Ino80 is essential for proximal-distal axis asymmetry in part by regulating Bmp4 expression

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Abstract

Background: Understanding how embryos specify asymmetric axes is a major focus of biology. While much has been done to discover signaling pathways and transcription factors important for axis specification, comparatively little is known about how epigenetic regulators are involved. Epigenetic regulators operate downstream of signaling pathways and transcription factors to promote nuclear processes, most prominently transcription. To discover novel functions for these complexes in axis establishment during early embryonic development, we characterized phenotypes of a mouse knockout (KO) allele of the chromatin remodeling Ino80 ATPase.

Results: Ino80 KO embryos implant, but fail to develop beyond the egg cylinder stage. Ino80 KO embryonic stem cells (ESCs) are viable and maintain alkaline phosphatase activity, which is suggestive of pluripotency, but they fail to fully differentiate as either embryoid bodies or teratomas. Gene expression analysis of Ino80 KO early embryos by in situ hybridization and embryoid bodies by RT-PCR shows elevated Bmp4 expression and reduced expression of distal visceral endoderm (DVE) markers Cer1, Hex, and Lefty1. In culture, Bmp4 maintains stem cell pluripotency and when overexpressed is a known negative regulator of DVE differentiation in the early embryo. Consistent with the early embryo, we observed upregulated Bmp4 expression and down-regulated Cer1, Hex, and Lefty1 expression when Ino80 KO ESCs are differentiated in a monolayer. Molecular studies in these same cells demonstrate that Ino80 bound to the Bmp4 promoter regulates its chromatin structure, which correlates with enhanced SP1 binding. These results in combination suggest that Ino80 directly regulates the chromatin structure of the Bmp4 promoter with consequences to gene expression.

Conclusions: In contrast to Ino80 KO differentiated cells, our experiments show that undifferentiated Ino80 KO ESCs are viable, but fail to differentiate in culture and in the early embryo. Ino80 KO ESCs and the early embryo up-regulate Bmp4 expression and down-regulate the expression of DVE markers Cer1, Hex and Lefty1. Based on this data, we propose a model where the Ino80 chromatin remodeling complex represses Bmp4 expression in the early embryo, thus promoting DVE differentiation and successful proximal-distal axis establishment. These results are significant because they show that epigenetic regulators have specific roles in establishing embryonic axes. By further characterizing these complexes, we will deepen our understanding of how the mammalian embryo is patterned by epigenetic regulators.

Keywords: Ino80, Chromatin remodeling, Bmp4, DVE, PD Axis, Gastrulation, Embryonic ectoderm

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Background

Mammalian embryonic development is best understood using the mouse model [1]. Fertilized eggs develop into a blastocyst by embryonic day 3.5 (E3.5). At this stage, the blastocyst is composed of both an inner cell mass (ICM), which is fated to become the embryo, and an extraembryonic trophoderm (TE). The blastocyst implants into the uterus at E4.5, and during this time the embryo forms the epiblast (EPI) and primitive endoderm (PrE). During implantation the EPI rapidly proliferates and expands to form an egg cylinder, which is composed of both embryonic ectoderm (EmE) and extraembryonic ectoderm (ExE) covered by visceral endoderm (VE). During egg cylinder expansion, the distal tip of the embryo differentiates into the distal visceral endoderm (DVE), establishing a proximal-distal (P-D) axis. During the transition from E5.5 to E6.0, the DVE migrates up the anterior portion of the embryo to create the anterior visceral endoderm (AVE) where it has organizer activity through secreting several inhibitors of the Tgfβ and Wnt family ligands. The secretion of these inhibitors constrains Tgfβ and Wnt signaling activity to the posterior portion of the embryo, which promotes the differentiation of the primitive streak and the anterior-posterior (A-P) axis.

Early mammalian development, and the establishment of asymmetric axes (P-D and A-P axes), require coordinated gene expression [2]. Gene expression is regulated by both transcription factors and epigenetic regulators. Both operate within chromatin, which is composed of nucleosomes at its fundamental level [3, 4]. Essential epigenetic regulators in eukaryotes are chromatin remodeling complexes. Chromatin remodeling complexes are usually multi-subunit enzymes that slide or evict nucleosomes, or exchange its histone subunits. These activities change chromatin structure by altering the position, occupancy or composition of nucleosomes [5]. In turn, changes in chromatin structure regulate access to the underlying DNA, that in turn influences nuclear processes like transcription.

Chromatin remodeling complexes are classified into the SWI/SNF, ISWI, CHD or INO80 families based upon the sequence homology of their ATPase subunit [5]. In mammals, the INO80 family is composed of the Srcap, p400, and Ino80 remodeling complexes. These complexes are large, 12-15 subunit, complexes that are unique among chromatin remodeling complexes because they catalyze histone exchange reactions [6]. In addition to histone exchange, Ino80 has significant nucleosome sliding activity, suggesting that it can either alter nucleosome position or occupancy, or change nucleosome composition in vivo. Presumably through these activities, Ino80 regulates a variety of nuclear processes which include transcription, DNA repair, DNA replication, and telomere structure [6].

While much has been done to characterize the nuclear functions of Ino80, little has been done to determine its functions in metazoan development. In plants, INO80 is essential for flowering and reproductive organ development, possibly through functions in homologous recombination and regulated transcription [7, 8]. In flies, INO80 mutants are late embryonic lethal, deregulate Hox gene expression, and manifest homeotic transformations [9]. In addition to Hox genes, INO80 regulates ecdysone response genes, which are essential for pupal development and molting [10]. Localization and nucleosome mapping studies in insect cells show that INO80 is widely distributed throughout the genome and remodels nucleosomes onto energetically unfavorable DNA sequences [11]. Similar to studies in plants and flies, mammalian Ino80 is also essential for development. Ex vivo studies with Ino80 knockout (KO) or shRNA knockdown (KD) pre-implantation embryos show that Ino80 maintains stem self-renewal by promoting the expression of pluripotency factors like Oct4 and stabilizing DNA replication forks [12, 13]. In utero, Ino80 KO embryos implant into the uterus but fail to develop to mid-gestation, possibly due to roles for Ino80 in regulating telomere structure or DNA damage repair [14]. Because Ino80 is a well-documented regulator of gene expression, developmental defects could also result from abnormal gene expression [13].

Building on these earlier studies, we show that Ino80 KO embryos fail to specify a DVE and a P-D axis in utero. Coincident with defects in the DVE, Ino80 KO embryos aberrantly express Bmp4 in the EmE, a known repressor of DVE specification [15]. Molecular studies using differentiating ESC models show that Ino80 is specifically localized to the Bmp4 promoter, remodels its chromatin structure, and regulates the binding of transcription activators to its DNA sequence. These results in combination suggest that Ino80 directly represses Bmp4 expression in the EmE through its chromatin remodeling activity to promote DVE specification and P-D axis establishment.

Results

Ino80 KO embryos fail to develop beyond the egg cylinder stage

To discover the functions of Ino80 in mammalian development, we created a conditional KO allele in mice using Cre-loxp technology [16]. With this strategy, essential exons are flanked by loxp sites that can be excised by tissue-specific Cre recombinase expression. The targeted Floxed-Neo allele (Neo) is designed to delete exons 2-4 of the Ino80 gene with Cre excision, deleting the initiating ATG (Additional file 1a). To create both the Ino80 Floxed allele (F) and the Ino80 deletion allele (KO), we crossed mice carrying our targeted allele to
mice constitutively expressing the Flp or Cre recombinases. Successful recombination and germ line transmission of the Floxed and KO alleles were identified by Southern blotting and PCR-based genotyping strategies (Additional file 1b-d).

To determine if our targeted allele resulted in a loss of Ino80 protein, we crossed a Tet inducible Cre expression system (M2-rTA and TetO-Cre alleles) into our Ino80 Floxed mice [17, 18]. Homozygous Ino80 Floxed (F/F) mouse embryonic fibroblasts (MEFs), with or without TetO-Cre, were isolated from mid-gestation E12.5 embryos. As expected, the addition of doxycycline to TetO-Cre, were isolated from mid-gestation E12.5 embryonic fibroblasts (MEFs), with or without TetO-Cre, were isolated from mid-gestation E12.5 embryos. Successful recombination and germ line transmission of the Ino80 Floxed allele resulted in the conversion of the Ino80 Floxed alleles to Ino80 KO alleles in two days (Additional file 2a). Over this same time course, Western blotting demonstrated a complete depletion of the Ino80 protein by day 6 (Additional file 2b). Coincident with the depletion of the Ino80 protein, we observed a loss of the exons 1–7 Ino80 transcript and the appearance of two smaller Ino80 transcripts (KO-1 and KO-2) by RT-PCR (Additional file 2c, d). Cloning and sequencing these smaller PCR products identified aberrant exon 1-5 and exon 1-6 splice events for the Ino80 transcript, with the deletion of exons 2-4 byCre-mediated recombination (Additional file 2d). In silico translation from the first ATG of each aberrant splice transcript and sequencing these smaller PCR products identified and KO-2) by RT-PCR (Additional file 2c, d). Cloning and sequencing these smaller PCR products identified aberrant exon 1-5 and exon 1-6 splice events for the Ino80 transcript, with the deletion of exons 2-4 by Cre-mediated recombination (Additional file 2d). In silico translation from the first ATG of each aberrant splice product results in an out of frame transcript and a random protein product (Additional file 2e). These results demonstrate that our Ino80 Floxed allele can be successfully excised by Cre recombinase, resulting in loss of the Ino80 protein.

A previous publication showed that MEFs depleted of Ino80 arrest the cell cycle and senesce [14]. To determine if depletion of Ino80 using our Ino80 Floxed allele resulted in similar phenotypes, we counted the number of viable fibroblasts over 12 days post-Cre expression by doxycycline exposure. As described previously [14], we observed that after doxycycline exposure Ino80 KO MEFs stopped proliferating and senesced as identified by an increased number of cells positive for endogenous β-gal activity, an indicator of senescence [19] (Fig. 1a-c). Reproducing previous phenotypes of Ino80 KO MEFs supports the conclusion that our Ino80 KO allele is a null allele.

Phenotypes for Ino80 KO embryos are largely uncharacterized. To identify the functions of Ino80 in mammalian development, we intercrossed mice heterozygous for the Ino80 KO allele. Genotyping litters of mice from this intercross identified no homozygous Ino80 KO mice, demonstrating that Ino80 is essential for mouse viability (Additional file 3). To stage the earliest point where Ino80 is required for development, we genotyped litters of embryos at mid-gestation (E13.5), gastrulation (E6.5), and peri-implantation (E4.5). Results from these experiments showed that Ino80 KO homozygous embryos were absent at E13.5, but present at the expected Mendelian ratios at gastrulation (E6.5) and prior to implantation (E4.5) (Additional file 3). These results demonstrate that Ino80 is essential for mammalian development post-gastrulation. A gross analysis of Ino80 KO embryos prior to (E5.5), during (E6.5), and post-gastrulation (E7.5) revealed that Ino80 is important for expansion of the egg cylinder during gastrulation (Fig. 1d). Quantitation of embryo length showed that the size of Ino80 KO embryos at E6.5 is similar to that observed for Ino80 KO embryos at E5.5 (Fig. 1e). Embryo reabsorption likely occurs after E7.5 because Ino80 KO embryos after E7.5 begin to show a loss in structural integrity (data not shown).

Ino80 KO embryonic stem cells are viable but exhibit an unstable pluripotent state

To characterize roles for Ino80 in the early embryo, we isolated Ino80 KO embryonic stem cells (ESCs) from the ICM of pre-implantation E3.5 Ino80 KO embryos. Collection and genotyping pre-implantation E3.5-day blastocysts from Ino80 KO heterozygous intercrosses showed that Ino80 KO homozygous blastocysts look morphologically similar to wild-type littermates by gross inspection (Fig. 2a). These same isolated blastocysts successfully attached and outgrew onto gelatinized plates in media containing serum and leukemia inhibitory factor (LIF). After seven days of outgrowth, we observed a similar expansion of trophoblasts and the ICM between wild-type and Ino80 KO blastocysts (Fig. 2a). From these outgrowths, we successfully cloned Ino80 KO ESCs under culture conditions that maintain ground state pluripotency (media containing serum + 2i + LIF) [20, 21]. Western blotting showed that the Ino80 KO ESCs do not have the Ino80 protein, further confirming that our Ino80 KO allele is a null allele (Fig. 2b). Unlike Ino80 KO MEFs, Ino80 KO ESCs proliferate and are not apoptotic when maintained in culture under conditions that promote ground state pluripotency (Additional file 4a, b). A microscopic analysis shows that Ino80 KO ESCs did not form prototypical undifferentiated colonies and exhibit slight cell scattering when maintained at ground state pluripotency (Fig. 2c). Cell scattering is more pronounced when the ESCs were passaged in a media formulation that maintains a metastable pluripotent state (serum + LIF) (Fig. 2c). The percentage of colonies that maintain alkaline phosphatase (AP) staining was equivalent between wild-type and Ino80 KO ESCs when maintained at ground state pluripotency (Fig. 2d, e). However, Ino80 KO ESCs did not form robust AP positive colonies if the colony formation assay was repeated when cells were maintained in a metastable state (Fig. 2d, e). The ability to form AP positive colonies
was recovered if cells maintained in the metastable state are returned to ground state conditions (Fig. 2e). These results demonstrate that \textit{Ino80}\textsuperscript{KO} ESCs are viable, AP positive, and morphologically show evidence of slight cell scattering when compared to wild-type controls when maintained at ground state pluripotency. However, \textit{Ino80}\textsuperscript{KO} ESCs lose AP staining and exhibit cell scattering when grown under conditions that maintain a metastable pluripotent state.

To further characterize phenotypes of \textit{Ino80}\textsuperscript{KO} ESCs grown under differing culture conditions, we measured transcript levels of several markers of pluripotency, stem cell differentiation, and lineage commitment. When maintained at ground state pluripotency, \textit{Ino80}\textsuperscript{KO} ESCs have equivalent expression of pluripotency markers \textit{Nanog}, \textit{Oct4}, \textit{Klf4}, \textit{Sox2}, and \textit{Essrb}. Markers of ectoderm, mesoderm, and endoderm lineages were also approximately equivalent between wild-type and \textit{Ino80}\textsuperscript{KO} ESCs (Fig. 2f). In contrast, we observed reduced pluripotency marker expression (\textit{Nanog}, \textit{Oct4}, \textit{Klf4}, and \textit{Essrb}) when \textit{Ino80}\textsuperscript{KO} ESCs were maintained in the metastable state for five days (Fig. 2g) [13]. These results are
consistent with the metastable pluripotent state, but not
ground state pluripotency requiring Ino80.

**Ino80 KO ESCs fail to differentiate using models of mammalian development**

To further characterize roles for Ino80 during mammalian development, we differentiated Ino80 KO ESCs using the teratoma model [22]. To create teratomas, both wild-type and Ino80 KO ESCs maintained at ground state were introduced into opposing flanks of NOD/SCID mice. Teratomas were allowed to grow until control tumors were approximately 1 cm in diameter. From five inoculations, we harvested four wild-type tumors and two Ino80 KO tumors. A histological analysis showed that control tumors formed a wide range of differentiated tissues from each of the three germ lines. Differentiated tissues include blood islands, keratin pearls, and neural rosettes from the ectoderm lineage, both striated and smooth muscle from mesoderm, and ciliated endoderm from the endoderm lineage (Fig. 3a, b). In contrast to the well-differentiated tissues observed in wild-type teratomas, we observed that Ino80 KO teratomas are composed of undifferentiated mesenchyme with a thin layer of epithelial cells on the outer periphery (Fig. 3a, b).

In addition to forming teratomas, we utilized the embryoid body ESC differentiation model [23]. To create embryoid bodies, ESCs maintained at ground state were dispersed in clumps and aggregated in serum containing

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**Fig. 2 Ino80 KO ESCs are viable, but differentiate when maintained in a metastable pluripotent state.**

a) Microscopic analysis of wild-type and Ino80 KO pre-implantation E3.5 blastocysts. Blastocysts were allowed to outgrow for seven days onto gelatinized plates in media containing serum + LIF. Wild-type and Ino80 KO are at identical magnification. Red arrows designate ESC colony outgrowth.

b) Western analysis of Ino80 protein expression in wild-type and Ino80 KO ESCs using a custom Ino80 antibody. Ponceau S was used as a loading control.

c) Microscope analysis of wild-type and Ino80 KO ESCs grown in media containing serum + 2i + LIF or serum + LIF. Wild-type and Ino80 KO are at identical magnification.

d) Alkaline phosphatase (AP) activity in wild-type and Ino80 KO ESCs after plating at clonal density and maintained in media containing serum + 2i + LIF or serum + LIF. Wild-type and Ino80 KO are at identical magnification.

e) Percent AP positive colonies quantified from panel d and after Ino80 KO ESCs passaged in serum + LIF were recovered in serum + 2i + LIF. At least 50 colonies were quantified per group.

f) Quantitative RT-PCR analysis of gene expression in wild-type and Ino80 KO ESCs maintained in serum + LIF. (N = 3 biological replicates; * = t test p ≤ 0.05). KO knockout, ESC embryonic stem cells, E embryonic day, LIF leukemia inhibitory factor.
media without 2i + LIF. ESC aggregates were then harvested at three-day intervals over the course of nine days. Histological analysis of day-9 wild-type embryoid bodies provided evidence of differentiated cell types including a well-defined endoderm, and an organized mesenchyme with evidence of cavitation as previously described [24] (Fig. 3c). In contrast, embryoid bodies derived from Ino80 KO ESCs were not well organized and lacked a continuous endoderm (Fig. 3c). We subsequently used RT-PCR to measure markers of pluripotency and differentiated tissues in the embryoid bodies. From these experiments, we observed that pluripotency marker repression during embryoid body differentiation is largely Ino80-independent (Fig. 3d). In contrast to pluripotency markers, we observed Ino80-dependent expression for several differentiation markers during embryoid body differentiation. Ino80 KO embryoid bodies showed enhanced expression of endoderm markers Gata6 and Gata4 [25], and the stem cell maintenance factor Bmp4 (Fig. 3d) [26]. The expression of these markers could be linked since Bmp4 positively regulates Gata4 and Gata6 expression [27]. In contrast to these changes, Ino80 KO embryoid bodies showed reduced expression of the ectoderm marker Fgf5 [28], the endoderm marker Hnf4a [29], and mesoderm markers Fgf8 [30] and T [31] (Fig. 3d). Because the expression of these...
markers is essential for embryonic development, Ino80-dependence of these same genes could contribute to a post-implantation lethal phenotype.

**Ino80 is expressed in the embryonic and extraembryonic ectoderm of post-implantation embryos**

In a first step towards understanding why Ino80 KO embryos fail to develop beyond E6.5, we determined where Ino80 is expressed during embryonic development and in the adult. Towards this end, we performed in situ RNA hybridization (ISH) using both sense and antisense probes to the Ino80 transcript in E5.5- to E7.5-day embryos. Whole mount ISH of E5.5- to E6.5-day embryos with sense and antisense probes showed widespread staining of embryos with the antisense probe, but not the sense probe (Fig. 4a). The most intense staining with the antisense probe was observed in both the EmE and ExE, and comparatively less staining was observed in the VE (Fig. 4a, b). The EmE contributes to all cell types of the embryo proper, whereas the ExE contributes to the placenta and umbilical cord [1]. To better visualize Ino80, we sectioned E6.5-day embryos either sagittal or in cross section. Inspection of these sections showed that staining with the antisense Ino80 probe is highest in the ExE and EmE, with less staining in the VE (Fig. 4c). Consistent with widespread expression in the early embryo, Northern blotting for Ino80 transcripts showed that it is expressed in all adult tissues analyzed (Additional file 5). The combination

![Fig. 4](image-url)
of these results supports the conclusion that Ino80 is widely expressed in many tissues of post-implantation embryo and the adult.

**Ino80 KO embryos have a defective distal visceral endoderm and fail to gastrulate**

To better understand why Ino80 KO embryos fail to develop, we used whole mount ISH to monitor marker expression for several essential cell types of the gastrulating embryo. We focused on markers previously analyzed from our studies of Ino80 KO embryoid bodies (Fig. 3d). In the early embryo, gastrulation occurs at E6.5 when the posterior of the embryo differentiates into mesoderm and definitive endoderm [1]. Initial ISH experiments documented a lack of mesoderm marker **T** expression in Ino80 KO embryos, suggesting that they do not differentiate mesoderm and likely do not gastrulate (Fig. 4d). Gastrulation requires the specification of the AVE and is the first organizer of the mammalian embryo. The AVE secretes inhibitors to Tgfβ and Wnt ligands including Cer, and its expression at E6.5 is a marker of the AVE [32]. At E6.5 Cer1 is not expressed in the Ino80 KO embryo, suggesting that Ino80 is required for AVE establishment (Fig. 4d). In addition to the AVE, signaling molecules that originate from the ExE are essential for AVE establishment and gastrulation [33]. One such molecule is the Tgfβ superfamily member Bmp4 [34]. ISH for Bmp4 transcripts showed that it is weakly expressed in the ExE in Ino80 KO E6.5 embryos (Fig. 4d). The results from these analyses demonstrate that Ino80 KO embryos do not gastrulate and have defects in specifying both the AVE and ExE.

To determine if defects in gastrulation result from defects at earlier stages of development, we measured tissue differentiation in Ino80 KO E5.5 embryos. From these experiments, we observed that Bmp4, which is normally expressed in the ExE [34], is abnormally expressed in the EmE in Ino80 KO E5.5 embryos (Fig. 4d). To further confirm EmE Bmp4 expression we performed a double ISH using Bmp4 and the EmE marker Otx2 [35]. From this experiment we show co-localization of Bmp4 and Otx2 expression in Ino80 KO E5.5-day embryos (Fig. 4e). To further monitor for defects in the EmE, we next measured Nodal expression. At E5.5 Nodal is expressed in the EmE where it promotes proliferation of the epiblast and differentiates the DVE [36]. In Ino80 KO E5.5 embryos, Nodal was normally expressed suggesting that the EmE is partially specified (Fig. 4d). Previous reports have shown that over expression of Bmp4 in the early embryo inhibits DVE differentiation [15]. To determine if Ino80 KO embryos have a defective DVE, we measured the expression of DVE markers Cer1, Hex1, and Lefty1 [37]. None of these markers is expressed in E5.5 Ino80 KO embryos, suggesting that Ino80 KO embryos do not specify a DVE (Fig. 4d). Defects in DVE specification could be due to a general defect in the VE because we observed reduced expression of the extraembryonic VE marker Hnf4a in Ino80 KO embryos (Fig. 4d) [29]. These results demonstrate that the DVE does not form in Ino80 KO E5.5 embryos, which could result from expression of the inhibitory molecule Bmp4 in the EmE and general defects in the VE.

Both the VE and the DVE originate from the PrE, which is specified during implantation at E4.5 [38]. To determine if defects in the DVE result from defects in the PrE, we stained Ino80 KO E4.5 embryos with the PrE markers, Lefty1 and Hex1. From these experiments, we observed equivalent expression of Lefty1 and Hex1, suggesting that the PrE of Ino80 KO embryos are specified at E4.5 (Fig. 4c). In addition to the PrE, the E4.5 embryo has an EPI that is the precursor to the E5.5 EmE [38]. We next determined if the EPI is specified by staining for Nanog, a marker of the ICM and EPI. From these experiments we observed equivalent expression of Nanog in the EPI, suggesting that it is specified in Ino80 KO embryos (Fig. 4d).

We next stained E6.5-day embryos using TUNEL to determine if defects in development are due to increased apoptosis with Ino80 KO. From these experiments we observed no increase of TUNEL positive cells between wild-type and Ino80 KO embryos, suggesting that the observed defects are not due to increased cell death (Fig. 4f, g) and likely are due to reduced cell proliferation.

In combination, these expression studies demonstrate that Ino80 KO embryos do not gastrulate. Defects in mesoderm differentiation coincide with a lack of the AVE and its organizer activity, and are not due to increased cell death. The lack of an AVE at E6.5 is due to the inability of Ino80 KO embryos to specify a DVE at E5.5. The lack of a DVE coincides with the abnormal expression of Bmp4 in the EmE, a known negative regulator of DVE differentiation [15]. Defects in the DVE are not likely due to defects prior to E5.5 because both the PrE and EPI appear to be normal in E4.5 Ino80 KO embryos.

**Ino80 represses Bmp4 expression during embryonic stem cell differentiation**

ISH experiments suggest that Ino80 represses Bmp4 expression during embryoid body differentiation and in the EmE of early post-implantation embryos (Figs. 3d and 4d, e). These results suggest a model where Ino80 directly represses Bmp4 expression in the EmE. To explore if Ino80 directly represses Bmp4, we used an in vitro model of early embryonic development. The differentiation of ESCs in monolayer by withdrawal of 2i + LIF in the presence of serum is a relevant model of early
embryonic development. When ESCs are grown under these conditions, they differentiate into EPI lineages including EmE and mesoderm precursors [39]. Under these conditions, we observe elevated expression of Bmp4 in Ino80 KO ESCs (Fig. 5a). Elevated Bmp4 expression coincides with repression of DVE markers Cer1, Hex, and Lefty1. Defects in DVE marker expression are not due to general defects in endoderm differentiation because we observed equivalent expression of endoderm markers Gata4 and Gata6 (Fig. 5a). In embryonic tissue lineages Bmp4 is regulated by two upstream enhancers (Fig. 5b) [40]. Chromatin immunoprecipitation (ChiP) experiments showed that Ino80 is localized to the Bmp4 promoter, but not the enhancers, when ESCs are differentiated for six days in serum containing media lacking 2i + LIF (Fig. 5c). Under these same conditions, we next used formaldehyde assisted isolation of regulatory elements (FAIRE) to detect differences in open chromatin at the Bmp4 promoter in Ino80 KO ESCs. From these studies, we detected increased open chromatin ~1.5 Kb upstream of the Bmp4 transcription start site in Ino80 KO ESCs (Fig. 5d, e). These changes in chromatin structure at ~1.5 Kb upstream of Bmp4 correlated with increased H3K4me3 and SP1 occupancy (Fig. 5f, g). H3K4me3 is a well characterized mark of active promoters [41], and SP1 is a transcription factor known to activate Bmp4 expression [42]. In combination, these experiments suggest that Bmp4 is a direct target of Ino80 chromatin remodeling activity with consequences to SP1 binding and Bmp4 expression. From these studies, we present a model where Ino80 normally represses Bmp4 expression in the EmE. When Ino80 is deleted, Bmp4 expression increases, DVE differentiation is repressed, and embryonic development is stopped (Fig. 5b).

Discussion

In the last few decades it has been discovered that chromatin remodeling complexes are essential for many aspects of mammalian development [43, 44]. Several chromatin remodeling complexes are essential for pre-implantation development including SWI/SNF (Brg1 KO) [45], NURD [46], and TIP60/p400 [47] complexes. In contrast, several other complexes including NURF (Bptf KO) [48] and CHD7 [49] have post-implantation phenotypes, suggesting that they regulate specific developmental pathways. Similarly, we showed that Ino80 KO embryos implant, but fail to develop beyond the egg cylinder stage. These findings contrast with the cell essential functions for Ino80 in differentiated MEFs, where it regulates telomere structures and DNA replication, which are known to occur between the cell types [50, 51].

To understand better how Ino80 regulates embryonic development, we derived Ino80 KO ESCs from pre-implantation embryos. We maintained Ino80 KO ESC under growth conditions that promote either ground state (serum + 2i + LIF) or metastable pluripotency (serum + LIF). Previous studies have shown that these culture conditions simulate growth conditions of the E3.5 blastocyst ICM (ground state) or E4.5 EPI and PrE (metastable state) [21]. Ino80 KO ESCs remain undifferentiated at ground state pluripotency, but begin to show signs of differentiation in the metastable state, including loss of robust AP staining, and down-regulation of pluripotency markers Nanog, Oct4, Klf4, and Esrrb. These results support a model where Ino80 is not required to maintain the pluripotent E3.5 blastocyst ICM in utero, but rather has essential functions in regulating cellular differentiation of the E4.5 peri-implantation embryo.

Our conclusions deviate slightly from previous studies which document that Ino80 is required to maintain ESC pluripotency and pre-implantation development [13]. Differences in methods used to achieve loss of function could explain these contrasting reports. The use of a siRNA KD approach could deplete maternal Ino80 transcripts, thus revealing pre-implantation phenotypes [13]. Conversely, a genetic KO approach would leave maternal Ino80 transcripts intact, mask any pre-implantation phenotypes, and allow the embryo to progress to post-implantation development (this study and [12, 14]).

In addition, the discrepancy could arise because Ino80 is required for Oct4, Nanog, and Sox2 expression in the metastable state (serum + LIF) [13], but not the ground state (2i + LIF) [this study]. The ability of 2i + LIF to mask the self-renewal defects of Ino80 KO ESC is not unprecedented as it has also been reported for Mbd3 KO ESC [46, 52]. In the metastable state, FGF4/MAPK signaling upregulates pro differentiation transcription factors promoting differentiation [53, 54]. It is plausible that depletion of Ino80, and the accompanying reduction in Oct4 expression [13], could synergize with aberrantly expressed pro differentiation factors to promote differentiation in the metastable state. Alternatively, increased H3K4me3 modified nucleosomes at the promoters of pluripotency regulators in the ground state [21] could recruit a redundant set of activating histone readers, masking the Ino80 dependency of Oct4 expression.

To characterize functions for Ino80 during early embryonic development we performed RNA ISH of several differentiation markers in peri- and post-implantation embryos. Our analysis documents that the DVE markers Cer1, Hex, and Lefty1 are not expressed in Ino80 KO E5.5 embryos. DVE establishment involves several Smad signaling pathways including those activated by the
Fig. 5  Ino80 localizes to Bmp4 promoter and regulates its expression during embryonic stem cell differentiation. 

a Quantification of Bmp4, Hex, Lefty1, Cer1, Gata6, and Gata4 expression by RT-qPCR during ESC differentiation in serum containing media without 2i + LIF in monolayer (Representative of N = 3 biological replicates; * = t test p ≤ 0.05).

b Cartoon showing the Bmp4 gene, location of upstream enhancers, and immediate downstream convergent gene Gm15217 position of PCR amplicons used in chromatin immunoprecipitation (ChIP) experiments are shown as black bars labeled 1-6.

c Localization of Ino80 to regulatory elements of Bmp4 by chromatin immunoprecipitation (ChIP). ChIP was performed with wild-type and Ino80 KO ESC after differentiation as a monolayer for six days in serum containing media lacking 2i + LIF. Position of PCR amplicons are shown in panel a (Representative of N = 3 biological replicates; * = t test p ≤ 0.05).

d Cartoon showing the Bmp4 promoter and the location of PCR amplicons used in FAIRE experiments are shown as black bars labeled 1-5.

e Changes in chromatin structure ~1.5 Kb upstream of Bmp4 in Ino80 KO ESC. FAIRE was performed with wild-type and Ino80 KO ESC after differentiation in serum containing media lacking 2i + LIF as a monolayer for six days. Position of PCR amplicons is shown in panel d (Representative of N = 3 biological replicates; * = t test p ≤ 0.05).

f Increased H3K4me3 occupancy at Bmp4 in Ino80 KO ESC. ChIP was performed with wild-type and Ino80 KO ESC after differentiation as a monolayer for six days in serum containing media lacking 2i + LIF. PCR amplicon 3 from panel b was used for ChIP (Representative of N = 3 biological replicates; * = t test p ≤ 0.05).

g Increased SP1 occupancy at Bmp4 in Ino80 KO ESC. ChIP was performed with wild-type and Ino80 KO ESC after differentiation in serum containing media lacking 2i + LIF as a monolayer for six days. PCR amplicon 3 from panel b was used for ChIP (Representative of N = 3 biological replicates; * = t test p ≤ 0.05).

h Model for Ino80 function during proximal-distal axis establishment. In wild-type embryos, Ino80 in part functions in the EmE to repress Bmp4 expression. Bmp4 repression in the EmE promotes DVE differentiation. In Ino80 KO embryos, Bmp4 is abnormally expressed in the EmE, which inhibits DVE differentiation. A lack of DVE prevents the AVE from forming, which subsequently prevents gastrulation. ESC embryonic stem cell, LIF leukemia inhibitory factor, FAIRE formaldehyde assisted isolation of regulatory elements, KO knockout, EmE embryonic ectoderm, DVE distal visceral endoderm, AVE anterior visceral endoderm.
Nodal and Bmp4 ligands [15]. Nodal was the first discovered ligand to promote the DVE [55], then later it was discovered that Bmp4 signaling repressed the DVE [15]. Bmp4 is normally expressed in the ExE, which is proximally located from the distal tip of the embryo, and future site of the DVE [34]. During egg cylinder expansion (E4.5-E5.5), the ExE is rapidly moved away from the distal tip of the embryo. During this period, the repressive effects of Bmp4 on the distal tip of the embryo subdue to the increased distance between the ExE and distal tip during egg cylinder expansion [34], and activating Nodal signals become dominant due to increased Nodal expression in the EmE [55, 56]. In this model, the combined loss of Bmp4 signaling and increased Nodal signaling to the distal VE, establishes the DVE [15]. From our studies, we observed that Nodal is Ino80-independent during embryoid body differentiation and embryonic development. In contrast to Nodal, we observe that Bmp4 is upregulated during embryoid body differentiation and in the E5.5 EmE. The abnormal expression of Bmp4 in the EmE could result in down regulation of DVE markers [15]. Alternatively, defects in the DVE could result from general defects in the VE as supported by reduced Hnf4a expression with Ino80 KO, or due to reduction in epiblast proliferation. Ino80 repression of Bmp4 expression in the EmE could be direct because it localizes to, regulates chromatin structure and transcription factor occupancy at, the Bmp4 promoter during ESC differentiation. In addition to these factors, we observed reduced Bmp4 expression in the ExE, suggesting that Ino80 could also be an activator of Bmp4 in extraembryonic tissues. These activities could also be direct, and the study of Bmp4 expression in differentiating Ino80 KO trophoblast stem cells could test this hypothesis [57].

Conclusions
From the data presented in this manuscript, we show that Ino80 is required to pattern the early embryo. Using several models of development, we showed that Ino80 KO embryos upregulate Bmp4 expression, and when measurable, we observed a coincident repression of the DVE markers Cer1, Hex, and Lefty1. Because Bmp4 represses the DVE, we propose that the lack of a DVE in Ino80 KO embryos is in part due to elevated Bmp4 expression in the EmE. Ino80 functions in regulating Bmp4 expression are likely direct because we observed specific Ino80 localization to its promoter in differentiating ESCs. We also observed evidence that Ino80 remodels chromatin and regulates SP1 binding at the Bmp4 promoter under these same conditions. These results support a model where Ino80 directly represses Bmp4 expression in the EmE, and its repression is essential for DVE establishment and ultimately a functional P-D axis (Fig. 5h).

Methods
Gene targeting and animal husbandry
The RPC121 mouse PAC library (MRC, UK, London) was screened using probe P2 internal to the Ino80 gene. Probe hybridization was performed using 0.25 M sodium phosphate pH 7.2, 1 mM EDTA, 7 % SDS at 65 °C overnight. Blots were washed three times for 30 minutes with 0.25XSSC, 0.1 % SDS at 65 °C. Five positive P1 clones were identified, and recombineering was used to move the chr2: 119,464,212-119,447,795 (mm10 assembly) DNA sequence from clone 386-C4 into DNDF-7. Recombineering was subsequently used to insert an upstream loxP site and a downstream frt-Neo-frt-loxP selection cassette.

Linearized Ino80 targeting vector was electroporated into R1 ESC (129 Sv X 129Sv-CP F1) on feeder layers. Neomycin resistant colonies were isolated and screened for successful Ino80 targeting using a BglII digest and probe P2, BglII digest and probe P3, and an EcoRV digest using probe P4. Primers used to amplify probes from genomic DNA are available in Additional file 6. Southern blotting was performed as described for probe hybridization. Ino80 was successfully targeted in 1 of 456 neomycin resistant clones. Germ line transmission of Ino80 Floxed-Neo ESCs from two founder chimera mice was confirmed by Southern blotting.

The Ino80 Floxed-Neo mice were crossed to Cre deleter (Jackson labs Cat#006054) and Flp deleter strains (Jackson labs Cat#011065) to create Ino80 Floxed and Ino80 KO alleles, respectively. Ino80 Floxed and Ino80 KO mice were then backcrossed to C57BL/6 J for five generations and confirmed Cre and Flp negative by PCR genotyping. Once backcrossed founder lines were established, the Ino80 Floxed mice were cryopreserved at Jackson Laboratory (Jackson labs Cat#027920). To create Ino80 doxycycline inducible KO MEFs, the Ino80 Floxed mice were crossed to TetO-Cre (Jackson Labs Cat#006234) and subsequently rtTA (Jackson Labs Cat#006965). MEFs were generated from an Ino80 F/F, TetO-Cre hemizygous, r'TTA homozygous X Ino80 F/F, and r'TTA homozygous cross. Genotypes of mice were determined by PCR using the Ino80 genotyping protocol or those provided by Jackson Labs for purchased alleles. The Ino80 genotyping protocol includes primers P1, P2, and P3 in a multiplex PCR reaction using recombinant Taq polymerase with Thermal Pol Buffer (NEB). PCR conditions were as follows: 94 °C for three minutes for one cycle, 94 °C for 30 sec, 54 °C for 60 sec, 72 °C for 45 sec repeated for 40 cycles. Primer sequences are available as Additional file 6.
The mice used in this study were housed in a specific pathogen-free facility at Virginia Commonwealth University, Richmond VA, USA. Mice were maintained on a 12-hour light/dark cycle, and provided a low fat diet and hypochlorinated water ad libitum throughout the duration of the study. All experiments and animal procedures were approved by the Animal Care and Use Committee of Virginia Commonwealth University under protocol AD10000372 and its modifications.

**Cell line isolation and maintenance**

To create Ino80 KO ESCs, we harvested and maintained E2.5-day embryos in KSOM media until the blastocyst stage. Expanded blastocysts were hatched using Tyrodes solution (Sigma T7188) and allowed to attach and outgrow on gelatinized plates for seven days in defined 2i + LIF media without serum (Millipore Cat# SF016-100). After outgrowth, the ICM was removed with a fine glass pipette, dispersed with 0.25% trypsin + EDTA, and expanded in 2i + LIF + 15% ESC grade serum (Life Technologies) media on gelatinized plates. Depending on the experiment, ESC lines were also grown in 2i + LIF growth media lacking serum (Millipore SF016-100), or a serum + LIF formulation (15% ESC grade FCS (Life Technologies), DMEM, nonessential amino acids, 2 mM glutamine, 10 μM β-mercaptoethanol, penicillin, and streptomycin, 1000 U/ml LIF (Life Technologies)). ESC lines were genotyped by PCR methods (see above) and routinely maintained in serum + 2i + LIF containing media on gelatinized plates. Ino80 KO at the protein level was confirmed by Western blotting using a custom rabbit polyclonal antibody and standard techniques. For gene expression studies, cells were harvested from culture using Tri-Reagent (Sigma) and total RNA was purified according to the manufacture’s protocol. Quantitative RT-PCR was performed as described below.

Primary MEFs were produced from E12.5-day embryos using standard methods. MEFs were maintained in DMEM containing 10% FCS, 2 mM glutamine, 1 mM nonessential amino acids, penicillin, and streptomycin. All experiments were performed using early passage (P0-1) fibroblasts. Ino80 deletion was achieved by adding 10 ng/ml doxycycline to the growth media for two days. Following doxycycline exposure, the medium was replaced and MEFs were maintained at subconfluence using standard tissue culture techniques. Efficiency of Ino80 deletion was confirmed by PCR as described above. Analysis of Ino80 transcripts was accomplished by RT-PCR using Superscript II (Life Technologies) followed by amplification using Phusion polymerase (NEB). PCR fragments were cloned into pTOPO Blunt Zero vector (Life Technologies) and sequenced.

To stain for β-galactosidase activity, MEFs were washed in phosphate buffered saline (PBS) and then fixed in 4% buffered formaldehyde for five minutes. Fixed cells were washed and then stained in 40 mM citric acid/sodium phosphate buffer, pH 6.0, containing 5 mM potassium ferrocyanide, 5 mM potassium ferriyanide, 150 mM sodium chloride, and 2 mM magnesium chloride overnight with gentle rocking. Cells were washed in PBS and blue cells counted with a microscope.

Annexin V + 7AAD staining of ESC lines was performed using AnnexinV-PE (BD Biosciences Cat#556421) using the protocol provided by BD Biosciences.

To stain for AP activity, ESC colonies were first briefly washed in PBS. Colonies were then fixed for five minutes at room temperature in PBS containing 0.2% glutaraldehyde, 0.02% NP-40, and 0.01% sodium DOC. Colonies were then washed for ten minutes at room temperature in 100 mM Tris-HCl buffer, pH 9.5, containing 100 mM NaCl and 10 mM MgCl₂. Colonies were then stained in BM purple AP substrate (Roche Cat# 11442074001) until the desired level of staining was achieved. Cells were then washed in PBS containing 0.1% tween-20 and 2 mM MgCl₂. The plates were then air dried.

To determine doubling time cells were plated at 5 × 10⁴ cells per well in a 24-well plate and incubated for the indicated time points, trypsinized, and counted using a hemocytometer. Doubling time was calculated using standard equations.

**Teratoma and embryoid body production**

To produce teratomas, 1 × 10⁶ ESC were injected into the flank of NOD/SCID mice (Jackson Labs Cat# 001303), after which the tumors were allowed to grow for three weeks. The mice were then sacrificed, and the tumors were removed and frozen in liquid nitrogen. Tumors were sectioned in OCT medium using a cryostat, fixed in 10% neutral buffered formalin, stained with hematoxylin and eosin, and visualized with light microscopy using standard techniques.

Embryoid bodies were created by transferring 10 × 10⁶ ESC into 13 × 100 mm polypropylene culture tubes with 3 ml DMEM containing 15% ESC grade FCS (Life Technologies), nonessential amino acids, 2 mM glutamine, 10 μM β-mercaptoethanol, penicillin and streptomycin (2i + LIF withdrawal growth conditions). Upon harvest, embryoid bodies were dispersed using Tri-Reagent (Sigma) and total RNA was purified according to the manufacture’s protocol. Total RNA was then processed for quantitative RT-PCR as described below. Day 9 embryoid bodies were fixed in 10% neutral buffered formalin overnight, washed in PBS, and then equilibrated in OCT before freezing and sectioning with a cryostat. Sections were stained with hematoxylin and eosin and imaged using light microscopy according to standard procedures.
RNA in situ hybridization and TUNEL assays

Embryos were dissected from timed pregnancies and photographed previously [48]. Alkaline phosphatase (AP) coupled anti-DIG Fab fragments and BM Purple substrate (Roche) were used to detect DIG labeled antisense probes as previously described [48]. For co-localization staining, DIG-Bmp4 and FITC-Otx2 antisense labeled probes were detected by AP conjugated anti-DIG Fab and anti-FITC Fab fragment, subsequently. The two probes were visualized by two different AP substrates; the Bmp4 detection used BM Purple (Roche) and Otx2 detection used INT/BCIP (Roche) after inactivating the first AP activity (Otx2) with 4 % formaldehyde for 30 min.

To measure apoptosis E6.5 embryos were fixed for two hours at 4C, washed with PBST and kept in 100 % a methanol until use. Embryos were treated with 5:1 methanol:30 % hydrogen peroxide for three hours. Then, they were washed in a methanol and treated with 20 µg/ul proteinase K for three minutes and fixed with 0.2 % glutaraldehyde in 4 % paraformaldehyde for 20 minutes at room temperature. Embryos were washed three times with PBST and treated with sodium borohydride for 20 minutes. Embryos were then washed with TdT buffer and apoptotic cells were detected by TUNEL staining using a fluorescein in situ cell death detection kit (Sigma). Apoptosis was calculated for each embryo by averaging the percentage TUNEL+ cells (brown nuclei) to all nuclei (DAPI staining) in three representative 50 um fields using microscopy.

Formaldehyde Assisted Isolation of Regulatory Elements (FAIRE)

FAIRE-qPCR was performed as described [58]. The qRT-PCR cycle threshold (Ct) value from the Input and FAIRE samples (control and Ino80 KO) were normalized to signal from the reference region in each sample. Next, the relative enrichment for site in the Bmp4 promoter in comparison to the reference primer set was calculated using the comparative ΔΔCt method. Primers used for FAIRE are available as Additional file 6.

Chromatin immunoprecipitation (ChIP)

ChIP was performed as described previously [48]. Antibodies used include Ino80 (VCU23, custom rabbit polyclonal), SP1 (Santa Cruz Biotech # sc-59), pan histone H3 (Abcam # ab1791), and H3K4me3 (Abcam # ab1012). DNA quantification was accomplished using SsoAdvanced universal SYBR Green supermix (Biorad) according to the manufacturer's established protocol. Data collection was performed using a 7900HT Fast Real Time PCR System (Applied Biosystems) and quantified as % Input DNA for Ino80 or SP1, or as a ratio of % Input H3K4me3/pan H3. The primers used for ChIP are available in Additional file 6.

Mouse Ino80 antibody VCU23 was raised in rabbits as a GST fusion to aa792-892 of Genbank # AAH59235.1. It was subsequently affinity purified by column chromatography using 6 x His-aa792-892. Specificity to Ino80 was confirmed by Western blotting using Ino80 control and KO ESC extracts. As a reference we observed identical results using the commercially available Ino80 antibody from ProteinTech (Cat# 18810-1-AP) (See Additional file 7).

Gene expression assays

Quantitative RT-PCR was performed on cDNA libraries converted using Superscript II (Life Technologies). cDNA libraries were amplified using SsoAdvanced universal SYBR green supermix (Biorad) according to the manufacturer’s established protocols. Primers used for gene specific quantitative PCR were previously described [48]. Data collection was performed using a 7900HT Fast Real Time PCR System (Applied Biosystems) and the ΔΔCt method was used to estimate transcript abundance relative to the normalization control Gapdh.

Northern blotting was performed as previously described [48]. A total of 1 ug of polyA+ RNA was resolved by 1 % formaldehyde agarose electrophoresis and transferred to Hybond N+ (GE Health Sciences) using the manufacturer’s suggested protocol. The Ino80 probe was amplified from ESC cDNA library using Taq polymerase, sequenced to confirm identity and labeled with P32 using random hexamer labeling. Detection was performed by phosphoimager.

Statistics

Standard deviation was used to calculate error bars throughout the study. P values were calculated using two tailed T tests. Number of replicates are designated in the figure legend.

Availability of supporting data

Data supporting the results of this article are available in Additional file 8.

Additional files

Additional file 1: Figure S1. Targeting and genotyping strategy for conditional and knockout Ino80 alleles. a Southern blotting strategy for screening embryonic stem cell clones successfully integrating the Ino80 targeting vector. Probes used in the screen are positioned at the site of hybridization, are designated in red, and are labeled P2, P3, and P4. Restriction enzyme sites are designated above the genome sequence. Loxp and Neo features of the targeting vector are shown in black. Exons are shown as vertical lines and designated by number in sequential order from 5’ to 3’. b Results of Southern blotting experiments to confirm successful integration through Cre- or Flp-mediated recombination of the Ino80 targeting vector. Wild-type targeted Ino80 FlxedNeo (Neo), Ino80
Floxed (Floxed) and Ino80 knockout (KO) alleles are shown. DNA fragment sizes obtained from restriction enzyme digestion and Southern blotting with P2, P3, and P4 probes are as expected from diagrams in panel a. c Diagram describing PCR-based genotyping strategy. Ino80 alleles as defined in panel a are shown in a cartoon format with exons and introns not to scale. Position of the Loxp sites (red dots) and the Neo selectable marker are shown relative to Ino80 exons 2-4, which are designated by numbers. Positions of primers P1, P2, and P3 are shown as arrows designating 5′-3′ orientation. PCR amplicons and their size are shown as black bars above amplifying primers. d Results from the PCR genotyping strategy using DNA from wild type and mice heterozygous for the Ino80 Floxed-Neo, Ino80 Floxed, and Ino80 KO alleles. (TIF 10353 kb)

Additional file 2: Figure S2. Molecular characterization of the Ino80 KO allele. A efficiency of Cre-mediated excision of Ino80 exons 2-4 from control Ino80 Floxed/Floxed (Cre-Neg) and Ino80 Floxed/Floxed, TetO-Cre (TetO-Cre) primary MEFs were monitored using the PCR genotyping strategy. Cells were incubated with 10 µg/ml doxycycline for two days, and then shifted to culture medium without doxycycline. DNA was taken for analysis prior to the addition of doxycycline (Day 0), and at two, four and ten days post-doxycycline addition. b Western blot analysis of Ino80 protein from total cell extracts prior to doxycycline addition (Day 0) and six days after initial doxycycline treatment with a custom Ino80 antibody. Ponceau S was used as a loading control. c Cartoon showing the position of RT-PCR primers used for characterizing the Ino80 KO allele transcript. Location of the forward primer in exon 1 and reverse primer in exon 7 are shown. The position of the loxp sites flanking exons 2-4 are shown as black boxes. The position of the initiating ATG is shown in exon 2. d RT-PCR results using primers shown in panel c on cDNAs converted from total RNA harvested from control Ino80 Floxed/Floxed (Cre-Neg) and Ino80 Floxed/Floxed, TetO-Cre (TetO-Cre) MEFS 6 days post-doxycycline treatment. PCR products were gel purified, cloned and sequenced. exon composition of PCR fragments is shown. e Predicted reading frames for the KO-1 and KO-2 transcripts initiating from the first available ATG. (TIF 10038 kb)

Additional file 3: Genotyping results from intercross of heterozygous Ino80 KO mice. Litters of mice or embryos at specified stages were collected and genotyped using the PCR-based genotyping strategy. Significant deviations from the expected genotype ratios were identified using a Chi-Square test (**p < 0.01). (XLSX 10 kb)

Additional file 4: Figure S4. Ino80 is not essential for the growth and viability of ESCs maintained in ground state pluripotency. A Growth curve of wild-type and Ino80 KO ESCs. 1.0 x 104 cells were seeded onto gelatine plates containing serum + 2i + LIF media. Numbers of trypan blue negative cells were counted every day over a four-day period. (Doubling Time, WT = 36 ± 4.5 hours, KO = 44 ± 4.2 hours; t test p < 0.05, N ≥ 3 biological replicates) b AnnexinV+ 7AAD staining of wild-type and Ino80 KO ESCs maintained in serum + 2i + LIF (N ≥ 3 biological replicates). (TIF 8494 kb)

Additional file 5: Figure S5. Ino80 is widely expressed in adult tissues. Abundance of Ino80 transcripts in a variety of adult tissues was determined by Northern blotting. Ethidium bromide stained RNA serves as a loading control. (TIF 1828 kb)

Additional file 6: Primers used in this study. (XLSX 12 kb)

Additional file 7: Figure S7. Western blotting confirming custom Ino80 antibody. Western analysis of Ino80 protein expression in wild-type and Ino80 KO ESCs using custom antibody YCU23 (a) and a commercially available Ino80 antibody (ProteinTech Cat 188810-1-AP) (b). In each case Ponceau S was used as a loading control. (TIF 6874 kb)

Additional file 8: Supporting data. (XLSX 36 kb)

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
QZ, Ze, and JW designed experiments. QZ, Ze, VP, SA, DB, and JW performed experiments. JW, wrote the manuscript. All authors read and approved the final manuscript.

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