The fossil record of ‘early’ tetrapods: evidence of a major evolutionary transition?

Paul Garner

According to evolutionary theory, the origin of tetrapods from a fish-like ancestor during the Devonian Period was one of the major events in the history of life on Earth. The ‘drying pond’ hypothesis was proposed to explain the selection pressures behind the transition. According to this hypothesis, the tetrapods evolved as fishes became progressively better adapted to terrestrial conditions during prolonged episodes of drought. Recently, however, the assumption that feet and legs evolved to facilitate life on the land has been called into question. The ‘earliest’ known tetrapods with feet and legs are now thought to have been aquatic animals; evolutionists therefore argue that feet and legs evolved in a shallow water environment and were only later co-opted for use on the land. This paper reviews the radical changes in thinking about the fish-tetrapod transition that have taken place in the evolutionary community. It also considers the chimeromorphic nature of Devonian tetrapods and fishes, and offers some critical comments on the evolutionary interpretation of their fossil record.

Evolutionists believe that tetrapods—i.e. vertebrates with four limbs—were the first animals to move on to the land, having evolved from a fish ancestor during the Devonian period (conventionally 408 to 360 million years ago). The fossil record of Devonian tetrapods is often presented as compelling evidence of this major evolutionary transition. Science writer Carl Zimmer has written a popular book, *At the Water’s Edge*, which purports to show how life came ashore (i.e. how fish evolved into tetrapods) and then went back to the sea (i.e. how land mammals gave rise to the whales). A more technical presentation was written recently by Jenny Clack, Reader in Vertebrate Palaeontology and Senior Assistant Curator of the University Museum of Zoology, Cambridge. Entitled *Gaining Ground: The Origin and Evolution of Tetrapods*, it begins with these words:

‘About 370 million years ago, something strange and significant happened on Earth. That time, part of an interval of Earth’s history called the Devonian Period by scientists such as geologists and paleontologists, is known popularly as the Age of Fishes. After about 200 million years of earlier evolution, the vertebrates—animals with backbones—had produced an explosion of fishlike animals that lived in the lakes, rivers, lagoons, and estuaries of the time. The strange thing that happened during the later parts of the Devonian period is that some of these fishlike animals evolved limbs with digits—fingers and toes. Over the ensuing 350 million years or so, these so-called tetrapods gradually evolved from their aquatic ancestry into walking terrestrial vertebrates, and these have dominated the land since their own explosive radiation allowed them to colonize and exploit the land and its opportunities. The tetrapods, with their limbs, fingers, and toes, include humans, so this distant Devonian event is profoundly significant for humans as well as for the planet.’

Indeed, according to the cladistic framework that now dominates evolutionary systematics, humans are not simply descended from fish—they are fish! Clack states: ‘Although humans do not usually think of themselves as fishes, they nonetheless share several fundamental characters that unite them inextricably with their relatives among the fishes … Tetrapods did not evolve from sarcopterygians [lobe-finned fishes]; they are sarcopterygians, just as one would not say that humans evolved from mammals; they are mammals.’

In this paper I will critically examine the fossil record of ‘early’ tetrapods and discuss the way in which older evolutionary views of their origin have been overturned in the last two decades. I will also consider the mosaic distribution of characters that we observe in Devonian tetrapods and fishes, the problems that it poses for evolutionary theory, and how it might be understood in a creationist framework.

**The ‘drying pond’ hypothesis**

Many evolutionary scenarios have been proposed to explain the origin of tetrapods. Most of them were developed to answer the question, ‘Why did fish leave the water and come onto the land?’ The early theories usually focused on the environmental setting and selection pressures behind the transition. Tetrapods were thought to have evolved during the Devonian, a period associated in many parts of the world with sediments stained red by iron oxide. Classic red beds, such as the Siluro-Devonian rocks of Europe (the Old Red Sandstone) and their North American equivalents (the Catskill and Escuminac formations), have often been
interpreted as the product of hot, semi-desert environments with seasonal wetness. This led many to speculate that an increasingly arid climate was a major influence on the evolution of air-breathing vertebrates. A classic paper by Barrell set the scene for much future discussion. He argued that the first tetrapods arose ‘under the compulsion of seasonal dryness’. Under such conditions, it was suggested, the air-bladder of certain fishes became progressively better adapted as an organ of respiration and the gills atrophied. The development of a new system of breathing allowed fishes to survive the drought conditions by moving between bodies of water. Those fishes with more limb-like appendages were better able to make the journey and this ultimately led to the evolution of limbs with digits. This became known as ‘the drying pond hypothesis’ and was popularized by the great vertebrate palaeontologist Alfred Sherwood Romer.

**‘Early’ tetrapods from East Greenland**

When Romer was popularizing the ‘drying pond’ idea, the earliest known tetrapods were *Ichthyostega* and *Acanthostega* from the Upper Devonian of East Greenland. *Ichthyostega* was first described by Säve-Söderbergh and then by Jarvik in a series of papers and a monograph. Although the anatomy of *Ichthyostega* is known in considerable detail, its body proportions are uncertain because the fossil material comes from more than one individual. *Ichthyostega* is about one metre long with a broad, flat head, short, barrel-shaped

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**Table 1. ‘Early’ tetrapods and so-called ‘near tetrapods’.** Most are represented by single specimens; *Acanthostega* is unique in that it represents a stratigraphic range. Givetian is a subdivision of the Middle Devonian, Frasnian and Famennian are subdivisions of the Upper Devonian, and Tournaisian is a subdivision of the Lower Carboniferous.

<table>
<thead>
<tr>
<th>Taxon</th>
<th>Stratigraphic unit</th>
<th>Age</th>
<th>Location</th>
<th>Material</th>
<th>Reference(s)</th>
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<td>Tournaisian</td>
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<td><em>Sinostega</em></td>
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<td>Ketleri Fm</td>
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<td>Latvia</td>
<td>Skull fragments, girdle fragments</td>
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<td><em>Acanthostega</em></td>
<td>Britta Dal Fm</td>
<td>Famennian</td>
<td>East Greenland</td>
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<td>Aina Dal Fm</td>
<td>Famennian</td>
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<td>Famennian</td>
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<td>Cloghnian Shale</td>
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<td>Scat Craig Beds</td>
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<td><em>Livoniana</em></td>
<td>Gauja Fm</td>
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<td>Latvia</td>
<td>Lower jaw fragments</td>
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**Figure 1.** Reconstruction of *Ichthyostega*, showing skull, vertebral column, and limbs, and its hind limb based on a specimen collected in 1987. Note the seven digits on the hind limb (from Clack).15
body, stocky legs, large pelvic and pectoral girdles, and a rib cage with broad, overlapping ribs (Figure 1). It is very evidently a tetrapod, with limbs rather than fins. Nevertheless, *Ichthyostega* has some fish-like characteristics, including a lateral line system and a tail with bony fin rays. Early reconstructions portrayed *Ichthyostega* as a semi-aquatic creature but most later ones depicted it as a predominantly terrestrial animal (e.g. Jarvik). As recently as 1988, a major vertebrate palaeontology text described *Ichthyostega* as a fairly typical land animal with the usual complement of five digits on the hind limb. The second Devonian tetrapod from East Greenland was *Acanthostega*. For many years this animal was known only from two partial skull roofs, but these were enough to mark it out as different from *Ichthyostega*.

**The search for evolutionary ancestors**

Evolutionists sought the ancestry of the tetrapods among the lobe-finned fishes. Although the lobe-fins are dominant in the fossil fish faunas of the Palaeozoic (conventionally 590 to 248 million years ago), they are represented today by only four surviving genera (the coelacanth *Latimeria* and three genera of lungfish). In 1892, Cope and others argued that tetrapods had evolved from the crossopterygians, the group of lobe-fins that includes the coelacanths. Various crossopterygians were proposed as the ‘model ancestor’, including *Sauripterus* and *Osteolepis*. However, most attention settled upon *Eusthenopteron*, from Escuminac Bay in Quebec, Canada. This is the fish that was commonly illustrated, in popular books on fossils, as hauling itself up onto Devonian riverbanks (e.g. Owen).

Nevertheless, there was evidently a substantial discontinuity in the fossil record between terrestrial vertebrates like *Ichthyostega* and their presumed ancestors. This was reflected in creationist treatments of the problem and acknowledged by evolutionists, such as Carroll who wrote:

‘We have not found any fossils that are intermediate between such clearly terrestrial animals and the strictly aquatic rhipidists described in the previous chapter.’

**Aquatic tetrapods challenge the ‘drying pond’ hypothesis**

Since 1990 our knowledge of ‘early’ tetrapods has been greatly expanded, with many new taxa described. Fossil material is now known from Scotland, Greenland, Latvia, the USA, Australia, Russia, and China (Table 1). Furthermore, our understanding of the Greenland tetrapods has been revolutionized by the discovery of new material. As a consequence, a major re-evaluation of tetrapod origins has taken place, and almost every aspect of the ‘drying pond’

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**Figure 2.** *Acanthostega* in a swimming posture (from Clack).
Devonian tetrapods has led modern-day evolutionists to reject this assumption. They now argue that the key tetrapod characters evolved for a shallow-water existence and were only later co-opted for terrestrial use. The new generation of Darwinists dismisses the ‘drying pond’ hypothesis as untestable story-telling, and increasingly relies on cladistics as an alternative framework for understanding the transition. The cladistic approach to the fish-tetrapod transition focuses on determining the sequence of acquisition of key tetrapod characteristics, from which inferences are drawn about the nature of the transition.53 We should recognize, however, that the cladistic methodology is inherently Darwinian and assumes from the outset the continuity of life. By its very nature, cladistics is insensitive to the discontinuities which creationists believe characterize living things.54

Other problems with the ‘drying pond’ hypothesis

The drying pond hypothesis has other problems.55 For instance, it is recognized that red beds are not necessarily indicators of arid climates:

‘The red bed problem has been extremely controversial, with marked differences of opinion, possibly due to the fact that the term “red bed” is a catchall for many sedimentary types produced under different conditions, the only common feature of them being the red color.’56

Modern red beds develop in the oxidizing conditions of the low latitude tropics (e.g. the Amazon Basin). Such environments are characterized by monsoonal rainfall, not arid conditions. Another problem is that, even if the red beds were laid down under conditions of semi-aridity, evolutionists cannot assume that the tetrapods arose in such environments, for the simple reason that many Devonian sediments are not red beds. Some are interpreted as river, lake, or near-shore sediments rich in organic matter, suggesting nearby forests.57

Furthermore, a survey of modern fishes that leave the water to spend time on land affords no support for the ‘drying pond’ hypothesis. There is no association between those that leave the water and those that possess digit-like fins. For example, eels undertake long journeys overland but they have nothing that could be described as digit-like appendages. Indeed, most of the fishes that possess digit-like structures are deep water species or habitual bottom dwellers, such as the Sargassum frogfish.

New views on tetrapod ancestry

There have also been changes of opinion about which group of fishes is closest to the ancestry of tetrapods. Eusthenopteron is no longer regarded as the model ancestor. Depictions showing this fish emerging onto dry land owed more to evolutionary presuppositions than evidence. Eusthenopteron was a rather undistinguished fish with no obvious adaptations to terrestrial life; tetrapod-like behaviour was attributed to it simply because there was no better candidate to fill the role of tetrapod ancestor. The true lifestyle of Eusthenopteron seems to have been that of a lurking aquatic predator, somewhat similar to the modern pike (Esox).

Attention is now focused on the formerly more obscure lobe-finned fishes, Panderichthys and Elpistostege. Until recently, these two genera were united in a family called the panderichthyids, but evolutionists now believe that they are not uniquely related to each other.59 Fossil material from Latvia and Canada shows that these fish were more tetrapod-like than other lobe-fins. Indeed, based on a partial skull roof, Elpistostege was originally described as a tetrapod.60 Although there has been dissent,6162 these genera are increasingly regarded by evolutionists as the closest known relatives of tetrapods.6365 The latest work by Ahlberg et al.43 indicates that Elpistostege is even more tetrapod-like than Panderichthys. These fish have crocodile-like skulls with dorsally placed eyes, straight tails, and slightly flattened bodies without dorsal or anal fins (see Figure 4). Like tetrapods, but unlike all other fishes, they also have frontal bones in the skull roof. Like Eusthenopteron, they seem designed for life as shallow-water predators.

Chimeromorphs pose problems for evolutionary theory

Creationists and evolutionists have observed that many organisms, both fossil and living, exhibit a mosaic distribution of character traits. Parker put it this way:

‘Each created kind is a unique combination of traits that are individually shared with members of other groups.’

Stephen Jay Gould called such organisms ‘mosaic forms’ or ‘chimeras’ while Kurt Wise calls them chimeromorphs. The duck-billed platypus (Ornithorhynchus anatinus), for instance, has features of both mammals (hair, milk production) and reptiles (egg-laying). Perhaps the best-
known fossil example is *Archaeopteryx*, which combines feathers with teeth and wing claws. In fact, a mosaic pattern of character distribution is seen in many other fossil organisms. For instance, Woodmorappe recently drew attention to the chimeric nature of the pakicetids, a group of terrestrial artiodactyls with a whale-like inner ear.

This observation seems to apply to the Devonian tetrapods and fishes considered in this article. For example, Daeschler *et al.* noted that:

‘Devonian tetrapods show a mosaic of terrestrial and aquatic adaptations.’

Some of the fishes possess tetrapod-like characters while the tetrapods have fish-like features. Evolutionists interpret mosaic organisms like these as evolutionary intermediates linking major groups. However, Wise makes an important point against this interpretation:

‘Although the entire organism is intermediate in structure, it’s the combination of structures that is intermediate, not the nature of the structures themselves. Each of these organisms appears to be a fully functional organism full of fully functional structures.’

Evolutionary theory might lead us to expect examples of intermediate structures, but there is nothing intermediate about, for example, the internal gills of *Acanthostega*, its lateral line system, or its limbs. They are fully developed and highly complex. What is unusual is their combination in a single organism. Intelligent design offers an alternative understanding of this widespread pattern. The Devonian tetrapods are thought to have lived a predatory lifestyle in weed-infested shallow water. They were therefore equipped with characteristics appropriate to that habitat (e.g. crocodile-like morphology with dorsally placed eyes, limbs and tails made for swimming, internal gills, lateral line systems). Some of these features are also found in fishes that shared their environment.

The mosaic pattern makes it difficult to identify organisms or groups of organisms that possess the ‘right’ combination of characters to be considered part of an evolutionary lineage. Consider the tetrapod-like lobe-fins *Panderichthys* and *Elpistostege*. Despite their appearance, these fish have some unique characters (such as the design of the vertebrae) that rule them out as tetrapod ancestors. At best, evolutionists can only claim that they are a model of the kind of fish that must have served as that ancestor. The same problem is encountered with the Devonian tetrapods. For example, *Ichthyostega* is described as ‘a very strange animal, and parts of it are like no other known tetrapod or fish’. Similarly, the shoulder girdles of the Devonian tetrapods ‘are not obviously halfway in structure between those of fishes and those of later tetrapods but have some unique and some unexpected features’.

Another example is *Livoniana*, a so-called ‘near tetrapod’ known from two lower jaw fragments. It possesses a curious mixture of fish-like and tetrapod-like characteristics, but it also has up to five rows of teeth, a feature not seen either in the fishes from which it is thought to be descended nor the tetrapods into which it is said to be evolving. That the mosaic distribution of characters can cause great confusion is exemplified by the recent discovery of *Psarolepis*, a fish from the Upper Silurian/Lower Devonian of China, which combines characters found in placoderms, chondrichthyans, ray finned fishes, and lobe-fins.

**Additional problems with ‘early’ tetrapod evolution**

Another problem is that the fossil record imposes tight constraints on the timing of the supposed transition. The earliest tetrapod fossils are found in late Frasnian sediments, but their presumed ancestors are hardly much older. To exacerbate the situation, the Frasnian ‘near tetrapods’ (*Obruchevichthys*, *Elginerpeton*, *Livoniana*) are already morphologically diverse at their first appearance. Thus Darwinists are compelled to postulate a rapid burst of evolution in which radical changes must have taken place:

‘*Panderichthys* and *Elpistostege* flourished in the early Frasnian and are some of the nearest relatives of tetrapods. But tetrapods appear only about 5 to 10 million years later in the late Frasnian, by which time they were widely distributed and had evolved into several groups, including the lineage leading to the tetrapods of the Famennian. This suggests that the transition from fish to tetrapod occurred rapidly within this restricted time span.’

Second, key morphological transitions, such as the purported change from paired fins to limbs with digits, remain undocumented by fossils. Where appendages are known they are clearly either fish-like fins or digit-bearing limbs, not at some transitional stage from one to the other. At one time it was claimed that the pectoral fins of rhizodonts, a group of lobe-finned fish, were remarkably similar to tetrapod limbs, but following the description of *Goolooongongia* from the Famennian of New South Wales, Johanson and Ahlberg have urged that they not be used as a model for the origin of tetrapod limbs. Furthermore, the pectoral fins of lobe-finned fish tend to be larger than the pelvic fins, whereas the
Devonian tetrapods were ‘rear-wheel drive’ animals with larger hind limbs than fore limbs. None of the recent fossil discoveries shed any light on this supposed reconfiguration.

Third, there are functional challenges to Darwinian interpretations. For instance, in fish the head, shoulder girdle, and circulatory systems constitute a single mechanical unit. The shoulder girdle is firmly connected to the vertebral column and is an anchor for the muscles involved in lateral undulation of the body, mouth opening, heart contractions, and timing of the blood circulation through the gills.41 However, in amphibians the head is not connected to the shoulder girdle, in order to allow effective terrestrial feeding and locomotion. Evolutionists must suppose that the head became incrementally detached from the shoulder girdle, in a step-wise fashion, with functional intermediates at every stage. However, a satisfactory account of how this might have happened has never been given.

**Conclusion**

Recent discoveries have undoubtedly advanced our knowledge of Devonian tetrapods and future creationist discussions of tetrapod origins must take this into account. It is no longer sufficient for creationists to contrast *Eusthenopteron* with *Ichthyostega* and point to the large morphological gap between them. We need to have more to say. Nevertheless, the presumed transition from fish to tetrapods remains contentious. The data and their interpretation are a source of lively debate and ongoing controversy:

‘In the not-too-distant past, there was almost no fossil material, and ideas were based largely on informed guesswork. Speculation was intense, and as is often the case, in inverse proportion to the amount of data. To be truthful, there is still not much real data, so that speculation is still active, and whatever is concluded today may be overturned by the discovery of a new fossil tomorrow. That in some sense is to be hoped for, because only in that way can guesses be falsified and tested as scientific hypotheses.’

A robust rationale for concluding that the Upper Devonian tetrapods evolved from a fish ancestor, or that they gave rise to Carboniferous tetrapod lineages, is lacking. It is hoped that this paper may stimulate creationists to develop a fuller understanding of these remarkable creatures and their ecological and geological context.

**References**

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55. Daeschler et al, Ref. 37, p. 641.


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