Hematologic Disorders of Fish

Tonya M. Clauss, DVM, MS\textsuperscript{a,*}, Alistair D.M. Dove, PhD\textsuperscript{a}, Jill E. Arnold, MS, MT\textsuperscript{b}
\textsuperscript{a}Veterinary Services and Conservation Medicine, Georgia Aquarium, 225 Baker Street, Atlanta, GA 30313, USA
\textsuperscript{b}National Aquarium in Baltimore, Pier 3, 501 East Pratt Street, Baltimore, MD 21202, USA

Most fish health research and medicine traditionally has focused on aquaculture and food fish species. As society recognizes the need for conserving and protecting its natural resources, public aquarium facilities, commercial ornamental fish producers, collectors, and hobbyists are following the lead of the aquaculture industry by improving fish health practices for popular display fish. The growth of domestic animal medicine as a discipline over the past few decades also has had an impact on fish medicine. Pets, including fish, often are believed members of the family and, as a result, more private practitioners are being consulted about pet fish health. Hematologic data not always have been used in evaluating fish health because of the difficulty of obtaining samples, the challenges involved in evaluating hemograms, and the lack of meaningful reference intervals to aid in interpretation. Hematologic evaluation can be useful in monitoring the health status of fish, as long as interpretation accounts for intrinsic and extrinsic factors that can influence the appearance of cells and the quantitative values obtained. Care must be taken when comparing data as many published reference ranges do not account for differences attributed to factors, such as age, gender, water quality, and season. Even the capture, handling, and anesthetics involved in obtaining blood samples from fish can have profound affects on the hemogram [1].

Hematologic disorders are marked by aberrations in structure or function of the blood cells or the mechanisms of coagulation. Although many other diseases may be reflected by the blood and its constituents, the abnormalities of erythrocytes, leukocytes, thrombocytes, and clotting factors are considered primary blood disorders. Diseases of fish may result in anemia, leukopenia, leukocytosis, thrombocytopenia, and other blood cell abnormalities.

* Corresponding author.
E-mail address: tclauss@georgiaaquarium.org (T.M. Clauss).
This article summarizes some of the salient features of hematologic analysis in teleosts (bony fishes) and elasmobranchs (sharks and rays) and outlines the ways in which the major types of diseases can present themselves in a fish hemogram. Blood sample collection and handling techniques for both classes of fishes are described in the literature [2–5] and are not discussed further in this review.

Normal erythrocytes

Erythrocytes of teleost fishes have similar appearance and ultrastructure to those of other nonmammalian vertebrates. The cells are oval to elliptic in shape with abundant pale, eosinophilic cytoplasm and centrally positioned oval to elliptic nuclei, which is moderately to deeply basophilic (Fig. 1). Elasmobranch erythrocytes are similar in appearance to those of teleosts but considerably larger (Fig. 2). A small amount of erythropoiesis occurs in the peripheral blood of both classes of fish, so it is common to find a small percentage of immature erythrocytes when examining a hemogram from a normal fish [6,7]. Moderate anisocytosis and polychromasia also is normal in many species of teleosts and elasmobranchs. Immature erythrocytes tend to be more rounded than oval with a blue-tinted cytoplasm and larger, more heterochromatic nucleus, thus a higher nucleus to cytoplasm ratio. Erythrocyte morphology, interspecific variation, and variations noted in response to intrinsic and extrinsic variables are summarized in detail in existing literature [4,5].

Techniques for laboratory evaluation of erythrocytes also are well described [4,5,8,9]. Examining morphology, determining the packed cell volume (PCV), and obtaining total erythrocyte counts and red blood cell indices, such as mean cell volume, mean corpuscular hemoglobin concentration, and mean corpuscular hemoglobin, all can be useful in diagnosing disease. PCV varies within and between species and seems to correlate with the normal activity level of the fish. For example, actively swimming species, such as tuna and other pelagic species, tend to have much higher PCVs than would a sedentary bottom dweller, such as a flatfish. Elasmobranchs have lower concentrations of larger cells and usually have lower PCVs. As with other hematologic parameters, PCV values may vary with age, gender, water quality, photoperiod, diet, and season [10–17]. In the absence of established reference ranges, an accepted PCV range for fish is 20% to 45% [17].

Normal leukocytes

Leukocytes of fishes are variable between species (see Figs. 1 and 2), such that initially it can be hard to identify some cell types. The main leukocytes can be identified using a comparative approach and process of elimination on a typical blood smear processed with a three-step Romanowsky-type
Fig. 1. Normal teleost blood cell morphology. (A) Mature erythrocyte morphology of golden trevally, *Gnathanodon speciosus*. (B) Mature erythrocyte morphology of ocean sunfish *Mola mola*, showing comparatively larger cells. (C) Eosinophil of a goatfish *Upeneus* sp showing marginal nucleus and large cigar-shaped eosinophilic granules. (D) Neutrophil of goatfish showing pale cytoplasm with sparse granulation (some basophilic in this case) and eccentric unsegmented nucleus. (E) Thrombocyte of ocean sunfish showing spindle shape typical of many teleost thrombocytes. (F) Monocyte of the arowana *Osteoglossum bichirrosum* showing typical indented nucleus and abundant vacuolated basophilic cytoplasm. (G) Mature lymphocyte (solid arrow) and immature erythrocyte (dashed arrow) of golden trevally. Note that the immature erythrocyte nucleus is essentially similar to that of mature erythrocytes and that, despite the similar basophilic nature, there is comparatively more cytoplasm in the immature erythrocyte. (H) Heterophils of the arowana showing pale basophilic cytoplasm and abundant tiny eosinophilic granules. These cells may fill the same roles as the neutrophils in (D) (see text for further discussion).

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Fig. 2. Normal elasmobranch blood cell morphology. (A) Mature erythrocyte morphology of bonnet head shark, *Sphyrna tiburo*, note throughout the plate that erythrocytes are much larger than those of teleosts shown in Fig. 1. (B) Buffy coat smear from the whale shark, *Rhincodon typus*, showing erythrocytes (E), a lymphocyte (L), heterophil (F), eosinophil (C), neutrophil (N) and thrombocytes (T). (C) Eosinophil or CEG (solid arrow) and two heterophils or FEG (dashed arrows) in the blood of a southern ray, *Dasyatis americana*. The granules of the CEG have vivid eosinophilic properties and are larger and cigar shaped. (D) Lymphocyte (arrow) of southern ray showing sparse basophilic cytoplasm, dense nucleus and blebbing of the plasma membrane. (E) Neutrophil of spotted wobbegong, *Orectolobus maculatus*. (F) Monocyte of the southern ray showing typical indented nucleus and abundant vacuolated basophilic cytoplasm. (G) CEG of the carcharhinid black tip shark *Carcharhinus melanopterus*. This cell type has larger granules similar to the eosinophil or CEG of other shark orders, but the tinctorial properties are more similar to the heterophil or FEG found in other orders. The same applies to carcharhinid FEGs, which display vivid eosinophilia, despite their smaller granules. The homologies of these two cell types with the CEG and FEG of other elasmobranch orders have not been established. (H) Thrombocytes in the blood of the black tip shark. Both granular (white arrow) and agranular (dashed arrows) thrombocytes occur in this species.
hematology stain. Evaluation of ultrastructure and thorough review of fish leukocytes in the literature has alleviated some of the confusion and difficulty associated with hemogram evaluation. Teleost and elasmobranch leukocytes are discussed briefly. In-depth summaries of identification and laboratory evaluation can be found in the literature [2,4,5,8,9,18].

Lymphocytes are the most common and probably most variable leukocyte in most healthy teleosts and elasmobranchs. These are small cells with densely basophilic nuclei and, usually, a very small amount of distinctly blue cytoplasm; the nucleus-to-cytoplasm ratio is high. They sometimes are rounded but often irregular in shape and often with “blebs” or blister-like outpocketings of the outer membrane. Reactive lymphocytes are larger and plasma cells do occur and can appear 2 to 3 times larger than regular lymphocytes [9].

The most common granulocyte of teleosts is the neutrophil. These typically are rounded cells that have nuclei that may or may not be segmented and their cytoplasm varies from extremely pale to gray or slightly blue, depending on the species [19]. The nucleus-cytoplasm ratio is low and cells tend to be similar to or larger than erythrocytes, rarely if ever smaller. Granulation also varies from indistinct to obvious, with the granules themselves appearing anywhere from glassy to slightly eosinophilic or basophilic. Some cell populations with more prominent eosinophilic granules can resemble heterophils of birds and reptiles or the heterophil or fine eosinophilic granulocyte (FEG) of elasmobranchs. Campbell [4] and Campbell and Ellis [5], however, conclude that there are enough functional studies to show that these cells fill the neutrophil role in the teleost leukocyte panoply. Although immature neutrophils occur in the circulating blood of teleosts, they do not resemble the classic band cell of the mammalian leukogram. In typical neutrophils and heterophils, the nucleus may be segmented or nonsegmented in mature cells depending on species and, sometimes, within a species. Neutrophils also are found commonly in elasmobranch species. The nuclei can be lobed or round. The cytoplasm is colorless with no visible granulation [8,9].

True eosinophils are less common in teleosts whereas basophils are rare. The granules of eosinophils tend to be larger and more distinct than those of the neutrophil/heterophil series, round or rod-shaped, and prominently eosinophilic; they are not easily overlooked. Basophils, when they occur, contain granules that stain so darkly as to obscure the nucleus [4]. In contrast, elasmobranchs show abundant eosinophilic cells, generally falling into two morphologic types. Consistent in sharks and rays is a cell that closely resembles the avian heterophils, with elongated, rod-shaped granules that stain a reddish color in Romanowsky-type stains. Nomenclature varies in the literature; this article uses the terminology described by Hine and Wain [20,21] as FEG. The second cell type, termed coarse eosinophilic granulocyte (CEG), is of equal size, but the granules are round and usually less abundant in the cytoplasm. The staining properties vary with species; most shark and ray CEGs stain orange to bright orange whereas the carhcharinid
CEG granules stain pale pink. The nuclei can be lobed or round in the FEG and CEG cells. Basophils rarely, if ever, are present in shark species but commonly found in rays. Elasmobranch leukocyte function warrants further study.

Monocytes of teleosts and elasmobranchs are similar to those of other vertebrates; they probably are the largest cells in any differential, irregular in shape, with an eccentric, large, heterochromatic nucleus (often indented or kidney shaped) and abundant slightly basophilic cytoplasm replete with vacuoles and other membrane-bound organelles.

Abnormalities of erythrocytes and leukocytes

Hematologic abnormalities involving the erythrocytes of fish include polycythemia, anemia, abnormal morphology, and nuclear or cytoplasmic inclusions. Like other vertebrates, fish leukograms are affected by disease, inflammatory processes, stress, nutrition, and physiologic and environmental factors. During disease processes, the leukogram of teleosts can show many of the patterns that are seen in mammals, although Campbell and Ellis [5] stress that the exact functions of piscine granulocytes are not established, so direct comparisons with mammalian processes are somewhat speculative.

Polycythemia and anemia

Fish with a PCV of 45% or greater generally are considered to have a relative polycythemia resulting from dehydration, especially when in conjunction with an elevation in serum osmolality, total protein, sodium, or chloride. Polycythemia also can be observed, however, in sexually mature males, in freshwater fish exposed to hypoxia, in stressed fish resulting from release of catecholamines, during splenic contraction, and with erythrocyte swelling [6,22,23].

Anemias are well documented in fish. PCVs less than 20% in teleosts usually are associated with anemia. Diagnosing anemias in elasmobranchs can be more challenging as some species have a normal PCV of 20% or less (for example, sandbar shark [Carcharhinus plumbeus] and the Port Jackson shark [Heterodontus portusjacksoni]) [2,9]. The PCV of some sharks varies between blood collection sites and this possibility should be taken into account when diagnosing anemia [24]. There are three primary types of anemia: hemorrhagic (blood loss), hemolytic (erythrocyte destruction), and hypoplastic (poor erythropoiesis). The basic descriptive terminology used for anemias in other animals applies equally to fishes and may refer to cell size (microcytic, normocytic, or macrocytic), hemoglobin concentration (hypochromic or normochromic), cell loss (hemolytic or hemorrhagic), and hemopoietic status (regenerative or nonregenerative). Causes of nonregenerative anemias include inflammatory disease, nutritional disorders, toxins, and renal or splenic disease with disruption or destruction of hematopoietic tissues.
The specific hematologic presentation of a given anemia often provides indications as to the etiology of the disease. Marked hemorrhage or hemolysis often results in microcytic anemia because regenerating immature erythrocytes make up the majority of cells in peripheral circulation and are smaller in size than mature erythrocytes. Marked polychromasia also may be noted. The presence of immature erythrocytes in circulation does not always signify a regenerative anemia; immature cells may be present in response to environmental stressors, such as hypoxia, toxins, or temperature change [18,25–27]. Hemorrhagic anemia in fish may be associated with trauma, cutaneous ulceration, parasitism (lampreys, leeches, and isopods) [5,28], nutritional deficiencies (vitamin K and B, inositol, and choline) [5,18], and viral or bacterial septicemia (Tables 1 and 2). When severe or chronic blood loss occurs, the net loss of iron may result in iron deficiency anemia. Hemolytic anemia in fish may be associated with hemolysin-producing bacteria (see Table 1), environmental toxins [29–31], viral infection [32], select nutritional deficiencies [33–35], and hemoparasites [3–5,36,37]. Marked regeneration often is noted with hemolytic anemia because the iron is not lost from the body and can be incorporated back into hemoglobin for hematopoiesis. Erythrocytes with pyknotic nuclei, erythroplastids, and erythrocyte fragmentation are associated with conditions that interfere with splenic removal of senescent erythrocytes from peripheral circulation [38]. Table 1 provides an overview of various causes and clinical presentations of anemia in teleosts.

Intrinsic and extrinsic induced variability

Stress factors in fish are as variable as they are with other animals. Extrinsic factors, such as handling, transport, poor water quality, and high population densities, initially may cause a significant stress response in most fish but with chronicity or acclimation the response may diminish. Stress response in fish is similar to that of higher vertebrates with a rapid release of catecholamines followed by the release of corticosteroids. A stress leukogram for most fish is manifested as a leukopenia with a lymphopenia and a relative granulocytosis. Hematologic changes associated with the stress response may persist for several days after the stressor is removed [39,40]. Juvenile teleosts have notably higher lymphocyte and total leukocyte counts compared with adults [15,22]. Noninfectious diseases also can manifest in the blood of fishes; these often are associated with husbandry practices and environment. Among husbandry-related diseases, a microcytic normochromic anemia can result from environmental stressors, such as increases in population density [41,42]. Higher lymphocyte and total leukocyte counts are found in fish from production systems with high densities, marginal water quality, and often elevated bacterial load [17]. Nitrite poisoning, also known as brown blood
Table 1
Some anemias and their etiologies in teleost fishes

<table>
<thead>
<tr>
<th>Etiology</th>
<th>Presentation</th>
<th>Citation</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Hemorrhagic</strong></td>
<td></td>
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<tr>
<td><strong>Bacterial</strong></td>
<td></td>
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<tr>
<td><em>Aeromonas</em> sp</td>
<td>External/internal lesions, septicemia</td>
<td>Campbell and Ellis [5]</td>
</tr>
<tr>
<td><em>Pseudomonas</em> sp</td>
<td>External/internal lesions, septicemia</td>
<td>Campbell and Ellis [5]</td>
</tr>
<tr>
<td><em>Yersinia</em> sp</td>
<td>Reticulocytosis, septicemia</td>
<td>Tobback et al [52]</td>
</tr>
<tr>
<td>Ammonia toxicity</td>
<td>Microcytic normo or hypochromic</td>
<td>Groff and Zinkl [18]</td>
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<tr>
<td><strong>Nutritional deficiency</strong></td>
<td></td>
<td></td>
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<tr>
<td>Vitamin K</td>
<td>Hypocoagulation, hypochromic</td>
<td>Groff and Zinkl [18]</td>
</tr>
<tr>
<td>Vitamin B</td>
<td>Hypocoagulation, hypochromic</td>
<td>Groff and Zinkl [18]</td>
</tr>
<tr>
<td>Inositol</td>
<td>Hypocoagulation, hypochromic</td>
<td>Groff and Zinkl [18]</td>
</tr>
<tr>
<td>Choline</td>
<td>Hypocoagulation, hypochromic</td>
<td>Groff and Zinkl [18]</td>
</tr>
<tr>
<td><strong>Parasites</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leech, lamprey, isopod</td>
<td>External lesions, pallor, macrocytic</td>
<td>Snderman [66], Nair and Nair [28]</td>
</tr>
<tr>
<td><strong>Iron deficiency</strong></td>
<td></td>
<td>Groff and Zinkl [18]</td>
</tr>
<tr>
<td>Viral (see Table 2)</td>
<td></td>
<td></td>
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<tr>
<td><strong>Hemolytic</strong></td>
<td></td>
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<tr>
<td><strong>Bacterial</strong></td>
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<tr>
<td><em>Flavobacterium columnare</em></td>
<td>Fragmentation, macrocytic hypochromic</td>
<td>Rehulka and Minarik [54]</td>
</tr>
<tr>
<td><em>Aeromonas</em> sp</td>
<td>Macrocytic, ± bacteremia, septicemia</td>
<td>Roberts and Ellis [49]</td>
</tr>
<tr>
<td><em>Pseudomonas</em> sp</td>
<td>Macrocytic, ± bacteremia, septicemia</td>
<td>Roberts and Ellis [49]</td>
</tr>
<tr>
<td><em>Vibrio</em> sp</td>
<td>Macrocytic, ± bacteremia, septicemia</td>
<td>Roberts and Ellis [49]</td>
</tr>
<tr>
<td><strong>Toxins</strong></td>
<td></td>
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<tr>
<td>Nitrite</td>
<td>Methemoglobinemia, hypochromic</td>
<td>Avilez et al [29]</td>
</tr>
<tr>
<td>Mercury contamination</td>
<td>Macrocytic normo or hypochromic</td>
<td>Elahee and Bhagwant [31]</td>
</tr>
<tr>
<td>Chlorine</td>
<td>Heinz bodies, methemoglobinemia</td>
<td>Buckley [30]</td>
</tr>
<tr>
<td><strong>Nutritional</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Folate deficiency</td>
<td>Pyknotic nuclei, erythroplastids, fragmentation</td>
<td>Plumb et al [34], Ferguson [33]</td>
</tr>
<tr>
<td>Vitamin E deficiency</td>
<td>Pyknotic nuclei, erythroplastids, fragmentation</td>
<td>Hibiya [45], Eiras [44]</td>
</tr>
<tr>
<td>Yeast excess</td>
<td>Poikilocytosis, microcytic hypochromic</td>
<td>Sanchez-Muiz et al [46]</td>
</tr>
<tr>
<td><strong>Hemoparasites</strong></td>
<td></td>
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<tr>
<td>Trypanoplasma</td>
<td>Microcytic hypochromic</td>
<td>Woo [37], Rowley [48]</td>
</tr>
<tr>
<td>(cryptobiosis, sleeping sickness)</td>
<td></td>
<td></td>
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<tr>
<td>Trypanosoma</td>
<td>Microcytic hypochromic</td>
<td>Woo [36]</td>
</tr>
<tr>
<td>Piroplasmds</td>
<td>Intracytoplasmic</td>
<td>Campbell and Ellis [5]</td>
</tr>
<tr>
<td>Hypoxia</td>
<td>Macrocytic normo or hypochromic</td>
<td>Elahee and Bhagwant [31]</td>
</tr>
<tr>
<td>Viral (see Table 2)</td>
<td></td>
<td></td>
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<tr>
<td><strong>Hypoplastic</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yersiniosis (chronic)</td>
<td>Septicemia, kidney/splenic necrosis</td>
<td>Tobback et al [52]</td>
</tr>
<tr>
<td><strong>Toxins</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ammonia</td>
<td>Cell fragility, impaired erythropoiesis</td>
<td>Groff and Zinkl [18]</td>
</tr>
<tr>
<td>Heavy metals</td>
<td>Impaired erythropoiesis, low hemoglobin</td>
<td>Groff and Zinkl [18], Noga [3]</td>
</tr>
</tbody>
</table>

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disease or new tank syndrome, occurs because of incomplete nitrification of ammonium waste in closed or high-density culture situations and can result in cyanosis and hemolytic anemia [5,29]. Ammonia and heavy metal toxicoses can result in various forms of nonregenerative anemia [3,18]. Chronic exposure to cypermethrin, a synthetic pyrethroid insecticide, may cause enlargement of erythrocytes and abnormal erythrocyte morphology yielding a macrocytic anemia [43].

Nutritional disorders, such as folic acid and vitamin E deficiency or toxicosis from rancid oils and environmental pollutants, may contribute to formation of abnormal erythrocyte nuclei and erythroplastids [44,45]. Deficiencies in dietary vitamin K and B vitamins, inositol, and choline, may cause coagulation disorders resulting in hemorrhagic anemia [18]. Folate deficiency in channel catfish (Ictalurus punctatus) results in chronic hemolytic anemia [34,35]. Vitamin C, iron, or copper deficiency can cause

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### Table 1 (continued)

<table>
<thead>
<tr>
<th>Etiology</th>
<th>Presentation</th>
<th>Citation</th>
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<tr>
<td>Nutritional Starvation</td>
<td>Microcytic hypochromic, abnormal nuclei</td>
<td>Rios et al [47]</td>
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<td>Vitamin B12 deficiency</td>
<td>Normocytic hypo or normochromic</td>
<td>Ferguson [33]</td>
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<td>Parasites</td>
<td>Microcytic hypochromic, impaired production (Cryptobiosis, sleeping sickness)</td>
<td>Woo [37], Rowley [48]</td>
</tr>
<tr>
<td>Trypanosoma</td>
<td>Microcytic hypochromic, impaired production</td>
<td>Woo [36]</td>
</tr>
<tr>
<td>Myxozoans</td>
<td>Normocytic hypochromic, poikilocytosis</td>
<td>Hoffmann and Lommel [73]</td>
</tr>
<tr>
<td>Kidney or splenic disease</td>
<td>Erythropoietic tissue destruction/ displacement</td>
<td>Campbell and Ellis [5]</td>
</tr>
</tbody>
</table>

### Table 2

Hematologic features of some viral diseases of teleost fishes

<table>
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<tr>
<th>Disease</th>
<th>Host</th>
<th>Presentation</th>
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<tr>
<td>IHNV</td>
<td>Salmonids</td>
<td>Hemorrhagic, hypochromic, nonregenerative anemia; loss of acid-base equilibrium</td>
<td>Amend and Smith [58,59]</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Egusa [62]</td>
</tr>
<tr>
<td>VHSV</td>
<td>Many species</td>
<td>Severe hemorrhagic and hemolytic anemia, relative monocytesis, lymphopenia, acute mortality</td>
<td></td>
</tr>
<tr>
<td>SVC</td>
<td>Cyprinids</td>
<td>Hemorrhagic hypochromic anemia, neutrophilia, monocytesis.</td>
<td>Egusa [62]</td>
</tr>
<tr>
<td>VEN</td>
<td>Marine species</td>
<td>Hemolytic anemia with prominent intraerythrocytic inclusions, typically a chronic progression</td>
<td>Hershberger et al [32]</td>
</tr>
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</table>
microcytic hypochromic anemia [18]. Rainbow trout fed diets containing certain peroxide-producing yeasts may develop microcytic hypochromic anemia with marked poikilocytosis [46]. Vitamin C and pyridoxine deficiencies may cause leukopenia, and biotin deficiency in cyprinids may result in leukocytosis [18]. Studies of a neotropical species, *Hoplias malabaricus*, revealed that food deprivation for extended periods may result in reversible leukopenia and thrombocytopenia [47].

**Inflammatory processes**

Granulocytosis often is associated with inflammation in teleost and cartilaginous fishes. Monocytes occur in low numbers in a differential white blood cell count but are actively phagocytic cells in piscine fishes [48]. A monocytosis is suggestive of an inflammatory response in teleost fishes. Neutrophilia often is associated with inflammatory processes, but neutrophils are not always phagocytic. An acute neutropenia may reflect extravasion of neutrophils at a focal site of significant insult. Some reports indicate that fish eosinophils are involved in inflammatory responses but have limited phagocytic capability [48, 49].

**Cell-mediated abnormalities**

Lymphocytosis may indicate a leukemic condition and typically are accompanied by cellular signs of degeneration, immaturity, or malignancy. Lymphocytes play an important role in humoral and cell-mediated immunity in fish [50]. Lymphocytosis may be suggestive of immunogenic stimulation, whereas lymphopenia may be suggestive of immunosuppressive conditions.

**Infection-induced abnormalities**

**Bacterial disease**

Gram-negative bacteria, such as *Aeromonas* spp and *Pseudomonas* spp, are common causes of septicemia and hemorrhagic anemia in freshwater fish [18]. *Aeromonas* spp, *Pseudomonas* spp, and *Vibrio anguillarum* produce hemolysins and are the most common causes of hemolytic anemia in fish [18, 49]. Yersiniosis of marine fish causes hemorrhagic septicemia characterized by anemia, leukocytosis, and reticulocytosis [51, 52]. A severe hemorrhagic anemia caused by cold-water vibriosis (Hitra disease) was documented in juvenile Atlantic salmon (*Salmo salar*) [53]. Brook trout (*Salvelinus fontinalis*) affected by columnaris disease (*Flavobacterium columnare*) developed macrocytic hypochromic anemia with evidence of fragmented erythrocytes noted on blood smears [54]. Fig. 3 shows a gram-positive septicemia caused by *Streptococcus iniae* in a yellow tang (*Zebrasoma flavescens*). In sharks, increases in granulocytes and decreases in lymphocytes can be associated with bacterial septicemia. Leukopenia with lymphopenia or a relative neutrophilia is
indicative of a stress response or sepsis [4,5]. Proliferative kidney disease (PKD)-affected fish are believed particularly sensitive to bacterial challenge because of the depression of granulocyte activity [55].

Viral disease

Of the myriad viral infections known from fishes, several are expressed prominently in blood, most often through the presence of intraerythrocytic inclusions or profound hemorrhagic or hemolytic anemia. Erythrocytic necrosis virus (VEN) causes hemolytic anemia and is associated with intracytoplasmic inclusions and nuclear changes in the erythrocyte of marine fish [32,56]. Erythrocytic inclusion body syndrome virus also results in intracytoplasmic inclusions but is distinct from VEN. It is associated with transient anemia in freshwater Pacific salmon in the United States but has had no adverse effects on other infected species [57]. Infectious hematopoietic necrosis virus (IHNV) infects a range of salmonid and other fish species, wherein it obliterates the lymphoid tissues of the spleen and head kidney [58,59]. Viral hemorrhagic septicemia (VHSV) has become a prominent emerging disease in the North American fisheries scene since an apparent epizootic started in the Great Lakes region in 2005 [60,61]. Spring viremia of carp (SVC) is a disease restricted to freshwater cyprinid fishes and like VHSV, it causes hemorrhagic anemia [62]. Channel catfish virus (Herpesvirus ictaluri) causes acute hemorrhage at the bases of the fins and in liver, kidney, and the gastrointestinal system [63,64]. Table 2 provides an overview of the clinical presentations associated with the viral diseases discussed.
Parasitic disease

Several parasites, mostly protistan, manifest in the blood of teleosts; these may be intraerythrocytic or occur in the plasma. *Trypanosoma* probably is the best-known extraerythrocytic protist in fish (Fig. 4). Fish trypanosomes essentially have the same appearance as those that cause sleeping sickness in humans (*T. brucei*) and trypanosomiasis in domestic animals, although they usually are larger (example, *T. mukasai* large form > 80 μm) [65] and occur at much heavier intensities than do mammalian kinetoplastids. Despite their higher parasitemia, they are infrequently pathogenic but are capable of causing fatal anemias [36]. Fish trypanoplasms are very similar in morphology to trypanosomes and are extraerythrocytic protists. *Cryptobia (Trypanoplasma) salmositica* [37] causes cryptobiosis in salmonids and may result in severe anemia because of destruction of hematopoietic tissue. *Trypanoplasma borreli* causes sleeping sickness in cyprinids with systemic illness and progressive anemia [48]. Species of trypanosomes and trypanoplasms with known life cycles are transmitted by leeches rather than arthropods.

The intraerythrocytic hemogregarines probably are the most common hemoparasites of teleosts [66,67]. Among elasmobranchs, they have been found in spiny dogfish *Squalus acanthias* [68], epaulette sharks (*Hemiscyllium ocellatum*) [69], and cat sharks (Fig. 5). Possible vectors are leeches and isopods. These usually occur as small to larger cellular inclusions in the cytoplasm of erythrocytes (see Diniz and colleagues for example [70]),

![Fig. 4. Protistan disease in a blood smear: trypanosmiasis caused by an undescribed *Trypanosoma* species, in straight-backed freshwater catfish, *Neosilurus hyrtilii* from Australia. These are typical trypanosomes in possessing a recurrent flagellum that originates from a darkly staining kinetoplastid organelle (solid arrow) and attached to the cell by an undulating membrane (dashed arrow).](image-url)
with or without a parasitophorous vacuole (a membrane-bound space surrounding an intracytoplasmic parasite and preventing direct contact with the cytoplasm) and with a distinct basophilic nucleus. They may appear similar to the large hemogregarines of birds and reptiles, such as *Haemoproteus*, occupying more than half of the cytoplasmic space and curving around the erythrocyte nucleus, or they may resemble the smaller, more basophilic piroplasms of mammals, such as *Babesia* spp, occurring singly or as pairs or tetrads, depending on the particular genus of parasites (see van der Straaten and colleagues for example [67]). Intraerythrocytic protist parasites of fishes (and amphibians) have been comprehensively reviewed by Davies and Johnston [71].

Many metazoan parasites cause hematologic changes, most commonly anemia, due to the consumption of host blood as part of the normal feeding biology of the parasite. They also may cause anemia by disruption of hematopoiesis. Elevated eosinophil counts are suggestive of an inflammatory response associated with antigenic stimulation or parasitic infections, such as metazoans. Leeches and lampreys are blood-sucking parasites that are large compared with most parasites and, therefore, can cause profound hemorrhagic anemia in fish. A study with captive blackeye thicklip (Labridae) showed that parasitic isopods (*Gnathia* sp) can significantly reduce hematocrit [72] as a result of the “tick-like” biology of female gnathiids, wherein they take periodic large blood meals before leaving the host to mate and molt in the substrate. Rainbow trout (*Salmo gairdneri*) affected by PKD, caused by the myxozoan parasite *Tetracapsuloides bryosalmonae*, develop a chronic normocytic hypochromic anemia with distinct poikilocytosis [73].

Fig. 5. Protistan disease in a blood smear: two hemogregarine morphotypes in the blood of the catshark, *Holohalaelurus punctatus*. One type (solid arrow) shows a fairly typical hemogregarine presentation, displacing the nucleus and with a distinct nucleus, whereas the other (dashed arrow) shows an unusual frothy appearance with the nucleus displaced to one pole. (Courtesy of Dr. Nico Smit, Department of Zoology, University of Johannesburg, South Africa.)
Thrombocytes and hemostasis

Thrombocytes play a large role in mediating the clotting response. They typically are small cells, ovoid, oblong, or spindle shaped (often all three in any given smear), with clear cytoplasm and a condensed basophilic nucleus. Some elasmobranch species have a second population of thrombocytes; in addition to the cell identical to that described for teleosts, there are thrombocytes with abundant FEGs in the cytoplasm. The clinical significance of these cells is not yet understood. Fish seem to rely primarily on extrinsic pathways of coagulation. Clot formation usually occurs within 5 minutes in teleosts whereas it may take greater than 20 minutes for some elasmobranchs to clot. Thrombocytopenia could have devastating affects on fish as not only are these cells responsible for blood clotting but also they are responsible for controlling fluid loss from surface wounds in fish. High levels of glucocorticoids decrease the number of thrombocytes and increase clotting times. Prolonged clotting times also is attributed to vitamin K deficiency [5].

Summary

Hematology of fishes lags behind that of other classes of vertebrates, but analysis of blood still can be informative about disease processes in teleosts and elasmobranchs. Although robust interpretation of fish hemograms often is hampered by a lack of reference values, this knowledge deficit represents an opportunity for expansion of clinical pathology studies among fishes. This article offers a summary of some of the more common hematologic abnormalities documented in fish and insight to some of the more obscure causes of hematologic variations and abnormalities that warrant further investigation and documentation. Practitioners are encouraged to obtain samples whenever possible as they can benefit individual animals and contribute to establishment of reference data.

References


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