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RESEARCH ARTICLE

The invisible fish: hydrodynamic constraints for predator-prey interaction in fossil fish *Saurichthys* compared to recent actinopterygians

Ilja Kogan^{1,2,*}, Steffen Pacholak^{3,4,*}, Martin Licht^{1,‡}, Jörg W. Schneider^{1,2}, Christoph Brücker³ and Sebastian Brandt⁵

ABSTRACT

Recent pike-like predatory fishes attack prey animals by a quick strike out of rest or slow movement. This fast-start behaviour includes a preparatory, a propulsive and a final phase, and the latter is crucial for the success of the attack. To prevent prey from escape, predators tend to minimise the duration of the interaction and the disturbance caused to surrounding water in order to not be detected by the prey's lateral line sensory system. We compared the hydrodynamic properties of the earliest fossil representative of the pike-like morphotype, the Triassic actinopterygian *Saurichthys*, with several recent pike-like predators by means of computational fluid dynamics (CFD). Rainbow trout has been used as a control example of a fish with a generalist body shape. Our results show that flow disturbance produced by *Saurichthys* was low and similar to that of the recent forms *Belone* and *Lepisosteus*, thus indicative of an effective ambush predator. Drag coefficients are low for all these fishes, but also for trout, which is a good swimmer over longer distances but generates considerable disturbance of flow. Second-highest flow disturbance values are calculated for *Esox*, which compensates the large disturbance with its extremely high acceleration performance (i.e. attacks at high speeds) and the derived teleostean protrusible mouth that allows prey catching from longer distances compared to the other fishes. We show CFD modelling to be a useful tool for palaeobiological reconstruction of fossil fishes, as it allows quantification of impacts of body morphology on a hypothesised lifestyle.

KEY WORDS: *Saurichthys*, Ambush predators, Fossil fish, Locomotion, CFD, Palaeontology

INTRODUCTION

Modern actinopterygian fishes exhibit a wide variety of body shapes, adapted for – and indicative of – a number of different lifestyles (e.g. Greenway, 1965). General appearance of the body,

shape and placement of fins, structure and position of the mouth or type of dentition are only some of the characters that correlate with requirements of the habitat, locomotor performance, feeding habits and even times of activity during the day and night cycle. Such correlations are corroborated by numerous field observations, experimental and theoretical studies (e.g. Hobson, 1979; Keast and Webb, 1966; Lauder, 2006; Licht, 2009; Wainwright and Richard, 1995; Webb, 1984). Nonetheless, attribution of behavioural types to extinct animals based on comparison with extant forms must remain to some extent speculative, as long as no individual parameters of the fossils are taken into consideration (see Boucot and Poinar, 2010; Fletcher et al., 2014).

The term 'pike-like predators' is used to describe actinopterygians with elongated heads, long and slender bodies, posteriorly placed dorsal and anal fins and symmetrical tails. These mostly piscivorous fishes are not designed for long-term continuous swimming, but attack their prey by quick strikes out of rest or slow movement (also called fast-start or ambush predation). Fishes partly matching this body plan go back to the Palaeozoic, but the first typical representatives of such a morphotype appeared in the fossil record around the Permian-Triassic boundary and belonged to the 'palaeopterygian' family Saurichthyidae Owen, 1860 (*sensu* Stensiö, 1925). Soon after the End-Permian mass extinction, species of the genus *Saurichthys* Agassiz, 1834 (Fig. 1) radiated all over the globe, invading both marine and freshwater ecosystems. Saurichthyids ranged in length from few centimetres to more than 1.5 m, and at least the larger species are known to have been piscivorous (Kogan et al., 2014). Becoming rare in freshwater environments, saurichthyids retained the role of high-level consumers in the marine realm until their last representatives died out during the Jurassic (e.g. Romano et al., 2012). From that time on, however, similar morphologies independently arose in several actinopterygian clades (Fig. 2), raising the question whether or not they can be related to a similar lifestyle.

In fishes, fast-start predation (lunging) follows a well-known scheme that can be subdivided in distinct stages. Weihs (1973) identified a preparatory phase (I), during which the predator, when detecting the prey, changes from a stretched out position to an L or S shape; a propulsive phase (II), where the predator accelerates by quick movements in its posterior body half; and a final stage (III) of gliding or continued swimming (also called variable stage by some workers, e.g. Harper and Blake, 1991). Prey is caught in phase III (Frith and Blake, 1995; Webb and Skadsen, 1980).

The energetic costs of the fast-starting hunting tactics are high and need to be compensated with an increased success rate (e.g. Harper and Blake, 1988), which can be achieved, for instance, when the predator approaches the prey quicker than it can escape. Besides

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List of abbreviations

<i>a</i>	deceleration	ms^{-2}
<i>A</i>	surface/frontal area	m^2
<i>c</i>	drag coefficient	
<i>d</i>	distance	m
<i>F</i>	force	N
<i>FR</i>	fineness ratio	
<i>h</i>	height	m
<i>l</i>	length	m
<i>m</i>	mass	kg
<i>p</i>	pressure	Pa
<i>r</i>	radius	m
ρ	density	kg/m^3
<i>t</i>	time	s
<i>u</i>	velocity	ms^{-1}
<i>V</i>	volume	m^3
<i>w</i>	width	m
<i>x</i>	regression coefficient	m^{-1}
<i>y</i>	dimensionless coefficient	

Units

Since the size of the investigated fishes varies within one order of magnitude, length units have been related to the total length of the respective fish to ensure comparability of the calculation results. These fish length-related values are marked with an asterisk in the text.

locomotor requirements to the fast-starting predator itself, perception of the approaching predator by the prey is crucial for the outcome of this interaction.

Fishes detect predators not only by vision, but also – and perhaps most importantly – by means of their lateral line sensory organ, which is sensitive to hydrodynamic signals (e.g. Bleckmann et al., 2004). To avoid being detected, an aquatic predator should therefore minimise flow disturbances caused by its movement. Experiments (e.g. Webb and Skadsen, 1980) demonstrate that fast-start predatory fishes reduce undulation of their body in the final phase of the strike and approach the prey largely without generating additional thrust.

As a first step towards the reconstruction of the possible hunting behaviour of the fossil actinopterygian *Saurichthys*, we examined the potential predator-prey interaction of this fish in comparison with several recent forms whose behaviour is known (Fig. 2, Table 1). For this purpose, we produced digital 3D surface models of a generalised *Saurichthys* and the recent pike-like actinopterygians *Esox lucius* (Linnaeus, 1758), *Belone belone* (Linnaeus, 1761), *Lepisosteus osseus* (Linnaeus, 1758) and *Ctenolucius hujeta* (Valenciennes in Cuvier and Valenciennes, 1849) and investigated their hydrodynamic properties in a digital (simulated) flow channel. Rainbow trout *Oncorhynchus mykiss* (Walbaum, 1792) was used as a control example for a predatory fish with a generalist shape.

We used computational fluid dynamics (CFD) modelling to determine flow velocities around and pressure distribution at the surface of the fish body. Simulations have been performed with rigid fish models held stationary in a constant current (corresponding to a fish gliding at a constant velocity). This represents an approximation for phase III (*sensu* Weihs, 1973) of the fast-start attack where acceleration is terminated and the predator attempts to catch prey without producing additional thrust, so the actual speed may be nearly constant for a short time between the propulsive phase and the interaction with prey. To simulate predation in riverine and marine/lacustrine environments, different turbulence intensities have been implemented into the calculations.

RESULTS**Pressure**

The distribution of pressures over the fish body in a moving fluid is visualised in Fig. 3. Areas of elevated pressure are nearly absent in *Belone* and *Saurichthys*, small in *Lepisosteus* and *Ctenolucius* and large in *Esox* and *Oncorhynchus*. In general, the highest pressures are recorded in the anterior part of the head and at the leading edges of the paired fins, thus in the body portions facing the current. However, elevated pressures are also found at the caudal peduncle and over the caudal fin.

Drag

All bodies moving through a fluid are confronted with resistance of the medium called drag. Total drag force F_{drag} is a combination of the pressure-induced drag F_{pressure} and the surface friction F_{friction} .

Whereas the total drag force F_{drag} increases with increasing speed and turbulence intensity (Table 2), the contribution of pressure-induced drag F_{pressure} and surface friction F_{friction} to total drag changes distinctly. Friction grows slower with higher velocities, so that drag is increasingly pressure-induced at higher swimming speeds (Table 2, Fig. 4). Among taxa, the percentage of friction varies from very low (*Esox*) to very high (*Lepisosteus* and *Belone*) (Fig. 4). The highest increase in pressure-induced drag compared to friction is noted for *Ctenolucius*.

A universally comparable drag measure is the drag coefficient c_{drag} . According to different conventions, it can be calculated based either on the projected frontal area ($c_{\text{drag,front}}$) or on the total body surface ($c_{\text{drag,surf}}$). Both coefficients are nearly speed-independent and are lowest for *Belone*, *Saurichthys* and *Oncorhynchus* (Table 1). Drag coefficients are highest for *Esox* and *Ctenolucius*, and mostly higher in fluvial systems than in open waters (Table 2).

Flow disturbance

To analyse the hydrodynamic effects of an approaching predator, we quantified the disturbance of flow caused by the body of a fish



Fig. 1. *Saurichthys deperditus* (Costa, 1862), a large Late Triassic saurichthyid from marine deposits of Austria.

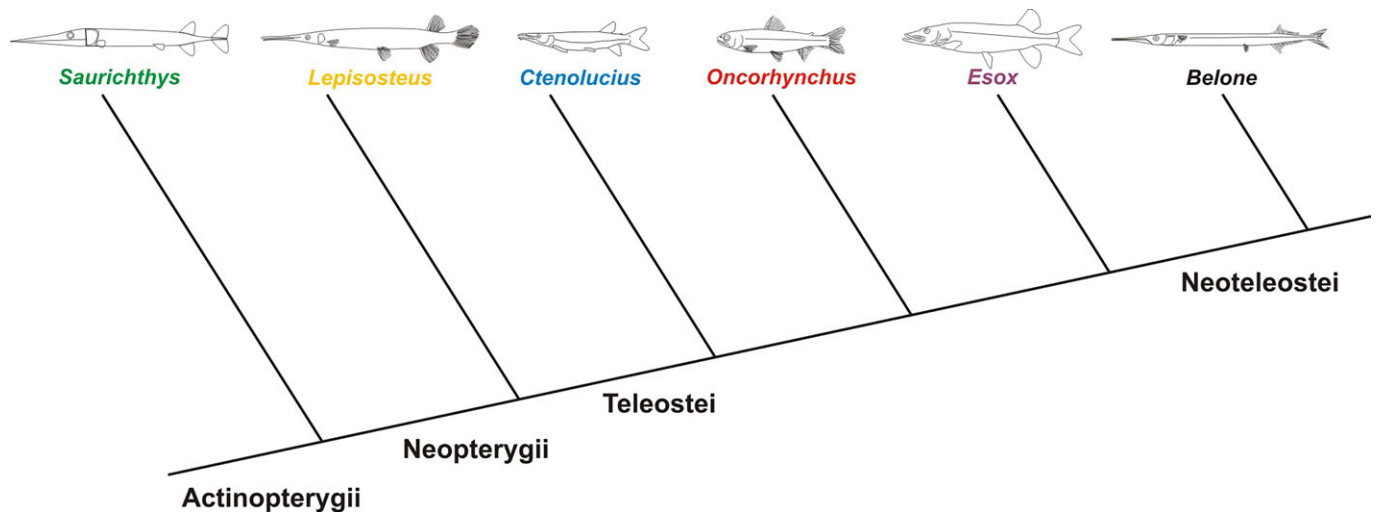


Fig. 2. Overview of the fish taxa incorporated in this study. Phylogenetic relationships are summarised in the cladogram. The extinct *Saurichthys* is the basalmost genus included. Colours used here for the fish names are found in the following figures as well.

moving through the fluid (Fig. 5). Flow disturbance can be expressed as variation of velocities in the flow channel where the fluid streams at a given inlet velocity u_{fish} (simulating the average swimming speed of the fish), while the fish body is held static (Fig. 5B), or as the circulation around the fish body when moving through a static fluid (Fig. 5C). Maximum disturbance is visualised in a contour plot generated by colouring areas of the fluid domain (shown in Fig. 5) where flow velocity u_{local} differs from the inlet velocity by at least 1%. Low velocity regions ($u_{\text{local}} \leq 99\% \cdot u_{\text{fish}}$) are coloured in blue and high velocity regions ($u_{\text{local}} \geq 101\% \cdot u_{\text{fish}}$) in red.

The dependency of flow disturbance on swimming speed and turbulence regime is shown in Table 3, using *Saurichthys* as an example. Disturbed area (expressed by the relative contour radius r^*) is smaller at higher turbulence intensities, but virtually no effect is documented on the distance d from the tip of the snout to the anterior margin of the contour.

The largest flow disturbance area in the frontal plane is shown by *Esox*, followed by *Oncorhynchus*. In both taxa, the contour expansion comprises more than 4.5 times their own radius. *Belone* generates the minimal relative contour radius (Fig. 6A, Fig. 7A). The greatest differences in the contour radius between the less turbulent lacustrine and the higher turbulent fluvial environment were recorded for *Lepisosteus* and *Esox* (Fig. 7A). With increasing velocity, the relative area of flow disturbance slightly decreases for *Lepisosteus* and *Ctenolucius*, but remains nearly constant for the other fishes. Relative snout distance d^* is lowest for *Belone* and highest for *Oncorhynchus*, followed by *Esox*

(Fig. 6B), and is nearly unaffected by velocity (Fig. 7B). *Saurichthys* plots in the lower half of both diagrams close to *Belone* and *Lepisosteus*.

Parametric studies

To evaluate the impact of environmental conditions on the hydrodynamic model, we performed additional calculations at different values of temperature, viscosity and density of the surrounding water. The temperature dependency of viscosity and density led to a triple parametric study that cannot be considered separately. In comparison with the initial calculations performed at 15°C, the results obtained for temperatures of 10°C and 20°C showed only minor deviation of less than 1%.

Additionally, a parametric study for turbulence intensity was made for several hunting domains (ideal-theoretic with nearly no turbulence, open marine/lacustrine, normal-fluvial and higher turbulent river). The results are compared in Table 4 and show only slight differences between ideal-theoretic and open marine/lacustrine habitat as well as between normal-fluvial and higher turbulent rivers at the considered parameters. Deviation remains constant irrespective of gliding velocity at which the calculations are performed.

Morphometric predictability

We found no correlation between the various length distances measurable in the fishes (Fig. 8) and their hydrodynamic properties as summarised above. In contrast, the fineness ratio defined here as the ratio of total length to maximum height (excluding fins) is a good indicator for flow disturbance, pressure distribution and drag

Table 1. Morphometric parameters (total fish length l_{fish} , relative head length l_{head}^* , fineness ratio FR , projected frontal area A_{front} and total body surface A_{surf}), Reynolds numbers Re and drag coefficients of the fishes incorporated in this study

Parameter	<i>Saurichthys</i>	<i>Belone</i>	<i>Lepisosteus</i>	<i>Ctenolucius</i>	<i>Esox</i>	<i>Oncorhynchus</i>
l_{fish} [m]	0.74	0.74	0.13	0.09	0.40	0.40
l_{head}^* [%]	31	22	31	22	26	19
FR	12	14	10	9	7	5
A_{front} [m ²]	0.004	0.001	0.0001	0.00006	0.003	0.005
A_{surf} [m ²]	0.124	0.074	0.003	0.002	0.056	0.097
Re [10 ³]	675–3380	634–3270	19–114	9–45	183–1760	182–1750
$C_{\text{drag,front}}$	0.345	0.359	0.421	0.751	0.684	0.262
$C_{\text{drag,surf}}$	0.011	0.007	0.014	0.023	0.041	0.013

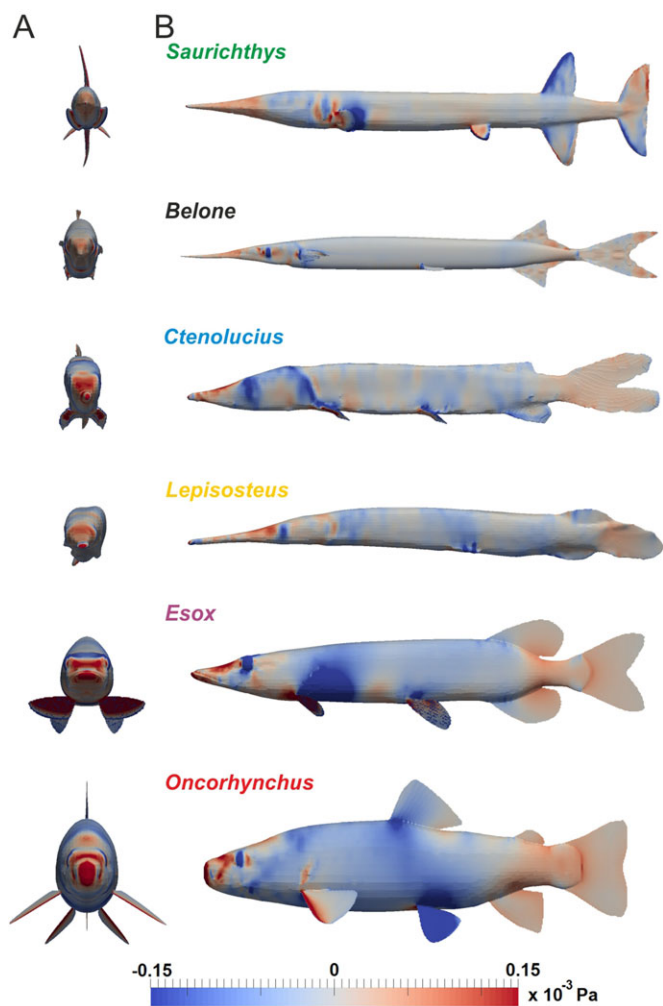


Fig. 3. Surface pressures. Distribution of surface pressures ($p=p_{\text{total}}-p_{\text{dyn}}$) over the fish bodies at $u_{\text{fish}}=1\text{ ms}^{-1}$ in anterior view (A) and lateral view (B). Regions of low pressure are coloured in blue, high pressure areas in red.

coefficient of the pike-like predators. Only the drag coefficient of trout is much lower than would be predicted from fineness ratio (Table 1).

DISCUSSION

Body shape and lifestyle of animals are known to be correlated (e.g. Greenway, 1965). Although the investigated extant fishes with a pike-like body shape belong to different phylogenetic lineages (Fig. 2), they not only share morphological traits such as

elongated skull, long and slender body, posteriorly positioned dorsal and anal fins, narrow caudal peduncle and symmetrical tail, but also exhibit considerable similarities in their hunting behaviour. They are considered fast-start predators, which attack mobile prey by quick, unexpected movements in order to prevent it from escape (Webb, 1984). Apart from the rapidity of the attack itself, this involves mechanisms for avoiding early discovery by the prey.

We assumed that successful fast-start predators should generate few flow disturbance to avoid being discovered by the prey’s lateral line sensory system. Furthermore, as body undulations increase friction and disturbance of flow, predators reduce or cease undulation in the final phase of the strike and approach prey without additional thrust (e.g. Weihs, 1973; Webb and Skadsen, 1980). Consequently, one may expect that the drag coefficient of such predators is low, permitting them to reach the target easily once the propulsive phase is terminated. Indeed, disturbed area (normalized by fish length) is small for *Belone*, *Saurichthys*, *Lepisosteus* and *Ctenolucius* but large for both *Oncorhynchus* and *Esox*. The drag coefficients are also low for *Belone*, *Saurichthys* and *Lepisosteus*, but surprisingly high for *Esox* and *Ctenolucius*. In contrast, the generalist *Oncorhynchus* is characterised by a very low drag coefficient. According to our predictions, *Belone* and *Lepisosteus* should be the best-adapted recent forms for fast-start predation.

Esox is, nevertheless, seen as the iconic fast-starting predator in literature and even eponymous to the pike-like morphotype. When more flow disturbance is caused by its less streamlined body shape, pike should attack more quickly to prevent prey from escape. During acceleration, drag plays a minor role compared with fish mass, mass of water accelerated with the fish (added mass) and rate of acceleration (Webb, 1975), and so the higher drag coefficient of pike has less impact on fast-starting than muscle mass percentage (lower in forms with extensive body armour), lateral body profile or size and position of fins (Domenici and Blake, 1997; Weihs, 1989). Extensive experimental studies demonstrated the fast-start performance of pike to be superior to that of *Lepisosteus* (Webb et al., 1992) and trout (Harper and Blake, 1990), despite the pike’s higher drag coefficient (see also Webb, 1988). Whereas in Harper and Blake’s (1991) experiments, the prey – goldfish – attempted escape in most cases but were rarely successful, only 15% of fathead minnows used by Webb and Skadsen (1980) showed escape movements at all. This means that in spite of its relatively high flow disturbance, pike is quick enough to be successful in predation. The protrusible jaws of teleost fishes, primarily not developed in *Saurichthys* and *Lepisosteus* and secondarily lost in *Ctenolucius* and *Belone*, might be crucial for predatory ability in *Esox*.

Table 2. Surface parameters (pressure-induced drag forces F_{pressure} , friction F_{friction} and total drag force F_{drag}) and the drag coefficient calculated based on the frontal area ($c_{\text{drag,front}}$) and the wetted surface area ($c_{\text{drag,surf}}$) for *Saurichthys* moving in marine/lacustrine and riverine environments at three different velocities

Parameter	$u_{\text{fish}}=1\text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=3\text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=5\text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=1\text{ ms}^{-1}$ riv.	$u_{\text{fish}}=3\text{ ms}^{-1}$ riv.	$u_{\text{fish}}=5\text{ ms}^{-1}$ riv.
F_{drag} [N]	0.69	5.51	14.69	0.73	5.87	15.64
F_{pressure} [N]	0.33	3.09	8.66	0.36	3.39	9.47
% of F_{drag}	48.40	56.05	58.96	50.00	57.69	60.54
F_{friction} [N]	0.35	2.42	6.03	0.36	2.48	6.17
% of F_{drag}	51.60	43.95	41.04	50.00	42.31	39.46
$c_{\text{drag,front}}$	0.354	0.316	0.303	0.375	0.336	0.323
$c_{\text{drag,surf}}$	0.011	0.010	0.009	0.012	0.011	0.010

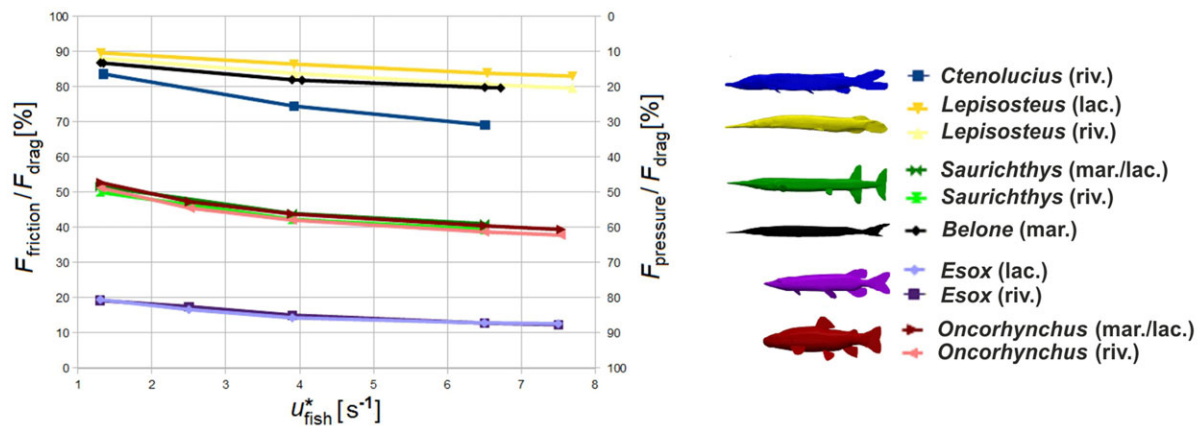


Fig. 4. Drag forces. Speed-dependent variation of the components of the total drag force applied on the fish bodies, friction (left scale) and pressure-induced drag (right scale) over standardised fish velocity (u^* in body length/second) in higher turbulent riverine (riv.) and less turbulent marine/lacustrine (mar./lac.) environments.

Although a predator as well, the lifestyle of trout includes more continuous swimming, which is reflected in its generalist body shape. Trout causes high flow disturbance with the overall greatest anterior extension, but its streamlined body enables it to travel over long distances at low energetic costs (e.g. Tytell, 2007).

In terms of flow disturbance and drag, the fossil fish *Saurichthys* occupies a position intermediate to the extant neopterygians *Lepisosteus* and *Belone*, both of which nearly comply with our predictions for a successful fast-start predator. The model-derived hydrodynamic properties allow to evaluate the adaptation of *Saurichthys* for the fast-start hunting behaviour, which previously could only be suggested based on general morphology (Lombardo and Tintori, 2005; Rieppel, 1985; Tintori, 1990). We demonstrate that a hypothetical generalised saurichthyid caused moderate flow disturbance, especially low directly in front of the snout, and was able to maintain a relatively long final phase of the strike thanks to

its low drag coefficient. Differences in squamation, vertebral column and fin morphology certainly had further impacts on the swimming behaviour of different *Saurichthys* species, which shall be quantified in forthcoming studies.

Environmental factors like water temperature, density and viscosity were found to have little effect on fish hydrodynamic properties when varied within a natural range. The only considerable deviation was caused by different turbulence intensities corresponding to open marine, lacustrine or riverine habitats. These especially influence drag forces as well as contour radius of the created flow disturbance. For instance, we noted a decrease of relative contour radius of *Saurichthys* with increasing turbulence but an increase of drag forces. So the prey may recognize the predator at a later moment but the predator needs a greater amount of energy to catch it in a turbulent environment. However, the general trends presented above remain constant throughout the turbulence regimes.

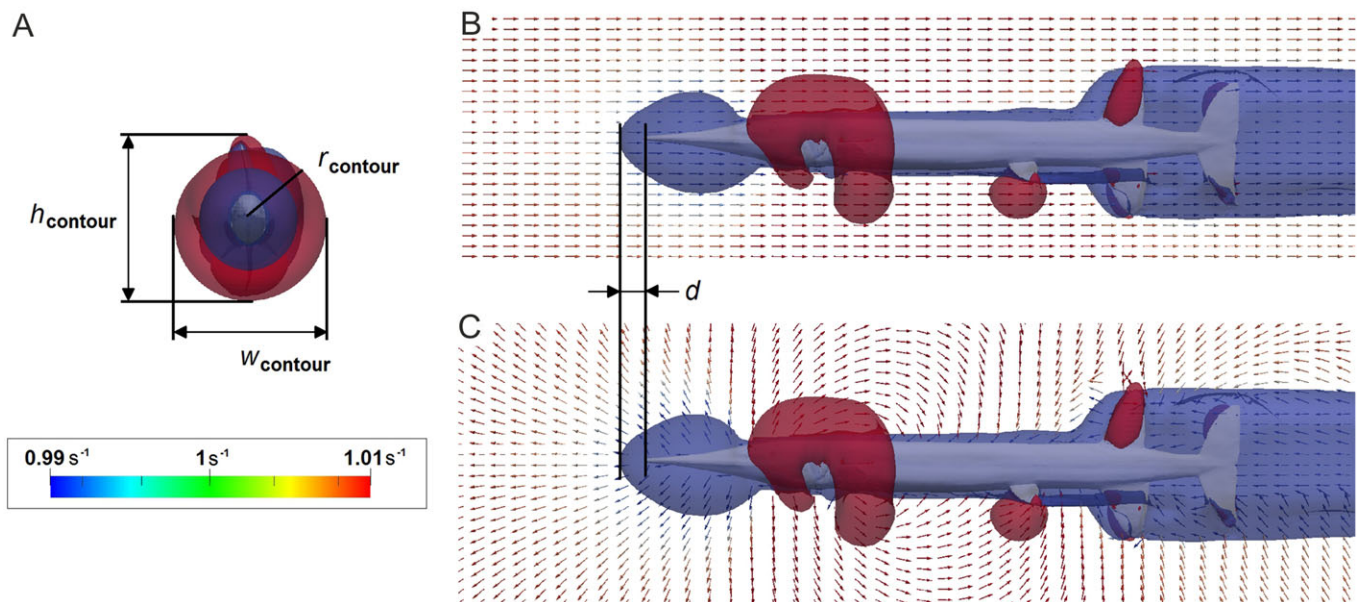


Fig. 5. Visualisation of the flow velocity disturbance around *Saurichthys* at $u_{\text{fish}} = 1 \text{ ms}^{-1}$. (A) flow disturbance in anterior view; (B) disturbance of a constant current streaming around a static fish in lateral view; (C) disturbance caused by the fish moving through a static water body in lateral view. d, anterior expansion of the contour (snout distance); h, height; r, radius; w, width.

Table 3. Contour parameters (relative radius r^* , anterior distance between snout and contour front d in metres and relative anterior distance d^*) for *Saurichthys* moving in marine/lacustrine and riverine environments at three different velocities

Parameter	$u_{\text{fish}}=1\text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=3\text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=5\text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=1\text{ ms}^{-1}$ riv.	$u_{\text{fish}}=3\text{ ms}^{-1}$ riv.	$u_{\text{fish}}=5\text{ ms}^{-1}$ riv.
r^*	3.80	3.80	3.79	3.69	3.68	3.69
d [m]	0.03	0.03	0.03	0.03	0.03	0.03
d^*	0.04	0.04	0.04	0.04	0.04	0.04

Our results show that numerical hydrodynamic modelling is a good tool to estimate the performance of fossil aquatic animals and to test hypotheses concerning their lifestyle. Furthermore, the procedure presented here is a low-cost method in comparison with experimental studies. Refinement of the calculation routines and inclusion of more morphological data as well as further taxa will lead to more detailed and quantifiable outcomes. Applied to fast-start predatory fishes, modelling of the acceleration phase and implementation of motion functions shall allow precise reconstructions of the complete strike movement, which is crucial to understand the evolution of predator-prey interactions.

MATERIALS AND METHODS

Considered taxa

Saurichthys

51 species are currently considered valid in the family Saurichthyidae, one of which is of Late Permian, 48 of Triassic and two of Early Jurassic age (see Romano et al., 2012, for an overview, and Maxwell et al., 2015; Tintori, 2013; Tintori et al., 2014; Werneburg et al., 2014 and Wu et al., 2015 for newest taxa). About one third of the Triassic species are documented by material sufficiently complete as to allow the reconstruction of the whole skeleton. For the purpose of this work, a generalised saurichthyid morphology was required, which is compiled from the best-known species of Early, Middle and Late Triassic. Besides several schematic drawings, whole-body restorations of Triassic saurichthyids have been published by Griffith (1977), Kogan et al. (2009) and Rieppel (1985).

Synapomorphic characteristics of the genus *Saurichthys* can be summarised as follows (Figs 1, 8): long and slender body (fineness ratio *sensu* Maxwell and Wilson, 2013, up to 20); elongate head usually accounting for 1/4 to 1/3 of the total body length; pelvic fins placed near the middle of the trunk; dorsal and anal fins symmetrical to each other and placed in the middle of the distance between the pelvis and the caudal fin; the latter externally and structurally symmetrical, having equal epaxial and hypaxial lobes separated by the vertebral column that proceeded straight to the posterior margin of the tail (abbreviate-diphycercal tail, Brough, 1936); narrow caudal peduncle, stiffened by interlocking scutes along the dorsal and ventral midlines; and a specialised, usually strongly reduced squamation (Gardiner, 1960; Griffith, 1959, 1962; Maxwell and Wilson, 2013; Rieppel, 1985, 1992; Romano et al., 2012; Stensiö, 1925; Woodward, 1888, 1890, 1895). The axial skeleton of *Saurichthys* lacks ossified vertebral centra and consists of a persistent notochord flanked by up to 200 pairs of neural arches dorsally and a usually lower number of haemal arches ventrally. In several species, the vertebral column is stiffened by long processes directed anteriorly and posteriorly to the neural arches (Tintori, 1990, 2013). Furthermore, many forms possessed unsegmented or rarely segmented fin rays, indicating increased stiffness of the fins (Romano et al., 2012; Schmid and Sánchez-Villagra, 2010).

Recent forms

Lepisosteus osseus (Ginglymodi: Lepisosteiformes) lives in rivers and lakes of North America and is one of the rare recent fishes whose body is covered by rhombic ganoid scales. Lepisosteids feed mainly on fish, but

can also include crustaceans, higher vertebrates and carrion in their diet (Kammerer et al., 2006). They approach their prey rather slowly before attacking it by a quick lunge (Kammerer et al., 2006; Porter and Motta, 2004).

Ctenolucius hujeta (Ostariophysi: Characiformes) is found in rivers of equatorial South America, where it preys on small fish and

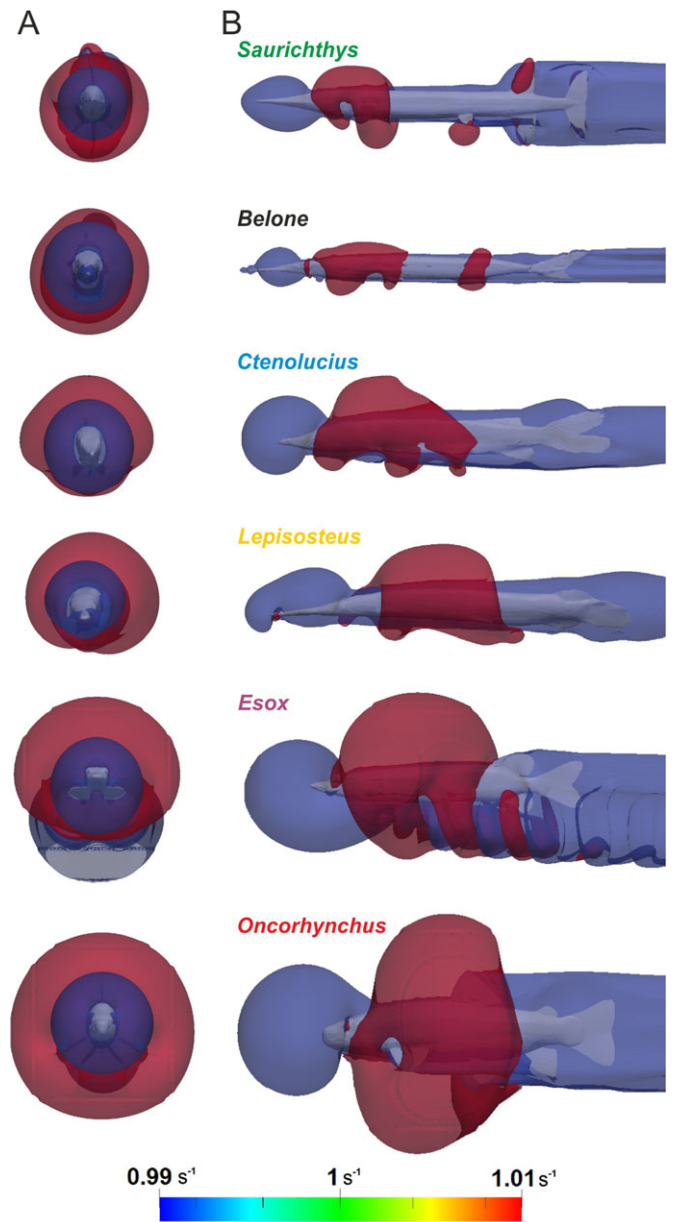


Fig. 6. Flow velocity disturbance around the fish bodies at $u_{\text{fish}}=1\text{ ms}^{-1}$. Blue regions indicate low velocity areas (99% of swimming velocity) and red ones symbolize high velocity areas (101% of swimming velocity) in anterior view (A) and lateral view (B).

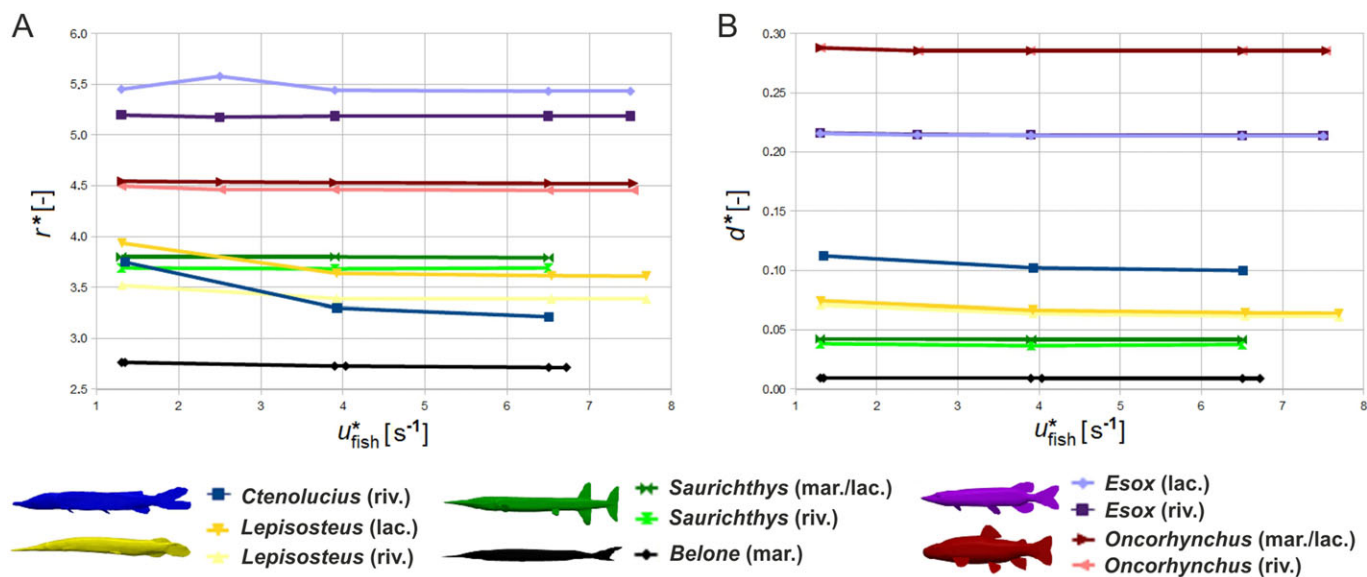


Fig. 7. Flow velocity disturbance. Speed-dependent variation of the radius (A) and the anterior expansion (B) of the area of disturbed flow velocity around the fish bodies. u^* , standardised fish velocity in body length/second; r^* , contour radius/fish radius; d^* , anterior contour expansion/fish length.

occasionally decapods. *Ctenolucius* has bright silver coloured skin and hides in small inlets waiting for prey that it catches with a quick strike (Vari, 1995).

Belone belone is a European representative of the needlefishes (Teleostei: Beloniformes) and lives in surface waters of the North Sea, the Black Sea, the Azov Sea, the Mediterranean and its adjacent regions. Belonids are piscivorous and use a similar hunting strategy, performing relatively quick strikes over short distances (Collette, 2003; Porter and Motta, 2004).

Esox lucius (Teleostei: Esociformes) is the emblematic ambush predator whose prey capture behaviour has been extensively studied (e.g. Harper and Blake, 1991; Frith and Blake, 1995; Rand and Lauder, 1981; Schrieffer and Hale, 2004). Species of *Esox* are found in fresh waters of the northern hemisphere and are essentially piscivorous (Bregazzi and Kennedy, 1980) and hide in vegetated areas of the water body waiting for prey that they attack by quick strikes.

Oncorhynchus mykiss (formerly called *Salmo gairdnerii*) is a widespread salmoniform teleost with a generalist morphology (Greenway, 1965), able to hold position in streams, to perform fast-start escapes, but also to migrate over long distances (Przybilla et al., 2010; Tytell, 2007; Webb, 1976). In nature, the diet of trout largely depends on age and available food, being composed in varying proportions of benthic macroinvertebrates, terrestrial insects, fish and even plants (Di Prinzio et al., 2013; Rikardsen and Sandring, 2006).

Morphometrics

For the *Saurichthys* model and the recent fishes investigated, we recorded several morphometric measurements, such as head length, total body length and maximum height (Fig. 8, Table 1). Head length is

measured from snout tip to the posterior margin of the operculum. Additionally, we considered the fineness ratio (*FR*) defined as the relationship between total length and maximum height (without fins). We furthermore calculated fish volume and mass out of the 3D geometry (Table 6).

Obtaining geometries
Saurichthys plastic model

The proportions mentioned above for *Saurichthys* are in good agreement with those of the recent garfish *Belone belone*, which therefore was used as a living control form. The lateral outline of *Saurichthys*, based on the published restorations and own observations (I.K., S.B.), was combined with shape data from three-dimensionally preserved saurichthyid skulls and supplemented with the garfish soft part morphology to obtain a true to scale *Saurichthys* trunk model executed in rigid foam (Fig. 9). Morphological details such as skull bones, muscles and scales have been formed in modelling resin and plastically applied in several steps. Special attention was paid to the configuration of the head, the squamation and the skin. Fins have been modelled separately, and their position was marked on the trunk model according to scientific reconstructions (Kogan et al., 2009; Rieppel, 1985). Negative silicone casts of all parts were prepared, allowing to reproduce the surfaces in epoxy resin. These could now be combined into a hollow body with an ultralight metallic core wire. Since the material used is thermally plastic, the obtained models can be deformed depending on the required position. For display purposes, models can be airbrushed in colours inferred from recent examples and covered with protective lacquer.

Digitalisation

The *Saurichthys* model was digitalised using a mobile MicroScribe 3D-Scanner from the Technical University of Dresden and the rendering software MicroScribe Utility for creating a point cloud of the scan data. Applying a Delaunay triangulation algorithm on this data field led to a stereo lithography (STL) surface consisting of a finite number of triangles (Pacholak et al., 2014). Specimens of *Belone*, *Lepisosteus* and *Ctenolucius* were scanned by CreaForm using a mobile HandyScan3D and were transformed the same way into STL data. The geometries of rainbow trout and northern pike were taken from the Online Toucan Virtual Museum of the Toucan Corporation Japan (<http://www.toucan.co.jp/3DCG/3ds/FishModelsE.html>) and converted from Tank3 Demo Files

Table 4. Parametric study of different hunting habitats for <i>Saurichthys</i> at low gliding velocity				
Parameter at $u_{fish}=1\text{ ms}^{-1}$	ideal	marine/lacustrine	riverine	turbulent river
r^*	3.87	3.80	3.69	3.57
d^*	0.04	0.04	0.04	0.03
F_{drag} [N]	0.69	0.69	0.73	0.82
$C_{drag,front}$	0.353	0.354	0.375	0.424
$C_{drag,surf}$	0.011	0.011	0.012	0.013

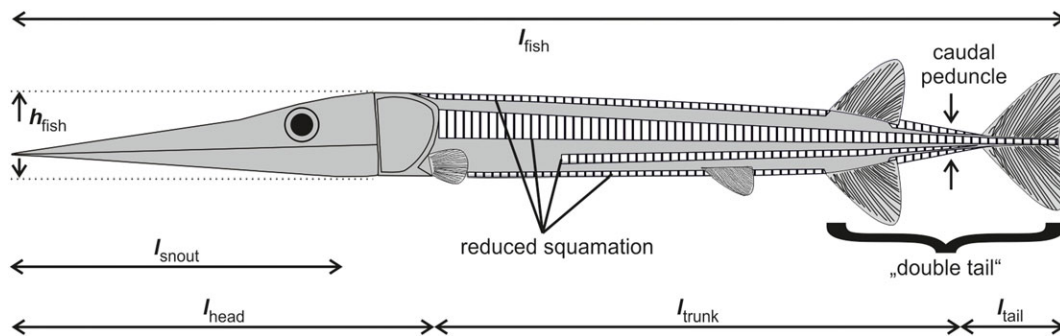


Fig. 8. General morphology of *Saurichthys* with some morphometric distances. h_{fish} , maximal height excluding fins; l_{fish} , total body length; l_{snout} , maximal length of the mouth gape; l_{head} , head length; l_{trunk} , trunk length from the hind margin of the operculum to the narrowest point of the caudal peduncle; l_{tail} , length from the narrowest point of the caudal peduncle to the hind margin of the caudal fin. The dorsal, anal and caudal fins form the 'double tail' *sensu* Weihs, 1989.

into STL with NURBS Modelling for Windows (Rhino), covering gill openings and closing the mouth, as no surface holes or unnatural rough edges are allowed to fulfil the circulation condition for fluid dynamics.

Additionally, deformations of scanned specimens like back bone bending caused by conservation or adverse storing could be undone by applying a reverse motion algorithm onto the surface models in MATLAB (The Mathworks Inc.). This method unbends a deformed vertebral column back to prone position under consideration of the length consistence criteria so that both the curved and the straightened

model have the same length after editing. Afterwards the unbent surface points are calculated using their relative position to the back bone.

Numerical methods

Steady gliding

The basic study of hydrodynamic properties of the fish bodies involves their interaction with the fluid at constant swimming speeds. We simulated gliding at constant speeds numerically by placing rigid surface models of the fishes into a digital flow channel. The latter approximates the fish's natural

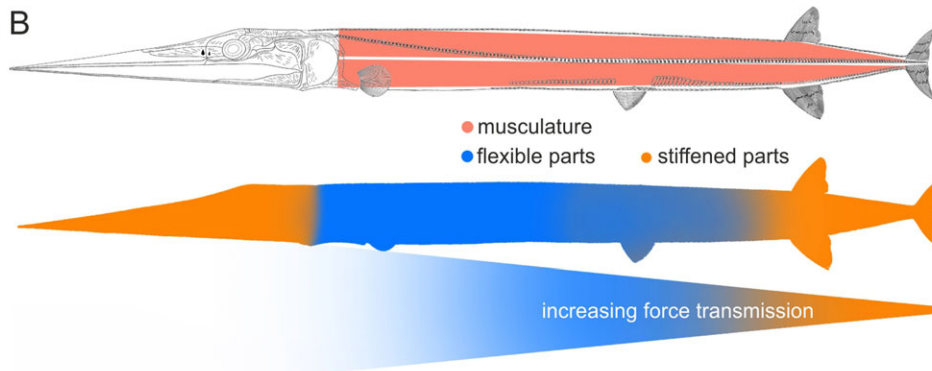


Fig. 9. Steps of plastic rebuilding of a generalised *Saurichthys*. (A) a Middle Triassic *Saurichthys curionii* (Bellotti, 1857) from Monte San Giorgio, Switzerland; (B) presumed distribution of muscles, soft and stiffened regions, based on Rieppel's (1985) reconstruction of *S. curionii*; (C) sketch of the cranial elements on a model of *Saurichthys* head; (D) modelling of the snout and head structures; (E) application of paired fins; (F) completed *Saurichthys* plastic model.

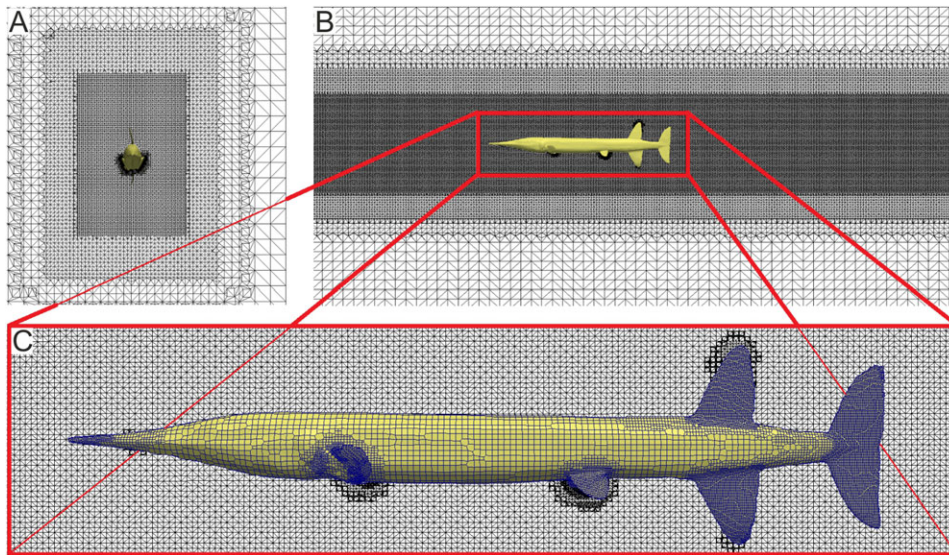


Fig. 10. CFD calculation mesh. Discretised flow channel around digitalised *Saurichthys* model with a relative size of five fish long, nine fish high and seven fish wide in anterior view (A) and lateral view (B). Each grid cell contains a spatial solution of the NS equation, so the resolution of the grid has a finer core near the surface of *Saurichthys*. (C) Discretisation of the model surface to calculate surface forces.

hunting environment that can be characterised by temperature, density and turbulence intensity of the surrounding water. Up to six different velocities were investigated for each fish species. When fluid parameters are held constant, the distribution of pressures and stream velocities over the fish body depends on body shape. To calculate the pressure-induced forces (lift and drag) and the viscous force (friction) out of pressure and stream velocity on the fish surface, the governing equations of fluid dynamics, the Navier–Stokes Equations (NS), mass conservation as well as momentum conservation:

$$\nabla \cdot \vec{u} = 0 \quad (1)$$

$$\frac{\partial \vec{u}}{\partial t} + (\vec{u} \cdot \nabla) \vec{u} = \frac{\eta}{\rho} \Delta \vec{u} - \frac{1}{\rho} \nabla p, \quad (2)$$

where p is pressure, $\vec{u}$ is fluid velocity, ρ and η are density and dynamic viscosity of water at time t , have to be solved numerically. Unfortunately, an analytic solution for NS with turbulence models (here we used the $k\text{-}\omega$ -SST model) describing finer vortex structures and velocity development still doesn't exist. So the circulation area surrounding the fish model had to be divided into a finite number of grid cells (shown in Fig. 10 for about 5 million cells totally) for a discrete spatial solution. Therefore an outer hexahedral mesh and a finer inner mesh adapted to the fish's shape was created. Surface layers were added along the fish surface to enhance the wall resolution. The spatial discretisation is also applied to the fish surface model to calculate surface pressures (Fig. 10C). In each cell, the NS is solved using numerical approximation functions for the pressure and velocity field. The according boundary conditions can be found in Table 5.

Calculations were performed by OpenFOAM using the finite volume method with second-order discretisation in space at a stationary case (e.g. Ferziger and Peric, 2002; Versteeg and Malalasekera, 2007) in species-specific flow channels spanning five times the length, nine times the height and about seven times the width of each fish, with a water

temperature set at 15°C and standard turbulence intensity values for a (less turbulent) marine/lacustrine and a (higher turbulent) fluvial environment, according to the natural habitats of the studied fishes (see Sukhodolov et al., 1998).

Post-processing

While only the distribution of pressures and flow velocities is directly obtained from the calculation, several other parameters can be derived from these results. These include the magnitude and shape of flow disturbance around the fish, friction and pressure-induced drag force and the drag coefficients.

The area within which flow velocity of the water is disturbed by 1% or more compared with u_{fish} (inlet velocity of the flow channel that corresponds to the average fish velocity) is expressed by its envelope (contour). To estimate flow disturbance around the fish body, the radius of the greatest expansion of the contour (shown in Fig. 3) $r_{\text{contour}} = \frac{1}{4}(h_{\text{contour}} + w_{\text{contour}})$ is divided by the mean fish radius $r_{\text{fish}} = \frac{1}{4}(h_{\text{fish}} + w_{\text{fish}})$:

$$r^* = \frac{r_{\text{contour}}}{r_{\text{fish}}} \quad (3)$$

where h is height and w width. The distance d between the tip of the fish snout and the anteriormost point of the contour is of greatest interest because this is the distance at which a frontally attacked prey would detect the approaching predator.

Drag is the force applied to any type of body moving through a fluid. It can be expressed as

$$F_{\text{drag}} = F_{\text{pressure}} + F_{\text{friction}}. \quad (4)$$

This drag includes pressure-induced drag forces F_{pressure} and surface friction F_{friction} . F_{pressure} follows from the distribution of pressure values p_i over the fish body, which in turn are calculated incrementally for small, user-defined

Table 5. Boundary conditions of the system of equations 1,2 for pressure p and velocity $\vec{u}$

Parameter	flow inlet	flow outlet	boundary	fish surface
pressure p	$\nabla p = 0$	$p = 0$	$\nabla p = 0$	$\nabla p = 0$
velocity $\vec{u}$	$\vec{u} = (0, u_{\text{fish}}, 0)$	$\nabla \vec{u} = 0$	$\nabla \vec{u} = 0$	$\vec{u} = \vec{0}$

Table 6. Volume V_{fish} and mass m_{fish} of the fishes/fish models used in this study

Parameter	<i>Saurichthys</i>	<i>Belone</i>	<i>Lepisosteus</i>	<i>Ctenolucius</i>	<i>Esox</i>	<i>Oncorhynchus</i>
V_{fish} [cm ³]	940	512	6.20	1.92	392	82
m_{fish} [kg]	1.06	0.58	0.007	0.002	0.44	0.93

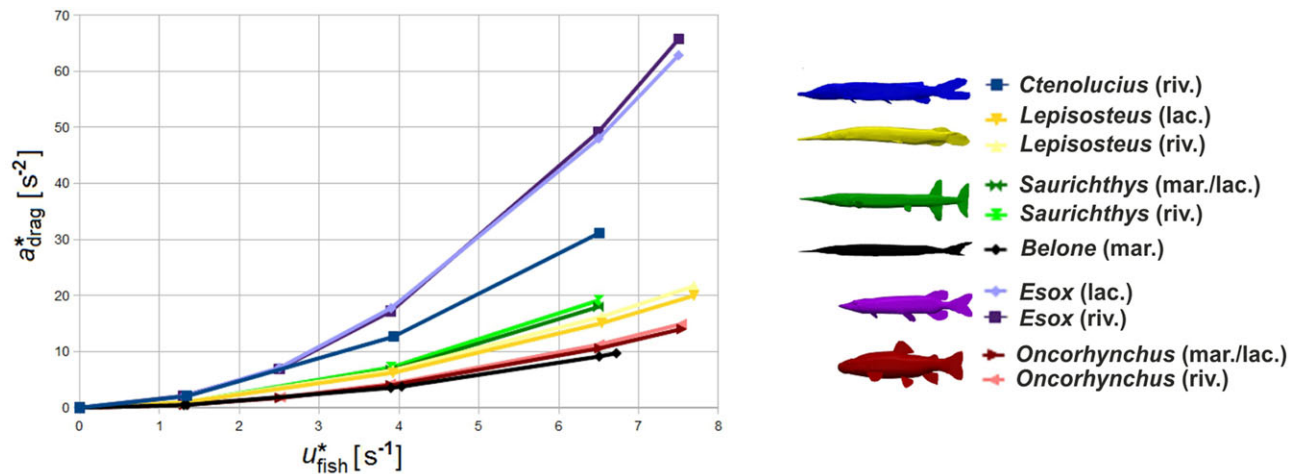


Fig. 11. Gliding abilities. Variation of the gliding deceleration a_{drag}^* over standardised fish velocity (u^* in body length/second) in higher turbulent riverine (riv.) and less turbulent marine/lacustrine (mar./lac.) environments.

surface areas A_i :

$$F_{\text{pressure}} = \sum_{i=1}^n p_i A_i. \quad (5)$$

F_{friction} is approximated considering the surface roughness and thus the fluid adhesion. Surface roughness is derived from the scan data and expressed by the grid mesh on the body surface; the actual scan resolution does not allow for incorporation of finer surface structures in the model. Remaining forces from surface waves or interactions with other objects can be neglected as well as the lift forces. It can be assumed that fishes are able to compensate the lift through control of their swim bladder.

Out of equation 5 the drag coefficient

$$c_{\text{drag}} = \frac{F_{\text{drag}}}{(\rho/2)u_{\text{fish}}^2 A} \quad (6)$$

is calculated, where $(\rho/2)u_{\text{fish}}^2$ is the dynamic pressure that is induced through the inlet velocity u_{fish} and fluid density ρ , and the specified area A is the projected frontal area (A_{front} for calculating $c_{\text{drag,front}}$) or the total wetted surface (A_{surf} to determine $c_{\text{drag,surf}}$), respectively.

Estimation of gliding ability

When predators cease undulating and approach their prey gliding in the final stage of the strike, it may be interesting to estimate how far they can glide before the movement is stopped by drag forces. Calculations of gliding distance have proven difficult in our simulation setting, but drag coefficient and gliding deceleration can be used as appropriate proxies.

Deceleration of a gliding movement without propulsion results from the division of drag forces F_{drag} by the fish mass m_{fish} : $a_{\text{drag}} = (F_{\text{drag}}/m_{\text{fish}})$. Taking into account that most fishes accomplish neutral buoyancy using their swim bladder, we assume that mean density of a fish with swim bladder must be equal to that of water and density of the fish excluding swim bladder should be significantly higher. Weighing the recent fish specimens at our hand led to establishing the following approximation for

the fish mass:

$$m_{\text{fish}} = 1.13 \cdot \rho_{\text{water}} V_{\text{fish}}, \quad (7)$$

where V_{fish} is the volume of the fish and ρ_{water} the density of the replaced water. Fish volumes are determinable from the available surface models (Table 6).

Deceleration is speed-dependent and can be derived from the model only for the initial velocities incorporated in the calculation. The relationship between velocity and deceleration a_{drag} shows an exponential behaviour and can be approximated by the regression:

$$a_{\text{drag}}(u) = x \cdot u_{\text{fish}}^y, \quad (8)$$

where x and y are variable components specific to each fish's body shape. Since $y \approx 2$ for all investigated species, x is given in m^{-1} .

With increasing swimming speed, *Esox*, *Ctenolucius* and *Saurichthys* exhibit the highest deceleration values (Fig. 11). Lowest gliding deceleration is noted for *Belone*. Deceleration is somewhat higher in the riverine environment. Calculation results for *Saurichthys* are given in Table 7.

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Table 7. Inertial gliding deceleration a_{drag} for *Saurichthys* moving in a marine/lacustrine and a riverine environment at three different velocities

	$u_{\text{fish}}=1 \text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=3 \text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=5 \text{ ms}^{-1}$ mar./lac.	$u_{\text{fish}}=1 \text{ ms}^{-1}$ riv.	$u_{\text{fish}}=3 \text{ ms}^{-1}$ riv.	$u_{\text{fish}}=5 \text{ ms}^{-1}$ riv.
$a_{\text{drag}} [\text{ms}^{-2}]$	0.65	5.19	13.84	0.68	5.53	14.74

Competing interests

The authors declare no competing or financial interests.

Author contributions

S.P., C.B., I.K., J.W.S. and M.L. designed the study. S.B. prepared the 3D model of a generalised *Saurichthys*. S.P. and I.K. performed the study. I.K., S.P., M.L. and J.W.S. analysed the results. All authors contributed to the discussion. I.K., S.P. and M.L. wrote the paper.

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References

- Agassiz, L. (1834). Abgerissene Bemerkungen über fossile Fische. *Neues Jahrb. Mineral. Geognosie Geol. Petrefaktenkunde* **1834**, 379–390.
- Bellotti, C. (1857). Descrizione di alcune nuove specie di pesci fossili di Perledo e di altre località lombarde. In *Studi Geologici e Paleontologici sulla Lombardia* (ed. A. Stoppani), pp. 419–438. Milano: Turati.
- Bleckmann, H., Mogdans, J., Engelmann, J., Kröther, S. and Hanke, W. (2004). Wie Fische Wasser fühlen: Das Seitenliniensystem. *Biol. Unserer Zeit* **34**, 358–365.
- Boucot, A. J. and Poinar, G. O., Jr. (2010). *Fossil Behavior Compendium*. Boca Raton: CRC Press.
- Bregazzi, P. R. and Kennedy, C. R. (1980). The biology of pike, *Esox lucius* L., in a southern eutrophic lake. *J. Fish Biol.* **17**, 91–112.
- Brough, J. (1936). On the evolution of bony fishes during the Triassic period. *Biol. Rev.* **11**, 385–405.
- Collette, B. B. (2003). Family Belontiidae Bonaparte 1832 – needlefishes. *Calif. Acad. Sci. Annotated Checklists of Fishes* **16**, 1–22.
- Costa, O. G. (1862). Studi sopra i terreni ad ittiolitti del Regno di Napoli diretti a stabilire l'età geologica dei medesimi. Parte I: Scisti bituminiferi di Giffoni. *Estratta dall'appendice agli atti della Real Accademia delle Scienze* **1862**, 1–44.
- Cuvier, G. and Valenciennes, A. (1849). *Histoire Naturelle des Poissons*, t. 22. Paris: P. Bertrand.
- Di Prinzio, C. Y., Miserendino, M. L. and Casaux, R. (2013). Feeding strategy of the non-native rainbow trout, *Oncorhynchus mykiss*, in low-order Patagonian streams. *Fish. Manage. Ecol.* **20**, 414–425.
- Domenici, P. and Blake, R. (1997). The kinematics and performance of fish fast-start swimming. *J. Exp. Biol.* **200**, 1165–1178.
- Ferziger, J. H. and Peric, M. (2002). *Computational Methods for Fluid Dynamics*. Berlin: Springer.
- Fletcher, T., Altringham, J., Peakall, J., Wignall, P. and Dorrell, R. (2014). Hydrodynamics of fossil fishes. *Proc. R. Soc. B Biol. Sci.* **281**, 20140703.
- Frith, H. R. and Blake, R. W. (1995). The mechanical power output and hydromechanical efficiency of northern pike (*Esox lucius*) fast-starts. *J. Exp. Biol.* **198**, 1863–1873.
- Gardiner, B. G. (1960). A revision of certain actinopterygian and coelacanth fishes, chiefly from the Lower Lias. *Bull. Br. Mus. Nat. Hist. Geol.* **4**, 239–384.
- Greenway, P. (1965). Body form and behavioural types in fish. *Experientia* **21**, 489–498.
- Griffith, J. (1959). On the anatomy of two saurichthyid fishes, *Saurichthys striolatus* (Bronn) and *S. curioni* (Bellotti). *Proc. Zool. Soc. Lond.* **132**, 587–606.
- Griffith, J. (1962). The Triassic fish *Saurichthys krambergeri* Schlosser. *Palaeontology* **5**, 344–354.
- Griffith, J. (1977). The Upper Triassic fishes from Polzberg bei Lunz, Austria. *Zool. J. Linn. Soc.* **60**, 1–93.
- Harper, D. G. and Blake, R. W. (1988). Energetics of piscivorous predator-prey interactions. *J. Theor. Biol.* **134**, 59–76.
- Harper, D. G. and Blake, R. W. (1990). Fast-start performance of Rainbow Trout *Salmo gairdneri* and Northern Pike *Esox lucius*. *J. Exp. Biol.* **150**, 321–342.
- Harper, D. G. and Blake, R. W. (1991). Prey capture and fast-start performance of northern pike *Esox lucius*. *J. Exp. Biol.* **155**, 175–192.
- Hobson, E. S. (1979). Interactions between piscivorous fishes and their prey. In *Predator-Prey Systems in Fisheries Management* (ed. H. E. Clepper), pp. 231–242. Washington, DC: Sport Fishing Institute.
- Kammerer, C. F., Grande, L. and Westneat, M. W. (2006). Comparative and developmental functional morphology of the jaws of living and fossil gars (Actinopterygii: Lepisosteidae). *J. Morphol.* **267**, 1017–1031.
- Keast, A. and Webb, D. (1966). Mouth and body form relative to feeding ecology in the fish fauna of a small lake, Lake Opinicon, Ontario. *J. Fish. Res. Board Can.* **23**, 1845–1874.
- Kogan, I., Schönberger, K., Fischer, J. and Voigt, S. (2009). A nearly complete skeleton of *Saurichthys orientalis* (Pisces, Actinopterygii) from the Madygen Formation (Middle to Late Triassic, Kyrgyzstan, Central Asia) – preliminary results. *PSF 17, Freib. Forsch.-H.* **C532**, 139–152.
- Kogan, I., Romano, C. and Wu, F. X. (2014). Triassic saurichthyid fishes as predators and prey – evidence from the fossil record. *J. Vertebr. Paleontol. Program and Abstracts* **2014**, 161.
- Lauder, G. V. (2006). Locomotion. In *Fish Physiology* (ed. D. H. Evans and J. B. Claiborne), pp. 3–46. Boca Raton: CRC Press.
- Licht, M. (2009). Is it possible to determine a special fish group by one habitat?: a short discussion. *Stud. Geol. Salmant.* **45**, 181–186.
- Linnaeus, C. (1758). *Systema Naturae, per Regna tria Naturae, secundum Classes, Ordines, Genera et Species, cum characteribus, differentiis, synonymis, locis*. Stockholm: Salvius.
- Linnaeus, C. (1761). *Fauna Svecica Sistens Animalia Sveciae Regni: Mammalia, Aves, Amphibia, Pisces, Insecta, Vermes*. Stockholm: Salvius.
- Lombardo, C. and Tintori, A. (2005). Feeding specializations in Late Triassic fishes. *Annali dell'Università degli Studi di Ferrara. Museologia Scientifica e Naturalistica, volume speciale* **2005**, 25–32.
- Maxwell, E. E. and Wilson, L. A. B. (2013). Regionalization of the axial skeleton in the 'ambush predator' guild – are there developmental rules underlying body shape evolution in ray-finned fishes? *BMC Evol. Biol.* **13**, 265.
- Maxwell, E. E., Romano, C., Wu, F. and Furrer, H. (2015). Two new species of *Saurichthys* (Actinopterygii: Saurichthyidae) from the Middle Triassic of Monte San Giorgio, Switzerland, with implications for character evolution in the genus. *Zool. J. Linn. Soc.* **173**, 887–912.
- Owen, R. (1860). *Palaeontology or A systematic Summary of Extinct Animals and Their Geological Relations*. Edinburgh: A. and C. Black.
- Pacholak, S., Hochstein, S., Rudert, A. and Brücker, C. (2014). Unsteady flow phenomena in human undulatory swimming: a numerical approach. *Sports Biomech.* **13**, 176–194.
- Porter, H. T. and Motta, P. J. (2004). A comparison of strike and prey capture kinematics of three species of piscivorous fishes: Florida gar (*Lepisosteus platyrhincus*), redbfin needlefish (*Strongylura notata*), and great barracuda (*Sphyræna barracuda*). *Mar. Biol.* **145**, 989–1000.
- Przybylla, A., Kunze, S., Rudert, A., Bleckmann, H. and Brücker, C. (2010). Entraining in trout: a behavioural and hydrodynamic analysis. *J. Exp. Biol.* **213**, 2976–2986.
- Rand, D. M. and Lauder, G. V. (1981). Prey capture in the chain pickerel, *Esox niger*: correlations between feeding and locomotor behavior. *Can. J. Zool.* **59**, 1072–1078.
- Rieppel, O. (1985). Die Triasfauna der Tessiner Kalkalpen 15. Die Gattung *Saurichthys* (Pisces, Actinopterygii) aus der mittleren Trias des Monte San Giorgio, Kanton Tessin. *Schweiz. Palaeontol. Abh.* **108**, 1–103.
- Rieppel, O. (1992). A new species of the genus *Saurichthys* (Pisces: Actinopterygii) from the Middle Triassic of Monte San Giorgio (Switzerland), with comments on the phylogenetic interrelationships of the genus. *Palaeontographica A* **221**, 63–94.
- Rikardsen, A. H. and Sandring, S. (2006). Diet and size-selective feeding by escaped hatchery rainbow trout *Oncorhynchus mykiss* (Walbaum). *ICES J. Mar. Sci.* **63**, 460–465.
- Romano, C., Kogan, I., Jenks, J., Jerjen, I. and Brinkmann, W. (2012). *Saurichthys* and other fossil fishes from the late Smithian (Early Triassic) of Bear Lake County (Idaho, USA), with a discussion of saurichthyid palaeogeography and evolution. *Bull. Geosci.* **87**, 543–570.
- Schmid, L. and Sánchez-Villagra, M. R. (2010). Potential genetic bases of morphological evolution in the Triassic fish *Saurichthys*. *J. Exp. Zool. B Mol. Dev. Evol.* **314B**, 519–526.
- Schrieffer, J. E. and Hale, M. E. (2004). Strikes and startles of northern pike (*Esox lucius*): a comparison of muscle activity and kinematics between S-start behaviors. *J. Exp. Biol.* **207**, 535–544.
- Stensiö, E. A. (1925). Triassic fishes from Spitzbergen 2. K. Sven. Vetenskapsakad. Handl. 3 Ser. **2**, 1–261.
- Sukhodolov, A., Thiele, M. and Bungartz, H. (1998). Turbulence structure in a river reach with sand bed. *Water Resour. Res.* **34**, 1317–1334.
- Tintori, A. (1990). The vertebral column of the Triassic fish *Saurichthys* (Actinopterygii) and its stratigraphical significance. *Riv. It. Paleont. Strat.* **96**, 93–102.
- Tintori, A. (2013). A new species of *Saurichthys* (Actinopterygii) from the Middle Triassic (Early Ladinian) of the Northern Grigna Mountain (Lombardy, Italy). *Riv. It. Paleont. Strat.* **119**, 287–302.
- Tintori, A., Huang, J.-D., Jiang, D.-Y., Sun, Z.-Y., Motani, R. and Chen, G. (2014). A new *Saurichthys* (Actinopterygii) from the Spathian (Early Triassic) of Chaoahu (Anhui Province, China). *Riv. It. Paleont. Strat.* **120**, 157–164.
- Tytell, E. (2007). Do trout swim better than eels? Challenges for estimating performance based on the wake of self-propelled bodies. *Exp. Fluids* **43**, 701–712.
- Vari, R. P. (1995). The Neotropical fish family Ctenoluciidae (Teleostei: Ostariophysi: Characiformes): supra and intrafamilial phylogenetic relationships, with a revisionary study. *Smithson. Contrib. Zool.* **564**, 1–97.
- Versteeg, H. and Malalasekera, W. (2007). *An Introduction to Computational Fluid Dynamics: The Finite Volume Method*. Harlow: Pearson.
- Wainwright, P. C. and Richard, B. A. (1995). Predicting patterns of prey use from morphology of fishes. *Environ. Biol. Fish.* **44**, 97–113.

- Walbaum, J. J.** (1792). *Petri Artedi sueci Genera piscium. Ichthyologiae pars III. Emendata et aucta a Iohanne Iulio Walbaum.* Grypeswaldiae: Ant. Ferdin. Röse.
- Webb, P. W.** (1975). Hydrodynamics and energetics of fish propulsion. *Bull. Fish. Res. Board Can.* **190**, 1-159.
- Webb, P. W.** (1976). The effect of size on the fast-start performance of rainbow trout *Salmo gairdneri*, and a consideration of piscivorous predator-prey interactions. *J. Exp. Biol.* **61**, 157-177.
- Webb, P. W.** (1984). Body form, locomotion and foraging in aquatic vertebrates. *Am. Zool.* **24**, 107-120.
- Webb, P. W.** (1988). 'Steady' swimming kinematics of Tiger Musky, an Esociform Accelerator, and Rainbow Trout, a Generalist Cruiser. *J. Exp. Biol.* **138**, 51-69.
- Webb, P. W. and Skadsen, J. M.** (1980). Strike tactics of *Esox*. *Can. J. Zool.* **58**, 1462-1469.
- Webb, P. W., Hardy, D. H. and Mehl, V. L.** (1992). The effect of armored skin on the swimming of longnose gar, *Lepisosteus osseus*. *Can. J. Zool.* **70**, 1173-1179.
- Weihs, D.** (1973). The mechanism of rapid starting in slender fish. *Biorheology* **10**, 343-350.
- Weihs, D.** (1989). Design features and mechanics of axial locomotion in fish. *Am. Zool.* **29**, 151-160.
- Werneburg, R., Kogan, I. and Sell, J.** (2014). *Saurichthys* (Pisces: Actinopterygii) aus dem Buntsandstein des Germanischen Beckens. *Semana* **29**, 3-35.
- Woodward, A. S.** (1888). Note on the early Mesozoic ganoid, *Belonorhynchus*, and on the supposed Liassic genus *Amblyurus*. *Ann. Mag. Nat. Hist. Ser. 6* **1**, 354-356.
- Woodward, A. S.** (1890). The fossil fishes of the Hawkesbury series at Gosford. *Mem. Geol. Survey N. S. W. Palaeontol.* **4**, 1-55.
- Woodward, A. S.** (1895). *Catalogue of the Fossil Fishes in the British Museum (Natural History)* **3**. London: British Museum (Natural History).
- Wu, F.-X., Sun, Y.-L., Hao, W.-C., Jiang, D.-Y. and Sun, Z.-Y.** (2015). A new species of *Saurichthys* (Actinopterygii; Saurichthyiformes) from the Middle Triassic of southwestern China, with remarks on pattern of the axial skeleton of saurichthyid fishes. *Neues Jahrb. Geol. Palaeontol. Abh.* **275**, 249-267.