

# Defective prelamin A processing and muscular and adipocyte alterations in *Zmpste24* metalloproteinase-deficient mice

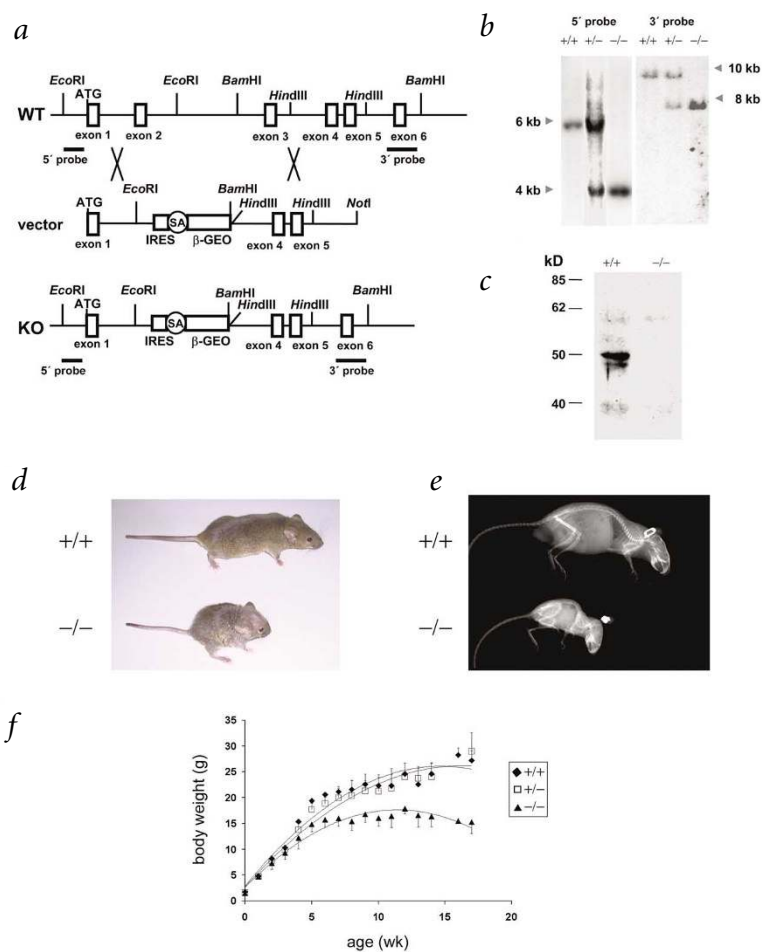
Alberto M. Pendás<sup>1\*</sup>, Zhongjun Zhou<sup>2\*</sup>, Juan Cadiñanos<sup>1\*</sup>, José M.P. Freije<sup>1</sup>, Jianming Wang<sup>2</sup>, Kjell Hultenby<sup>3</sup>, Aurora Astudillo<sup>4</sup>, Annika Wernerson<sup>3</sup>, Francisco Rodríguez<sup>1</sup>, Karl Tryggvason<sup>2</sup> & Carlos López-Otín<sup>1</sup>

\*These authors contributed equally to this work.

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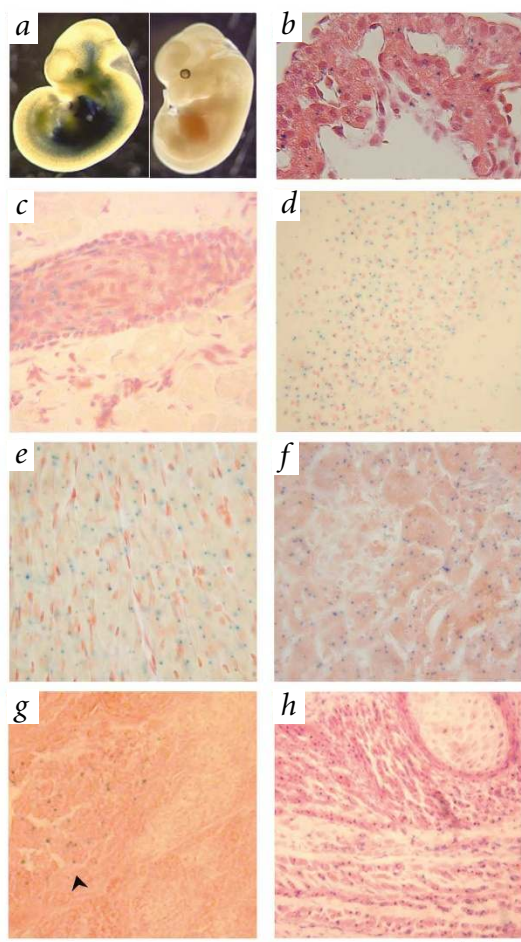
The mouse ortholog of human FACE-1, *Zmpste24*, is a multi-spanning membrane protein widely distributed in mammalian tissues<sup>1,2</sup> and structurally related to Afc1p/ste24p, a yeast metalloproteinase involved in the maturation of fungal pheromones<sup>3</sup>. Disruption of the gene *Zmpste24* caused severe growth retardation and premature death in homozygous-null mice. Histopathological analysis of the mutant mice revealed several abnormalities, including dilated cardiomyopathy, muscular dystrophy and lipodystrophy. These alterations are similar to those developed by mice deficient in A-type lamin<sup>4</sup>, a major component of the nuclear lamina<sup>5</sup>, and phenocopy most defects observed in humans with diverse congenital laminopathies<sup>6–8</sup>. In agreement with this finding, *Zmpste24*-null mice are defective in the proteolytic processing of prelamin A. This deficiency in prelamin A maturation leads to the generation of abnormalities in nuclear architecture that probably underlie the many phenotypes observed in both mice and humans with mutations in the lamin A gene. These results indicate that prelamin A is a specific substrate for *Zmpste24* and demonstrate the usefulness of genetic approaches for identifying the *in vivo* substrates of proteolytic enzymes.

To clarify the function of *Zmpste24*, and to identify *in vivo* substrates targeted by this proteinase, we generated *Zmpste24*-null mice (Fig. 1a). After heterozygote intercrossing, we obtained *Zmpste24*-null, heterozygous and wildtype mice in the expected mendelian ratio. We verified homozygosity with respect to the mutated allele by Southern blot (Fig. 1b), and lack of *Zmpste24*



**Fig. 1** Generation of *Zmpste24*-deficient mice. **a**, Schematic representation of the wildtype *Zmpste24* locus (WT), targeting vector and targeted allele (KO). Positions of restriction enzyme sites and probes used for Southern-blot analysis are shown. **b**, Southern-blot analysis of genomic DNA from three littermate progeny of *Zmpste24* heterozygote crosses. Probing of *EcoRI*-digested DNA revealed fragments of 6 kb and 4 kb for wildtype and disrupted alleles, respectively. Probing of *BamHI*-digested DNA revealed fragments of 10 kb and 8 kb for wildtype and disrupted alleles, respectively. **c**, Extracts of kidneys from wildtype and *Zmpste24*<sup>-/-</sup> mice were examined with a monoclonal antibody directed against the C terminus of *Zmpste24*. **d**, Photograph of two littermate progeny of a *Zmpste24* heterozygote cross, at 3 mo. **e**, Radiograph of a *Zmpste24*<sup>-/-</sup> mouse at 3 mo, compared with a wildtype control. **f**, Cumulative plot of body weight versus age. Dots represent mean values, and error bars indicate s.e.m.

<sup>1</sup>Departamento de Bioquímica y Biología Molecular, Facultad de Medicina, Instituto Universitario de Oncología, Universidad de Oviedo, 33006 Oviedo, Spain. <sup>2</sup>Division of Matrix Biology, Department of Medical Biochemistry and Biophysics, Karolinska Institutet, Stockholm, Sweden. <sup>3</sup>Department of Pathology, Karolinska Institutet, Huddinge University Hospital, Huddinge, Sweden. <sup>4</sup>Servicio de Anatomía Patológica, Hospital Central de Asturias, Oviedo, Spain. Correspondence should be addressed to C.L.-O. (e-mail: CLO@correo.uniovi.es).



**Fig. 2** Expression analysis of *Zmpste24* during mouse development and in adult tissues. **a**, Whole-mount staining of E10.5 embryos showing X-gal staining in heart, skin and central nervous system (left). **b**, Section of the same embryo showing X-gal staining in cardiomyocytes. **c–h**, X-gal staining of organ cryostat sections from an adult heterozygous mouse. **c**, Skin showing staining in keratinocytes of a hair follicle and in dermal fibroblasts. **d**, Hippocampus of the brain. Most of the staining localized to neuronal cells. **e**, Heart showing staining in cardiomyocytes. **f**, Kidney showing staining mainly in epithelial cells of proximal/distal tubules. **g**, Pancreas showing positive staining in endocrine (arrow), but not exocrine, tissue. **h**, Paravertebral muscle. Sections were counterstained with either eosin (**b,f,g,h**) or safranin (**c–e**). Scale bars represent 50  $\mu$ m in **b–g** and 200  $\mu$ m in **h**.

took advantage of the *lacZ-neo<sup>r</sup>* fusion expressed under the control of the *Zmpste24* promoter as a consequence of the targeting strategy. We saw extensive X-gal staining as early as embryonic day (E) 10.5, with the strongest activity being detected in the heart (Fig. 2a). We also carried out staining on tissue sections from mice at different stages of postnatal development, from 3 to 24 weeks of age (Fig. 2b–h). The intensity of the X-gal staining increased with age. We observed strong signals in cardiomyocytes, proximal and distal tubules and collecting ducts of kidney, neuronal cells of brain and the basal layer of epidermis, as well as hair follicles and fibroblasts of dermis. We also detected staining in pancreatic islets, smooth muscle cells of intestine, urinary bladder, blood vessels, seminiferous tubules of the testes, epithelial cells of the thyroid and skeletal muscle cells (Fig. 2 and data not shown).

As *Zmpste24* is highly expressed in the heart and kidney, we carried out serum and blood analyses to explore the possible causes of death. Blood glucose and serum triglyceride levels did not differ between *Zmpste24*-null mice and controls prior to the stage in which the mutants began to lose weight. Levels of both biochemical markers decreased in mutant animals that were losing weight, however (see Web Table A online). The serum levels of aspartate aminotransferase, creatine kinase, glutamate dehydrogenase and creatinine were normal in *Zmpste24*-null mice until 8 weeks of age. In end-stage mutants (approximately 20 weeks), however, we noticed a significant rise in serum aspartate aminotransferase, creatine kinase and glutamate dehydrogenase levels, whereas serum creatinine levels remained normal (see Web Table A online). These results suggest damage of the cardiac muscle. In agreement with serological data, we observed a significant thinning of the ventricular wall and, in some cases, a dilatation of both right and left ventricles. Microscopic analysis revealed foci of muscle degeneration, infiltration of inflammatory cells and interstitial fibrosis (Fig. 3a,b). We found an increased number of lipid droplets in hepatocytes surrounding the hepatic veins in liver from null mice, suggesting a chronic cardiac dysfunction (Fig. 3c). Taken together, these abnormalities observed in the heart suggest that dilated cardiomyopathy and heart failure probably caused the death of the *Zmpste24*-null mice.

In addition to the heart pathology, muscular fibers of the paravertebral region and the deltoid and quadriceps were dystrophic, showing a significant increase in abnormal small round fibers (Fig. 3d–e). Moreover, about one-third of the *Zmpste24*-null mice over 16 weeks of age started to lose their fur, whiskers and sometimes eyelashes. Microscopic analysis revealed that mutant mice at 16 weeks had completely lost the subcutaneous fat layer (Fig. 4). The epidermis and hair follicles were atrophic. There was an increased number of apoptotic bodies in the basal layer of the epidermis and in hair follicles, indicating the involvement of apoptosis in the pathogenesis of these skin-related lesions (Fig. 4). Notably, we saw thymic hypoplasia with loss of corticomedullary demarcation in young null mice. Finally, there was a

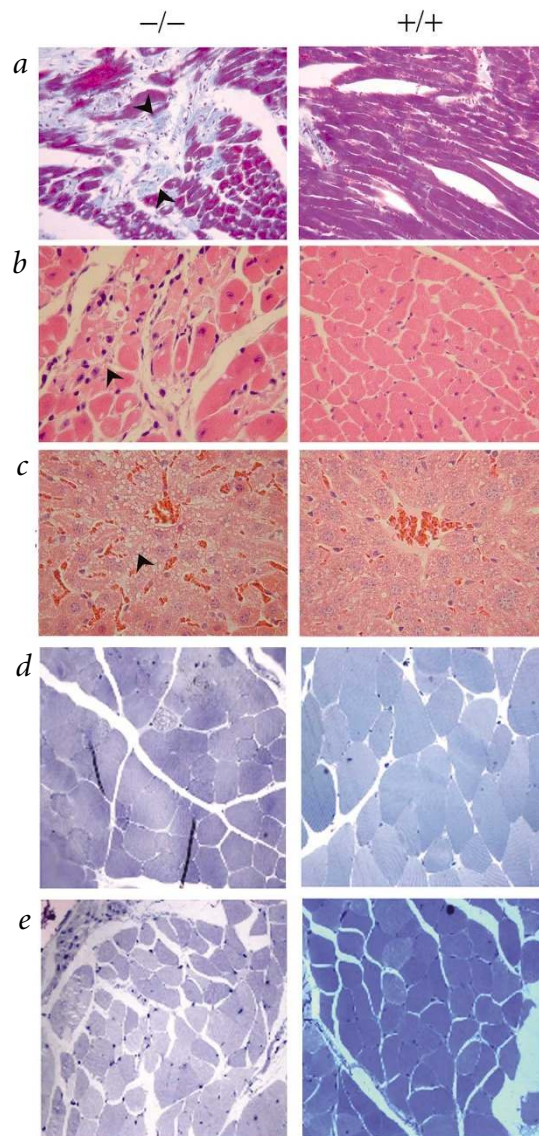
protein by western blot (Fig. 1c). Heterozygous mice were morphologically indistinguishable from their wildtype littermates. At birth, *Zmpste24*-null mice were also indistinguishable from their heterozygous or wildtype siblings and seemed healthy, until four weeks of age. Subsequently, null mice could be distinguished from wildtype littermates by size and weight. Within four to six weeks, their growth rate was reduced; by seven weeks, they stopped growing, despite normal feeding habits. At this stage, null mice were 30% smaller and had gained 40% less weight than their wildtype or heterozygous littermates (Fig. 1d–f). After two months of age, homozygous null mice began to progressively lose weight, showing an abnormal posture characterized by a hunched position and scoliosis (Fig. 1e). They became less mobile and began to slap and splay their hind paws while walking. When lifted by their tails, they reflexively overextended their hind limbs, trembled and were unable to bow upwards, unlike wildtype mice. In addition, most *Zmpste24*-null mice died prematurely, having an average lifespan of 20 weeks. We dissected animals between 14 and 20 weeks of age and found that the relative weight of the heart was greater in the mutants than in their littermate controls ( $0.53\% \pm 0.028\%$  versus  $0.44\% \pm 0.014\%$ , mean  $\pm$  s.e.m.;  $P < 0.05$ ). The relative weight of the kidneys was also higher in the *Zmpste24*-null mice than in controls ( $P < 0.05$ ).

To analyze the molecular alterations underlying the observed abnormalities in *Zmpste24*-null mice, we first studied *Zmpste24* expression during mouse development and in adult tissues. Northern-blot analysis identified a transcript of 3.5 kb in all tissues, indicating widespread expression of this gene (data not shown). To dissect more precisely the spatial and temporal expression pattern of *Zmpste24* during mouse development, we

**Fig. 3** Histology of heart, liver and muscle from *Zmpste24*-null (left) and wildtype (right) mice. **a**, Gomori trichrome staining showing foci of muscle degeneration and fibrosis in mutant heart (left). Arrows indicate foci of fibrosis of the heart. Collagen was stained blue-green. **b**, Higher magnification ( $\times 400$ ) of a heart section stained with hematoxylin and eosin showing muscle degeneration, lymphocytic infiltration, interstitial fibrosis and intracellular vesicles in the mutant heart (left). Arrow indicates a degenerating cardiac muscle and infiltration of lymphocytes. **c**, Histology of liver ( $\times 400$ ) showing congestion and fatty degeneration of liver in the mutant mouse (left). Arrow indicates intracellular lipid accumulation. **d,e**, Semi-thin sections of quadriceps (**d**) and paravertebral (**e**) muscles at  $\times 400$  from mutant (left) and wildtype (right) mice. Scale bars represent 50  $\mu\text{m}$  in **a** and **c**, and 25  $\mu\text{m}$  in **b,d** and **e**.

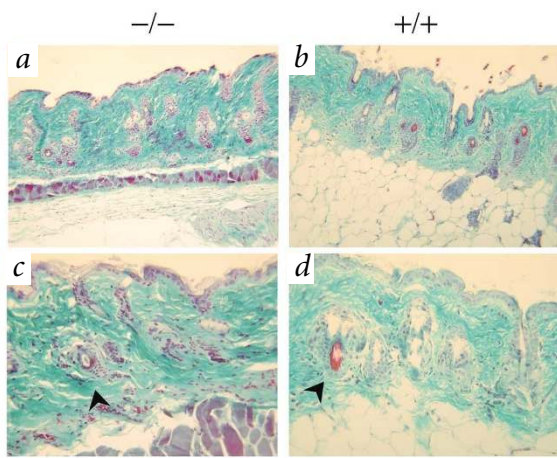
growth plate dysplasia in the distal femur and proximal tibia (data not shown). Mice deficient in *Zmpste24* have previously been generated, but no phenotypic abnormalities or putative substrates targeted by this enzyme in mammalian cells were reported<sup>9</sup>. Apparently, these null mice were analyzed only during early development, which resulted in failure to detect the many phenotypes described here that are shown later in mouse development. Nevertheless, we cannot rule out the possibility that putative differences between those mice and the *Zmpste24*-null mice described here may be caused by the different targeting strategies used for inactivating the gene. The abnormalities that we observed in *Zmpste24*-null mice, including postnatal growth retardation, shortened lifespan, loss of fat layer and muscular dystrophy, are very similar to those developed by mice lacking A/C-type lamin<sup>4</sup>, a major protein component of the nuclear lamina<sup>5</sup>. Moreover, mutations in the gene encoding lamin A are responsible for a series of rare and seemingly unrelated human diseases that resemble the phenotype observed in the *Zmpste24*-null mice. These genetic disorders include dilated cardiomyopathy with conduction-system disease, Dunnigan-type familial partial lipodystrophy, Emery-Dreifuss muscular dystrophy and limb girdle muscular dystrophy (type 1B)<sup>6–8</sup>.

The finding that the *Zmpste24*-null phenotype in mice seemed to recapitulate the diverse pathological features associated with lamin A deficiencies in both human and mice led us to hypothesize that lamin A could be an *in vivo* substrate of *Zmpste24*. The lamin A protein is synthesized as a precursor that undergoes a complex maturation process, initiated by the farnesylation of a cysteine residue present in a conserved CaaX motif (C, cysteine; a, aliphatic; X, any residue) at the carboxy-terminal end of this protein. This step is followed by removal of the aaX tripeptide and methylation of the carboxyl group of the S-prenyl-cysteine<sup>10</sup>. These modifications are shared by other mammalian proteins,

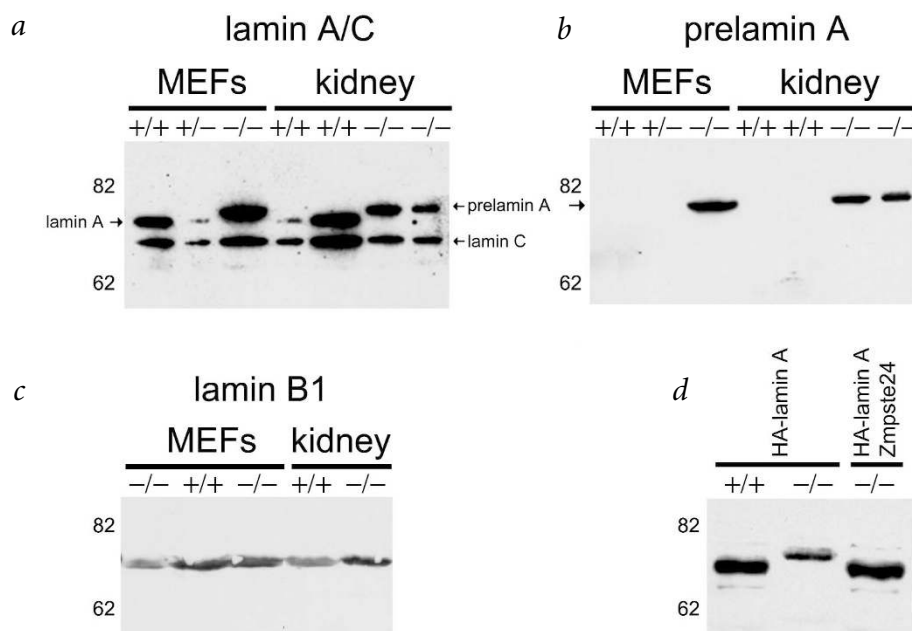


including the Ras family of oncoproteins and some heterotrimeric G proteins<sup>11</sup>. Farnesylated prelamin A undergoes a second endoproteolytic cleavage characteristic of this protein, however, resulting in the release of 15 amino acids from its C-terminal region, including the farnesylated cysteine residue<sup>12</sup>. The enzyme responsible for this specific endoproteolytic maturation step of the lamin A precursor has not been identified.

To evaluate whether *Zmpste24* is involved in prelamin A processing, we first examined by western blot the maturation stage of this protein in cells and tissues from *Zmpste24*-null mice. Murine embryonic fibroblasts (MEFs) and tissue extracts from mutant mice lack the mature lamin A band (72 kD) present in wildtype extracts. Instead, we observed a band of the expected molecular mass for prelamin A (74 kD; Fig. 5a). We confirmed this finding by hybridizing the same blot with an antibody



**Fig. 4** Masson trichrome staining of skin and subcutaneous tissue, showing lipodystrophy. The *Zmpste24*-null mouse (left) shows complete loss of subcutaneous adipose tissue and severe atrophic epidermis and hair follicles (arrow), as compared with the wild type (right). **a,b**, Lower magnification ( $\times 200$ ). **c,d**, Higher magnification ( $\times 400$ ). Scale bars represent 100  $\mu\text{m}$  in **a** and **b**, and 50  $\mu\text{m}$  in **c** and **d**.

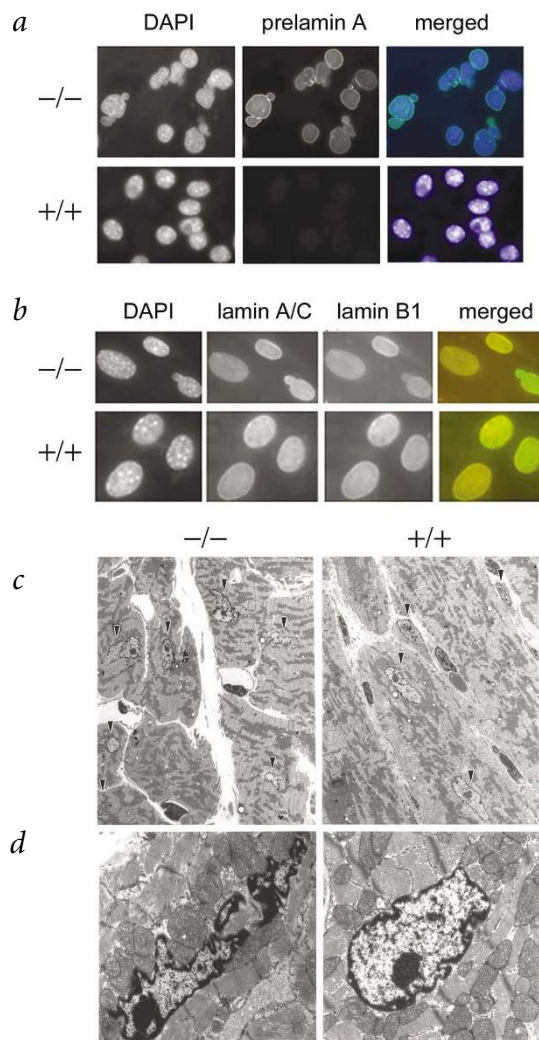


**Fig. 5** Loss of *Zmpste24* prevents prelamin A processing. **a–c**, Extracts of MEFs and kidney from wildtype and mutant mice were probed with anti-lamin A/C (**a**), anti-prelamin A (**b**) or anti-lamin B1 antibodies (**c**). **d**, Wildtype and mutant MEFs were transfected with a hemagglutinin-tagged lamin A construct alone or together with a construct expressing *Zmpste24* cDNA. Extracts of the transfected cells were analyzed with an anti-hemagglutinin monoclonal antibody.

against the C-terminal tail, which is proteolytically removed during prelamin A maturation (Fig. 5b). High accumulation of prelamin A was seen in *Zmpste24*<sup>-/-</sup> cell extracts, but was absent in wildtype controls. By contrast, we did not see molecular weight changes in either lamin C (Fig. 5a), a splice variant of lamin A without the CaaX box, or in lamin B1 (Fig. 5c), a CaaX-prenylated lamin that does not undergo the second proteolytic maturation event characteristic of lamin A. To demonstrate a direct role of *Zmpste24* in the processing of prelamin A, we transfected wildtype and *Zmpste24*<sup>-/-</sup> MEFs with an hemagglutinin-tagged murine prelamin A cDNA, alone or in combination with a *Zmpste24* expression construct. When we analyzed the transfected cells by western blot, we detected bands of 72 kD and 74 kD in wildtype and *Zmpste24*-null mice, respectively, confirming the lack of prelamin A processing in the mutant cells (Fig. 5d). Co-transfection with the *Zmpste24* expression plasmid resulted in the appearance of the 72 kD band in *Zmpste24*<sup>-/-</sup> MEFs, indicating that re-expression of the proteinase is sufficient to revert the deficiency in prelamin A processing observed in these cells (Fig. 5d). These results indicate that *Zmpste24* is necessary for the *in vivo* maturation of prelamin A.

To examine whether the observed deficiency in prelamin A maturation causes nucleoskeleton abnormalities, we carried out immunofluorescence analysis of nuclear lamina components in MEFs and newborn fibroblasts of both wildtype and *Zmpste24*-null mice. Prelamin A was associated with the nuclear envelope in mutant cells (Fig. 6a). Co-localization of lamin B1 and lamin A/C confirmed this observation. Prelamin A, however, could not be readily detected in wildtype nuclei under the same conditions. By contrast, we did not observe differences in lamin B1 and emerin (two additional intermediate filament components of the nuclear lamina) between *Zmpste24*<sup>-/-</sup> cells and wildtype controls (Fig. 6b and data not shown). We obtained similar results in

wildtype and mutant MEFs transfected with a prelamin A cDNA that was tagged with hemagglutinin at its amino-terminal end (data not shown). These labeling experiments also revealed changes in the nuclear morphology of cultured mutant cells. Whereas nuclei of MEFs and neonatal fibroblasts are circular or ovoid, those of mutant cells are often of irregular shape and show herniation-like nuclear blebs (Fig. 6a,b). Electron microscopy



**Fig. 6** Nuclear envelope architecture in wildtype and *Zmpste24*-deficient cells. **a,b**, MEFs were analyzed by immunofluorescence with anti-prelamin A (**a**), or with anti-lamin A/C and anti-lamin B1 antibodies (**b**) and counterstained with DAPI. **c**, Electron microscope micrographs of cardiac muscle fibers and nuclei of cardiac muscle cells (arrowhead) showing irregular shape of the nuclei in mutant mice (left) compared with wildtype mice (right). **d**, Higher magnifications of the nuclei of cardiac muscle from mutant end-stage (left) and wildtype (right) mice. Scale bars represent 5  $\mu$ m in **c** and 1  $\mu$ m in **d**.



analysis also confirmed the abnormalities in nuclear architecture of mutant cardiac muscle cells (83% abnormal shape in mutant nuclei versus 8% in wildtype; see Methods; Fig. 6c,d). Taken together, these results indicate that although uncleaved prelamin A is able to assemble normally in the nucleus during interphase, the integrity of the nuclear envelope becomes partially compromised in *Zmpste24*<sup>-/-</sup> cells. In addition, and consistent with the proposed role of *Zmpste24* in prelamin A processing, immunofluorescence analysis of hemagglutinin-*Zmpste24* transfected cells revealed that this protease localizes at the endoplasmic reticulum and at the nuclear envelope (data not shown).

The links between prelamin A accumulation and tissue alterations could operate through two distinct but not mutually exclusive mechanisms. First, the altered structure of the nuclear lamina could affect directly mechanically stressed tissues, such as the muscle fibers of the heart and skeleton, where lamin B1 is not expressed<sup>13</sup>. In addition to this structural function, there is growing evidence that lamins are implicated in DNA replication, chromatin organization, regulation of transcription and cell division. These effects on nuclear dynamics may account for many of the clinical features and tissue-specific alterations observed in human laminopathies<sup>14–16</sup>. Our studies of *Zmpste24*-deficient mice confirm and extend these proposals. We provide further structural evidence of the occurrence of marked nuclear abnormalities associated with these diseases in mutant mice. The finding that proteins interacting with prelamin A and lamin A, like emerin, are incorporated into the nuclear lamina in mutant cells suggests that beyond structural alterations, downstream regulatory effects must also contribute to the observed phenotypes. Notably, the *Zmpste24*-null mice described here have a milder and later-acting phenotype than that observed in lamin A/C-deficient mice<sup>4</sup>. These *Zmpste24*-null mice can be considered a multiple knockout model, owing to the simultaneous deficiency of lamin A and lamin C and to the functional inactivation of emerin, which is mislocalized in lamin A/C-null cells<sup>4</sup>. Our *Zmpste24* knockout model thus resembles more precisely the phenotype observed in individuals with hereditary laminopathies and may provide a useful tool to further dissect the mechanisms by which lamin alterations compromise muscular and adipocyte function, leading to these progressive and life-threatening diseases.

## Methods

**Gene targeting of *Zmpste24*.** We isolated PAC clones enclosing *Zmpste24* from a 129 Sv library (HGMP Resource Centre, UK). This gene consists of 1,475 bp of coding sequence, organized in 10 exons distributed along a genomic segment of 38 kb (Fig. 1a). To create *Zmpste24*-null mice, we designed a targeting vector to replace exons 2 and 3 with a promoter-less IRES  $\beta$ -geo containing a splice acceptor sequence. Expression of the *lacZ* reporter gene was thus driven by the endogenous *Zmpste24* promoter, and the *Zmpste24* coding sequence was interrupted after codon 39. After electroporation of *NotI*-linearized targeting in 129 Ola embryonic stem cells, Southern-blot analysis of 120 neomycin-resistant colonies revealed homologous recombination in six independent clones. We used two of these heterozygote clones to generate chimeric mice. Five highly chimeric males were bred to C57BL/6 females; two gave germline transmission of the *Zmpste24* targeted allele, as confirmed by PCR genotyping and Southern-blot analysis. Animal experimentation was done in accord with the guidelines of the Universidad de Oviedo and the Karolinska Institutet regarding the care and use of laboratory animals.

**Histology, whole-mount staining and  $\beta$ -gal histochemistry.** We fixed samples with cold 4% paraformaldehyde in PBS, processed them into serial paraffin sections and stained them with hematoxylin and eosin or with the Masson's trichromatic staining. We stained epoxy semithin muscle sections

with toluidine blue following standard procedures. We carried out whole-mount embryo staining and histochemical analysis of  $\beta$ -galactosidase reporter activity in frozen sections as described<sup>17</sup>.

**Electron microscopy.** We killed mice by cervical dislocation and immediately placed the heart in 2% glutaraldehyde and 0.5% paraformaldehyde in 0.1 M sodium cacodylate buffer containing 0.1 M sucrose and 3 mM  $\text{CaCl}_2$ , pH 7.4. We cut small pieces from the myocardium and stored them in the fixative overnight at 4 °C. We rinsed samples in 0.15 M sodium cacodylate buffer containing 3 mM  $\text{CaCl}_2$  and post-fixed them in 2%  $\text{OsO}_4$  in 0.07 M sodium cacodylate buffer containing 3 mM  $\text{CaCl}_2$  for 2 h. Specimens were dehydrated and embedded in LX-112, and we examined them in a Tecnai 10 transmission electron microscope at 80 kV.

**Evaluation of nuclear morphology in heart muscles.** We defined normal nuclei as spool-shaped or rounded, and pathological nuclei as irregular and more elongated. We randomly took electron micrographs and blindly selected counting areas at low magnification. We counted the number of 'pathological nuclei' at  $\times 4400$  and divided by the total number of nuclei in controls and knockouts, respectively. We analyzed a total of 60 wildtype and 104 mutant nuclei.

**Expression constructs.** To produce hemagglutinin-lamin A (epitope-tagged at the N terminus), we subcloned a PCR fragment (*BglII*-*XbaI*) encoding mouse prelaminA in-frame into pcDNA3-HA (Invitrogen). Human *FACE-1* cDNA was subcloned into pcDNA3.

**Western blot and immunofluorescence analysis.** We separated in SDS-polyacrylamide gels total protein extracts homogenized in SDS lysis buffer containing Tris-HCl (60 mM, pH 6.8), 1% SDS, 8% glycerol. We transferred proteins to nitrocellulose and incubated them overnight at 4 °C with the different primary antibodies polyclonal anti-prelamin A (sc-6214, Santa Cruz Biotechnology), polyclonal anti-lamin B1 (sc-6217, Santa Cruz Biotechnology), monoclonal anti-lamin A/C (Manlac-1, provided by G. Morris), monoclonal anti-*FACE-1* (Daiichi) and monoclonal anti-HA (Roche). We used a secondary antibody (rabbit anti-goat or goat anti-mouse) coupled to peroxidase for detection of the reaction with Supersignal Chemiluminiscent Substrate (Pierce). We derived MEFs from embryos at E13.5 as previously described<sup>18</sup> and transiently transfected them with Eugene-6 (Roche). For immunofluorescence, we fixed cells in cold paraformaldehyde, rinsed them in PBS and permeabilized them with 0.2% Triton X-100. Finally, we incubated cells with the corresponding primary antibodies diluted in 15% FCS in PBS. The rhodamine and FITC-conjugated secondary antibodies were from Santa Cruz Biotechnology. We counterstained nuclei with DAPI.

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*Note: Supplementary information is available on the Nature Genetics website.*

## Competing interests statement

The authors declare that they have no competing financial interests.

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1. Freije, J.M.P. *et al.* Identification and chromosomal location of two human genes encoding enzymes potentially involved in proteolytic maturation of farnesylated proteins. *Genomics* **58**, 270–280 (1999).
2. Tam, A. *et al.* Dual roles for Zmpste24p in yeast a-factor maturation: NH<sub>2</sub>-terminal proteolysis and COOH-terminal CAAX processing. *J. Cell Biol.* **142**, 635–649 (1998).
3. Boyartchuk, V.L., Ashby, M.N. & Rine, J. Modulation of ras and a-factor function by carboxyl-terminal proteolysis. *Science* **275**, 1796–1800 (1997).
4. Sullivan, T. *et al.* Loss of A-type lamin expression compromises nuclear envelope integrity leading to muscular dystrophy. *J. Cell Biol.* **147**, 913–920 (1999).
5. Fisher, D.Z., Chaudhary, N. & Blobel, G. cDNA sequencing of nuclear lamins A and C reveals primary and secondary structural homology to intermediate filament proteins. *Proc. Natl Acad. Sci. USA* **83**, 6450–6454 (1986).
6. Bonne, G. *et al.* Mutations in the gene encoding lamin A/C cause autosomal dominant Emery-Dreifuss muscular dystrophy. *Nature Genet.* **21**, 285–288 (1999).
7. Fatkin, D. *et al.* Missense mutations in the rod domain of the lamin A/C gene as causes of dilated cardiomyopathy and conduction-system disease. *N. Engl. J. Med.* **341**, 1715–1724 (1999).
8. Shackleton, S. *et al.* LMNA, encoding lamin A/C, is mutated in partial lipodystrophy. *Nature Genet.* **24**, 153–156 (2000).
9. Leung, G.K. *et al.* Biochemical studies of Zmpste24-deficient mice. *J. Biol. Chem.* **276**, 29051–29058 (2001).
10. Sinensky, M. *et al.* The processing pathway of prelamin A. *J. Cell Sci.* **107**, 61–67 (1994).
11. Zhang, F.L. & Casey, P.J. Protein prenylation: molecular mechanisms and functional consequences. *Annu. Rev. Biochem.* **65**, 241–269 (1996).
12. Weber, K., Plessmann, U. & Traub, P. Maturation of nuclear lamin A involves a specific carboxy-terminal trimming, which removes the polyisoprenylation site from the precursor; implications for the structure of the nuclear lamina. *FEBS Lett.* **257**, 411–414 (1989).
13. Maniatis, S. *et al.* Distribution of emerin and lamins in the heart and implications for Emery-Dreifuss muscular dystrophy. *Hum. Mol. Genet.* **8**, 353–359 (1999).
14. Cohen, M., Lee, K.K., Wilson, K.L. & Gruenbaum, Y. Transcriptional repression, apoptosis, human disease and the functional evolution of the nuclear lamina. *Trends Biochem. Sci.* **26**, 41–47 (2001).
15. Wilson, K.L., Zastrow, M.S. & Lee, K.L. Lamins and disease: insights into nuclear infrastructure. *Cell* **104**, 647–650 (2001).
16. Morris, G.E. & Maniatis, S. Heart to heart: from nuclear proteins to Emery-Dreifuss muscular dystrophy. *Hum. Mol. Genet.* **8**, 1847–1850 (1999).
17. Hogan, B., Beddington, R., Costantini, F. & Lacy, E. *Manipulating the Mouse Embryo: a Laboratory Manual* (Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York, 1994).
18. Palmero, I. & Serrano, M. Induction of senescence by oncogenic Ras. *Methods Enzymol.* **333**, 247–256 (2001).