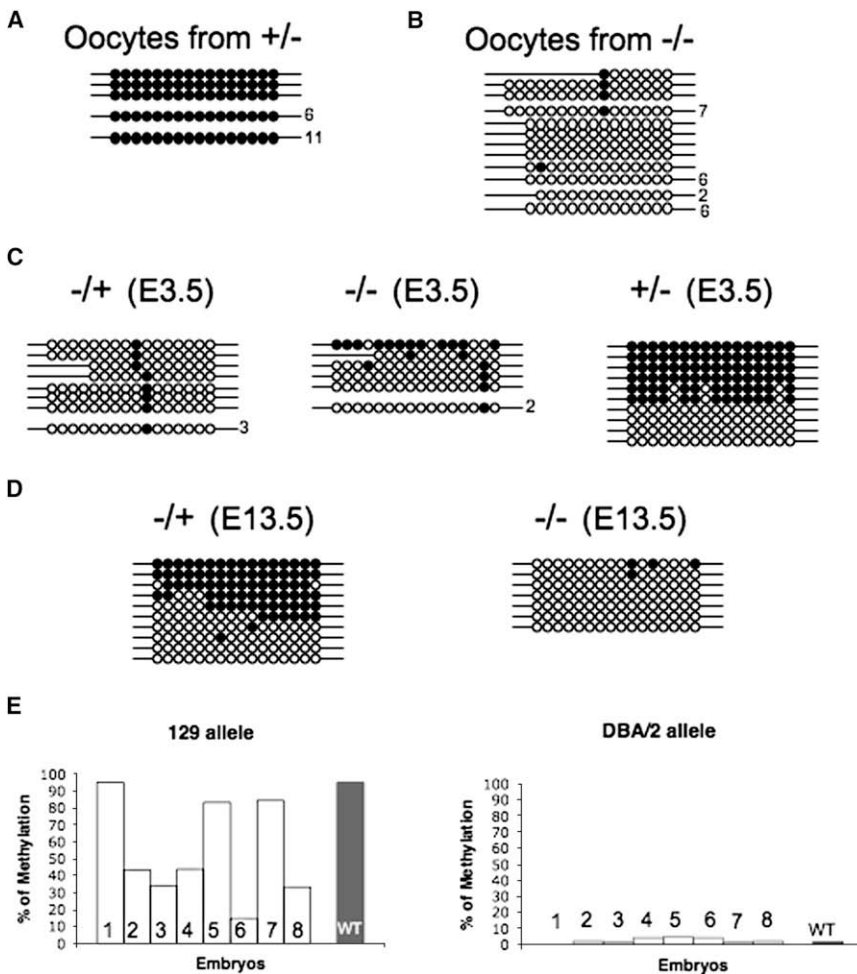


Figure 4. Differential Methylation at the *Snrpn* DMR Is Not Established in the Absence of the Maternal *Zfp57* and Can Be Acquired in the Presence of the Zygotic *Zfp57*

(A–D) Genomic DNA samples from oocytes or embryos in the 129 Sv/Ev background were subjected to bisulphite sequencing. A total of 16 differentially methylated CpG sites at the *Snrpn* DMR are shown for (A)–(D). Filled oval, methylated CpG site; open oval, unmethylated CpG sites; line with ovals, a unique clone. A unique clone was assigned based on unconverted C residues in the sequences. The number of sequences is shown for a nonunique clone. The high conversion rate fails to distinguish whether these nonunique sequences represent clonal or individual products. (A) Two different pools of unfertilized oocytes isolated from heterozygous female mice. (B) Two different pools of unfertilized oocytes isolated from homozygous female mice. (C) Pooled E3.5 embryos. -/+ (E3.5), 29 heterozygous E3.5 embryos from the crosses between homozygous female mice and wild-type male mice. -/- (E3.5), ten homozygous E3.5 embryos from the crosses between homozygous female mice and homozygous male mice. +/- (E3.5), 16 heterozygous E3.5 embryos from the crosses between wild-type female mice and homozygous male mice. (D) Genomic DNA samples were made from littermate E13.5 embryos from the cross between a homozygous female mouse and a heterozygous male mouse. (E) Acquisition of DNA methylation imprint occurred on the maternally derived *Snrpn* DMR. Histograms are shown for the levels of methylation at the CpG sites of this DMR in E11.5–E13 embryos from the cross between a homozygous female mouse in the 129 Sv/Ev background and a wild-type male mouse in the DBA/2 background. Unfilled bars, eight littermate heterozygous embryos (1–8) from this cross. Filled

bar, seven littermate wild-type (WT) embryos from the cross between a wild-type 129 Sv/Ev female mouse and a wild-type DBA/2 male mouse. Vertical axis, percentage of methylated CpG sites analyzed. Left panel, maternally derived 129 Sv/Ev allele; right panel, paternally derived DBA/2 allele.

both maternal and zygotic functions of *Zfp57* in embryos. These data indicate that ZFP57 plays a role in the postfertilization maintenance of genomic imprints at a large subset of the imprinted regions. Indeed, it was just published online that zygotic mutations in human *Zfp57* also affect differential DNA methylation at a subset of imprinted regions (Mackay et al., 2008). Variable effects of this mutation in different individuals are consistent with the defects we observe in the zygotic mutant mice. Another maternal effect gene, *Stella/PGC7*, appears to play a partially protective role in the maintenance of the other two paternally derived methylation imprints at *H19* and *Rasgrf1* as well as a role in the maintenance of three maternally derived methylation imprints at *Peg1*, *Peg3*, and *Peg10* (Nakamura et al., 2007). In contrast to *Zfp57*, *Stella/PGC7* does not affect *Dlk1-Dio3*, *Snrpn*, and *Peg5* or play a role in the germline establishment of imprints. The overlapping effect of *Zfp57* and *Stella* at *Peg1* and *Peg3* suggests that their maintenance functions may not be mutually exclusive.

Interestingly, differential DNA methylation at the *Snrpn* DMR was established in the presence of the zygotic *Zfp57* in postimplantation embryos even though germline methylation was either

not established or lost in oocytes and remained absent at E3.5 (Figure S13). Since ZFP57 can bind to the *Snrpn* DMR in ES cells, it is likely that ZFP57 can bind to the unmethylated *Snrpn* DMR directly during methylation acquisition in embryos. This acquisition of methylation only occurs at the maternally derived *Snrpn* DMR. This suggests that there may be a DNA methylation-independent genomic imprint that can act as imprinting memory at the *Snrpn* ICR. Previously, it has been shown that the imprinting of some placental genes occurs in the absence of maintenance methylation, likely involving repressive histone modifications (Lewis et al., 2004; Umlauf et al., 2004). It will be interesting to determine whether repressive histone modifications play a role in the heritable methylation-independent germline mark at *Snrpn*. Future experiments will address the many intriguing questions arising from this finding, including the nature of the heritable memory and whether there is a bias for the grandmaternal over the grandpaternal maternal allele. Since some heterozygous embryos derived from null female mice had no or only partial DNA methylation around midgestation, whereas others displayed a fully methylated maternal allele, it is possible that differential methylation at the *Snrpn* DMR in the presence of the zygotic

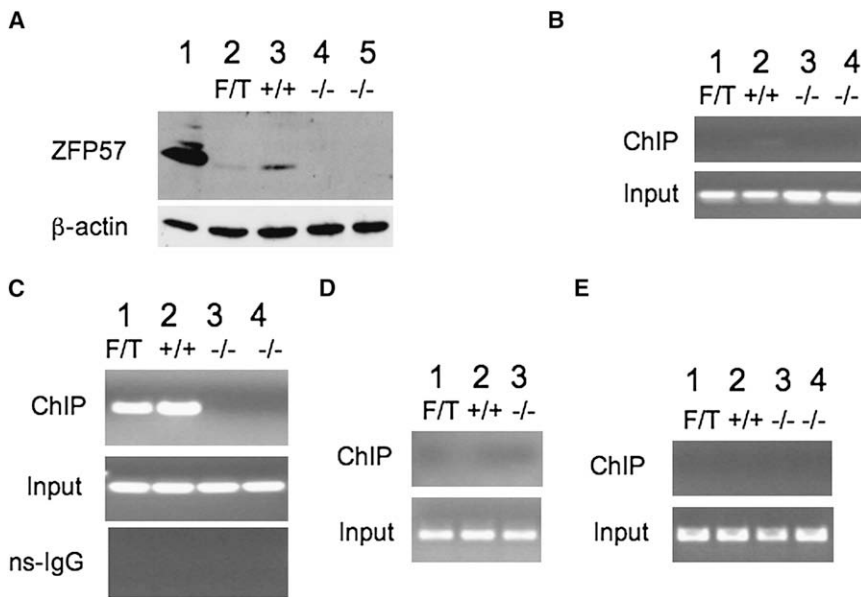


Figure 5. Endogenous ZFP57 Can Bind to the *Snrpn* DMR Region

(A) Western blot analysis of affinity-purified anti-ZFP57 polyclonal antibodies with total cell lysate samples. Lane 1, ZFP57 was overexpressed in COS cells; lane 2, an ES clone containing one floxed allele (F) and one targeted allele (T) at the *Zfp57* locus; lane 3, wild-type TC1 ES cells; lanes 4 and 5, two independent *Zfp57* null ES clones carrying two deleted alleles. Please see Figure S2 and Supplemental Data for the description of the targeted, floxed, and deleted alleles.

(B–E) Chromatin immunoprecipitation (ChIP) was performed with approximately one million ES cells. No ChIP PCR product was observed in the negative control samples without the addition of any antibodies (data not shown). (B) The first-round PCR product at the *Snrpn* DMR region. (C) The second-round PCR product at the *Snrpn* DMR region. (D) The second-round PCR product at a region approximately 40 kb upstream of the *Snrpn* DMR region. (E) The second-round PCR product at the *H19* DMR region. ChIP, ChIP PCR product; Input, PCR product from the input/starting samples; ns-IgG, ChIP PCR product from the control

samples when rabbit nonspecific IgG (ns-IgG) antibodies were added during immunoprecipitation. (B–E) Lane 1, an ES clone containing one floxed allele (F) and one targeted allele (T) at the *Zfp57* locus; lane 2, wild-type TC1 ES cells; lanes 3 and 4, two independent *Zfp57* null ES clones carrying two deleted alleles.

Zfp57 occurs over time in embryos, and all progeny from this cross may acquire full differential methylation at the maternal allele in late stages of development.

The effects of ZFP57 on imprint methylation are consistent with the lethality of embryos described for the range of genetic crosses: (1) loss of just zygotic *Zfp57* causes partial neonatal lethality as well as partial loss of differential DNA methylation; (2) loss of only the maternal *Zfp57* does not appear to cause any lethality or any loss of differential DNA methylation due to the rescue of the maternal effect by zygotic *Zfp57*; (3) eliminating both the maternal and zygotic *Zfp57* results in complete loss of

differential DNA methylation as well as a highly penetrant embryonic lethality around midgestation. This lethality is consistent with phenotypes expected from the cumulative effects of loss of imprinting of a subset of imprinted genes, although *Zfp57* may also play a role at some unidentified nonimprinted loci.

How might this KRAB zinc finger protein be influencing the epigenetic state of imprinted domains? Interestingly, it was recently reported that the KRAB domain can trigger heritable de novo DNA methylation when targeted to a reporter transgene specifically in early mouse embryos (Wiznerowicz et al., 2007). This is consistent with an endogenous role for the KRAB zinc finger protein ZFP57 in

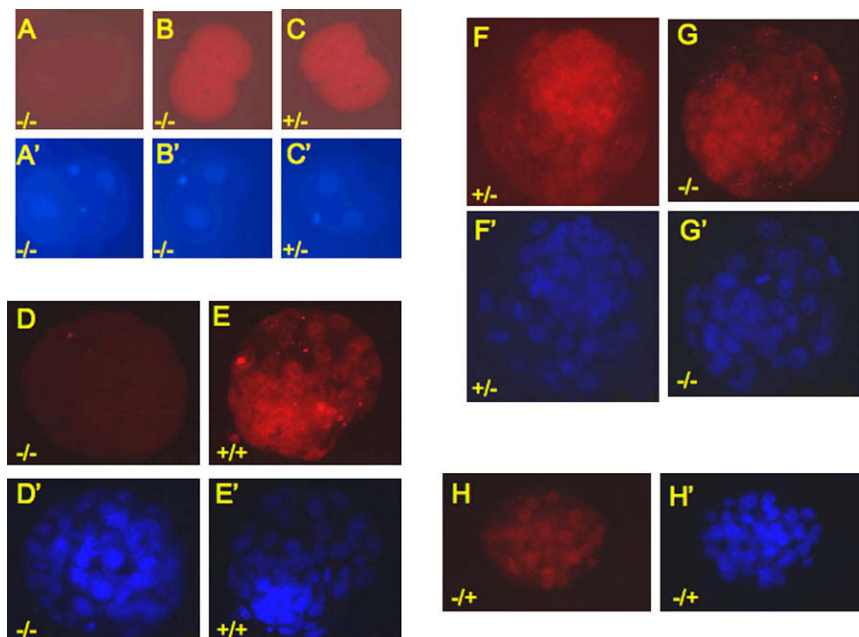


Figure 6. Maternal ZFP57 Is Present in Preimplantation Embryos

(A–H) Preimplantation embryos were stained with affinity-purified polyclonal antibodies against ZFP57; (A'–H') they were also incubated with Hoechst dye to illuminate genomic DNA. (A and A') homozygous E1.5 embryo and a (D and D') homozygous E3.5 embryo were derived from the crosses between homozygous female mice and homozygous male mice. Littermate (B and C) E1.5 embryos and (F and G) E3.5 embryos were derived from the crosses between heterozygous female mice and homozygous male mice. (B and G) A homozygote (–/–). (C and F) A heterozygote (+/–). (E) The wild-type E3.5 embryo was derived from the cross between a wild-type female mouse and a wild-type male mouse. (H) The heterozygous (–/+) E3.5 embryo was derived from the cross between a homozygous female mouse and a wild-type male mouse.

mediating DNA methylation at some imprinted domains. We therefore suggest a model in which ZFP57 might be directly involved in targeting methylation to certain ICRs: in the female germline at *Snrpn* and at several imprinted ICRs after fertilization. This implies that the maintenance of imprints during early development is a regulated event and more than a mere protection against active and passive demethylation. It is possible that ZFP57 can also bind to the methylated *Snrpn* DMR and other imprinted regions. Future experiments focusing on the functions of ZFP57 and associated complexes may elucidate a novel pathway of epigenetic control involving zinc finger proteins and lead to a better understanding of the underlying mechanisms of the establishment and maintenance of DNA methylation.

EXPERIMENTAL PROCEDURES

Targeting at the *Zfp57* Locus

A genomic fragment of *Zfp57* was used to make the targeting construct (see [Supplemental Data](#)). The linearized construct was electroporated into ES cells, and candidate targeted ES clones after G-418 drug selection were screened by Southern blot. A confirmed targeted ES clone was used to generate germline-transmissible chimera mice, and corresponding floxed and deleted alleles at the *Zfp57* locus were obtained.

Bisulphite Mutagenesis and PCR Primers

The protocol for bisulphite mutagenesis is described by [Arnaud et al. \(2003\)](#). PCR primers used for the *Snrpn* DMR and the IG-DMR are the same as those used by [Lucifero et al. \(2002\)](#). For other primers, please see [Supplemental Data](#). Prior to cloning for bisulphite sequencing, sample aliquots were tested by combined bisulphite restriction analysis (COBRA) for later comparison with sequenced products to rule out cloning bias. For the oocyte-specific *Snrpn* bisulphite analysis, two biological replicates on pools of oocytes derived from heterozygote ($n = 311$ from 12 females, and $n = 174$ from 8 females) and homozygote females ($n = 312$ from 7 females, and $n = 185$ from 8 females) were subjected to both bisulphite sequencing and COBRA analysis. For E3.5 embryos, numbers are indicated in the legend to [Figure 4](#); additional pooled embryos are assessed by COBRA analysis as described in [Figure S8](#).

Methylation-Sensitive Southern Blot Analysis

DNA was isolated, double digested, blotted, and hybridized as described by [Takada et al. \(2002\)](#), including probes for the *Gtl2* promoter and IG-DMR. For the probe of the *Snrpn* DMR, please see [Supplemental Data](#). For Southern blotting, a total of 17 heterozygotes and 13 homozygotes from heterozygous female mice plus 16 heterozygotes and 16 homozygotes from null female mice were generated. DNA was used for multiple probe hybridizations with sample overlap between the *Gtl2*-DMR and IG-DMR and the *Snrpn* and *H19* DMRs. In addition, there was overlap between samples used for Southern blot analysis and for COBRA analysis. For example, of 7 heterozygotes from null female mice analyzed for the *Snrpn* DMR by Southern blotting, 4 (plus an additional 18) were assessed by COBRA. All genotype-specific methylation analysis by Southern blotting was confirmed by COBRA analysis, although not necessarily on all the same samples. Genotype-specific methylation was consistent for each genotype when using these two different approaches.

Northern Blot Analysis

The polyA⁺ northern blot used in [Figure 1D](#) was generated with RNA isolated from adult organs and used previously ([Chester et al., 1998](#)) and was repeated with fresh tissues, which gave the same results (data not shown). Either total RNA or polyA⁺ RNA derived from midgestation embryos was used for northern blots on the imprinted genes shown in [Figure 2](#). Probes were derived from the corresponding cDNAs.

Generation and Purification of Anti-ZFP57

Antibodies and Immunostaining

A portion of *Zfp57* cDNA encoding the N-terminal portion of ZFP57 protein (Met1 to Arg240) was amplified by PCR and was cloned into the NcoI and

BamHI sites of the expression vector pQE60 (QIAGEN). Hexahistidine-tagged protein was induced by the addition of IPTG to the transfected bacterial clones and was purified under denaturing conditions according to the manufacturer's manual. The purified denatured protein was directly injected into the rabbits, and anti-ZFP57 sera were generated by following a standard protocol (Cocalico Biologicals, Inc., USA). Rabbit anti-ZFP57 polyclonal antibodies were purified with the hexahistidine-tagged ZFP57 recombinant protein attached to a Nickel-NTA affinity column ([Gu et al., 1994](#)). After elution with 4 M MgCl₂, the purified polyclonal antibodies were dialyzed against water for 1 hr and then against Tris-buffered saline solution (25 mM Tris [pH 8.0]) exhaustively at 4°C.

Immunofluorescence Staining in Embryos

Embryos were harvested from superovulated female mice and fixed in 4% paraformaldehyde. They were stained with the purified anti-ZFP57 antibodies by following a previously published protocol ([Payer et al., 2003](#)).

Coimmunoprecipitation

Mouse monoclonal antibodies against the myc epitope were used to immunoprecipitate the exogenously transfected myc epitope-tagged ZFP57 in COS cells. Purified rabbit polyclonal antibodies against ZFP57 were used to pull down the endogenous ZFP57 in ES cells.

Chromatin Immunoprecipitation

Chromatin immunoprecipitation was conducted according to the protocol recommended by Upstate (USA). For PCR amplification of immunoprecipitated DNA, please see information regarding PCR primers in [Supplemental Data](#).

SUPPLEMENTAL DATA

Supplemental Data include fourteen figures, two tables, Supplemental Results, Supplemental Experimental Procedures, and Supplemental References and can be found with this article online at [http://www.developmentalcell.com/supplemental/S1534-5807\(08\)00338-9](http://www.developmentalcell.com/supplemental/S1534-5807(08)00338-9).

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