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Skeletal Biology and Disease Modeling in Zebrafish

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ABSTRACT
Zebrafish are teleosts (bony fish) that share with mammals a common ancestor belonging to the phylum Osteichthyes, from which their endoskeletal systems have been inherited. Indeed, teleosts and mammals have numerous genetically conserved features in terms of skeletal elements, ossification mechanisms, and bone matrix components in common. Yet differences related to bone morphology and function need to be considered when investigating zebrafish in skeletal research. In this review, we focus on zebrafish skeletal architecture with emphasis on the morphology of the vertebral column and associated anatomical structures. We provide an overview of the different ossification types and osseous cells in zebrafish and describe bone matrix composition at the microscopic tissue level with a focus on assessing mineralization. Processes of bone formation also strongly depend on loading in zebrafish, as we elaborate here. Furthermore, we illustrate the high regenerative capacity of zebrafish bones and present some of the technological advantages of using zebrafish as a model. We highlight zebrafish axial and fin skeleton patterning mechanisms, metabolic bone disease such as after immunosuppressive glucocorticoid treatment, as well as osteogenesis imperfecta (OI) and osteopetrosis research in zebrafish. We conclude with a view of why larval zebrafish xenografts are a powerful tool to study bone metastasis. © 2021 American Society for Bone and Mineral Research (ASBMR).

KEY WORDS: ZEBRAFISH; SKELETON; REGENERATION; METASTASIS; GLUCOCORTICOID

Introduction

Zebrafish have become an important model organism to study the development and disease of the skeleton in basic and preclinical research. The potential of these teleost fish lies in their small size, ease of care, genetic amenability, and high regenerative capacity. Moreover, thanks to the transparency of embryonic and larval stages, osteogenesis and osteoblast activity can be monitored in much detail, using available transgenic and mutant lines affecting specific cells or tissues.1,2 This is combined with long-term in vivo imaging feasibility of embryonic, larval, and also adult individuals,3,4 which distinguishes zebrafish from other vertebrate models such as rodents, in which intravital imaging can be challenging (Table 1). Importantly, the zebrafish genome contains orthologues of about 82% of human disease-related genes,5,6 including those affecting the skeleton. Both tissue-specific overexpression via site-specific recombinases (Cre)7,8 and gene-specific knockout via clustered regularly interspaced short palindromic repeats (CRISPR-Cas9)9,10 are available in zebrafish, along with antisense oligonucleotide gene knockdown approaches11,12 and ideal conditions to carry out forward and reverse genetic screens.13,14,15 Furthermore, single-nucleotide genome editing can be performed by CRISPR-Cas9–mediated knock-ins.16,17 Finally, drugs can be administered to zebrafish in various ways including dissolving chemicals directly in zebrafish water/media,18 the preferred method in drug screening (Table 1).

The above descriptions illustrate the potential of zebrafish to study skeletal biology and disease. Excellent reviews have been published on diverse aspects of zebrafish as a skeletal research model.19-22 Here, we aim to introduce zebrafish to the wider bone research community, by presenting essential information on their skeletal architecture and patterning, cell types, and matrix mineralization (which is loading dependent), along with introducing a variety of zebrafish assays to study bone regeneration. Furthermore, we highlight the utility of larval xenografts to demonstrate the power of zebrafish in bone metastasis research.

Architecture of the Zebrafish Skeleton

The skeleton of vertebrates is generally divided into the exoskeleton and endoskeleton.23,24 The prominent parts of the zebrafish skeleton are as follows: (i) the craniofacial skeleton and
including parietal bones, jaw bones, and opercles (bones covering the gills); and (ii) the axial skeleton comprising the vertebral column, ribs, intermuscular bones, as well as unpaired dorsal, anal, and caudal fins (44,45) (Fig. 1A). Zebrafish are considered sexually mature at around 90 days, corresponding to a standard length (SL; measured from snout to the last caudal vertebra in adult individuals) of 1.5 to 2.0 cm. Zebrafish undergo continuous growth associated with an increase in skeletal volume, resulting in a body length of 3 to 4 cm, and typically live for around 3 years (though they can reach 5 years). (46)

The adult zebrafish skull is composed of 74 craniofacial bones, considerably more than the 22 bones of the mammalian skull. (47) Nevertheless, a number of bony structures in zebrafish have clear homologs in mammals, including the anterior part of the neurocranium, which resembles the mammalian palate, (48) and the cranial vault, which is conserved between zebrafish and mammals. (49) As with mammals, the zebrafish

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**Table 1. Comparison of Skeletal Features and Experimental Tractability in Humans, Rodents, and Zebrafish**

| Skeletal feature                  | Human          | Rodent         | Zebrafish     | References |
|----------------------------------|----------------|----------------|---------------|------------|
| Bone types                        | Dermal         | Dermal         | Dermal        | (5)        |
|                                   | Compact        | Compact        | Compact       |            |
|                                   | Spongy         | Spongy         | Tubular (spongy) |          |
| Skeletal cell types               | Chondrocytes   | Chondrocytes   | Chondrocytes  | (5,6)      |
|                                   | Osteoblasts    | Osteoblasts    | Osteoblasts   |            |
|                                   | Osteoclasts    | Osteoclasts (multinucleate) | Osteoclasts (multinucleate and mononucleate) |
|                                   | Osteocytes     | Osteocytes     | Osteocytes    |            |
| Ossification types                | Endochondral   | Endochondral   | Endochondral  | (5,7)      |
|                                   | Intramembranous| Intramembranous| Intramembranous|            |
|                                   | Perichondral   | Perichondral   | Perichondral  |            |
| Development                       | In utero       | In utero       | Extrauterine  |            |
| Average brood size                | n/a            | 6–8            | 100–150       | (8)        |
| Mineralization begins             | 4–5 weeks      | about 2 weeks  | 3–4 dpf       | (9,10)     |
| Skeletal maturity reached         | Up to 30 years | 4–5 months     | 2–4 months    | (11)       |
| Direction of loading              | Axial          | Orthogonal     | Axial         | (12)       |
| Repair after fracture             | Yes            | Yes            | Yes           | (13,14)    |
| Regeneration after amputation     | Limited (digit tip) | Limited (digit tip) | Yes         | (15–17)    |
| Gene conservation (versus humans) | 100%           | ~85%           | ~75%          | (18)       |
| Bone marrow                       | Yes            | Yes            | No            | (19)       |
| Dynamic histomorphometry          | Tetracycline   | Alizarin red stain | Alizarin red stain | (20–22) |
|                                   |                | Calcein green  | Calcein green |            |
| Visualization of cell dynamics of bone | No          | Limited (intravital imaging) | Transparency in mutants, larval stages and fin regenerates, adult scale regeneration | (4,15,23,24) |
| BMD assessment                    | CT (fixed/live); high-resolution peripheral quantitative CT (HRpQCT, fixed/live); ultrasound (live); DXA (live) | µCT (fixed); DXA (live) | µCT (fixed) | (25–27) |
| Drug screening                    | In vitro       | In vitro       | In vivo       | (11)       |
| Mosaic CRISPR mutagenesis time    | n/a            | 3 months       | 5 days to 3 months | (28,29) |
| Stable CRISPR mutagenesis time    | n/a            | 9 months       | 6–9 months    | (30)       |
| Ex vivo/in vitro study of bone    | In vitro differentiation of stem cells | In vitro differentiation of stem cells | Ex vivo scale culture | (23,31) |
|                                   | Ex vivo culture of 1° cells | Ex vivo culture of 1° cells | Ex vivo culture of 1° cells |            |

1° = primary; n/a = not applicable; dpf = day(s) post fertilization.
skull features skeletal joints, including fibrous joints (e.g., skull sutures), and articular joints in the jaw.\(^{(50)}\) Regarding spinal morphology, zebrafish share a similar number of vertebrae (30 to 32 in zebrafish versus 33 in humans) and a physiological curvature with humans: kyphosis in the abdominal region where the ribs protect the viscera, and lordosis in the caudal region.\(^{(51)}\) In craniocaudal order, the zebrafish vertebral column is composed of a Weberian apparatus consisting of four vertebrae connecting the swim bladder to the ear (important for the transmission of sound), 10 abdominal vertebrae (also known as precaudal vertebrae or trunk vertebrae) that are articulated with rod-shaped rib segments, transitioning into 14 caudal vertebrae, and three caudal fin vertebrae.\(^{(52)}\) (Fig. 1A). The spinal cord passes through the neural arches that extend dorsally from each vertebra, similarly to the mammalian spinal canal. Caudal vertebrae also have hemal arches that extend ventrally and enclose the caudal artery and vein.

The three-dimensional (3D) morphology of individual zebrafish vertebral body centra is characterized by its hourglass shape (Fig. 1B). In contrast to mammalian vertebrae that contain trabecular bone and carry bone marrow (BM), zebrafish vertebral bodies do not accommodate red BM, because adult hematopoiesis takes place in the kidney (Table 1). Instead, they are filled with vacuolated notochord cells and become surrounded by adipose tissue.\(^{(53,54)}\) Micrometer-thin trabecular struts are found surrounding the narrow center of individual vertebrae.\(^{(55)}\)

Although assessment of vertebral trabecular bone, a readout for bone fragility in mammals, is limited in zebrafish, morphology of the vertebral centra is commonly used as an indicator for vertebral bone quality. 3D morphometric parameters, e.g., vertebral body length (VBL), bone volume (BV), bone volume per tissue volume (BV/TV), vertebral cross-sectional thickness (V.Th), and eccentricity (i.e., roundness) are extracted from micro-computed tomography (μCT) scans of the zebrafish spine (Fig. 1B), allowing the quantification of changes, e.g., due to altered musculoskeletal activity,\(^{(55,56)}\) aging,\(^{(57)}\) and disease.\(^{(58)}\) In addition, thickness and volume of hemal and neural arches are used to assess vertebral morphology in zebrafish mutants.\(^{(25)}\) One advantage of the small-sized zebrafish is the possibility of analyzing the complete skeleton at high resolution (e.g., a whole-body μCT scan at a pixel size of 1 μm\(^2\)), more rapidly than it is possible in larger rodent species. This provides the possibility to both evaluate the bulk 3D morphology of the entire organism and to simultaneously characterize tissue morphology at high resolution,\(^{(59)}\) which has been done in deformed osteoarthritic vertebral bodies.\(^{(60)}\) Furthermore, longitudinal histological sections or whole-mount

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**Fig 1.** The zebrafish skeleton. (A) μCT image of the craniofacial and axial skeleton, including the vertebral column composed of the Weberian apparatus, abdominal (also referred to as precaudal or thoracic), caudal, and caudal fin vertebrae. Hox gene expression patterns are indicated. Similar to the mammalian skeleton, ribs in zebrafish are articulated to the abdominal vertebrae and protect the inner organs. (B) Close-up view of abdominal and caudal vertebrae. Sagittal views of two adjacent double-cone shaped vertebrae (left) display the neural arches extending dorsally and encompassing the neural canal. Extending ventrally from the abdominal vertebrae ribs are articulated, while the caudal vertebrae extend to hemal arches which encompass the caudal artery and vein. In frontal view (right), the unmineralized vertebral center is revealed which contains notochord and vacuolated soft tissue (not displayed). The ring-shaped vertebral endplate regions are connected by an IVL (not shown) and correspond to the vertebral growth zone in zebrafish. Parameters, including the VBL, V.Th, and BV/TV, provide valuable measures to quantify the vertebral morphology and structure. BV/TV = bone volumetric fraction; IVL = intervertebral ligament; VBL = vertebral body length; V.Th = vertebral thickness.
**Fig 2.** Assessment of matrix composition in zebrafish bone. (A) Raman spectroscopy allows mapping the mineral-related and protein-related properties of regenerating caudal fin bone (here, mineral-to-matrix ratio at 7 dpa) proximal and distal to the amputation plane (arrow). In the Raman spectra of regenerating parts (gray curve), protein-related peaks including amide I, amide III, and hydroxyproline, are more pronounced in respect to the mineral-related phosphate peak, reflecting a lower mineral-to-matrix ratio compared to native bone tissue (black curve). (B) qBEI can be used to assess the mineral density distribution in the zebrafish skeleton (here: sagittal plane of the vertebral column and close-up of the endplate region). A histogram generated from calibrated pixel intensities within a region of interest allows extracting the Ca-mean, Ca-peak, and Ca-width, as well as areas with Ca-low and Ca-high. Ca-high = high degree of mineralization; Ca-low = low degree of mineralization; Ca-mean = mean Ca content; Ca-peak = peak Ca content; Ca-width = heterogeneity of the Ca distribution; dpa = days post amputation; qBEI = quantitative backscattered electron microscopy.

stains, eg, Alizarin red staining, enable the display of the complete skeleton at microscopic resolution.\(^{20,45,61}\)

Similar to mammals, vertebral bones in zebrafish are interconnected by soft tissue, facilitating movement and increasing the range of locomotion. In mammals, the intervertebral disk (IVD) is composed of fibrocartilaginous cartilage (annulus fibrosus) and nucleus pulposus. In contrast, the intervertebral soft tissue in zebrafish is characterized by a ring-shaped ligament (intervertebral ligament [IVL]) connecting the outermost circular edges of two adjacent vertebrae. Although the physiological role of the IVD in humans is damping compressive forces from gravitational loading, the zebrafish spine is loaded axially due to compressive forces from swimming through viscous water and direct muscle forces transmitted by tendons attached to the vertebrae.\(^{12}\)

Compared to tetrapods (eg, mice, dogs), where gravitational forces apply orthogonally to the spine, zebrafish can be considered advantageous in terms of loading regime. Thus, they provide a valuable tool to study spine and intervertebral tissue degeneration and effects of altered locomotion patterns on the bone-tendon-muscle unit, which can be assessed in small-sized zebrafish using histological and advanced X-ray imaging approaches.\(^{62}\) (Fig. 2).

**Patterning of the Zebrafish Axial and Fin Skeleton**

As in other vertebrates, axial patterning in the zebrafish skeleton manifests in the vertebral column which is regionalized into different types of vertebrae along the anteroposterior axis.\(^{32}\) Studies in mice and chicken revealed the morphological diversity and axial position of the different types of vertebrae that are sensitive to positional cues encoded by regional Hox gene expression.\(^{63,64}\) Although Hox mutations in humans lead to early developmental lethality, HOX-related axial skeletal defects have been described, in addition to limb and craniofacial defects, arthritis, and diverse types of cancer.\(^{65,66}\) Regionalization of the zebrafish axial skeleton is under the same mechanistic control, and relies on spatial (and temporal) collinearity; ie, the correspondence between the physical sequence of the genes on the chromosome and their anteroposterior boundaries (and timing) of expression\(^{63,67}\) (Fig. 1A). Although there are 39 Hox genes distributed in four clusters in the mammalian genome, zebrafish possess 48 genes in seven clusters. These differences have come about as a result of genome duplication and gene loss during vertebrate evolution.\(^{68,69}\) Accordingly, Hox expression domains and axial structure regionalization are only partially equivalent between zebrafish and tetrapods (true, eg, for Hoxc6 and Hoxd12 but not for Hox9).\(^{64,67,70}\) Nevertheless, studying homeotic transformations in embryonic and larval zebrafish may be a useful tool to reveal the variety of Hox protein regulatory functions.

In fish, the axial skeleton includes the vertebral column and associated median fins (dorsal, anal, and caudal), whereas the paired pectoral and pelvic fins are located ventrolaterally in the abdominal region. Positioning of the dorsal and anal fin has not been investigated in zebrafish, but hypothetically may be determined as in sharks by the expression domain of Hox and Tbx genes.\(^{71}\) The development of the pectoral fin shows high similarity with tetrapod limb development especially at early stages, regarding Hox expression\(^{72-74}\) and inductive signals such as retinoic acid (RA)\(^{75-78}\) (Fig. 3A-F). During gastrulation, RA is responsible for limb field positioning.\(^{79}\) Subsequently, RA
Early patterning of limb and pectoral fin buds is comparable. (A–C) Early zebrafish fin bud patterning. (A) Fin induction at 12 hpf: RA from the somites activates signals that lead to activation of tbx5 in the lateral plate and Fgf signaling in the distal bud, establishing the AER. (B) Early expression of Hox genes and shh at 30 hpf. (C) Expression of Hox genes and shh at 60 hpf. Note that the hoxd11 expression domain in the posterior fin bud region remains restricted to this region at 30 hpf and 60 hpf. Dashed line = boundary of fin bud proper and fin fold. (D–F) Early mouse limb bud patterning. (D) Similar induction signals as described in A for the fin bud occur in the mouse at E9. (E) Early expression of Hox genes and shh in E10.5 mouse limbs shows a similar pattern observed as in B for the fin bud. While hoxd9 and hoxd10 extend through the whole bud, hoxd11, hoxd12, hoxd13, and shh are restricted to the posterior domain. (F) Hox expression domain in E12.5 mouse limbs differs from those observed in C in the fin bud; hoxd11, hoxd12, and hoxd13 domains extend more anteriorly; and shh remains restricted to the posterior region. Left to right = proximodistal axis, top down = anterior–posterior axis. E10.5 = embryonic day 10.5; E12.5 = embryonic day 12.5; E9 = embryonic day 9; hpf = hours post fertilization.

Types of Ossification and Osseous Cells

In zebrafish, ossification first occurs 3 to 4 days post fertilization (dpf), progressively leading to the formation of a mature skeleton by 2 to 4 months (dependent on zebrafish size, Table 1). Many elements form by intramembranous and perichondral ossification, with some elements formed by endochondral ossification (Table 1). Depending on the anatomical region, the zebrafish skeleton is formed by either of the three ossification types. Intramembranous ossification is the major form of ossification in the zebrafish skeleton and occurs in elements such as the cranial roof, the opercles covering the gills, and most vertebrae. Endochondral ossification, in which a cartilage template is successively replaced by bone, gives rise to only a few elements including the neural arches of vertebrae 1 to 5. Perichondral ossification occurs on a chondral surface without replacing the cartilage template, and takes place in the lower zebrafish jaw. Although mammalian vertebrae exclusively form via a cartilage intermediate, zebrafish centrae, directly mineralize from the notochord sheath, an ossification layer around the notochord, followed by intramembranous bone formation. Notably, the notochord sheath is also responsible for segmentation of the zebrafish spine prior to centra formation.
Based on the evolutionary conservation of skeletal genes across vertebrates, zebrafish bone contains the same osseous cells that are found in mammals, namely osteoblasts, osteoclasts, and osteocytes (Table 1). Like in mammals, bone apposition is performed by osteoblasts derived from osteoprogenitor cells. In the zebrafish skeleton, a large fraction of bone surface is covered by osteoblasts, whose morphology can vary greatly and is dependent on their location and function.\(^{(5)}\) In the spine, osteoblastic bone formation leads to an increase in vertebral volume and length in anterior and posterior directions, which elongates the spine. This bone formation can be assessed easily in the intervertebral growth region, more specifically on the circular vertebral body endplates.\(^{(85)}\) Bone can be analyzed in the zebrafish spine by using the static histomorphometry protocols that are commonly applied to rodent bones or human biopsies, including the number of osteoblasts per bone perimeter (N. Ob/B.Pm), osteoid surface per bone surface (OS/BS), and osteoid thickness (O.Th).\(^{(56,58)}\) However, zebrafish bone can also be labeled with fluorescent dyes like calcein or Alizarin red at consecutive time points, providing the opportunity to perform dynamic histomorphometry and determine the bone formation rate (BFR) and bone mineral apposition rate (MAR).\(^{(56,58)}\) Table 1. In contrast to rodent models or humans, where dyes are applied by injection or ingestion, zebrafish are more commonly stained by bathing them in the dye solution.

Bone resorption in zebrafish is performed by mononucleated and multinucleated osteoclasts. Although mononucleated osteoclasts are present at the early stages of development and are associated with shallow resorption patterns, multinucleated osteoclasts create resorption lacunae typically described for mammalian osteoclasts later in life.\(^{(60)}\) Similar to mammals, both types of osteoclasts express tartrate-resistant acid phosphatase (TRAP).\(^{(60,96)}\)

Concepts of bone modeling and remodeling described for mammalian bone can also be applied to zebrafish. While bone modeling is defined as the process of adapting bone shape by bone formation and resorption in response to increased or reduced loading at different surfaces, remodeling is carried out in the same location to maintain bone matrix quality and to repair microcracks.\(^{(58)}\) Remodeling, which involves the orchestration of both osteoclasts and osteoblasts by mechanosensitive osteocytes to renew bone and repair microcracks, is less pronounced in zebrafish. Osteons are essentially not present. Yet remodeling processes in zebrafish are suggested to occur site-dependently and to be linked to the demand of lifelong growth.\(^{(6)}\)

Zebrafish bone, in contrast to bone of the medaka fish (Oryzias latipes, another teleost), is generally osteoic, although the vertebral bones do not contain osteocytes in early juvenile stages and the bony fin rays and scales remain anosteocytic throughout life.\(^{(97)}\) Although mammalian osteocytes are the main orchestrator of remodeling, only a few studies have focused on the mechanosensing capabilities of the osteocyte lacunar network and dendrite characteristics in zebrafish. There is, however, evidence for a relationship between the morphology of the osteocyte network and bone formation in a zebrafish OI model, in which altered bone formation is associated with drastically reduced numbers of osteocyte lacunae.\(^{(58)}\) This information illustrates some important differences but also many commonalities between zebrafish and mammalian bone biology (Table 1).

### Bone Matrix and Mineralization in Zebrafish

Vertebrate bone is composed of a soft matrix containing mainly collagen I and noncollagenous proteins and hardens by incorporation of carbonated bone apatite. Although the degree of mineralization varies among vertebrate species, the basic macromolecular and elemental composition of bone matrix is conserved between mammals and zebrafish. The presence of phosphate, carbonate, amide I and III, proline, hydroxyproline, and phenylalanine in zebrafish bone has been demonstrated with vibrational spectroscopy.\(^{(58,88–91)}\) These analyses have shown the typical fingerprints of collagenous matrix with embedded carbonated bone apatite. In growing fin bone, amorphous calcium phosphate is suggested to transform into more crystalline mineral during tissue maturation.\(^{(88)}\) In addition to calcium (Ca) and phosphorous (P), magnesium is one of the main minerals stored in zebrafish bone.\(^{(92)}\) Moreover, trace elements including strontium and zinc are involved in bone formation, similar to the mammalian situation. In particular, these elements have been detected in scales\(^{(93)}\) and vertebral bone matrix\(^{(94)}\) using X-ray fluorescence microscopy and energy dispersive X-ray spectroscopy, respectively.

Clearly, some differences apply in terms of mineral metabolism between teleosts and terrestrial mammals. In contrast to tetrapods, which depend on both dietary P and Ca intake for maintaining the Ca–P–based bone matrix, zebrafish live in a Ca-rich environment and absorb Ca through their gills. However, dietary P intake is required, and reducing the P levels either through genetic manipulation or through a reduced diet leads to nonmineralizing matrix.\(^{(95,96)}\)

As in mammals, the composition and organization of the bone matrix is crucial to providing fracture resistance in zebrafish. Specifically, the sum of collagen-related and mineral-related structural properties (collagen alignment, enzymatic and nonenzymatic crosslinking, size and orientation of mineral particles) and compositional properties (carbonate-to-phosphate ratio), as well as the degree of mineralization in terms of overall bone mineral density (BMD) or local Ca content and distribution, determine the mechanical properties at the tissue level. This translates into fracture risk at the whole bone level.\(^{(58,94)}\) Several techniques have been adapted from mammalian bone analyses to determine the degree of mineralization in micrometer-sized zebrafish bone (Table 1). μCT based on calibration with hydroxyapatite phantoms has been used to assess BMD in the spine of zebrafish.\(^{(25)}\) Indeed, variations in BMD due to mutations support the presence of similar mineralization pathways in zebrafish and mammalian models. The macromolecular composition of pathologic bone matrix in zebrafish carrying mutations in \(\text{colla1a1}\) has been analyzed by vibrational spectroscopy and demonstrated alterations in the mineral-to-matrix and carbonate-to-phosphate ratios.\(^{(58)}\) Moreover, remineralization in regenerating zebrafish fin bones after amputation can be monitored using vibrational spectroscopy, allowing the comparison of new bone quality with the quality of native bone (Fig. 2A). Another indicator for a well-mineralized bone matrix is the Ca content and the distribution of Ca within the bone matrix. These parameters are commonly assessed in rodents and human biopsies using two-dimensional (2D) quantitative-backscattered electron microscopy (qBEI)\(^{(97)}\) (Fig. 2B). More recently, qBEI has also been adapted to zebrafish vertebral bone, in which an increase in mean Ca content by ~8% has been linked to increased musculoskeletal activity.\(^{(56)}\)

Increased mineralization may lead to increased bone fracture resistance.\(^{(96,99)}\) Although whole-bone mechanical testing of zebrafish bones, eg, of individual vertebrae, is challenging (though possible)\(^{(12)}\) due to their small size (VBL = 500 μm), nanoindentation techniques are valuable tools to assess mechanical properties of zebrafish vertebral bone at the tissue level. As an example, rising Ca/P ratios have been associated with
an increase in elastic modulus in zebrafish vertebral bone during aging.\(^\text{[94]}\) However, the opposite effect can be observed as well. In case of disturbed collagen and consecutive mineral particle deposition (eg, in chihuahua\(^\text{[85]}\), liliput\(^\text{[91]}\), and stöpse\(^\text{[100]}\) zebrafish mutants), the elastic modulus of vertebral bone is reduced despite a higher degree of mineralization, stressing the importance of well-organized matrix mineralization to withstand fracture. Notably, zebrafish bone matrix elastic modulus values lie in a similar range as in mammals (up to 24 GPa)\(^\text{[101]}\), highlighting the similarities in bone matrix composition and mechanical properties between both. This substantiates the use of zebrafish as a model to study the effects of genetic alterations and external stimuli on bone matrix quality.

Problems in matrix mineralization such as those present in phosphate homeostasis disorders have been phenocopied in zebrafish. Phosphate is an essential mineral for hydroxyapatite formation in bone\(^\text{[102]}\), and its lack (hypophosphatemia) is provoked in mutants, in which essential genes for phosphate regulation are altered. The no bone (nob) mutant, in which the gene ectonucleotide triphosphate diphosphohydrolase 5 (entpd5) is affected, forms no mineralized bone at all.\(^\text{[103]}\) In another mutant, dragonfish (dgf\(^\text{[\text{\textsuperscript{10}}]}\)), the gene ectonucleotide pyrophosphatase/phosphodiesterase 1 (enpp1) is mutated, leading to reduced pyrophosphate levels and ectopic mineralization of the axial and craniofacial skeleton.\(^\text{[103]}\) The dgf mutant shows altered expression of genes related to phosphate homeostasis and bone mineralization, such as fgf23, solute carrier family 34 member 1a (slc34a1a, also known as npt2a), entpd5, and secreted phosphoprotein 1 (spp1), and can thus also be used to model generalized arterial calcification of infancy (GACI) and pseudoxanthoma elasticum (PXE).\(^\text{[104]}\) Restored skeletal mineralization is observed in double mutant nob/dgf zebrafish, indicating a reciprocal regulation of Enptd5 and Enpp1.\(^\text{[104]}\) This illustrates the usefulness of zebrafish mutant analyses and state-of-the-art technology to understand mineralization defects in vertebrates.

### Response of the Zebrafish Skeleton to Loading

Although loading of the zebrafish skeleton differs from that of terrestrial animals, due to the supportive buoyancy of water, the skeleton does respond to mechanical loading (Table 1). This comes from the action of muscle contraction on the skeleton, and reaction forces from swimming through a viscous medium.

During early fetal life all vertebrates develop in an aqueous environment, whether in utero, in ovo, or in water (Table 1). During this time, biomechanical stimuli acting on the developing skeleton are caused by the action of muscle on skeletal tissues.\(^\text{[105]}\) It has been demonstrated in mice, chicks, and humans that restriction of fetal movement leads to altered mineralization and to failure of joint and eminence morphogenesis.\(^\text{[106–108]}\) In zebrafish, genetically or pharmacologically induced paralysis leads to altered chondrocyte maturation\(^\text{[109]}\) and abnormal joint morphogenesis through changes in chondrocyte proliferation and migration.\(^\text{[110,111]}\) Finite element (FE) modeling of the larval jaw, in which the structure is subdivided into smaller and simpler entities, allows modeling of the loading effects on tissue deformation, and has demonstrated that altered joint shape impacts the pattern of biomechanical strain.\(^\text{[112,113]}\) It is well established that biomechanical loading of the joint is a key risk factor for osteoarthritis. Moreover, mutants such as col1a1a2 and pro4 exhibiting altered joint shape go on to develop osteoarthritis in these joints.\(^\text{[90,113,114]}\) Given the recent identification of loci associated both with osteoarthritis and altered joint shape in presumptomatic humans\(^\text{[115,116]}\) and the need for functional screening platforms, this raises the prospect of using zebrafish to screen for osteoarthritis susceptibility genes implicated in joint development and maintenance (Fig. 4).

Mineralization of the vertebral column and fins of zebrafish occurs much later than the onset of locomotion, with first mineralization of vertebral centra observed at around 7 dpf (SL = 3.8 mm), and vertebral arches and fin rays at around day 14 (SL = 5–6 mm).\(^\text{[152,117]}\) Vertebral bone formation can be triggered by increased physiological musculoskeletal loading in adult zebrafish, demonstrating that zebrafish bone is susceptible to positive bone modeling according to Wolff’s law.\(^\text{[96]}\) Zebrafish subjected to swim training for 9 hours a day from 5 to 14 dpf exhibit premature ossification of fin and vertebral column bone.\(^\text{[118]}\) In terrestrial species osteocytess function as mechanosensors, directing the remodeling activity of osteoblasts and osteoclasts, through the regulation of the glycoprotein Sclerostin (SOST).\(^\text{[119]}\) However, anosteocytic fish also model bone in response to load.\(^\text{[120]}\) Swim training of osteocytic zebrafish and anosteocytic medaka led to strikingly similar patterns of new bone formation in both species, mediated by sost expression by chondrocytes and osteoblasts in regions of high strain modeled by FE on individual vertebrae.\(^\text{[121]}\) Interestingly, zebrafish vertebral motion analysis together with FE predict patterns of bone failure during loading,\(^\text{[122]}\) which could be used to test bone performance in mutants.

Gravity plays an important role in the loading of the skeleton of terrestrial animals, and prolonged exposure to microgravity (weightlessness or gravity close to zero) leads to decreased bone density in humans.\(^\text{[123]}\) Perhaps surprisingly, gravity also impacts the skeleton of teleost fish. Several studies on medaka have been performed using an aquatic habitat on the International Space Station (ISS). These have demonstrated that medaka lose bone density following exposure to microgravity, show transcriptional changes to skeletal genes\(^\text{[123]}\) and increased osteoclast activity.\(^\text{[126]}\) Although zebrafish have not been reared on the ISS, they have been exposed to increased gravitational forces (hypergravity). Exposure to 3g to 9g during zebrafish larval development led to altered chondrocyte maturation\(^\text{[125]}\) and changes to mineralization and the transcription of skeletal genes.\(^\text{[126]}\) These studies illustrate the versatility of zebrafish models to study loading effects on bone.

### OI and Osteopetrosis

The use of genetic tools, bone imaging, and pharmacological treatments in zebrafish models has increased our understanding of the pathophysiology of congenital skeletal diseases, and has been reviewed in detail.\(^\text{[39]}\) OI is the term given to a collection of rare genetic bone collagenopathies, which are characterized by suboptimal skeletal development, aberrant bone architecture, and high fracture incidence.\(^\text{[127]}\) There are multiple subtypes of OI, with varying degrees of severity; type I OI is the mildest form, which is underpinned by reduced production of normal type 1 collagen, whereas other types are the result of mutations which alter the molecular structure of type 1 collagen.\(^\text{[128]}\) The majority of OI cases result from mutations in COL1A1/2. The chihuahua zebrafish mutant, identified through forward genetic screening, was the first of several lines to accurately model OI in zebrafish.\(^\text{[45,129]}\) Heterozygous chihuahua zebrafish possess a dominant mutation in col1a1, resulting in gross...
skeletal deformities and molecular abnormalities in bone mineralization OI. Some rarer forms of OI arise from mutations in collagen-processing genes such as BMP1, PLOD2, CRTAP, and P3H1, or osteoblast-related genes such as SP7, for which there are stable zebrafish mutant lines. Although there is no cure for OI, many patients are administered antiresorptive bisphosphonate drugs to increase BMD. Zebrafish have been used to explore the role of bisphosphonates on bone in OI. For example, sustained treatment of the frilly fins (bmp1a) mutant with alendronate reduced the frequency of fractures in caudal lepidotrichia but could not rescue defects in fracture repair post injury.

At the opposite end of the spectrum, genes associated with high bone mass (osteopetrosis) have been modeled in...
zebrafish. The *panther* (*csf1ra*) mutant zebrafish line exhibits low levels of osteoclast activity and osteopetrosis, due to the lack of colony stimulating factor 1 receptor alpha, which promotes the differentiation of myeloid lineage cells.\(^{114,145}\) Studies of the *panther* zebrafish line have helped to demonstrate the need for effective intercellular signaling by osteoblasts and osteoclasts throughout skeletal development and for maintaining bone architecture.\(^{143,144,146}\) More recently, integrative studies have used pedigree analyses to identify rare mutations in high bone mass genes such as *CLCN7* and *SMAD9*, followed by functional validation in zebrafish.\(^{111,147,148}\) Collectively, these studies have implicated the CIC-7/CTSK/TGF-β/SMAD (CHLORIDE ION CHANNEL 7/CATHEPSIN K/TRANSFORMING GROWTH FACTOR β/SMAD) signaling axis in osteopetrosis.

Human epidemiological analyses and genomewide association studies (GWAS) continue to rapidly identify new loci associated with skeletal health (Fig. 4A).\(^{149,150}\) yet these gene candidates require functional validation. In order to keep up with this pace, follow-up studies using zebrafish have evolved away from traditional forward genetic screening methods and toward modern genome editing tools such as CRISPR-Cas9 (Fig. 4B).\(^{151}\) Emerging phenomics-based deletion approaches in mosaic zebrafish (crisprants) now present an efficient model for the rapid validation of novel GWAS-derived genes related to bone disease.\(^{29,40}\) (Fig. 4B, C, Table 1). In many cases generating knockouts by the creation of indels with CRISPR-Cas9 can be informative, providing information on the nature of the putative associated genes and the likelihood that they are indeed causal. A potential limitation is reached when knock-in of the specific genetic change(s) identified in humans is required, because gene editing using homologous recombination or base editing, while possible,\(^{152-154}\) is less efficient than the generation of frameshift alleles through non-homologous end joining. Although stable mutant line generation by CRISPR-Cas9 knockin\(^{36,37}\) will be the gold standard to prove altered function of a gene due to a point mutation, mosaic deletion in crisprants will be one of the ways to deal with the rich information obtained from GWAS.

**The Regenerative Capacity of the Zebrafish Skeleton**

Zebrafish regenerate various organs such as the retina, brain, and pancreas, which has promoted the use of zebrafish as a regeneration model.\(^{155}\) In teleosts and mammals, derivatives of dermal skeletal tissue are represented in terms of teeth, whereas endoskeletal tissue is represented in terms of bone and cartilage. In contrast to mammals, however, the dermal skeleton of zebrafish also encompasses fin rays and scales, which have the capability to regenerate throughout life (Table 1). The presence of such tissues facilitates the study of skeletal features that do not exist in humans, including continuous tooth replacement\(^ {156,157}\) and regeneration of scales and fins.\(^ {15}\) Moreover, calvaria and jaw bone regenerate in zebrafish (Fig. 5A-E, Table 2). Zebrafish fins quickly and completely regenerate after profound amputation, a process which is regulated by various intercellular signaling pathway events.\(^ {173}\) In fact, regenerated fins are nearly indistinguishable from uninjured fins, as long as the endoskeletal elements (eg, hyurpals) are retained during the amputation procedure.\(^ {5,174-176}\) Notably, regenerative capacity of the caudal fin is not altered even after repeated amputations.\(^ {177}\) The skeleton of the nonmuscular part of the caudal fin, predominantly used for regeneration studies because of its accessibility, is of dermal origin. The most prominent feature of this part is the presence of segmented bony fin rays (lepidotrichia), arranged in two concave hemirays. Bony fin rays contain lose mesenchyme (intra-ray fibroblasts), arteries and nerves and are covered by a sheet of osteoblasts and overlaying epidermis. Inter-ray mesenchyme connects individual bony fin rays to each other.

Control over lepidotrichia segment length and joint formation, which significantly differs from endoskeletal joint formation, is crucial in regulating fin growth and regeneration. Although *qdf5*, an endoskeletal joint marker, is not expressed in lepidotrichia segment joints,\(^ {178-180}\) distinct lepidotrichial joint markers such as *exv1* and the gap junction gene *cx43* have been identified.\(^ {180,181}\) Notably, joint formation and segment length are regulated independently during growth of the fin.\(^ {182}\) Mutation of *cx43* leads to the *sof* (short fin) phenotype resulting from shorter fin ray segments due to an increased rate of joint formation.\(^ {182,183}\) In contrast, longer fins in *long fin* (*lof*) and *rapunzel* mutants display normal-length segments, but more in number.\(^ {182,184}\) In another *long fin* (*alf*) mutants with altered potassium channel function fins are longer and segment length is variable.\(^ {185}\) These phenotypes are reestablished during regeneration after fin amputation. We direct the interested reader to excellent expert reviews on this topic.\(^ {41,186}\)

Regeneration occurs in a series of events. After amputation, epithelial cells at the wound site migrate to form a multilayered epithelium termed the wound epidermis.\(^ {187}\) Subsequently, mesenchymal cells from the stump migrate distally and proliferate to form the blastema,\(^ {188}\) which is a mass of cells restoring the missing structures. The blastema matures and subdivides into a distal signaling center (the most distal blastema),\(^ {189,190}\) and a more proximal, highly proliferative growth and patterning zone.\(^ {191}\) When blastemal organization is complete, regenerative outgrowth proceeds for approximately 2 weeks at high speed by coordinated cell proliferation and differentiation along the proximodistal axis, and is followed by slower growth until completion after 3 to 4 weeks post amputation depending on the amputation level.\(^ {192,193}\)

The bony fin rays are the stabilizing elements of the fins and much progress has recently been made in describing their regeneration. During blastema formation, mature osteoblasts lining the inner and outer surface of the stump hemirays differentiate by losing *bglap* (*osteocalcin*) and *sp7* (*osterix*) expression, a process that is regulated by RA and NF-κB signaling.\(^ {194,195}\) They migrate distally to form part of the blastema, where they then upregulate the immature osteoblast marker *runx2*. This is followed by redifferentiation in a proximal to distal sequence (most mature at proximal position: *osterix*/osteocalcin/sp1 expression, followed by *runx2/osterix* expression, and *runx2* most distally).\(^ {15,196}\) Lineage restriction of osteoblasts has been demonstrated during the regeneration process.\(^ {15,166,167}\) Notably, mature stump osteoblasts are not the only source of osteoblasts in regeneration, and several other cell populations contribute to regenerated bone in zebrafish and medaka.\(^ {197}\) Thus, bone regeneration in teleost fin rays is highly plastic.\(^ {41}\)

Human bone traumata rarely include profound tissue removal, but often present as fractures or tissue necroses. In addition to a cryoinjury model, in which necrotic bone fragments get displaced from the wound margin,\(^ {199}\) bony fin ray fractures
are increasingly used to elucidate mechanisms of vertebrate bone repair (Table 1). To date, two main fin fracture models have been described: a crush injury model that affects several bony fin rays\(^{(13)}\) and a milder fracture model which only affects a single bony fin ray in usually one hemiray segment\(^{(14,200)}\) (Fig. 5E). Although there are differences between these two models, fracture repair in both involves a thickening of the tissue surrounding the lesion, reminiscent of the callus formed during mammalian fracture repair (Fig. 5E, Table 2). The molecular mechanisms and cellular dynamics underlying zebrafish fracture repair and fin regeneration are surprisingly similar. For example, dedifferentiation, migration and redifferentiation of osteoblasts also occur during zebrafish fracture healing\(^{(13,14)}\). Thus, it remains to be investigated which events truly distinguish fin fracture healing from fin regeneration after amputation. One important difference between mammalian and zebrafish fracture healing, however, concerns the contribution of osteoblast precursors from the BM. These BM-derived stromal cells take part in mammalian long-bone fracture repair\(^{(201)}\) but are absent in zebrafish, which lack BM proper. In contrast, osteoblast dedifferentiation has not been described in mammals\(^{(202)}\) with the exception of digit tip regeneration during which bone cells acquire a blastemal state to regenerate the tissue,\(^{(16,17)}\) and in bone explants.\(^{(203)}\) Future work needs to evaluate the relevance
of osteoblast dedifferentiation for mammalian fracture repair and bone homeostasis.

In both mammals and in zebrafish, skull injuries (trepanations) are repaired by intramembranous ossification without callus formation (Table 2).[170] In zebrafish, trepanations have been performed by drilling a hole in the os frontale and/or os parietale[14,200] (Fig. 5A). As with fin injury paradigms, trepanation of zebrafish calvariae induces mature osteoblasts to dedifferentiate,[14] a mechanism that has not been tested for in mammalian skull injury models (Table 2). In contrast, calvarial bone healing in mammals might be mediated by stem cells at the sutures.[172]

Zebrafish bones have some special gene expression features best exemplified by reference to jaw regeneration, which produces a hybrid cartilage-bone cell type. This cell type first produces cartilage and later switches to bone matrix mineralization.[160] Interestingly, hypertrophic chondrocytes give rise to osteoblasts in developing zebrafish cartilage bone.[204] A phenomenon also observed in mammalian mandibular condyle[205] and long-bone development.[206,207] Besides this peculiarity, zebrafish jaw regeneration occurs by formation of a transient cartilage callus and involves a periostal origin of bone-forming cells, similar to what has been reported in mammalian fracture repair.[160–162] Surgical removal of jaw tissue in zebrafish, however, leads to wound epidermis and blastema formation.[160,162,163] as well as activation of signaling cascades known from appendage regeneration in zebrafish and other nonmammalian vertebrates.[208,209] Although jaw regeneration in zebrafish bridges wide skeletal gaps, it may result in a malformed shape with ectopic ossification of cartilage in the mandibular symphysis (Fig. 5B).[162,163] Unlike other bone regeneration models in zebrafish, jaw regeneration involves callus formation and progenitor sources equivalent to bone healing in nonregenerative species. In humans, jaw regeneration is limited and methods for repairing the missing jaw tissue include distraction of the remaining mandibular bone or the use of implants, tissue transplants or, more recently, stem cells, although these therapies are not completely successful.[210] Hence, investigating differences between appendage and jaw regeneration in different species could lead to novel approaches for regenerative therapy in humans.

Another telost bone structure with high regenerative capacity is the elasmoid scale, a dermal bone embedded in the skin (Fig. 5C). The scale is covered by osteoblasts on both sides and osteoclasts along mineral grooves (radii).[211] As for appendages, the first stage of regeneration is wound reepithelialization.[164] Rapid reconstruction of the scale occurs by proliferation of a pool of de novo osteoblasts, shape changes, and cell death, resulting in three spatially distinct osteoblast populations.[213,165] Osteoblasts then deposit collagen fibrils, and mineralization of the scale proceeds. Osteoclasts remodel the scale to its final shape.[31,164,165,211]

Zebrafish have a high capacity to regenerate different skeletal tissues, and plasticity of bone forming cells may be the key to maintaining this ability. Understanding and identifying conserved regenerative mechanism in vertebrates that potentially have been lost in mammals will be important to modulate human bone regeneration in a clinical setting in the future.

### Table 2. Overview on Zebrafish Bone Injury Paradigms and Mechanisms of Bone Restoration

| Bone injury | Ossification | Callus formation | De novo | Dedifferentiation-redifferentiation (in vivo) | References |
|-------------|--------------|-----------------|---------|---------------------------------------------|------------|
| Zebrafish   |              |                 |         |                                             |            |
| Skull       | Intramembranos | No              | ?       | Yes                                         | (14)       |
| Jaw         | Intramembranos | Yes             | Yes     | ? (yes)                                     | (160–163)  |
| Scale       | Intramembranos | No              | Yes     | ?                                           | (23,164,165)|
| Fin amputation | Intramembranos | No              | Yes     | Yes                                        | (15,166–169)|
| Fin fracture | Intramembranos | Yes             | ?       | Yes                                        | (13,14)    |
| Mammals     |              |                 |         |                                             |            |
| Tibia       | Endochondral  | Yes             | Yes     | ?                                           | (161,170)  |
| Femur       | Endochondral  | Yes             | Yes     | ?                                           | (171)      |
| Skull       | Intramembranos | No              | Yes     | ?                                           | (170,172)  |

Metabolic and hormonal problems cause a wide range of human disorders that frequently involve bone.[212] Several related bone disease models, eg, mimicking osteoporosis-like phenotypes, have been developed in zebrafish,[213] often by using unique anatomical bone structures such as exoskeletal zebrafish fin rays or scales.[214]

Zebrafish have been increasingly used to study the adverse effects of glucocorticoids (GCs), which cause GC-induced osteoporosis (GIO) in patients undergoing immunosuppressive treatment, on bone (Table 3). GC are mainly produced in the zebrafish interrenal gland within the head kidney,[228] the equivalent of the adrenal cortex in mammals.[229] Like humans,[230] zebrafish have two GC receptor (GR) isoforms (gene nr3c1, nuclear receptor subfamily 3 group C member 1), GRα and GRβ, which are nuclear hormone receptors, with a similar effective GRα/GRβ ratio and predominantly nuclear localization.[231,232] In contrast to humans, the zebrafish GRβ does not function as a dominant negative inhibitor of the GRα isoform and is not transcriptionally active.[233,234] Furthermore, zebrafish lack a direct homologue of Hydroxysteroid 11-beta dehydrogenase 1 (11β-HSD1)[235] and therefore do not reduce 11-ketosteroid, which means that they are not able to activate GC from inactive precursors and thus exhibit a limited tissue-specific action.[236] However, stress axis signaling is otherwise conserved in zebrafish.[237] Importantly, zebrafish react to stress by producing the corticosteroid cortisol, like humans, whereas rodent species rely on corticosterone production upon stress.[238,239] Moreover, active forms of GC can be used to circumvent the
lack of 11β-HSD1, which is why GC effects on bone can be well studied in zebrafish.\(^{(237)}\)

In vivo manipulation of GR signaling is technologically advanced in zebrafish models, and a variety of zebrafish mutants and transgenic reporter lines have been generated to analyze GR signaling activity and GC resistance (Table 4). Several GR mutants exist today, such as the hypomorph \(gr^{+357/244}\) and the null mutant \(nr3c1/ia^{30/ia^{30}}\)\(^{(245)}\). Remarkably, zebrafish null mutant larvae have increased levels of whole-body cortisol, but nevertheless show relatively normal morphology and are viable through adulthood, in contrast to mice.\(^{(245)}\) A variety of transgenic zebrafish reporter lines, such as the Tg(GRE:EGFP,myl7:TagBFP) line, also known as SR4G,\(^{(240)}\) the Tg(GRE:Luciferase) line, which is glucocorticoid-responsive in vivo zebrafish luciferase activity” (GRIZLY) assay,\(^{(242)}\) and the Tg(9xGRE-HSV:U23:EGFP) line,\(^{(241)}\) are useful to monitor GR activity across zebrafish tissues in vivo. GC resistance can be visualized with the help of the Tg(pomc:GFP) line, in which the reaction to GC treatment can be monitored via the decrease of pomc expression in the pituitary gland. A forward-genetic screen enabled the identification of GC-resistant zebrafish mutants, such as loopless (lp), lacking pomc suppression.\(^{(243)}\) Notably, ubiquitous GR knockout in zebrafish leads to increased muscle mass,\(^{(247)}\) which could mechanically impact bone, too. Zebrafish can also be utilized to investigate the function of the mineralocorticoid receptor (MR) that is evolutionarily linked and cooperating with the GR.\(^{(248)}\) Recently, studies on the MR and GR showed that both receptors differentially regulate transcription, protein deposition, and proteolysis during larval development and that their combined activation is responsible for growth suppression.\(^{(249)}\)

The mentioned tools can be used to model the effects of excess GC on bone. In embryos and larvae, models mimicking impaired bone formation as observed during GIO were established by using prednisolone\(^{(219–221)}\) and dexamethasone.\(^{(215–217)}\) Inhibited bone formation and mineralization linked to reduced expression of osteoblast-specific genes\(^{(215–217,219,220)}\) and increased osteoclast activity\(^{(217,221)}\).

**Table 3. Overview on Bone Inhibitory Effects of Excess GC Levels in Zebrafish**

| Zebrasfish model | Compound screen possible? GC | Effect | Reference |
|------------------|------------------------------|--------|-----------|
| Development      | +                            | Dexamethasone | Inhibition of bone formation, downregulation of osteoblast-specific genes, reduced mineralized matrix in skull | (215–218) |
|                  |                              | Prednisolone | Delayed and reduced mineralization, enriched NF-κB and focal adhesion signaling pathway, increased osteoclast activity | (219–222) |
| Fin fold regeneration | +                          | Beclomethasone | Inhibited regeneration, less proliferating cells, inhibition of neutrophil migration | (223,224) |
| Scales           | +                            | Dexamethasone | Decreased size and circularity in regenerating scales, suppressed osteoclast activity | (225) |
|                  |                              | Prednisolone | Enhanced matrix breakdown, increased osteoclast activation | (226) |
| Adult fin regeneration | –                          | Prednisolone | Regenerates remain shorter, osteoblast proliferation and differentiation reduced, no increased osteoclast activity | (159,227) |
| Skull regeneration | –                           | Prednisolone | Complete, but slow regeneration | (219) |

**Table 4. Tools to Study GR Signaling and GC Resistance in Zebrafish**

| Tool | Use/details | Reference |
|------|-------------|-----------|
| Reporter lines | Tg(6xGRE:EGFP,myl7:TagBFP)\(^{mn48}\) | Monitors GR activity, high-resolution model for physiological and stressed conditions | (240) |
|   | Tg(9xGRE-HSV:U23:EGFP)\(^{ia^{30}/ia^{30}}\) | Monitors GR activity, high responsiveness to GC treatment, endogenous stimuli, and molecular manipulation | (241) |
|   | Tg(GRE:Luciferase)\(^{ib6}\) | Monitors GR activity, can be monitored in a specially developed GRIZLY assay | (242) |
|   | Tg(pomc:GFP)\(^{184}\) | Monitors the reaction to GC treatment via pomc expression; visualization of GC resistance | (243) |
| Mutants | \(gr^{+357}\) | Hypomorph GR mutant | (244) |
|   | \(nr3c1/ia^{30/ia^{30}}\), \(nr3c1/co^{402/co^{402}}\) | GR null mutants, hypercortisolemic, failing cortisol stress response | (245,246) |
|   | loopless \(lp^{u6377}\) | GC resistant, lack pomc expression | (243) |
|   | \(nr3c2/co^{402/co^{402}}\) | MR null mutant, delayed and dysregulated cortisol response | (246) |

GRIZLY = glucocorticoid-responsive in vivo zebrafish luciferase activity.
were the most prominent effects observed. In 2-week-old, prednisolone-treated zebrafish, mRNA levels of *osterix* (*sp7*), osteocalcin (*bglap*), and *entpd5* were clearly reduced, whereas some matrix metalloproteases were induced. Excess GC levels have anti-regenerative effects in zebrafish. Larval fin fold regeneration was inhibited after prednisolone and beclo

methasone dipropionate treatment, a process that was mediated by *cripto-1*, a Nodal signaling co-factor and Activin-signaling antagonist. Larval fin fold regeneration was not inhibited with the selective GR agonist ginsenoside Rg1 in adults. Prednisolone treatment caused poor scale regeneration resulting from enhanced matrix breakdown due to increased osteoclast activity, and impaired fin regeneration also due to alterations in vesicular transport mechanisms. Studies with beclo

methasone suggest that a temporary activation of the GR during blastema formation is sufficient to block regeneration. Notably, antosteostaticogenic effects were observed in fin regeneration and fin ray fracture healing following prednisolone treatment in zebrafish (Geurtzen and colleagues and personal observations) and medaka. Vertebral bone volume and skull regeneration after trepanation were remarkably unaffected by prednisolone treatment.

Zebrafish are known for their use in drug screening approaches. High-throughput screens on larvae to identify anti-osteoporotic compounds have identified compounds such as RU486 and tanshinol. Another antosteoporotic drug, the flavonoid icaritin, which protects against Rankl-induced bone resorption, was identified in medaka embryos. Zebrafish scales are a useful tool for drug screening purposes and in vivo imaging. Scales are small and abundant, have transparent anatomical structures and can be cultured for up to 72 hours (Table 1). Transgenic reporter lines labeling osteoblasts are useful in this context. In terms of GC-related research, the scale model confirmed the bone protective function of the bisphosphonate alendronate and the vitamin D (vitD) homologue alfacalcidol. The scale model also holds promise to further elucidate close interactions of bone resident cells. Fractured zebrafish scales revealed a previously unrecognized cellular mechanism of osteoblast–osteoclast communication via osteoblast-derived extracellular vesicles promoting osteoclast differentiation.

Low bone quality and increased fracture risk are a frequent consequence of diabetes mellitus. Both type I and type II diabetes mellitus have been modeled in zebrafish. Type I diabetic zebrafish, which were injected with the diabetogenic drug streptozocin, killing pancreatic beta-cells, showed impaired fin regeneration due to decreased proliferative potential in the regenerate. In type II diabetic zebrafish, which were incubated in high glucose, scales showed an imbalance in bone metabolism inducing an osteoporosis-like phenotype. Moreover, scale assays revealed that the antioxidant liquiritigenin counteracts osteoporotic complications in hyperglycemic fish. It will be interesting to investigate bone quality of other bones in diabetic zebrafish, such as the vertebrae and skull, and to characterize fracture repair in one of the available fracture models in the future.

As in mammals, vitD is suggested to play a role in increasing the available plasma phosphate in fish. Overexpression of *hand2*, which stimulates vitD inactivation, led to altered regeneration of bones in the zebrafish pectoral fin, which could be partially rescued by administration of vitD. Likewise, vitD receptor inhibition suppressed fin regeneration, whereas vitD analogue treatment promoted fin regeneration. In vivo compound screening with larval zebrafish revealed dose-dependent increases in the formation of mineralized bone following vitD and calcitriol administration. Furthermore, diabetic zebrafish treated with paricalcitol, a vitD analog, showed improved bone regeneration and mineralization due to enhanced osteoblast differentiation and insulin expression. These studies confirmed the importance of vitD for bone health in zebrafish. Altogether, the described work strengthened the importance of zebrafish for studying hormonal and metabolic bone disorders in nonmammalian species.

**Larval Models to Study Bone Metastasis**

Zebrafish are increasingly used to study bone disease related to secondary tumor formation. Bone metastases are common in cancer patients and associated with a poor prognosis. The composition of bone extracellular matrix with its embedded cells, growth factors, and chemokines makes it an attractive site for cancer cell homing. Crosstalk between bone and cancer cells leads to an imbalance between osteoblast-mediated bone formation and osteoclast-mediated bone resolution resulting in dysregulated bone remodeling (reviewed in Guise). Such interactions can promote bone metastasis formation.

Notably, although osteoblasts and osteoclasts have been shown to impact bone metastasis, many studies demonstrate that the mammalian BM is a particularly favorable environment for tumor cells (reviewed eg, in Schmid-Alliana and colleagues). BM is not a mineralized tissue, but is of special interest to clinicians seeing cancer patients developing bone metastasis. In zebrafish larvae, bone formation and hematopoiesis take place in separate locations, which allows scientists to reveal the distinct roles of bone mineral and BM-like attractants for cancer cells. Mammalian BM expresses cancer cell attractants which is also true for the zebrafish caudal hematopoietic tissue (CHT) (Fig. 6). The cytoarchitecture and processes observed in the CHT resemble those of mammalian BM (Fig. 6A,B), which is why the CHT is often referred to as a BM-like niche. In zebrafish larvae, blood formation takes place in this vascular plexus in the ventral tail region of the growing zebrafish. Experimentally, it has been observed that some cancer cells home to the CHT leaving the circulation. This, together with the fact that cancer cells can be injected easily into zebrafish embryos and larvae, led to the establishment of zebrafish xenografts to study bone metastasis formation.

The CHT contains different cell types, including endothelial, lymphoid, myeloid, and mesenchymal stromal cells (MSCs) (Fig. 6A). Hematopoietic stem cells (HSCs) migrate to the zebrafish CHT niche in response to increased chemokine (*C-X-C* motif ligand 12a) expressed by MSCs. Although the CXCL12/CXCR4 (*C-X-C* Motif Chemokine Receptor 4) axis was already shown to exert an important role in cancer-endothelial cell adhesion, invasive activity modulation, and cancer cell proliferation in mice, a growing number of zebrafish studies highlight the contribution of this signaling pathway to metastasis formation. The CHT also expresses *gata2B*, the orthologue of *Gata2* that regulates HSC maintenance within the BM and is associated with distant metastatic progression of prostate cancer.

The CHT is highly vascularized and its endothelial cells can form a stem cell pocket to sustain HSCs. Endothelial cells in the CHT express transcription factor EC (*tfec*), *kit* ligand (*kitlg*), oncostatin M (*osm*), *kruppel-like factor 6a* (*klf6a*), *C-C motif chemokine ligand 25b* (*ccl25b*), *chemokine (C-X-C motif) receptor 1* (*cxcr1*), and *chemokine (C-X-C motif)
ligand 8a \((\text{cxcl8a})\)\(^{(278)}\) (Fig. 6A), which are responsible for HSC colonization and whose orthologues are partially expressed in BM and might contribute to metastasis formation\(^{(292-298)}\).

Myeloid cells (neutrophils and macrophages), whose involvement in bone metastasis has been established in murine models\(^{(298,299)}\) populate the CHT. In zebrafish, myeloid cell depletion results in breast cancer cell invasion in the proximity of the CHT\(^{(284)}\).

To induce metastasis in zebrafish, a variety of methods such as mutagenesis by carcinogens, targeted mutagenesis of tumor suppressor genes, and tissue-specific overexpression of oncogenes are available\(^{(300-302)}\). Xenografting, ie, the transfer of living cells between species, is a straightforward way to investigate the mechanisms underlying bone metastasis. The benefits of zebrafish larval xenografts include the possibility of high throughput, the ease of tracking of fluorescently labeled cancer cells due to optical clarity of larvae\(^{(303)}\) and relatively fast micrometastasis development\(^{(304,305)}\). Cancer cells can be injected into the bloodstream through the duct of Cuvier \((\text{DoC})\), heart, and posterior cardinal vein of 2 dpf larvae in which adaptive immunity is absent\(^{(276,306)}\).

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**Fig 6.** The CHT as an attractive site to study bone metastasis (A) Zebrafish larvae at 3 to 5 dpf contain developing bone (opercle and cleithrum) and CHT which acts as a BM-like niche. (Left) The opercle and cleithrum are among the first dermal bones to develop and can be imaged in live larvae due to their lateral, superficial positioning. (Right) The CHT is located laterally to the dorsal aorta in the posterior region of the larval zebrafish and is the site of early hematopoiesis, akin to mammalian BM. The cellular components of the CHT include endothelial cells, MSCs, HSCs, myeloid cells such as macrophages and neutrophils, lymphoid cells, and erythroid cells. Molecular components of the CHT listed have been previously shown to play a role in bone metastasis in mammalian models. Metastasis-related factors expressed in cancer cells uncovered in this model are shown on the right. (B) Illustration of the BM niche in a mouse long bone containing similar cell types as the CHT. dpf = days post fertilization; MSCs = mesenchymal stromal cells; HSCs = hematopoietic stem cells.
Homing of cancer cells to new sites requires extravasation and resembles micrometastasis formation, which can be observed within a few days post injection at sites such as the CHT, fin fold, and trunk.\(^{284,305,307}\)

The larval xenograft model has been employed to investigate bone metastasis of prostate cancer, breast cancer, and multiple myeloma (MM) in the CHT and studied regarding the molecules and signaling pathways involved\(^{276,308-312}\) (Fig. 6A). Human prostate cancer cells, resected from the CHT several days post–DoC injection, were found to upregulate stemness (eg, NANOGL, OCT4, Cripto, C-X-C Motif Chemokine Ligand 2 [CXCL2]) and mesenchymal (eg, Vimentin, Twist, and Zinc Finger E-Box Binding Homeobox 2 [ZEB2]) markers while reducing E-cadherin.\(^{308}\) Moreover, the CHT microenvironment enhanced Activin A expression, which correlates with increased bone metastasis risk in patients, in prostate cancer cells.\(^{309}\)

The zebrafish xenograft model has been used to study micrometastasis formation of triple-negative breast cancer (TNBC). High CXCR4 expressing TNBC cells progressively extravasated and invaded the CHT, more than cells displaying low CXCR4 mRNA levels.\(^{283}\) Importantly, CXCR4 sustained tumor metastasis in a Cxcl12-dependent manner, which resembles the situation in human and murine models of breast cancer.

Several other signaling pathways were found to drive breast cancer metastases in the zebrafish CHT (Fig. 6A). Similar experiments demonstrated the importance of SMAD6\(^{310}\) and αv integrin\(^{311}\) for invasion of TNBC. Depletion of αv integrin caused a decrease of SNAIL, SLUG, N-Cadherin, and Vimentin expression, and resulted in a dramatic decrease of invasion and metastases.

At the time patients are diagnosed with MM, various BM-lytic lesions can often be observed.\(^{312}\) In zebrafish, MM cell homing to the CHT increased gene expression in cancer cells related to cytokine and chemokine-mediated signaling (including IL-6), cell adhesion, and angiogenesis,\(^{276}\) signals known from the mammalian BM microenvironment.\(^{313-315}\) Reduced expression of Very Late Antigen 4 (VLA-4), PTK2 protein tyrosine kinase 2 (PTK2 or FAK), and CXCR4 led to impaired homing of MM cells to the CHT,\(^{276}\) demonstrating the similarity of CHT metastasis to processes observed in mammalian BM.\(^{316-318}\)

An advantage of zebrafish xenografts is the possibility to test drug regimes on human tumor material in a semi–high-throughput manner. Tumor cells can also be pretreated with chemical compounds and injected into zebrafish to observe their effect on tumor growth and progression.\(^{283}\) Examples of drugs tested in zebrafish xenotransplants include R-406 inhibiting SYK kinase for retinoblastoma\(^{319}\) and prostate cancer metastasis,\(^{320}\) gomesin and gomesin-like homologue for melanoma,\(^{321}\) and IT1t, a CXCR4 antagonists, for breast cancer metastasis.\(^{283}\) These and other examples demonstrate the power of zebrafish screens to identify promising drugs as potential cancer metastases treatments. Together with the high-resolution in vivo imaging this will further promote zebrafish as a preclinical animal model in cancer research.

**Conclusions**

Zebrafish are increasingly used in the field of skeletal disease and regeneration research because of their ease of genetic manipulation, convenient drug treatment options, and in vivo imaging possibilities. High-throughput genetic and drug screening can be performed in zebrafish larvae, which can also be used as xenograft recipients to study bone metastasis. There are many similarities between mammalian and zebrafish physiology, which allow the investigation of hormonal bone disease in zebrafish. Zebrafish bones regenerate remarkably well, and high plasticity of bone-forming cells might be key to this ability. Understanding the pathogenesis of skeletal alterations and regeneration in zebrafish will help to improve therapeutic approaches in a clinical setting in the future.

**Disclosures**

All authors declare no competing financial interest.

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Kristin Dietrich: Writing-original draft; writing-review & editing. Imke Fiedler: Visualization; writing-original draft; writing-review & editing. Anastasia Kurzyukova: Visualization; writing-original draft; writing-review & editing. Alejandra Lopez Delgado: Visualization; writing-original draft; writing-review & editing. Lucía R. Rodríguez-Delgado: Visualization; writing-original draft; writing-review & editing. Karina Geurtzen: Writing-original draft; writing-review & editing. Chrissy Hammond: Writing-original draft; writing-review & editing. Björn Busse: Writing-original draft; writing-review & editing. Franziska Knopf: Conceptualization; writing-original draft; writing-review & editing.

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