Dual Functions of ASCIZ in the DNA Base Damage Response and Pulmonary Organogenesis

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Abstract

Zn²⁺-finger proteins comprise one of the largest protein superfamilies with diverse biological functions. The ATM substrate Chk2-interacting Zn²⁺-finger protein (ASCIZ; also known as ATMIN and ZNF822) was originally linked to functions in the DNA base damage response and has also been proposed to be an essential cofactor of the ATM kinase. Here we show that absence of ASCIZ leads to p53-independent late-embryonic lethality in mice. Asciz-deficient primary fibroblasts exhibit increased sensitivity to DNA base damaging agents MMS and H₂O₂, but Asciz deletion or knock-down does not affect ATM levels and activation in mouse, chicken, or human cells. Unexpectedly, Asciz-deficient embryos also exhibit severe respiratory tract defects with complete pulmonary agenesis and severe tracheal atresia. Nkx2.1-expressing respiratory precursors are still specified in the absence of ASCIZ, but fail to segregate properly within the ventral foregut, and as a consequence lung buds never form and separation of the trachea from the oesophagus stalls early. Comparison of phenotypes suggests that ASCIZ functions between Wnt2-2b/β-catenin and FGF10/FGF-receptor 2b signaling pathways in the mesodermal/endodermal crosstalk regulating early respiratory development. We also find that ASCIZ can activate expression of reporter genes via its SQ/TQ-cluster domain in vitro, suggesting that it may exert its developmental functions as a transcription factor. Altogether, the data indicate that, in addition to its role in the DNA base damage response, ASCIZ has separate developmental functions as an essential regulator of respiratory organogenesis.

Introduction

Pathways that maintain genome integrity by responding to spontaneous DNA damage are crucial for normal development and ageing, and act as tumor suppressors to prevent the onset of cancer [1]. While DNA damage signaling is rather generic in that structurally diverse lesions eventually lead to activation of one or both of the central checkpoint kinases ATM or ATR [2], DNA repair pathways are believed to be highly lesion-specific [1]. In addition to environmental DNA damage, eukaryotic cells incur a high level of spontaneous DNA damage as a consequence of normal metabolism, most notably abasic sites that are generated as repair intermediates of the base excision repair (BER) pathway with an estimated incidence of ~10,000 per cell per day [3]. Abasic sites can emanate from various base modifications (e.g. oxidation, methylation), which for experimental purposes are most commonly generated by treatment with methylmethane sulfonate (MMS) or H₂O₂, and key BER enzymes for their repair include apyrimidinic/apurinic endonuclease (APE) and DNA polymerase beta (Polβ) [4,5].

The importance of the BER pathway is indicated by findings that absence of any of the BER genes acting downstream of abasic sites results in embryonic or perinatal lethality in mice [6,7]. However, some of the key BER enzymes also seem to have DNA damage-independent functions; for example, APE1 has a separate role as a redox regulator of several transcription factors [8].

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ASCIZ is a DNA damage response protein that has been proposed to be a regulator and stabilizing co-factor of the ATM kinase, mutations of which lead to a syndrome involving neurological and immune dysfunctions, tumour predisposition, and X-ray hypersensitivity. To study ASCIZ function in vivo, we have generated a knockout mouse model lacking this gene. Here we show that ASCIZ has a specific role in mediating cell survival in response to DNA base damage, but it is not required for stabilization and regulation of ATM. Strikingly, Asciz knockout mice fail to survive to birth and have tissue-specific defects in embryonic development. In particular, Asciz null embryos fail to develop lungs and undergo an early arrest in tracheal development. The precursor cells that normally form the lung are present in our embryos, but they fail to segregate from the foregut. These observations indicate that ASCIZ plays an important and previously unrecognized developmental role that is most likely unrelated to its function in mediating responses to DNA damage. Our study delineates the function of ASCIZ in DNA damage survival and highlights an exciting new function of the protein in controlling the early stages of lung development.

Similarly, increased apoptotic cell death during development of Polß-null mice can be suppressed by deletion of p53 (TRP53), indicating that this part of the phenotype is indeed due to defective base damage repair. On the other hand, the perinatal lethality of these mice that is associated with defective neuronal and lung development as a DNA damage-independent defect is not rescued by p53 deletion [9–11].

While the DNA damage processing enzymes of the BER pathway are clearly defined, new accessory factors that regulate the activity or stability of Polß and other BER enzymes keep emerging, including the non-histone DNA-binding protein HMGB1 [12], arginine methyl-transferases [13], and ubiquitin ligases [14]. We recently identified ASCIZ (=ATM substrate Cbk2-interacting Zn2+-finger) as a new Zn2+-finger (ZnF) protein with roles in the DNA base damage response. In human cells, ASCIZ forms DNA damage-induced nuclear foci specifically in response to DNA damaging agents that generate lesions repaired by the BER pathway (MMS and H2O2) in a manner that is enhanced by the BER inhibitor methoxyamine, and Asciz deletion by siRNA leads to increased MMS sensitivity [15]. Likewise, Asciz deletion in the chicken DT40 B lymphocyte line leads to increased sensitivity to MMS and H2O2, but not to ionizing radiation (IR), UV irradiation and other DNA lesions, as well as increased erroneous repair of enzyme-generated DNA base damage consistent with a role in the BER pathway [16]. Moreover, Asciz deletion suppresses the dramatic MMS hypersensitivity of Polß-deficient DT40 cells [16], reminiscent of the protective effect of simultaneous deletion of the relevant upstream methyl-purine-glicosylase (MPG) in Polß-deficient murine embryonic fibroblasts [17]. ASCIZ contains a large number of conserved ATM/ATR kinase phosphorylation sites in an SQ/TQ cluster domain [18], and consistent with its original classification as an ATM substrate, ASCIZ was subsequently re-identified as an ATM-interacting protein (thus also called ATM-IN) [19]. It was proposed that ASCIZ acts as an essential co-factor of ATM that was required for ATM stability (and vice versa ATM was required for ASCIZ stability) as well as for ATM activation by some stimuli, though surprisingly not by canonical DNA damaging ATM activators such as IR [19].

To better understand the role of ASCIZ in vivo, we have here generated a mouse line that lacks the vast majority of the Asciz protein-coding sequence in the germline. Our results confirm that Asciz-deficient cells are specifically hypersensitive to DNA lesions that are processed by the BER pathway, but challenge the proposed interdependence between ASCIZ and ATM levels. In contrast to Atm-deficient mice that overall develop normally [20], Asciz deletion results in late embryonic lethality with severe respiratory defects reminiscent of mouse mutants in Wnt2/2b and FGFl0 signaling pathways. The data indicate that Asciz has an unexpected DNA damage-independent developmental function as an essential regulator of pulmonary organogenesis.

Results

Generation of Asciz gene-targeted mice

Human and mouse Asciz have a similar gene structure where exons A–C encode the N-terminal ZnF region of about 220 amino acid residues, and exon D encodes the bulk of the protein (601 of 823 or 818 residues) including the nuclear localization signal, core domain and SQ/TQ cluster domain (Figure 1A, 1B; NCBI Gene ID 23300). Because there is evidence for expression of alternative isoforms that differ in the number of N-terminal ZnFs (http://www.uniprot.org/uniprot/043313), we integrated loxP sites flanking exon D into the murine Asciz locus to remove the majority of the protein-coding sequence (Figure 1B). Germine deletion of this exon after crossing with PGK-Cre knock-in mice, followed by outcrossing of PGK-Cre (all on a pure C57BL/6 background), was confirmed by Southern blot and PCR genotyping (Figure 1C). In over 600 offspring from Asciz+/−/− heterozygote intercrosses genotyped at weaning (+3 weeks of age), we failed to detect any homozygous Asciz-deleted mice (Figure 1C and Table 1). However, homozygous Asciz-deleted embryos were readily detectable even at relatively late stages of gestation (Figure 1D; and more detail below).

Western blotting of head extracts confirmed the absence of ASCIZ protein in Asciz−/−− embryos, and a ~50% reduction of protein levels in heterozygotes compared to wildtype (WT) littermates (Figure 1E). Levels of other DNA damage response proteins (including ATM) appeared to be normal in Asciz-deficient embryos (Figure 1E and below). In Northern blots using a probe for the non-deleted exon C, the residual exon D-deleted Asciz transcript was present in homozygous targeted embryos at ~15% of wildtype (WT) mRNA levels (Figure S1), indicating that the mutated mRNA is highly unstable. Using Asciz null embryo lysates as an antibody specificity control, we found that ASCIZ is ubiquitously expressed in adult mice, with overall similar levels relative to the loading control in all tissues except for somewhat higher levels in the brain, cerebellum and testes (Figure 1F).

Absence of Asciz leads to p53-independent late-embryonic lethality

The absence of homozygous Asciz−/− mice at weaning prompted us to investigate the development of ASCIZ-deficient embryos in more detail. Asciz−/− embryos were recovered at near-Mendelian ratios at all time points analysed (Table 1). Based on peripheral circulation scored during uterine dissections, Asciz−/− embryos appeared to lose viability around embryonic day 16.5 post conception (E16.5) (Table 1), at which point they were considerably growth-retarded compared to littermates (Figure 2A, 2B). Embryonic lethality due to DNA damage response gene deletions can often be suppressed by p53 deletion [6]. To test if p53 status affects the essential requirement for Asciz, we intercrossed compound
Asciz deficiency leads to modestly accelerated senescence and BER-like DNA damage hypersensitivity in primary fibroblasts

To monitor DNA damage sensitivity of primary Asciz-deficient cells, we isolated murine embryonic fibroblasts (MEFs) from viable Asciz+/− embryos and matched WT littermate controls between
In these assays, genotypes were minimal) with a panel of DNA damaging agents. passage MEFs (P2–P3, when proliferation differences between damage in primary cells. To corroborate this, we treated early-indicated a role of ASCIZ in the response to oxidative base senescence of MEFs under these conditions is believed to involve senesced earlier than the matched WT cultures (Figure 3B). As inhibition of APE1 and Polß [5]. In addition, MMS hypersensi-which blocks the single-nucleotide BER pathway through partial than that of WT cells co-treated with methoxyamine (Figure S2), indicating that absence of ASCIZ only partially impairs BER. Altogether, these results are consistent with a role of Asciz as an accessory factor in the BER pathway in primary cells.

Because we had originally identified ASCIZ based on its interaction with Chk2 [15], and because ASCIZ was later proposed to be sometimes required for ATM activation [19], we tested if the MMS hypersensitivity of Asciz−/− MEFs could be due to defective ATM signaling. However, there was no reduction in MMS-dependent ATM activation (detected by an antibody against mouse pS1987-ATM; human pS1981-ATM) and phosphorylation of key ATM/ATR targets γH2AX and pS18-pS3 in Asciz-deficient MEFs compared to WT littermate controls (Figure S3A, S3B). MMS-induced pS3-S18 phosphorylation was completely abolished by the highly specific synthetic ATM kinase inhibitor KU55933 [22] (Figure S3C), indicating that it is a genuinely ATM-dependent process, whereas H2AX phosphorylation under these conditions was ATM-independent and thus likely ATR-mediated. Because our antibody for Chk2 phosphorylation on T68, widely considered to be one of the most ATM-specific phosphorylation sites, did not detect this site in MEF extracts (data not shown), we monitored Chk2-T68 phosphorylation in human U2OS cells following Asciz depletion by siRNA treatment. However, Chk2 was still efficiently phosphorylated in response to MMS (as well as IR) in Asciz-depleted cells (Figure S4A, S4B). Altogether, these data indicate that the increased MMS sensitivity of Asciz-deficient cells is not caused by impaired ATM signaling.

Table 1. Genotypes of Asciz−/− intercross litters.

| age   | Asciz+/+ | Asciz+/− | Asciz−/− (%) | Asciz+/− alive | Asciz−/− exencephaly |
|-------|----------|---------|--------------|----------------|----------------------|
| weaning | 270       | 418     | 0            | 0              | -                    |
| E12.5  | 9         | 9       | 4 (18)       | 0              | 1                    |
| E16.5  | 25        | 36      | 13 (18)      | 8              | 5                    |
| E15.5  | 13        | 15      | 8 (24)       | 6              | 2                    |
| E14.5  | 29        | 60      | 22 (20)      | 18             | 7                    |
| E13.5  | 10        | 25      | 9 (20)       | 8              | 3                    |
| E12.5  | 32        | 53      | 13 (13)      | 12             | 2                    |
| E11.5  | 23        | 32      | 15 (21)      | 15             | 2                    |
| E8.5–10.5 | 29  | 44      | 30 (29)      | n.d.           | n.d.                 |

n.d., not determined.

**Figure 2. Late gestational growth defect of Asciz−/− embryos.** (A) Embryo weights of WT, heterozygotes and Asciz−/− embryos at the indicated times post-conception. Data are the mean ± standard error; 22–55 embryos were analysed per timepoint (KO: n = 8 (E12.5), 12 (E14.5), 8 (E15.5), 11 (E16.5), and 4 (E18.5)). (B) Crown-rump length of embryos determined by histomorphometry. Data are mean ± standard error, n = 3–9 per data point. *p<0.05, **p<0.01, ***p<0.001.

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E12.5–E14.5 (i.e., before growth retardation was apparent). Standardized proliferation assays using a 3T3 protocol under normoxic conditions (20% O2) revealed a modest premature senescence phenotype of Asciz−/− deficient MEFs compared to WT controls, with growth arrest after ~20% fewer population doublings (Figure 3A). When normalized to the maximum population doublings within each litter, Asciz−/− MEFs always senesced earlier than the matched WT cultures (Figure 3B). As senescence of MEFs under these conditions is believed to involve an oxygen-induced DNA damage response [21], these results indicated a role of ASCIZ in the response to oxidative base damage in primary cells. To corroborate this, we treated early-passage MEFs (P2–P3), when proliferation differences between genotypes were minimal) with a panel of DNA damaging agents. In these assays, Asciz−/− deficient MEFs were significantly more sensitive to agents such as UV or hydroxyurea (HU) whose damage is repaired by other pathways (Figure 3E, 3F). MMS hypersensitivity of Asciz−/− MEFs was less pronounced than that of WT cells co-treated with methoxyamine (Figure S2), which blocks the single-nucleotide BER pathway through partial inhibition of APE1 and Polß [5]. In addition, MMS hypersensitivity of ASCIZ-deficient cells was further enhanced by methox-

yamine (Figure S2), indicating that absence of ASCIZ only partially impairs BER. Altogether, these results are consistent with a role of Asciz as an accessory factor in the BER pathway in primary cells.

During analyses of DNA damage signaling in Asciz-depleted human cells (Figure S4B and data not shown) and initial protein blots of embryo extracts (Figure 1E), we could not detect any meaningful reduction of ATM levels in Asciz−/− cells. Because this contradicted the recent report that ASCIZ and ATM levels were mutually dependent on each other [19], we explored this discrepancy first in additional embryos. Again, we saw no reduction of ATM protein levels in any of the Asciz−/− or heterozygous samples compared to WT littermate controls (Figure 4A, left panel, and data not shown). Likewise, we also did not see a reduction of ATM levels in human cells after almost complete depletion of ASCIZ by siRNA-treatment (Figure 4B, left panel, compare control siRNA treated to si-Asciz treated U2OS cells, lanes 1 and 2; and data not shown), or in two independently
generated ASCIZ knockout clones [16] in the chicken DT40 B cell line (Figure 4C, left panel).

We also revisited the proposal that ATM was in turn required for ASCIZ stability [19]. In contrast to the severe reduction of ASCIZ levels in an ataxia telangiectasia (AT) fibroblast line reported by Kanu and Behrens (2007), we did not detect any loss of ASCIZ in another human AT patient-derived fibroblast cell line (Figure 4B, left panel, AT2221JE) that is considered to be bona fide ATM-deficient [23] compared to control fibroblasts (Figure 4B, left panel, GM847) or the isogenic AT cell line reconstituted with WT Atm (Figure 4B, left panel, AT2221JE+ATM [23]). We expanded this analysis to a panel of seven independent human AT-patient derived lymphoblastoid cell lines [24]. When adjusted for loading (actin), there was no reduction of ASCIZ levels in four of these lines that contained no or extremely low levels of residual ATM protein (AT1ABR, L3, AT4ABR, AT5ABR) compared to two independent healthy donor control cell lines (C3ABR, C35ABR) (Figure 4B, right panel); ASCIZ levels were also unaffected in two further AT lines that contained intermediate ATM protein levels (AT1ABR, AT33ABR), and there was only a modest reduction of ASCIZ levels in a third AT cell line with intermediate ATM levels (AT32ABR). Similarly, there was also no reduction of ASCIZ protein levels in brain lysates of Atm null mice [20] compared to WT littermates (Figure 4A, right panel), or in an Atm-deleted chicken DT40 clone [25] compared to the WT control (Figure 4C, right panel). Taken together, these data demonstrate that ASCIZ and ATM are not required for each other’s stability in three different vertebrate species.

**Asciz is essential for pulmonary organogenesis**

Because of the overall relatively mild DNA damage phenotypes of Asciz-deficient primary cells (Figure 3) and the absence of a p53-effect on viability (Table 2), we wondered whether the underlying cause for the late gestational lethality of Asciz−/− embryos could be DNA

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**Table 2. Genotypes of offspring from Asciz+/− p53+/− intercrosses at weaning.**

| Genotype         | p53+/+ | Asciz+/− | Asciz−/− |
|------------------|-------|---------|---------|
| p53+/+           | 36    | 69      | 0       |
| p53+/−           | 49    | 107     | 0       |
| p53−/−           | 16    | 25      | 0       |

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Figure 3. Cellular phenotypes of Asciz-deficient primary fibroblasts. (A) Cumulative population doublings (PDs) of Asciz-deficient and matched WT littermate MEFs in a standardized 3T3 assay. Data are mean ± standard error, n=6. (B) Data from panel A normalized to the PD maximum in the relevant litter. (C–F) DNA damage sensitivity assays. Cells were treated with the indicated doses of MMS, H2O2, 2 mM HU or 50 J/m2 UV, and viability was determined by propidium iodide exclusion and FACS. *p<0.05, **p<0.01, ***p<0.001.

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damage-independent, and performed histological analyses of litters between E12.5 and E18.5. The most striking defect at all time points was the complete absence of lungs in all Asciz-deficient embryos analyzed (n>30; Figure 5A–5C) and apparent lack of tracheal tissue in all but one of these (Figure 5A, 5C). As expected, all five Asciz−/− embryos analyzed again lacked developing pulmonary epithelium (Figure 6B, 6D, Figure S3, and data not shown). One Asciz null embryo contained a very short incompletely separated tracheal stump that ended bluntly where it would normally connect to the main bronchi (Figure 6B). Interestingly, the other Asciz null embryos contained single centrally located bud-like structures that emerged from the ventral oesophagus near the level where the trachea bifurcates into bronchi in the relevant WT littermates (Figure 6D, Figure S5); the central location suggested that this bud-like structure represented tracheal primordium.

Two of the Asciz−/− whole-mount embryos and littermate controls were sectioned at the level of the truncated trachea (Figure 7B, 7B′) or tracheal bud-like structure (Figure 7D, 7D′) for immunofluorescence staining with the respiratory marker Nkx2.1.

The combined trachea and lung defects in Asciz−/− embryos were interesting because both organs originate at the same time but presumably independently of each other from the common respiratory endoderm in the ventral foregut [27,28]. Shortly after specification of respiratory precursors that are characterized by expression of the Nkx2.1 transcription factor, bilateral lung buds and, just rostral of these, a central tracheal primordium emerge from the ventral foregut around E9.5 in the mouse. The junction of these primordia marks the bifurcation of the trachea into the two main bronchi, and the lung buds expand caudally into the surrounding mesoderm to form the bronchial tree and pulmonary epithelium by branching morphogenesis, whereas the trachea septates from the common foregut lumens in an upwards “unzipping” motion [27,28]. Thus, in a simplified view, the origin of the respiratory system can be traced back to the projection of the tracheo-bronchial bifurcation onto the ventral oesophagus.

To more clearly assess the developing trachea and lungs in three dimensions, we performed optical projection tomography (OPT) on whole-mount E-cadherin stained embryos. WT embryos showed clear separation of oesophagus and trachea at the larynx, bifurcation of the trachea into two bronchi and advanced branching of the developing pulmonary epithelium (Figure 6A, 6C). As expected, all five Asciz−/− embryos analyzed again lacked developing pulmonary epithelium (Figure 6B, 6D, Figure S3; and data not shown). One Asciz null embryo contained a very short incompletely separated tracheal stump that ended bluntly where it would normally connect to the main bronchi (Figure 6B). Interestingly, the other Asciz null embryos contained single centrally located bud-like structures that emerged from the ventral oesophagus near the level where the trachea bifurcates into bronchi in the relevant WT littermates (Figure 6D, Figure S3); the central location suggested that this bud-like structure represented tracheal primordium.

ASCIZ is required for separation of respiratory progenitors from the ventral foregut

The combined trachea and lung defects in Asciz−/− embryos were interesting because both organs originate at the same time but presumably independently of each other from the common respiratory endoderm in the ventral foregut [27,28]. Shortly after specification of respiratory precursors that are characterized by expression of the Nkx2.1 transcription factor, bilateral lung buds and, just rostral of these, a central tracheal primordium emerge from the ventral foregut around E9.5 in the mouse. The junction of these primordia marks the bifurcation of the trachea into the two main bronchi, and the lung buds expand caudally into the surrounding mesoderm to form the bronchial tree and pulmonary epithelium by branching morphogenesis, whereas the trachea septates from the common foregut lumens in an upwards “unzipping” motion [27,28]. Thus, in a simplified view, the origin of the respiratory system can be traced back to the projection of the tracheo-bronchial bifurcation onto the ventral oesophagus.

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Two of the Asciz−/− whole-mount embryos and littermate controls were sectioned at the level of the truncated trachea (Figure 7B, 7B′) or tracheal bud-like structure (Figure 7D, 7D′) for immunofluorescence staining with the respiratory marker Nkx2.1. The tracheal stump in the mutant stained homogenously with Nkx2.1 (Figure 7B, bottom panel), similar to the trachea in the WT littermate (Figure 7A), and the ventral part of the tracheal bud-like structure in the other Asciz−/− embryo was also enriched for Nkx2.1 (Figure 7D′) with staining intensity similar to the separated trachea in the matched WT littermate control (Figure 7C′). Interestingly, in stark contrast to the WT oesophagus, some ectopic Nkx2.1-positive cells remained in the ventral part of the oesophagus in the mutant where the trachea had partially separated (Figure 7B, top panel).

We also analysed these sections for expression of p63, a p53-like transcription factor that is normally highly expressed in the oesophagus, but also present in basal cells of the trachea [29]. Under our staining conditions at the developmental stages studied here, p63 seemed only to be present in the oesophagus but not in the trachea in WT embryos (Figure 7A′, 7C). However, p63-
positive cells were readily detectable in the ventral part of the tracheal bud-like structure in the Asciz$^{2/-}$ embryo (Figure 7D), suggesting defective partitioning of specified cells between trachea and oesophagus. As ectopic p63 expression can result from increased Sox2 levels [29,30], a transcription factor involved in foregut separation that is normally highly expressed in the oesophagus and dorsal part of the trachea but downregulated in the ventral part of the developing trachea, we also monitored Sox2 expression in these sections. WT tracheas (Figure 7A, 7C, bottom panels) and the partially separated Asciz$^{2/-}$ trachea (Figure 7B, bottom panel) exhibited the expected dorsally polarized Sox2 expression pattern; in contrast, Sox2 was still expressed at high levels throughout the ventral part of the bud-like structure in the Asciz$^{2/-}$ embryo (Figure 7D). Thus, while impaired local down-regulation of Sox2 could contribute to the Asciz$^{2/-}$ phenotype, it is interesting to note that most of the ectopic Sox2-positive cells in the tracheal bud-like structure were still able to downregulate p63. We also observed aberrantly high Sox2 levels in the ventral foregut in Asciz$^{2/-}$ embryos around E10.25, i.e. before oesophagus and trachea were separated in the matched littermate control with appropriately down-regulated Sox2 (Figure S6), indicating that impaired dorso-ventral patterning of Sox2 expression is not merely a secondary consequence of impaired foregut separation in our mutant.

Altogether, these analyses indicate that Asciz-deficient mice are able to initially specify the respiratory endoderm, based on Nkx2.1 expression, but then fail to remodel the endoderm in a manner required for initiation of lung budding and efficient separation of the trachea.

The ASCIZ SQ/TQ-cluster domain has the propensity to activate transcription

When ASCIZ was originally isolated in a yeast two-hybrid screen [15], we noticed during vector-swapping control experiments that ASCIZ could very strongly activate yeast two-hybrid reporter genes on its own once it was fused to the Gal4 DNA-binding domain (Gal4-DBD). As a large proportion of genes that regulate foregut development function as transcription factors (e.g., Sox2, p63, Nkx2.1 mentioned above), and because the modular domain composition of ASCIZ resembles some ZnF transcription factors (see below), we revisited the yeast reporter system to explore the potential of ASCIZ to function as a transcriptional regulator. Both the four-ZnF 823-residue and the two-ZnF 667-residue splice isoforms of human ASCIZ were able to activate the GAL1-HIS3 and GAL2-ADE2 reporter genes in these one-hybrid assays (Figure 8A). Importantly, similar dual luciferase reporter assays in human U2OS cells using the 667-residue isoform demonstrated that ASCIZ also has an intrinsic ability to
activate gene expression in mammalian cells when tethered to promoters (Figure 8B). Interestingly, truncation analysis revealed that the SQ/TQ-cluster domain - but not the ZnF or core domains - of ASCIZ was sufficient for reporter gene activation (Figure 8A).

Discussion
DNA damage and ATM-related functions of ASCIZ

Here we have shown that Asciz is essential for pulmonary organogenesis during embryonic development in mice, and required for proper DNA base damage responses in primary cells. Although the lung defect is mechanistically most likely unrelated to defective DNA damage responses, the overall phenotype - MMS and H2O2 hypersensitivity and embryonic lethality - is consistent with a role of ASCIZ as an accessory BER factor downstream of glycosylases, as proposed by previous work in human and chicken cells [15,16]. Although Asciz null embryos die a few days earlier and their lung defect is considerably more severe than in case of Polß-deficient embryos, the latter also seem to have a very comparable late gestational growth retardation [10,11], and furthermore, the essential requirement for Polß is also not suppressed by deletion of p53 [9]. Likewise, embryos deficient in Yb-1, another protein recently linked to accessory functions in the BER pathway [31,32], also share overall similar late embryonic growth retardation and lethality, frequent exencephaly and modestly increased cellular oxidative stress-induced senescence phenotypes [33].

In contrast to similarities with BER-related genes, the phenotype of Asciz-deficient mice differs fundamentally from the phenotype of Atm-deficient mice. For example, the key phenotype of Asciz-deficient mice - embryonic lethality with absence of lungs - is not shared by Atm-null mice [20], and the key phenotype of Atm-deficiency - dramatically increased ionizing radiation sensitivity - is not shared by Asciz-deficient cells [16,19]. Consistent with normal ATM protein levels in human, mouse or chicken cells in the absence of ASCIZ, ATM signaling was also unaffected in our Asciz-deficient MEFs or Asciz-depleted human cell lines (Figures S3, S4, and data not shown), including in response to HU, hypotonic NaCl and chloroquine, that required ASCIZ for ATM activation according to Kanu and Behrens [19]. Thus, the completely different phenotypes and absence of ASCIZ effects on ATM stability and activation question the classification of ASCIZ as an “essential co-factor” and regulator of ATM [19]. It is not clear why the other group obtained different results, as our gene targeting strategy was identical to theirs. Kanu and Behrens did not provide genetic background information for their mice, but given that we consistently observed unimpaired ATM levels in Asciz-deficient human, chicken or mouse cells, it seems unlikely that the differing effects could be mouse strain-dependent. As we have confirmed normal ATM levels directly in freshly prepared tissue extracts, we can also exclude the possibility that we may have missed differences in protein levels as a result of variable cell culture conditions. Likewise, given that we did not see a meaningful correlation between ATM and ASCIZ levels in numerous independent AT cell lines, including isogenic AT cell controls reconstituted with WT Atm, as well as genuine mouse and chicken Atm gene deletions (Figure 4), we can only speculate that the previously reported dramatic loss of ASCIZ may be a peculiarity of that particular AT cell line, possibly due to increased genome instability of AT cells. Considering that the positions of 15 potential ATM phosphorylation sites are exactly conserved from chicken to human and mouse ASCIZ, we favour a model where DNA damage-related functions of ASCIZ may be modulated by its direct phosphorylation by ATM. Indeed, our preliminary data that ASCIZ can be directly phosphorylated by ATM in vitro and

Figure 6. Defective pulmonary and tracheal development in Asciz-null embryos. Optical projection tomography of whole-mount E-cadherin stained of E11.5 (A, B) and E12.5 (C, D) littermates. Stippled boxes indicate the approximate plane of sections chosen for immunofluorescence analysis in Figure 7. Panels are arranged with the oesophagus on top.
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Figure 7. Expression analysis of markers of foregut development. Sections from the levels indicated in Figure 6 were stained with the indicated antibodies. All panels are oriented with the oesophagus or dorsal foregut on top. A’–D’ are sections adjacent to the ones shown in A–D. In the merged panel on the left, nuclei are counterstained with DAPI. Scale bars = 20 μm.
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that its MMS-induced focus formation in vivo seems to be at least partially regulated by ATM (to be reported elsewhere in detail) are consistent with a functional interaction between the two proteins.

Role of ASCIZ in lung development

As early lung development is unlikely to be specifically affected by DNA damage signaling, and severe tracheal atresia in Asciz null embryos was surprising, particularly as there are very few mouse mutants with comparable respiratory defects [reviewed in [27,28,34,35]]. Speculation and early development of the respiratory tract is regulated by extensive signaling crosstalk between the foregut endoderm and surrounding mesoderm [27,28], and mouse mutants have revealed major signaling pathways involved in these processes (Figure 9). Double-knockout mice lacking the Gli2 and Gli3 transcription factors of the hedgehog pathway also seem to lack lungs as well as the trachea; however, they also lack the oesophagus indicating a more severe foregut defect [36]. These defects are considerably less severe in sonic hedgehog (Shh) null embryos [37]. Foregut development appears overall normal in Wnt2/Wnt2b double-null embryos as well as Shh-Cre driven conditional β-catenin (Ctnnb1) KO mice, but these never establish complete lung and tracheal agenesis [38,39]. Mice lacking FGF-10 [40,41] or its cognate FGF-receptor 2b [42] also lack lungs, but seem to contain a grossly normal trachea (and are also characterized by a complete absence of limbs in contrast to the Asciz−/− phenotype). Conversely, FoxG1-Cre driven conditional Bmp4 deletion results in selective tracheal agenesis, where the main bronchi and primitive lungs emerge directly from the oesophagus [43]. Based on these comparisons (Figure 9), the Asciz−/− phenotype is less severe than the complete respiratory precursor defect with absence of Nkx2.1 expression and combined agenesis of lungs and trachea in Wnt2/2b and Shh-Cre/β-catenin mutants, but more severe than the respiratory tract defect in Figf10 or FGF-receptor 2b mutants with selective pulmonary agenesis yet preserved tracheal development. Genetically, these data thus suggest a crucial regulatory function for ASCIZ in the regulation of respiratory organogenesis at a level between endodermal β-catenin and mesodermal FGF10 signaling pathways (Figure 9). As FGF10 has been proposed to regulate the downregulation of Sox2 expression in respiratory precursors [30], our finding of impaired dorso-ventral patterning of Sox2 expression in Asciz−/− embryos before foregut separation (Figure S6), and when tracheal separation stalls early (Figure 7D), are also consistent with a role of ASCIZ upstream of FGF10.

The signaling pathways discussed here ultimately regulate developmentally important gene expression programs during specification, morphogenesis and differentiation of the respiratory system. ASCIZ is a predominantly nuclear protein [15,19], its ZnF structure is generally reminiscent of transcriptional regulators [44], and we have shown here that ASCIZ has the propensity to function as a transcriptional activator via its SQ/TQ cluster domain (Figure 8). In some regards, ASCIZ can be considered as a mirror image of the Sp1 transcription factor. Whereas ASCIZ contains a ZnF domain at the N-terminus and an extended SQ/TQ cluster towards the C-terminus, Sp1 that also is essential for murine development [45] contains an SQ/TQ rich N-terminal transcription activation domain and a triple-ZnF domain at the C-terminus. While Sp1 has been extensively studied as a transcription factor, it is now becoming apparent that it also has transcription-independent roles as an ATM substrate that relocates into DNA damage-induced foci [46,47], somewhat similar to our original interest in ASCIZ. Altogether, these
analyses make it tempting to speculate that ASCIZ may regulate pulmonary development as a transcription factor.

In conclusion, we have shown here that ASCIZ has dual functions with a role in the response to DNA lesions that are repaired by the BER pathway, as well as pleiotropic functions during murine embryonic development, most notably as a member of a very select group of essential regulators of respiratory organogenesis. Nks2.1-positive respiratory precursors seem to still be specified in the absence of Asciz, but then fail to properly segregate within the foregut. Impaired foregut separation in Asciz<sup>−/−</sup> embryos seems to correlate with an inability to downregulate Sox2 expression in the ventral foregut, but the exact mechanism responsible for this defect remains to be determined. Asciz null embryos die a few days before birth rather than perinatally from an acute inability to breathe, indicating that additional developmental defects beyond the respiratory system may contribute to the lethality. Our study provides a basis to further investigate how exactly ASCIZ regulates respiratory organogenesis and possibly other developmental processes by expanding the analysis to tissue-specific or temporally regulated conditional knockout systems.

**Methods**

**Ethics statement**

All mouse procedures were approved by the St. Vincent’s Hospital Animal Research Ethics Committee.

**Generation and genotyping of Asciz gene-targeted mice**

The mouse Asciz gene contains four exons spread over ~15 kbp on chromosome 8 and was targeted in C57BL/6 ES cells by integrating loxP sites on both sides of exon D (Figure 1B) using standard homologous recombination, ES cell and blastocyst manipulation techniques as a contracted service by Ozgene Pty Ltd, Perth. A diagnostic ScaI restriction site was integrated just 3′-terminal of the downstream loxP site. Gene targeting was confirmed by Southern blotting using 5′- and 3′-probes located outside the targeting vector. The 5′-probe can be used for Southern blot analysis of ScaI-digested DNA to distinguish between the WT (22.5 kbp), targeted (11.5 kbp), and Asciz-KO (7.6 kbp) alleles (Figure 1B). The germline Asciz KO allele was generated by crossing the targeted line with C57BL/6 mice containing a PGK-Cre knockin in the ROSOM locus, followed by two C57BL/6 backcrosses to remove PGK-Cre and for embryo transfer into the St. Vincent’s Hospital Biological Resources Centre. Thus, the Asciz KO line is on a pure C57BL/6 genetic background. Animals were housed in SPF microisolators. Genotyping can also be performed by PCR using primers F1 (5′-CATGGAATTGT-TAAAAGCTC-3′), F2 (5′-CCGACTGGGGATGTAGTCAG-3′) and R1 (5′-AAAAGATAGAATAGCTACAC-3′), which result in bands of 170 bp for the WT and 220 bp for the KO allele.

Asciz<sup>−/−</sup> mice were crossed with germline p53-targeted mice (deletion of exon 2–10 [48,49]) to generate compound heterozygous, and offspring were genotyped at weaning using primers above and 3′-probes located outside the targeting vector. The 3′-probe can be used for Southern blot analysis of ScaI-digested DNA to distinguish between the WT (15 kbp) and Asciz-KO (9 kbp) alleles (Figure 1B), which result in bands of 170 bp for the WT and 220 bp for the KO allele. Recovery of p53<sup>−/−</sup> mice at weaning was approximately half of the expected Mendelian ratios, a known phenomenon for null homozgyosity on inbred backgrounds [50].

**Embryo analyses**

The time of pregnancies was defined as E0.5 on the morning vaginal plugs were observed in Asciz<sup>−/−</sup> intercrosses. Embryos were dissected from the uterus in cold PBS, weighed after blotting off excess fluid and immediately fixed for histology, or processed for protein extraction or MEF isolation, and genotyped by PCR using yolk sac or tail DNA. For histology, whole embryos were fixed in Bouin’s solution or paraformaldehyde, processed to paraffin and sagittal sections were stained using haematoxylin-eosin and scanned using a Zeiss Mirax Digital Slide Scanner by the Australian Phenomics Network Histopathology and Organ Pathology Service, University of Melbourne, or manually processed and photographed as described [51].

**Cell lines and MEF cultures**

Human and chicken cell lines were cultured as described [15,16,24,32]. MEFs were prepared by dissecting embryos in cold PBS, heads and internal organs were removed, and remaining corpses were sliced into smaller pieces and trypsinized cells were cultured in Dulbecco’s Modified Eagles medium containing 10% fetal calf serum for 2 days, trypsinized, Coulter-counted, and re-seeded at 10<sup>6</sup> cells per 10 cm dish. This passage, defined as P1, was incubated for 3 days, trypsinized, counted and re-seeded at 10<sup>6</sup> cells per 10 cm dish (P2), and this process was repeated for 8 passages. For DNA damage sensitivity assays, 5×10<sup>4</sup> MEFs (P2–P3) were seeded per 35 mm well, grown in Dulbecco’s modified Eagle’s medium and treated as indicated in the figure legend, and after 10 hours cell viability was determined by propidium iodide exclusion using flow cytometry. Each set of DNA damage sensitivity experiments was performed in parallel with MEFs from at least three independent embryos per genotype.

**Optical projection tomography (OPT)**

Staged embryos were stained for OPT [53] with an antibody to E-cadherin (ECCD2, Invitrogen, 1/200 dilution) as described [54], with 48 hour primary and secondary antibody incubations interspersed with extensive 12 hour washes to remove unbound antibody. Samples were imaged on a Biophotonics 3001 OPT machine (Biophotonics, UK) and datasets reconstructed by NRecon (Skyscan, Belgium) and visualized using Drishti (http://anusf.anu.edu.au/Vizlab/drishti/). Embryos were rescued from agarose after imaging, processed to paraffin and sectioned, or directly prepared for cryo-sectioning. After antigen retrieval in citrate buffer sections were stained with antibodies to Nkx2.1 (anti-TTF1, Zymed, 1/200), p63 (Abcam, 1/200), and Sox2 (Chemicon) to examine differentiation.

**Biot analyses**

Southern, Northern and Western blots were performed as described [15,51]. Antibodies against ASCIZ [13], available from Millipore and chicken ATM [52] were described before. Other antibodies: Actin (MAB1501, Millipore), ATM (5G2, Abcam), ATR (sc-1857, Santa Cruz Biotechnology), human p53 (sc-126, Santa Cruz Biotechnology), mouse p53 (1C12, Cell Signaling Technology, PML (sc-5621, Santa Cruz), XRCC1 (sc-11429, Santa Cruz), γH2AX (05-636, Millipore), pS1981(mouse: pS1987)-ATM (200-301-400, Rockland; or 10H11.E12, Cell Signaling Technology), pS15(mouse: pS18)-p53 (9284, Cell Signaling Technology), pT68-Chk2 (2661 or 2197, Cell Signaling Technology).

**Transcription reporter assays**

For yeast assays, ASCIZ constructs were cloned in pAS2.1 and transformed into PJ69-4A, except the isolated SQ/TQ cluster domain that was cloned into the low-level expression vector pGBT9 because its high level expression was toxic in yeast. One-hybrid reporter assays were performed essentially as described previously.

**DNA damage sensitivity experiments**

Sensitivity experiments was performed in parallel with MEFs from at least three independent embryos per genotype.
for two-hybrid assays in our laboratory [55,56] except that plates were supplemented with leucine. For mammalian dual luciferase reporter assays, the 667-residue ASCIZ isoform was cloned into pCDNA3-Gal4DBD for transient transfection of U2OS cells with equal amounts of the reporter vectors pFR-Luc and pRL-CMV for use with the Dual-Luciferase Reporter Assay kit (Promega) according to the manufacturer’s instructions and measurement of luminescence using a Polarstar Optima (BMG Labtechnologies).

Supporting Information

Figure S1 Instability of the residual Asciz mRNA in Asciz null embryos. Northern blot analysis of E14.5 head extracts of 4 separate WT and Asciz null embryos probed with exon C- or D-specific probes and Gapdh as loading control. Markers on the left indicate (from top to bottom) 10 kb, 8 kb, 6 kb, 4 kb and 3 kb. Note that 2 bands of ~5.5 kb and ~3 kb are detected with both Asciz probes in the WT, indicating alternative splicing. The similar size of the main band of the exon D-deleted transcript to the 5.5 kb WT mRNA is likely due to read-through from the exon C splice donor junction (in the absence of an exon D splice acceptor) to a poly-adenylation signal downstream of the loxP site. Image quant phosphoimager density units for these bands are: WT, 21545 ± 1282; KO, 2986 ± 1032.

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Figure S2 Asciz-deficiency only partially impairs base excision repair. Primary MEFs (5–6 embryos per genotype; independent preparations from those shown in Figure S3) were pretreated with 6 mM methoxymethyl (MOA) for 2 hours and then with 0.005% MMS for 18 hours as indicated before propidium iodide exclusion assay by FACS.

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Figure S3 Unpaired ATM signaling in Asciz−/− MEFs. (A) Western blot analysis of WT and Asciz−deficient primary MEF cultures treated with 0.01% MMS or 40–120 μg/ml chloroquine (CHQ) for 4 hours, probed with the indicated antibodies. (B) Western blot analysis of WT and Asciz−deficient primary MEF cultures treated for 4 hours with 0.01% MMS, 2 mM HU or 20 μg/ml bleomycin, or for 1 hour with 50 mM NaCl, probed with the indicated antibodies (top panels); identical experiments except that MMS treatment was for only 1 hour (bottom panels). (C) Western blot analysis of WT and Asciz−deficient primary MEF cultures treated with 0.025% MMS for 3 hours and 15 μM KU55933 (ATMi with pretreatment for 2 hours before MMS addition) as indicated.

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Figure S4 Unpaired ATM signaling in Asciz−depleted human U2OS cells. (A) U2OS cells were treated with GL2 control or Asciz siRNA si579 [15] and treated with 0.02% MMS for the indicated times. (B) U2OS cells were treated with GL2 control or two separate Asciz siRNAs as described and lysed 1 hour after 2 Gy gamma irradiation, and blotted with the indicated antibodies. The arrow points to the position of ATM in the pS1981-ATM blot, the more abundant upper band represents cross-reactivity of the antibody with near-identical phosphorylation sites in a larger protein, possibly 53BP1. Blots above and below the lines are from separate experiments. Note that an older ASCIZ antibody batch was used for this experiment that crossreacts with a ~100 kDa band just below ASCIZ not observed with the new antibody batch in the other figures.

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Figure S5 Additional embryo analyses. E12.5 WT and Asciz−/− littermates were stained with E-cadherin for whole-mount optical projection tomography similar to Figure 6.

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Figure S6 Analysis of marker expression before foregut separation. Cryo-sections of E10.25 WT and Asciz−/− littermates stained with the indicated antibodies. Panels are oriented with the dorsal foregut on top.

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Author Contributions

Conceived and designed the experiments: SJ IS BvD AH CJM SVK HO MK LAC TJC KiY YT ST MFL JH. Wrote the paper: JH.

BvD NT AH KH JLN CJM SVK HO MK JH. Analyzed the data: SJ IS BvD AH JLN CJM SVK HO MK JH. Contributed reagents/materials/analysis tools: SJ IS BvD AH JLN CJM SVK HO MK JH.

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