Knock-in human FGFR3 achondroplasia mutation as a mouse model for human skeletal dysplasia

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Achondroplasia (ACH), the most common genetic dwarfism in human, is caused by a gain-of-function mutation in fibroblast growth factor receptor 3 (FGFR3). Currently, there is no effective treatment for ACH. The development of an appropriate human-relevant model is important for testing potential therapeutic interventions before human clinical trials. Here, we have generated an ACH mouse model in which the endogenous mouse Fgfr3 gene was replaced with human FGFR3G380R (FGFR3ACH) cDNA, the most common mutation in human ACH. Heterozygous (FGFR3ACH/+ ) and homozygous (FGFR3ACH/ACH) mice expressing human FGFR3G380R recapitulate the phenotypes observed in ACH patients, including growth retardation, disproportionate shortening of the limbs, round head, mid-face hypoplasia at birth, and kyphosis progression during postnatal development. We also observed premature fusion of the cranial sutures and low bone density in newborn FGFR3G380R mice. The severity of the disease phenotypes corresponds to the copy number of activated FGFR3G380R, and the phenotypes become more pronounced during postnatal skeletal development. This mouse model offers a tool for assessing potential therapeutic approaches for skeletal dysplasias related to over-activation of human FGFR3, and for further studies of the underlying molecular mechanisms.

Gain-of-function point mutations in fibroblast growth factor receptor 3 (FGFR3) cause a variety of congenital skeletal dysplasias inherited as an autosomal dominant trait. These skeletal dysplasias are characterised by varying degrees of skeletal deformities ranging from least to most severe as follows: hypochondroplasia (HCH), achondroplasia (ACH), severe achondroplasia with developmental delay and acanthosis nigricans, and thanatophoric dysplasia (TD) types 1 and 2. Achondroplasia (ACH) (OMIM 100800) is the most common form of genetic short-limbed dwarfism in human. ACH is characterised by short stature with disproportionately short limbs, macrocephaly, characteristic faces with frontal bossing, midface hypoplasia, and exaggerated thoracolumbar kyphosis¹. Over 99% of individuals affected with ACH have the same point mutation, G380R, in the transmembrane domain of FGFR3 protein (FGFR3G380R)²,³. The clinical features of heterozygous ACH are consistent among patients, and homozygous ACH causes severe skeletal deformities that lead to early death. The majority of ACH cases (over 80%) occur spontaneously through mutations in sperm related to advanced paternal age⁴.

FGFR3 is expressed mainly in proliferating chondrocytes in the developing long bones⁵ and has been proven to be a negative regulator of endochondral bone growth⁶. Morphometric examination revealed the shortening of growth plates in ACH patients⁷. Several ACH mouse models have been established to study the roles of FGFR3 in skeletal development and disease. Three ACH mouse models express murine ACH mutation (Fgfr3G380R), introduced using knock-in⁸,⁹ or transgenic¹⁰ approaches, and one ACH mouse model transgenically expresses human FGFR3G380R¹¹. These mouse models share some ACH phenotypes. However, some phenotypes have not been fully described or examined in these models.

Currently, there is no effective treatment for skeletal dysplasias caused by activating mutations of FGFR3. Several potential therapeutic strategies targeting either the over-activated FGFR3 or its downstream effects are currently under development. An FGFR3-binding peptide¹², a C-type natriuretic peptide analogue¹³, a soluble

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form of human FGFR3, parathyroid hormone, and statins have been shown to improve bone growth in genetically manipulated ACH or TD I mouse models in vivo or ex vivo. The effects of statins have been examined using the in vitro human-relevant model of chondrocytes differentiated from induced pluripotent stem cells from either TD I or ACH patients. The C-type natriuretic peptide analogue has reached clinical trials. It is important to develop a human-relevant in vivo model to provide a robust system for testing potential therapeutic interventions before human clinical trials.

In this report, we developed a human-relevant ACH mouse model by replacing mouse Fgfr3 with human FGFR3 cDNA containing the FGFR3G380R ACH mutation. The clinical phenotypes and histology of bone abnormalities were characterised in the mutant mice. This FGFR3ACH mouse model closely recapitulates human ACH. As such, it offers a valuable tool for assessing potential therapeutic approaches designed to target the over-activation of human FGFR3.

Results
Generation of FGFR3ACH and FGFR3WT mice. To generate FGFR3ACH mice, we used a gene-targeting approach to replace the mouse Fgfr3 with human FGFR3 cDNA carrying the ACH mutation (FGFR3ACH) under the full control of the endogenous mouse Fgfr3 promoter, intron 1, and 5′ and 3′ untranslated regions (Fig. 1A).

Skeletal abnormalities in newborn FGFR3ACH mice. The features of human ACH patients can be readily identified clinically and radiologically at birth. At birth, there were no obvious differences in appearance between FGFR3ACH+ or FGFR3ACH/ACH mice, collectively termed FGFR3ACH, and their WT littermates (Supplemental Fig. 1A). We therefore analysed the bone structure of newborn mice. The newborn FGFR3ACH mice showed proximal limb shortening with relatively normally sized trunks (Fig. 2A). Femur length was reduced by 15% in FGFR3ACH mice and 42% in FGFR3ACH/ACH mice compared with WT mice (Fig. 2C). A closer view of the skull structure revealed the skull was rounded and the calvarial bones were distorted in FGFR3ACH mice, due to a positional shift and compression of the frontal and parietal bones (Fig. 2B). The jugal limitans, i.e., the cranial suture that separates the frontal and nasal bones, was absent in FGFR3ACH mice (Fig. 2B). The metopic sutures, which line the midline between the two nasal bones, were unilaterally fused or partially absent in FGFR3ACH mice (Fig. 2B). Thus, newborn FGFR3ACH mice exhibited premature suture closure and abnormal skull shapes. Furthermore, a shorter intervetebral distance between cervical vertebrae (Supplemental Fig. 1B) and a narrower rib cage (Supplemental Fig. 1D) were observed in FGFR3ACH newborns. These phenotypes are similar in many respects to the skeletal deformities in human ACH newborns, and the bone abnormalities are more evident in FGFR3ACH/ACH mice than in FGFR3ACH+ mice.

Pronounced skeletal abnormalities in FGFR3ACH mice during postnatal development. The dwarfism phenotypes gradually became evident in FGFR3ACH mice. Dominant short stature (Fig. 3A), rounded head (Fig. 3B,D), short snout (Fig. 3B,C), and kyphosis (humpback) (Fig. 3C) phenotypes could be readily observed in FGFR3ACH mice at 10 days to 1 month of age. All FGFR3ACH/ACH mice developed kyphosis phenotypes at around 2 weeks of age, and about 90% of FGFR3ACH+ mice developed kyphosis phenotypes before 1 month of age. In addition, protrusion of the lower incisors was observed in FGFR3ACH+ mice (Fig. 2B). Thus, newborn FGFR3ACH mice exhibited premature suture closure and abnormal skull shapes. Furthermore, a shorter intervetebral distance between cervical vertebrae (Supplemental Fig. 1B) and a narrower rib cage (Supplemental Fig. 1D) were observed in FGFR3ACH+ newborns. These phenotypes are similar in many respects to the skeletal deformities in human ACH newborns, and the bone abnormalities are more evident in FGFR3ACH/ACH mice than in FGFR3ACH+ mice.

Two-dimensional micro-computed tomography (micro-CT) was used to examine the skeletal abnormalities in FGFR3ACH mice. The skeletal bone revealed dwarfism, rounded skulls, and severe curvature of the cervical and upper thoracic vertebrae in FGFR3ACH mice (Fig. 4A–C). FGFR3ACH/ACH mice exhibited more severe phenotypes compared with those of FGFR3ACH+ mice (Fig. 4A–C). Furthermore, these phenotypes became more pronounced in older mice (based on comparison among the phenotypes of 1-, 4-, and 12-month-old mice in Fig. 4A–C). Close observation of the skulls and vertebrae of FGFR3ACH mice revealed shortened snouts and dome-shaped skulls (Fig. 4D–F), and almost completely folded upper thoracic vertebrae in FGFR3ACH/ACH and older FGFR3ACH+ mice (Fig. 4G–I). The severities of these phenotypes were more consistent among FGFR3ACH/ACH mice, as compared with FGFR3ACH+ mice, as shown by the smaller variation in the body lengths of FGFR3ACH/ACH mice compared with that of FGFR3ACH+ mice (Fig. 3F). This is relevant because the variation in the severities of the short snout, rounded-head, and kyphosis phenotypes is represented in the body length.

Patients with ACH present with rhizomelic (short-limbed) dwarfism. This phenotype was reproduced in the FGFR3ACH+ mice, which showed a 22% shortening of femur length along with a 7.1% shortening of body length at 1 month of age, compared with the corresponding measurements in WT mice (Table 1). The results suggested that the limbs were disproportionately shortened relative to body length in FGFR3ACH+ mice. Furthermore, the
femurs were short, curved, and thick with widened diaphyses and flared metaphyses in FGFR3ACH mice (Fig. 4J), which are very similar to phenotypes observed in ACH patients.

Altered chondrocyte proliferation and differentiation in FGFR3ACH mice. Femur length is significantly reduced in FGFR3ACH mice (Fig. 2C and Table 1). To examine defects in the long bones of FGFR3ACH mice more closely, we performed a histological analysis of the distal femur from WT and FGFR3ACH mice at different developmental stages. The epiphyseal structure was similar between the WT and FGFR3ACH mice at birth (Fig. 5A). The secondary ossification centre was readily formed in WT mice at 1 week of age, whereas its formation was markedly delayed in FGFR3ACH mice (Fig. 5A,B), suggesting a delay in chondrocyte terminal differentiation. In endochondral ossification, chondrocytes sequentially transit through resting, proliferating, and hypertrophic

Figure 1. Generation of ACH mice and human FGFR3 WT controls by introducing human FGFR3G380R cDNA or WT FGFR3 into the murine Fgfr3 locus. (A) Strategy for the generation of the targeting vector and depiction of the final chromosomal structure of the murine Fgfr3 locus after the introduction of the human FGFR3G380R cDNA via gene targeting. (B) Mouse embryonic stem cell clones containing the targeted allele were identified by Southern blot analysis. (C) The neomycin resistance cassette in the identified stem cells was removed by Flp/FRT excision and analysed by PCR amplification and EcoRI digestion. A 528 bp PCR product was present in the stem cells without the neomycin resistance cassette, and the 328 bp and 254 bp fragments produced by EcoRI digestion of the PCR product could be detected. (D) PCR amplification analysis of genomic DNA isolated from WT and FGFR3ACH mice. A 1067 bp PCR product was amplified from the murine Fgfr3 locus. A 506 bp PCR product was amplified from the human FGFR3G380R targeted allele. (E) The mRNA expression of targeted human FGFR3G380R and endogenous mouse Fgfr3 in the heterozygous FGFR3G380R, homozygous FGFR3G380R, and WT mice was determined by RT-PCR using sequence-specific primers. ACH/+; the heterozygous FGFR3/ACH mice; ACH/ACH; the homozygous FGFR3ACH/ACH mice; +/+; wild type littermates.
stages. The FGFR3ACH mice showed good development of each stage. However, the growth plates were significantly shorter in FGFR3ACH mice with a shorter proliferative zone at 2, 4, and 8 weeks of age (Fig. 5B,C). This was caused by a reduction in the number of proliferative chondrocytes, indicating that chondrocyte proliferation was compromised in FGFR3ACH mice. Despite the shorter proliferative zone, the arrangement of chondrocyte columns in the growth plate remained normal in FGFR3ACH mice before 2 weeks of age. The disturbed arrangement of chondrocyte columns in FGFR3ACH mice can be appreciated at 4 and 8 weeks of age, and their arrangement was disrupted by an increased amount of space between the columns (Fig. 5B). We further showed that FGFR3ACH mice had higher FGFR3 phosphorylation in chondrocytes of growth plates (Fig. 5D) and the primary chondrocytes had lower proliferation rates compared with those from WT mice (Fig. 5E), suggesting that FGFR3 activation inhibited chondrocyte proliferation in FGFR3ACH mice.

Altered bone formation in FGFR3ACH mice. Low bone density has been reported in adult ACH patients39, which may have clinical relevance and lead to subsequent bone damage. The development of the long bones is coordinated between chondrogenesis and osteogenesis. Reduced growth of the longitudinal trabecular bone was observed in the distal femoral metaphysis of FGFR3ACH mice at several stages of postnatal development (Fig. 5A, stained in blue). Furthermore, the expression of osteocalcin, which is associated with the early stages of matrix ossification, was increased in the chondrocytes of the hypertrophic zone of the distal femur of FGFR3ACH mice at 2 weeks of age (Fig. 5F). A reduced hypertrophic zone was observed in FGFR3ACH mice at 8 weeks of age (Fig. 5B,C). These results indicate that the bone-forming process was disturbed in FGFR3ACH mice. To determine the structure

Figure 2. Skeletal defects and changes in bone architecture in newborn FGFR3ACH mice. (A) Lateral view of skeletal preparations of FGFR3ACH-/+ (ACH/+), FGFR3ACH/ACH (ACH/ACH), and WT mice. Cartilage was stained with Alcian blue and bone was stained with alizarin red. Scale bar: 30 mm. (B) Dorsal view for comparison of the skulls. The white arrowheads indicate the jugum limitans, and the white arrow indicates the metopic suture. N, nasal bones; F, frontal bones; Pa, parietal bones. (C) Femur length was significantly decreased in FGFR3ACH/ACH and FGFR3ACH+ mice. WT, n = 3; ACH/+ , n = 6; ACH/ACH, n = 3.
of trabecular bone, we performed a micro-CT analysis. Three-dimensional images of the distal femoral metaphysis showed a lower bone volume with thinner and fewer trabecular bones and larger intertrabecular spaces in newborn and 1-year-old FGFR3ACH mice compared to WT mice (Fig. 5G). A histomorphometric analysis of bone formation showed that the trabecular bone volume (BV/TV), trabecular thickness (Tb.Th), and trabecular number (Tb.N) were decreased, along with an increased trabecular separation (Tb.Sp) and structure model index (SMI) in the distal femoral metaphysis of FGFR3ACH mice compared with WT mice at birth and at 1 year of age (Table 2). Furthermore, we observed fewer osteoblasts and osteoclasts in the femurs of FGFR3ACH mice at 1 year of age (Fig. 5H), suggesting that the bone turnover rate might be altered in FGFR3ACH mice.

Discussion
In this study, we describe an ACH mouse model (FGFR3ACH), which expresses human FGFR3G380R, the most common mutation in human ACH patients. Our mouse model recapitulates the main human ACH phenotypes and offers a valuable tool for studying pathological conditions and testing potential therapeutic approaches designed to target the over-activation of human FGFR3 and its downstream signaling pathway in vivo. In addition to the postnatal phenotypes observed in previous ACH mouse models8,10,11, we showed that FGFR3ACH mice shared most of the key clinical phenotypes observed in human ACH patients at birth, including rhizomelic dwarfism, rounded skull, and midface hypoplasia. Furthermore, we described craniosynostosis and low bone density in newborn FGFR3ACH mice. These phenotypes became more pronounced during postnatal skeletal development and were correlated with the dose of FGFR3ACH. Kyphosis, which is common in human ACH during postnatal
development, develops postnatally in FGFR3ACH mice. Although previously generated ACH mouse models share some ACH phenotypes, some phenotypes have not been fully observed or described in these ACH mouse models. Here, we described the full range of ACH abnormalities in this mouse model. A comparison of skeletal phenotypes between human achondroplasia and ACH mouse models is summarised in Table 3.

Previously, two ACH mouse models were generated by either targeting8 or transgenically expressing10 murine \(\text{Fgfr}^{\text{G374R}}\) (an ortholog of the human mutation \(\text{FGFR}^{\text{G380R}}\), \(\text{FgfrACH}\)). Another ACH mouse model transgenically expresses human \(\text{FGFR}^{\text{G380R}}\) under the control of the mouse \(\text{Fgfr}^{\text{A}}\) promoter region11. All these ACH mouse models exhibit some human ACH phenotypes during postnatal development. However, rhizomelic (short-limbed) dwarfism was not obvious, and dwarfism phenotypes were not described at birth in both \(\text{Fgfr}^{\text{A}}\) models. In addition, the transgenic model expressing \(\text{FgfrACH}\) under the control of the collagen II promoter presented some specific defects that are not usually observed in human ACH patients due to misexpression of \(\text{Fgfr}^{\text{A}}\)10. The transgenic human \(\text{FGFR}^{\text{G380R}}\) mice presented the rhizomelic dwarf phenotype at birth11. However, thoracic kyphosis was not described for this model11. Moreover, homozygous \(\text{FGFR}^{\text{G380R}}\) transgenic mice expressed more severe phenotypes than the two \(\text{Fgfr}^{\text{A}}\) mouse models or our \(\text{FGFR}^{\text{G380R}}\) model, and died shortly after birth. The severe phenotypes in homozygous \(\text{FGFR}^{\text{G380R}}\) transgenic mice might be because the mice express both endogenous \(\text{Fgfr}^{\text{A}}\) and human \(\text{FGFR}^{\text{G380R}}\). Most homozygous ACH patients are stillborn or die during the neonatal period. Although our \(\text{FGFR}^{\text{G380R}}\) mice exhibited a higher mortality rate than the WT mice at birth and before 4 weeks of age, 60% of \(\text{FGFR}^{\text{G380R}}\) mice survived after 1 month in our study. The homozygous knock-in or transgenic \(\text{Fgfr}^{\text{A}}\) mice survived after birth, but their neonatal survival rate was not reported. The human and mouse \(\text{FGFR}^{\text{G380R}}\) genes share 92.7% identity and 94.9% similarity. The various phenotypes and different levels of severity observed in different ACH mouse models might be caused by the different promoter chosen and various copy numbers of activated \(\text{FGFR}^{\text{A}}\) in the transgenic models, the different knock-in approaches used in the KI models, and the different genetic backgrounds of the mice assayed. Having access to various ACH mouse models provides the opportunity for choosing particular ACH phenotypes of interest for further studies. Nevertheless,

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**Figure 4.** Radiographs of \(\text{FGFR}^{\text{G380R}}\) (ACH/+) (ACH/ACH), and WT mice (WT). \(\text{FGFR}^{\text{G380R}}\) mice showed profound rhizomelic dwarfism, a rounded skull, and kyphosis (arrow) in the cervico-thoracic spine. The severity of these phenotypes correlated with the dose of \(\text{FGFR}^{\text{G380R}}\) and postnatal development. Lateral view of whole skeleton at the age of 1 month (A), 4 months (B), and 1 year (C). The enlarged view shows rounded skull at the age of 1 month (D), 4 months (E), and 1 year (F). The enlarged view shows kyphosis (arrow) in the cervico-thoracic spine in \(\text{FGFR}^{\text{G380R}}\) and \(\text{FGFR}^{\text{A}}\) aged 1 month (G), 4 months (H), and 1 year (I). The enlarged view shows short, curved, and thick bones with widened diaphyses and flared metaphyses in the femurs of \(\text{FGFR}^{\text{G380R}}\) mice at several developmental stages (J).

| Body length (mm, n = 14) | Femur length (mm, n = 9) |
|--------------------------|--------------------------|
| WT 72.69 ± 4.905         | 12.33 ± 0.5             |
| ACH/+ 67.52 ± 4.332***   | 9.61 ± 0.8529***        |
| Percentage decreased 7.11% 22.06% |

**Table 1.** Body length and femur length of 1-month-old mice. WT: wild-type littermates; ACH/+; \(\text{FGFR}^{\text{G380R}}\) mice. Each value is expressed as the mean ± SD (n values shown in column headers). ***p < 0.001.
Figure 5. Histological, immunohistochemical, and micro-computed tomography analysis of distal femoral growth plates and chondrocyte proliferation in FGFR3ACH/ACH mice. The insets show magnified views. (A) Masson's trichrome staining of distal femoral growth plates at the ages of 1 day, 1 week, 2 weeks, 4 weeks, and 8 weeks. Square brackets: the region of trabecular bone. Scale bar, 500 μm. The insets show magnified views in (B). Femoral growth plates show the typical zonal structure of proliferating chondrocytes (PZ) and hypertrophic chondrocytes (HZ) in FGFR3ACH/ACH mice. Delayed formation of secondary ossification centres was apparent in FGFR3ACH/ACH mice at 1 week of age. Growth plates were shorter with more compacted proliferative cells in FGFR3ACH/ACH mice. An increased amount of space (yellow arrows) between the chondrocyte columns can be appreciated in FGFR3ACH/ACH mice at 4 and 8 weeks of age. (C) Quantitative measurements of the heights of the PZ and HZ of the distal femoral growth plates of FGFR3ACH (ACH) mice and WT mice at 2, 4, and 8 weeks of age. Data represent mean values ± SD. *p < 0.05; ***p < 0.001. (D) Immunohistochemistry staining of phospho-FGFR3 in growth plate chondrocytes showed increased abundance of phospho-FGFR3 in 16-day-old FGFR3ACH/ACH mice. (E) An increased growth rate of chondrocytes from FGFR3ACH/ACH analysed using an iCELLigence™ real-time cell analysis system. The data are presented as changes in the cell index over time. (F) Immunohistochemical analysis of osteocalcin showed increased abundance of osteocalcin in the hypertrophic zone in 2-week-old FGFR3ACH/ACH mice. (G) Vertical (left) and transverse (right) views of trabecular bone in the distal femoral metaphysis of WT and FGFR3ACH/ACH mice at 1 day and 1 year of age assessed using micro-computed tomography analysis. (H) Masson’s trichrome staining of the distal femurs of WT and FGFR3ACH/ACH mice at 1 year of age. The yellow arrows indicate the osteoblasts and osteoclasts. The insets show examples of the magnified views of osteoblasts (OB) and osteoclasts (OC).

|     | BV/TV (%) | Tb.Th (mm) | Tb.Sp (mm) | Tb.N (1/mm) | SMI     |
|-----|-----------|------------|------------|-------------|---------|
| newborn |           |            |            |             |         |
| WT (n = 3) | 43.16 ± 1.85 | 0.076 ± 0.007 | 0.086 ± 0.006 | 5.718 ± 0.35 | 1.487 ± 0.24 |
| ACH/ACH (n = 3) | 15.76 ± 3.93*** | 0.058 ± 0.011 | 0.17 ± 0.045 | 2.74 ± 0.71*** | 2.436 ± 0.26** |
| 1 year old |           |            |            |             |         |
| WT (n = 4) | 5.22 ± 0.51 | 0.079 ± 0.010 | 0.383 ± 0.047 | 0.661 ± 0.087 | 2.569 ± 0.20 |
| ACH/ACH (n = 3) | 2.74 ± 1.08* | 0.069 ± 0.003 | 0.378 ± 0.075 | 0.402 ± 0.169 | 2.87 ± 0.37 |

Table 2. Structural parameters of distal femur trabecular bone in newborn mice. WT: wild-type littermates; ACH/ACH: FGFR3ACH/ACH mice. BV/TV: trabecular bone volume/tissue volume; Tb.Th: trabecular thickness; Tb.Sp: trabecular separation; Tb.N: trabecular number; SMI: structure model index. Each value is expressed as the mean ± SD (all groups n = 3). *p < 0.05; ***p < 0.01; **p < 0.001.
the FGFR3ACH mice generated in this study exhibited obvious and homogenous clinical phenotypes of ACH. Furthermore, these mice expressed human FGFR3G380R alone without interference by endogenous Fgfr3, making them a useful model for further evaluating potential treatments targeting the over-activation of human FGFR3 and its downstream signaling.

The biological basis of dwarfism phenotypes in ACH patients involves a specific defect in endochondral ossification and longitudinal bone growth. The molecular consequences of over-activation of FGFR3 on chondrocyte proliferation and differentiation in the growth plate have been well established. Here, we demonstrated that over-activated human FGFR3G380R has inhibitory effects on chondrocyte proliferation and maturation in an in vivo mouse model. FGFR3 mutations might also affect membranous ossification. Gain-of-function mutations of FGFR3 (P250R and A391E) have been shown to cause human Muenke syndrome and Crouzon syndrome with acanthosis nigricans. Affected patients undergo premature fusion of the cranial sutures. Furthermore, TD patients also frequently exhibit severe craniosynostosis phenotypes. ACH patients present craniofacial phenotypes suggesting the possibility of craniosynostosis defects. Recent report showed that ACH is associated with craniosynostosis and suggested that craniosynostosis may be under-reported. An increased understanding of premature fusion of the cranial sutures and craniofacial phenotypes in ACH at early stages will facilitate improved treatment of these defects. For the first time, we have demonstrated here that the cranial sutures undergo premature fusion in newborn ACH mice, which offers the opportunity to better understand the development of cranial anomalies in ACH, and further study membranous ossification.

During the postnatal and adult stages, bone undergoes continuous remodelling through the coordinated processes of bone formation and bone resorption. Low bone density has been reported in adult ACH, which may have clinical relevance and lead to subsequent bone damage, such as increased fragility and risk of fracture. Enhanced osteoblast differentiation has been observed in long bone growth plates of Fgfr3ACH mice at the age of 15 days, suggesting advanced ossification at an early stage. Recent studies in Fgfr3ACH mice revealed enhanced osteogenic differentiation in cultured bone marrow stromal cells, and this was associated with decreased bone mass at 2 months of age. Here, we demonstrated that the expression of osteocalcin in the chondrocytes of the hypertrophic zone of the distal femur in FGFR3ACH mice at 2 weeks of age, suggesting enhanced osteoblast differentiation in FGFR3ACH mice. Recently, a study revealed that bone density was reduced in the majority of ACH and HCH patients in the age range of 10–33 years, which is indicative of osteopenia. Here, we provide direct evidence showing a low bone density in newborn and adult FGFR3ACH mice. Furthermore, we observed fewer osteoclasts and osteoblasts in the femur of FGFR3ACH mice at 1 year of age. Both endochondral ossification and bone remodelling regulate bone mass in adults, suggesting that altered bone remodelling might also contribute to the lower bone mass of adult FGFR3ACH mice. Potential changes in bone structure should be further evaluated in neonatal ACH patients to determine whether an adequate diet and exercise may help to prevent any such osteopenia at early developmental stages. Our mouse model offers a good opportunity for testing interventions for early onset osteopenia in ACH.

The mouse is a convenient animal model for the study of human genes and diseases, which has led to the development of treatments for many serious diseases and conditions. However, mice are not always reliable as preclinical models for human disease. Many drugs have shown promising results in preclinical trials in mice but later failed in human clinical trials. As such, there is a need to develop reliable preclinical mouse models of human diseases for clinical research and drug development. Mice expressing a mutated version of a human gene known to be associated with a specific human disease and faithfully mimicking the disease phenotypes can be useful for studying disease pathology, conducting preclinical research, and testing compound efficacy in vivo. We have generated an ACH mouse model that faithfully and comprehensively recapitulates the disease phenotypes, enabling the evaluation of potential treatments targeting the over-activation of human FGFR3 and its downstream signaling.

Methods
(Full experimental details are provided in the Supplemental Data).

### Table 3. Similarity of skeletal features found in human achondroplasia and observed in achondroplasia mouse models. Tg mFgfr3ACH, transgenic mice expressing mouse Fgfr3G380R using the type II collagen promoter and enhancer sequences. KI mFGFR3ACH, gene targeting mouse Fgfr3G380R. Tg hFGFR3ACH, transgenic mice expressing human FGFR3G380R using the mouse Fgfr3 promoter. KI hFGFR3ACH, gene targeting human FGFR3G380R as described in this report. ND, not described; NS, not significant. The time point when the specific phenotype was first observed in each ACH mouse model.

| Skeletal feature            | Tg mFgfr3ACH | KI mFgfr3ACH | Tg hFGFR3ACH | KI hFGFR3ACH |
|----------------------------|--------------|--------------|--------------|--------------|
| Rhizomelic dwarfism at birth| NS           | NS           | At birth     | At birth     |
| Large head with frontal bossing, mid-face hypoplasia at birth| 21 days      | 10 days      | At birth     | At birth     |
| Craniosynostosis at birth  | ND           | ND           | ND           | At birth     |
| Low bone density in adolescent and adult | ND           | ND           | ND           | At birth     |
| Homozygous ACH patients are stillborn or die during the neonatal period | ND           | ND           | Die soon after birth | A higher mortality rate at birth |
| Thoracic kyphosis by 4 months | 1 month      | 5 weeks      | ND           | 1 month      |
| Narrow growth plate        | 10 days      | 1 month      | At birth     | 1 month      |
Mice. Mice were housed in a temperature- and humidity-controlled room with a 12-h light/12-h dark cycle under specific pathogen-free conditions. All animal protocols were approved by the Institutional Animal Care and Utilization Committees, Academia Sinica, Taiwan (Protocol #14-12-795). The investigation conformed to the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health.

Gene targeting and generation of chimeric mice. To generate a targeting vector for the expression of human FGFR3<sup>G380R</sup>, we adopted a highly efficient recombination-based method, as previously described<sup>27</sup>. The human mutant FGFR3 cDNA (NM_000142) encoding the FGFR3<sup>G380R</sup> protein was used to replace the WT allele of Fgfr3 in 129 Sv mice. The stem cells carrying the targeting vector without the neomycin resistance cassette were injected into C57BL/6J blastocysts<sup>28</sup>. The resulting chimeric mice were crossed with 129 Sv females to enable germline transmission. Heterozygotes were used to continue the strain and to provide experimental pairs.

Quantitative reverse transcriptase polymerase chain reaction (qRT-PCR). Total RNA from the left hind-limb of neonatal mice was isolated, converted to cDNA, and subjected to qRT-PCR. Expression data were normalized to Gapdh mRNA levels. Gene-specific primer sequences are listed below.

- Mouse Fgfr3: forward sequences, 5′-TGCGGTGCTTCCACAGA-3′; reverse sequences, 5′-ACTTGGACCTCTCCGTG-3′.
- Human FGFR3: forward sequences, 5′-GCTTGAACACAGGTCGTG-3′; reverse sequences, 5′-CAGTCCCTCCA TTCTCTG-3′.
- Gapdh: forward sequences, 5′-CCAGAACATCATCCCTGCAT-3′; reverse sequences, 5′-GTTCAGCTCTGGGATGACCTT-3′.

Bone length measurement. Femurs of mice were dissected and the flesh was removed. The lengths of the femurs were measured using a millimetre-scale calliper ruler.

Micro-CT. Two-dimensional imaging of whole mice, skulls, shoulder joints, and hind limbs was performed on euthanised mice using a Skyscan 1076 system (Bruker, Brussels, Belgium). The distal femur metaphyses of fixed trabecular bones were analysed by three-dimensional micro-CT using a Skyscan 1076 3D system in the Taiwan Mouse Clinic, following their standard protocol (as described in a previous report)<sup>29</sup>. The following scanning parameters were chosen: image pixel size: 9 μm, X-ray voltage: 50 kV, X-ray current: 140 μA, filter: Al 0.5 mm, exposure: 3300 ms, rotation step: 0.8°, frame averaging: 2, tomographic rotation: 180°. Cross-sections were reconstructed using NRecon software (Bruker). The parameters were as follows: smoothing: 0, ring artefacts reduction: 6, beam-hardening correction: 20%, change dynamic image range: 0.015–0.07.

Histology, histochemistry, and immunohistochemistry. Sections of fixed bone tissues were prepared and examined with Masson’s trichrome stain. For immunohistochemistry, the sections were incubated with phospho-FGFR3 antibody (Cell Signaling, Danvers, MA, USA) or anti-osteocalcin antibody (Millipore, Billerica, MA, USA) and then visualised using 3,3′-diaminobenzidine.

Primary chondrocyte culture and cell proliferation assay. The primary chondrocytes were isolated and cultured as previously described<sup>30</sup> with several modifications as described in the Supplementary Data. The proliferation of primary chondrocytes (we did not use subcultured cells) was assessed using an iCELLigence™ real-time cell analyser (Acea Biosciences, San Diego, CA, USA, distributed by Roche Diagnostics, Basel, Switzerland). The changes in adhesion and spreading of the cells were continuously recorded for 15 days using the iCELLigence™ system. Data were expressed as a graph of cell index values during the exponential phase.

Skeletal preparation. Skinned and eviscerated newborn mice were fixed and stained with Alcian blue 8GX (Sigma–Aldrich, St. Louis, MO, USA) and alizarin red (Sigma–Aldrich).

Statistical analysis. A two-tailed Student’s t-test was used to test for differences between groups. A p value less than 0.05 was considered to be statistically significant (∗p < 0.05, ∗∗p < 0.01, ∗∗∗p < 0.001). Chi-square goodness-of-fit tests (2 degrees of freedom) were used to test for departures from Mendelian expectations for the genotypes of FGFR3<sup>G380R</sup> mice generated from heterozygous breeding pairs.

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Author Contributions

Study design: Y.C.L. and Y.T.C. Study conduct: Y.C.L., I.W.S. and S.D.C. Data collection and analysis: Y.C.L., I.W.S., Y.J.P. and S.D.C. Data interpretation: Y.C.L., I.W.S., S.D.C. and Y.T.C. Manuscript preparation: Y.C.L. Revision of manuscript content: Y.C.L. Approval of final version of the manuscript: Y.C.L., I.W.S., S.D.C. and Y.T.C. Y.C.L. takes responsibility for the integrity of the data analysis.

Additional Information

Supplementary information accompanies this paper at http://www.nature.com/srep

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