One step further in biomechanical models in palaeontology: a nonlinear finite element analysis review

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ABSTRACT

Finite element analysis (FEA) is no longer a new technique in the fields of palaeontology, anthropology, and evolutionary biology. It is nowadays a well-established technique within the virtual functional-morphology toolkit. However, almost all the works published in these fields have only applied the most basic FEA tools i.e., linear materials in static structural problems. Linear and static approximations are commonly used because they are computationally less expensive, and the error associated with these assumptions can be accepted. Nonetheless, nonlinearities are natural to be used in biomechanical models especially when modelling soft tissues, establish contacts between separated bones or the inclusion of buckling results. The aim of this review is to, firstly, highlight the usefulness of non-linearities and secondly, showcase these FEA tools to researchers that work in functional morphology and biomechanics, as non-linearities can improve their FEA models by widening the possible applications and topics that currently are not used in palaeontology and anthropology.

Subjects Anthropology, Evolutionary Studies, Paleontology, Zoology, Biomechanics
Keywords Finite element analysis, Palaeontology, Anthropology, Non-linear methods, Functional morphology

INTRODUCTION

Computational biomechanics represents the application of computational tools in mechanical problems to study biological systems. During the last decade, computational methods such as finite element analysis (FEA) have been widely used in the field of palaeontology to study biomechanical behaviour of a vast array of fossil species (Rayfield, 2007). However, almost all works published in the field have applied the most basic FEA capabilities, i.e. liner materials in static structural problems, where we can easily define the relationship between physical parameters by means of linear equations. This kind of equations are easy to solve using direct solvers as they exhibit a low computational cost that is a direct response of the size of the finite element mesh. A larger number of nodes in a finite element mesh involves more unknowns in the equation and, consequently, more time is required to solve the mathematical system of equations. Nevertheless, mathematical nonlinearities are natural in physical laws and the assumption of linearity and staticity is a
simplification of reality to make the problem easy to solve. This is because a nonlinear system is characterized with an output that is not proportional to the change of the input. The inclusion of this complexity entails an increase in the computational cost of solving the equations and the need of using iterative solvers. Consequently, palaeontologists have primarily used linear approximations and static problems because these are easier to calculate, computationally faster and solutions can be superposed on each other, hence avoiding an iterative process. In addition, most palaeontologists do not have a deep background on how FEA problems are defined and solved, and many are not aware of the potential of non-linear modelling. For example, including non-linearities in paleoanthropological models can improve the results obtained when modelling skulls by considering sutures, thus making the behaviour of the model closer to reality (Tanaka et al., 2000). They can also be useful in understanding the failure of slender and thin bones or when trying to understand the taphonomic deformations that affected some fossil material by carrying out retrodeformation procedures. Nowadays, most of both commercial and non-commercial FEA packages can solve non-linearities and some examples are already published in living species or in biomedical studies involving humans. Therefore, it makes sense to explore the possibilities that non-linear FEA could provide to the palaeontological and anthropological communities, as these approaches would allow them to explore a broader range of scientific questions, including topics that are currently unsolved or not modelled with enough accuracy. Consequently, the aim of this review is to, firstly, showcase when non-linearities can be useful in functional morphology and secondly, to introduce these methods to researchers that work biomechanics as a way to improving their FEA models by showing them examples that currently are not used in this field. Therefore, this review can be of interest for palaeontologists that seek new ideas in their research, functional morphologists that want to be one step beyond in their research and other researchers who work in the life sciences or in computational mechanics that want to know the state-of-the-art in non-linear FEA applied to biomechanical models.

SEARCH METHODOLOGY

The literature cited in this text is based on a personal selection made by the author to reliably characterise the methods described in the text. A previous search in the Google Scholar database was done to select the appropriate references for the examples. Different keywords were used in each analysed case to fit the expected search. The final selection of the references was based on covering- if possible- diverse animal families, different morphologies, or different fields.

DISCOVERING ALL THE FEA ELEMENTS: SOLIDS, SHELLS, PLATES, BEAMS, SPRINGS, AND TRUSSES

Finite element analysis (FEA) is the mathematical way to solve problems of elasticity in complex geometries by dividing the geometry in small elements where the equations are easy to solve (Zienkiewicz & Taylor, 1981). The equations of elasticity relate the external forces applied in a body to understand how it deforms and how the inner forces are
distributed inside them (Timoshenko & Gere, 1961). The underlying premise of the
method is that a complex geometry can be subdivided into a mesh consisting of a finite
number of elements in which the respective equations are approximately solved
(Marcé-Nogué et al., 2015). This method has been widely used in palaeontology,
anthropology and functional morphology mostly because we can easily digitize bony
structures (Lautenschlager, 2016) to then apply FEA to the obtained geometries. It cannot
be omitted that the generated models are not literal representations of reality, but they still
may be useful for answering scientific questions (Anderson et al., 2012). Following the idea
of simplification there are different kinds of elements that we can use when we are creating
a FEA model (Fig. 1). The use of some of these elements will result in a greater degree of
simplification from reality than others because different simplifications are assumed in
terms of geometry and mechanical behaviour, but also because it involves a reduction of
the complexity of the mathematical equations and the computational time.

The elements **beam** and **spring** or **truss** are used when the original geometry is a line or
can be assumed as a line and the model is defined either in the 3D space or in 2D.
The primary difference between these elements is that beam elements follow beam theory
(Timoshenko, 1955), which enables the calculation of the loads and deflection of beams
subjected to outer forces (including bending, shear, torsion and axial forces). Springs and
truss elements, in contrast, are only designed to handle tensile and compressive forces in
the axial direction of the element. Examples can be found when simplifying the skull of
reptiles and mammals to a beam model (Preuschoft & Witzel, 2002). However, they are not
widely used because these models can be solved most of the time by hand without the need
of a computer. However, the use of springs or truss are widely extended as a complement of
the model when it is necessary to include tendons, ligaments, or other complementary
biological structures of the main model. For example, FEA models of the carpal bones
include spring elements to model the presence of ligaments between bones (Gíslason et al.,
2017).
Shell and plate elements are commonly used when the geometry can be assumed as a surface with constant thickness and the model is defined in 3D space. The difference between shells and plates are that shells are used in curved surfaces and plates only in plane surfaces. Mechanically speaking, both shells and plates can handle bending, but shells develop membrane forces whereas plates do not. This means that shell elements include the membrane effects of resistance to compressive and tensile forces, whereas plates do not. In most of the biological models modelling bone structures, shells have been the preferred option in front of plates. An example of shell elements can be found in works analysing carpal bones (Püschel et al., 2020a) or talar morphologies (Püschel et al., 2020b) because they have a tiny layer of cortical bone with cancellous bone inside where the cortical bone can be assumed as a surface or when modelling something thin as dragonfly wings (Rajabi et al., 2016a).

Another assumption that may further reduce the dimensions of the problem may be simplifying to a surface that lies in a 2D plane using plane elements. I suggest calling them as plane elements because these elements are not really in 2D since they have a constant thickness and use the equations of plane elasticity. When solving the equations of elasticity, plane elasticity refers to the study of specific solutions of the elastic problem in bodies that are surfaces with a constant thickness that are lying in a plane and the forces you apply should lie in this plane. Examples of plane models can be widely found in studies focused on mammal mandibles (Lautenschlager et al., 2020; Marcé-Nogué et al., 2020) or in dinosaurs and other fossils (Neenan et al., 2014; Ma et al., 2021). Plane models also can be useful when modelling other morphologies such as trilobites (Esteve et al., 2021), claws (Patiño, Pérez Zerpa & Fariña, 2019), beaks (Miller et al., 2020) or teeth (Ballell & Ferròn, 2021).

It is important to point out the differences between shell, plate, and plane elements. First, shell elements are not lying in a plane whereas plane and plate elements are. Secondly, plate element allows forces that are not in the plane, like perpendicular forces, supporting bending whereas plane elements do not. This difference can be seen in previous FEA modelling studies of several temnospondyl amphibians (Fortuny et al., 2012) or crocodylomorphs (Pierce, Angielczyk & Rayfield, 2009) where the forces applied are perpendicular to the flat surface of the skull during bilateral cases where plate elements where used.

Finally, solid elements are used when the geometry is a volume, and the model is built in the 3D space. They have been the most widely used in palaeontology and anthropology because they can be easily created from after digitizing a real geometry using CT scanning, photogrammetry, laser scanners, among others. Examples can be found in FEA models of mandibles which have been modelled in 3D (e.g. (Zhou et al., 2019)), unlike the simpler plane models described above. Solid elements can also be found in models of skulls (Zhou et al., 2017), teeth (Benazzi et al., 2012) and a broad range of postcranial (Püschel & Sellers, 2016; McCabe et al., 2017; Bucchi et al., 2020) and other biological structures (Nagel-Myers et al., 2019; Bicknell et al., 2021; Klunk et al., 2021; Krings, Marcé-Nogué & Gorb, 2021).
In general, a nonlinear system is a mathematical system in which the change of the output variable is not proportional to the change of the input variable and, consequently, the equations cannot be written as a linear combination of the unknown variables (Kim, 2015). Therefore, the equations of nonlinear systems are more difficult to solve. A common strategy to deal with them is to approximate the system by linear equations performing multiple iterations to converge to the correct solution (Fig. 2). On the contrary, problems are linear when the output variable is proportional to the change of the input variable. Linearities are found in elastic materials (i.e., following the Hooke’s Law) or when using the small strain theory. This theory is applied when deformations are much smaller than the body dimensions. Therefore, the deformed and undeformed configurations of the body under analysis are assumed to be the same. The equations of continuum mechanics are considerably simplified when applying this assumption by ‘linearising’ (i.e., making linear) the problem to be solved. Non-linearities can be originated by different phenomena in these systems:

1. **Material non-linearity**: When a material is non-linear, the strain it experiences is not proportional to the stress applied i.e., the material does not conform to Hooke’s Law. This situation occurs in plastic or hyperelastic materials where the relationship between stress and strain does not follow a lineal proportion.

2. **Large deformation non-linearity**: The so-called finite strain theory, large strain theory, or large deformation theory is used when strains are large enough to invalidate the assumptions of the small strain theory, which is the theory commonly used in linear elastic problems. In this case, the deformed and undeformed configurations of the body under analysis are notably different, requiring a clear distinction between them in the formulation that, consequently, also affects the relation between stress and strain in the constitutive equation. This theory is common in elastomers and soft tissues and needs to be used when modelling hyperelastic materials.

3. **Large displacement non-linearity**: Also called as geometrical non-linearity, assumes small strains but large rotations and displacements. In the geometrically linear case, the
forces are applied in the undeformed geometry when solving the model whereas in the geometrically nonlinear cases, the applied forces depend on the deformed upcoming geometry. It involves an iterative solution accounting for the displacements and needs to be considered when analysing buckling.

4. **Non-linear contacts:** Separate surfaces of two bodies are in contact without overlapping in such a way that they become mutually tangential. Depending on the relationship between these two surfaces, contacts that allow the separation in the perpendicular direction require a nonlinear solution because there are unknowns at the start of the solving process i.e., where and which force is applied.

The mathematical methods applied to solve general nonlinear functions are all iterative starting from an initial estimation. The solution is obtained by solving iteratively a linearization of the non-linear system in different steps towards the convergence of the solution. Different methods are available depending on the procedure of calculating the increment of the steps: the Newton-Raphson method, the incremental secant method or the incremental force method among others (Kim, 2015). Therefore, the computational cost of the solving procedure of a nonlinear FEA model is now not only affected by the size of the mesh, but also affected by the number of iterative resolutions before convergence.

**Non-linear materials: hyperelasticity and plasticity**

Non-linear materials are materials in which the constitutive equation that defines their behaviours establishes a relationship between stress and strain that is not proportional to a constant. Typical material non-linearities can be found in phenomena such as plasticity and hyperelasticity. Plasticity describes the deformation of a material undergoing non-reversible changes of shape in response to applied forces. In a typical stress-strain curve for a plastic material there is a linear elastic region which satisfies Hooke’s law and a plastic region before fracture that can also follow a linear law or can be defined using different linear sections (Fig. 3). The transition from elastic behaviour to plastic behaviour is called yield and a non-linear solution is required because the solver needs to discover if the body is in the plastic region or not. The total strain is defined by $\varepsilon_{\text{total}} = \varepsilon_{\text{elastic}} + \varepsilon_{\text{plastic}}$ and the value of stress will depend on the value of this total strain. In a biological context,
plasticity can be found in trabecular bone formulations to capture tension-compression asymmetry in the yield strength (Gupta et al., 2007) or, more generally, in studies where a permanent deformation or plasticity is assumed in cortical bone or other biological materials such as dentin, enamel or nacre (An, 2016). In materials such as dentine or enamel, plasticity represents the extremes of the loading environment rather than everyday behaviour.

A hyperelastic material is one that shows extreme elastic behaviour, in that it can return to its original shape even after experiencing very high strains. They are ideal elastic materials in which the stress-strain relationship is non-linear because derives from a strain energy function instead of Hooke’s law (Fig. 3). Moreover, these materials use the large deformation theory already mentioned above. However, the response of the material is not plastic because deformations are fully recoverable. Typical formulations of hyperelastic materials are, among others, phenomenological descriptions of observed behaviour in Mooney–Rivlin and Ogden formulations or equations describing the underlying structure of the material in the Neo–Hookean model (Ogden, 1984). Hyperelastic formulations are common in soft tissues such as ligaments or tendons (Shearer, 2015). Specifically, they can be found in the periodontal ligament (PDL) (Bucchi et al., 2019), muscles such as the pelvic floor (Stansfield et al., 2021), the abdominal muscle (Tu et al., 2019) or generic muscular tissues (Hedenstierna, Hallidin & Brolin, 2008), skin (Ito et al., 2022), corneas (Shan et al., 2010), cartilage (Pataky, Koseki & Cox, 2016), the temporomandibular joint (Sagl et al., 2019) or when modelling blood vessels (Vorp, 2007).

Sometimes the equations that are defined to control soft tissue behaviour include a viscous term (Huang et al., 2017). Viscoelasticity describes the variation of material response within time containing an elastic and a viscous part. The viscous part can describe creep, when stress remains constant and the deformation increases with time, or relaxation, when the deformation remains constant and stress decreases over time. On the other hand, the elastic response is instantaneous and can be defined using a linear material (Booker & Small, 1977) or a nonlinear hyperelastic material (Kulkarni et al., 2016).

More complex models, including fibres in their formulation, exist for the arterial vessels (Gasser, Ogden & Holzapfel, 2006) or the intervertebral discs (Noailly, Planell & Lacroix, 2011) among others. Despite the complexity of these formulae, which combines the overlay of the stiffness in the preferred directions of the fibres with the hyperelastic formulation of the matrix, the constitutive equation is also nonlinear, and it must be solved following an iterative procedure.

**Non-linearities in geometry: buckling**

In a linear problem, the equations of equilibrium are formulated in the original undeformed state and are not updated with the deformation. This is common in most engineering problems because the deformations are small enough to avoid differentiating the original geometry and the deformed one. However, there are cases where the deformation cannot be ignored, and we need to include large displacement non-linearities due to the geometrical update during the application of forces: This is the case of buckling.
Buckling implies a sudden change in shape of a body under load because the loss of stability when this load reaches certain critical value (Fig. 4). If a body—such as a column under compression or a plate under shear, for example—is subjected to a gradually increasing load, when the load reaches the critical value, the body may suddenly change shape. Although buckling appears before failure, it can be decisive in the ergonomics of certain biological bodies, limiting the range of forces under which they are able to remain functional. Buckling is caused by nonlinearities in the geometry and can be approached by a linearisation that drives to a bifurcation problem of eigenvalues. Therefore, the linear buckling analysis is done in parallel to a linear elastic analysis. Otherwise, the full nonlinear solution of the point of collapse can be obtained by increasing the load in smaller steps with an iterative method while the geometry is updated to its deformed state. This latter is significantly more computationally expensive but might be more accurate than the linear buckling. In a biomechanical context, buckling can be found when study slender bodies such as the swordfish rostrum (Habegger et al., 2020), the weevil rostrum (Matsumura et al., 2021) or even in bones under compression such as the vertebrae (Williams et al., 2021). Buckling is also considered in humans as a cause of fractures of postcranial bones (Lee et al., 2009).

**Non-linearities in contacts**

Contacts between two bodies are divided between **linear contacts** and **non-linear contacts**. Linear contacts can be included in a linear elastic model without modifying the solving mode and keeping the direct solution. It also involves a low computational cost that is simply a function of the size of the finite element mesh (namely, the number of elements and nodes). However, the inclusion of non-linear contacts changes the solving mode to a non-linear solution with an iterative solver, increasing the computational cost of the analysis. Contacts can be described according to the relationship between the two separate surfaces of each body that become mutually tangential in five general different types.
according to how they can move perpendicularly to each other and how they can move in the tangential plane. In other words, if they are allowed to separate and slide (Fig. 5).

1. **Bonded contacts**: when separation and sliding is not allowed. It is a linear contact.
2. **No-separation contact**: when separation is not allowed but sliding in the tangential plane is allowed. It is a linear contact.
3. **Frictionless contact**: when separation and sliding is allowed. It is a non-linear contact.
4. **Rough contact**: when separation is allowed but sliding in the tangential plane is not allowed. It is a non-linear contact.
5. **Frictional contact**: when separation is allowed but sliding in the tangential plane is controlled by a friction coefficient. It is a non-linear contact.

Frictional contact can be understood as an intermediate status, where sliding in the tangential plane is not free but is allowed and bonded contact is used when we have two bodies that are perfectly joined but they are created or defined as separate bodies during the FEA modelling. For example can be used for defining all the pieces involving a teeth such as the cortical bone, dentine, enamel, pulp and the PDL. (Benazzi et al., 2013; Bucchi et al., 2019)

In general, contacts are found in FEA models involving more than one body and the definition of each contact depends on the nature of its behaviour. It can be found in models when studying the carpal bones of the wrist (Gíslason et al., 2017; Püschel et al., 2020a) or the feet (Ito et al., 2022), the ossicles of the auditory system (Marcé-Nogué & Liu, 2020), the intervertebral discs and the vertebræ of the spine (Guan et al., 2019), all the tissues in the hip (Fleps et al., 2018) or the patella (Fitzpatrick & Rullkoetter, 2012), the mandible, the tempomandibular joint and the skull (Sagl et al., 2019) or the interaction between the
bodies in the wings of dragonflies (Rajabi et al., 2016b), bees and wasps (Eraghi et al., 2021), among many others. Therefore, contacts can be used to establish relationship between bones or soft tissues. Contacts are also useful when studying occlusal forces during mastication to model the interaction between teeth and food (Skamniotis, Elliott & Charalambides, 2019) or even the impact of eggshells with the floor (Sellés et al., 2019).

**SUMMARY: IDEAS FOR PALAEOONTOLOGISTS**

FEA is no longer an incipient technique in the fields of palaeontology, anthropology, and evolutionary biology. Instead, it is nowadays a well-established technique within functional virtual morphology toolkit that has been used in more than 750 biological and evolutionary publications between 2005 and 2020 (Tseng, 2021). Most of this works present FEA models without non-linearities. This is not necessarily a problem by itself if the linear approach is sufficient to answer the scientific question of interest. Indeed, many engineering problems can be solved without trespassing the threshold of the linear models. Therefore, this text does not want to spread an incorrect idea regarding the use of supposedly more accurate non-linear models. In fact, the use of linear and not expensive computational approaches without nonlinearities can be certainly useful to understand the behaviour of many biological systems under analysis. For instance, most of the FEA works that include fossils have focused on the study of skeletal elements that can be successfully modelled using linear elastic material properties and solved using a static analysis under small strains and displacements, *i.e.* without the need of non-linearities. However, although linear models can be used in a broad range of functional works, the aim of this text is to highlight the value of non-linearities when they can be of utility, or they are needed to improve the knowledge we have in fields such as palaeontology and anthropology.

**Non-linear soft tissues**

Little is known about soft tissue properties in fossils. The direct examination of fossil soft tissues and preserved blood cells is of little value when studying palaeontological remains due to the degradation or the contamination from modern remains (van Dongen et al., 2017). The reconstruction of soft tissues from fossils is an issue that it is unresolved but can be approached through investigating extant relatives to infer the palaeo-physiology of extinct taxa, *e.g.*, via the phylogenetic bracketing approach (Witmer, 1995). Therefore, any FEA models can potentially include an inference regarding soft tissues properties. As an example, cranial sutures are deformable joints between adjacent bones bridged by collagen fibres and there are several works on fossil taxa that have include soft tissues modelling sutures in *Tyrannosaurus rex* skull (Cost et al., 2020), australopithecines (Dzialo et al., 2014) or in dicynodonts (Jasinoski, Rayfield & Chinsamy, 2009), as well as FEA models of current lizard species (Dutel et al., 2021), Sphenodon (Curtis et al., 2013) or even some mammals (Bright & Gröning, 2011). All of these examples used linear material properties to characterize the elastic behaviour of soft tissues which can be an appropriate simplification if this is validated experimentally (Bright & Gröning, 2011). However a recent diagnosis suggested that the lack of sutures or and inappropriate modelling can
result in inaccurate results of stress, strain or deformation (Rayfield, 2019) although it is not clear how the soft tissue can be accurately predicted in fossils (Broyde et al., 2021). It is at this point that the researcher needs, at least, to be aware that a more accurate modelling of these soft tissues should be done using nonlinear material properties, which in turn involve an increase in the computational cost of the model. Unfortunately, soft tissue models in living animals have not been extensively documented, with the exception of biomedical contexts (Tanaka et al., 2000). Hopefully, studies focused on experimentally testing non-linear models in diverse biological taxa will be carried out in the near future.

**Plasticity in retrodeformations**

Retrodeformation is very common in fossil taxa as the process that produces the original form of the taxon prior to fossil diagenesis when this has been recovered in any deformed way. Deformation in fossils is produced due to a multiple array of taphonomic and tectonic processes. Overburden stress due to the weight of the overlying sediments linearly compacts the fossil from above causing the fossil to break and/or warp. Other causes of fossil deformation include tectonic stresses and sediment cracking. Under the action of these loads, the fossil can break in a brittle manner or can be distorted plastically, preserving the structure of the fossil due to the lack of breakage. Fossils under plastic deformation, where forces applied during time modify the original shape of the bone structure may be restored. Although there are several techniques to virtually restore deformed specimens available without using mechanical equations (Lautenschlager, 2016), it has sense to use methods from mechanics such as FEA that involve forces if one want to infer which was the actual process that drove the fossil to be deformed (Arbour & Currie, 2012; Di Vincenzo et al., 2017). Modelling this phenomenon would require including the nonlinear plastic behaviour of bone, because retrodeformation is a permanent deformation in cortical bone. In this case FEA could be inversely applied by defining the plastic behaviour of the fossil material and then, setting the forces applied at the fossil as the unknowns of the problem. It would allow to answer the question of which forces do we need to apply deformed bodies to recover its original form.

**Buckling in slender bones**

In palaeontology there are a lot of slender structures that are susceptible to be analysed using buckling. Probably the most common and useful case would be in bones under compression such as the leg bones of large, heavy dinosaurs and mammals. This is because mass is considered as one of the main factors affecting the morphology and osteological adaptation of these bones (Etienne et al., 2020). To understand how these bones are adapted to the heavy weight that they needed to support, evaluation of the maximal stress as a measure of bone strength is not the only informative metric (Hutchinson, 2021). This is because bone may fail without involving fracture. Alternatively, bone could fail through buckling if it is not stiff enough (Currey, 2012). In this case, buckling needs to be considered, because it can cause the collapse of the legs before the fracture of the bone. Usually, buckling reduces the capacity of the strength of the structure because it appears in
a lower value than the yield stress and the fracture stress that defines the strength of the material.

If we assume that the leg bones in heavy dinosaurs as slender columns like those from a building, Euler’s critical load is defined as the compressive load at which the column will suddenly buckle (Timoshenko, 1955). This equation can give clues about the relationship between geometrical factors such as the length of the bones or how they are joined to the articulations. Given that the length, material, or boundary conditions cannot be modified from the original model, Euler’s critical force will depend on the second moment of area or moment of inertia. Increasing the value of the critical force implies a modification of the cross-section of the bone through more inertial geometries. Therefore, if we assume the cross-section of leg bones as an annulus, thicker annulus will increase the inertia. But also, if the thickness is kept constant, a broader annulus will increase the inertia of the cross section. This simple consequence can be obtained assuming leg bones with a straight morphology not close enough to the reality, but very useful for the purpose of study. However, in case of analysis of the real and irregular geometry of the bones, the simple formula of Euler cannot be used but the problem of buckling can be solved via computational methods by means of FEA solving an eigenvalue problem. Few works are paying attention to it, discarding the effect of buckling in the morphology of the long bones in living mammals (Brassey et al., 2013). Considering than an eigenvalue problem in a FEA model is not increasing the computational cost of the analysis too much, it would be worth to more exhaustively test if the leg bones of heavy dinosaurs or mastodontic extant mammals are affected by buckling, as was suggested in horses (Currey, 2003). Another case where buckling could be a concern is in bones with a high aspect ratio where the walls are substantially thinner (De Margerie et al., 2005). This could be the case of wing bones (Palmer & Dyke, 2012), and hence buckling should be also explored.

**Bone grouping using contacts**

Functional implications of fossil bones have been widely studied in fossil taxa using FEA models (Richmond et al., 2005). Depending on the purpose, bones can be studied alone or as a group and the main difference between these two cases is the absence or presence of contacts. When separation between bones is not desired, for example in the analysis of teeth, considering the bonding of the cortical bone, dentine, enamel, pulp and the PDL (Benazzi et al., 2013), the contacts used are linear and it does not require an increase in the computational cost of the solving process. This is something that can be considered when creating FEA models because it allows the inclusion of several bones in the model without nonlinearities.

On the other side, nonlinear contacts allow separation between the bones. Although the inclusion of this contacts implies an iterative solution through convergence, it may be necessary to implement when a group of bones need to be studied together such as the carpal bones of the wrist (Gíslason et al., 2017) or the bones of the foot (Ito et al., 2022). This has been done extensively in biomechanical models of living primates; therefore, it should be considered in other FEA works in the field of the palaeontology and anthropology. In fact, the literature is full of biomechanical analysis of kinematics and
dynamics of solid bodies where bones of fossils are grouped to study its performance (Sellers et al., 2017; Bishop et al., 2021). Therefore, it makes sense when creating FEA models, to include more than one bone in the model if it can be useful for the desired analysis despite increasing the computational cost of the solution. Also when the contact between bones is through articular cartilage, the contact can be defined between cartilages that are also in contact with the bone (Püschel et al., 2020a).

Models with shells, plates, beams, springs, and trusses

Finally, although this is not related with the use of a nonlinear iterative solving, the use of other kind of elements other than solid elements can have a great advantage when dealing with nonlinear models. This is because they provide a useful way to reduce the number of elements and nodes of the FEA mesh and, consequently, a reduction of the time spent solving the equation in each iteration. Hence, a nonlinear model will particularly benefit from the use of these elements.

The use of shell elements to model cortical bone in morphologies that can be assumed as thin and with a constant thickness, such as carpal bones or talar morphologies, require a lower number of elements and nodes because there is only one mesh layer. Using solid elements in the same morphology would need at least four or five layers of elements would along the thickness to properly build an adequate mesh to accurately capture the results. This was used in an analysis of carpal bones (Gíslason et al., 2017) to model both the cortical bone and the articular cartilage, reducing significantly the number of elements to allow a smooth non-linear solution, due to the presence of non-linear contacts. The same example uses non-linear spring elements to model the behaviour of the ligaments. This decision is also in favour of not increasing the number of nodes and elements of the model, because spring or truss elements can be defined using only one element with the origin and final nodes. In this manner, the model avoids the inclusion of a three-dimensional geometry modelled with solid elements for each ligament, which would exponentially increase the number of nodes and elements in the mesh and consequently, increase the computational cost of the solution.

When creating FEA models of fossils and considering the inclusion of some of the non-linearities it is a good option to evaluate if the use of simpler elements can reduce computational cost. Although researchers should be aware of the potential ramifications of simplifying their models, it is also true that any model will necessarily not represent a literal representation of reality. Instead, the requirements necessary to answer the research question of interest should always be kept in mind when making decisions about model complexity.

Summarizing, this text highlighted the usefulness of non-linearities in FEA palaeontological and anthropological models in spite of increasing their complexity and the computational costs. Nowadays, most of the commercial and non-commercial FEA packages include the resolution of non-linear problems in their capabilities, and they also documented with tutorials and examples how to deal with them. Therefore, the main aim of this review is to provide a road map for the next generation of palaeontologists, anthropologists, and functional morphologists by showing them unexplored ways that
could a profound impact in finite element analysis and how they can explore these methods further.

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Competing Interests
The author declares that they have no competing interests.

Author Contributions
Jordi Marcé-Nogué conceived and designed the experiments, performed the experiments, analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the article, conceived the text, and approved the final draft.

Data Availability
The following information was supplied regarding data availability:
This is a literature review; there is no raw data or code.

REFERENCES
An B. 2016. Constitutive modeling the plastic deformation of bone-like materials. *International Journal of Solids and Structures* 92–93(6):1–8 DOI 10.1016/j.ijsolstr.2016.05.003.
Anderson PSL, Bright JA, Gill PG, Palmer C, Rayfield EJ. 2012. Models in palaeontological functional analysis. *Biology Letters* 8(1):119–122 DOI 10.1098/rsbl.2011.0674.
Arbour VM, Currie PJ. 2012. Analyzing taphonomic deformation of ankylosaur skulls using retrodeformation and finite element analysis. *PLOS ONE* 7(6):e39323 DOI 10.1371/journal.pone.0039323.
Ballell A, Ferrón HG. 2021. Biomechanical insights into the dentition of megatooth sharks (*Lamniformes: Otodontidae*). *Scientific Reports* 11(1):1232 DOI 10.1038/s41598-020-80323-z.
Benazzi S, Kullmer O, Grosse IR, Weber GW. 2012. Brief communication: comparing loading scenarios in lower first molar supporting bone structure using 3D finite element analysis. *American Journal of Physical Anthropology* **147**(1):128–134 DOI 10.1002/ajpa.21607.

Benazzi S, Nguyen HN, Kullmer O, Hublin J-J. 2013. Unravelling the functional biomechanics of dental features and tooth wear. *PLOS ONE* **8**(7):e69990 DOI 10.1371/journal.pone.0069990.

Bicknell RDC, Holmes JD, Edgecombe GD, Losso SR, Ortega-Hernández J, Wroe S, Paterson JR. 2021. Biomechanical analyses of Cambrian euarthropod limbs reveal their effectiveness in mastication and durophagy. *Proceedings of the Royal Society B: Biological Sciences* **288**(1943):20202075 DOI 10.1098/rspb.2020.2075.

Bishop PJ, Falisse A, De Groote F, Hutchinson JR. 2021. Predictive simulations of running gait reveal a critical dynamic role for the tail in bipedal dinosaur locomotion. *Science Advances* **7**(39):1–14 DOI 10.1126/sciadv.abi7348.

Booker JR, Small JC. 1977. Methods for the numerical solution of the equations of viscoelasticity. *International Journal for Numerical and Analytical Methods in Geomechanics* **1**(2):139–150 DOI 10.1002/nag.1610010203.

Bucchi C, Marcé-Nogué J, Galler KM, Widbiller M. 2019. Biomechanical performance of an immature maxillary central incisor after revitalization: a finite element analysis. *International Endodontic Journal* **52**(10):1508–1518 DOI 10.1111/iej.13159.

Bucchi A, Püschel TA, Lorenzo C, Marcé-Nogué J. 2020. Finite element analysis of the proximal phalanx of the thumb in Hominoidea during simulated stone tool use. *Comptes Rendus Palevol* **19**(2):26–39 DOI 10.5852/cr-palevol2020v19a2.

Cost IN, Middleton KM, Sellers KC, Echols MS, Witmer LM, Davis JL, Holliday CM. 2020. Palatal biomechanics and its significance for cranial kinesis in tyrannosaurus rex. *The Anatomical Record* **303**(4):999–1017 DOI 10.1002/ar.24219.

Currey JD. 2003. How well are bones designed to resist fracture? *Journal of Bone and Mineral Research* **18**(4):591–598 DOI 10.1359/jbmr.2003.18.4.591.

Currey JD. 2012. The structure and mechanics of bone. *Journal of Materials Science* **47**(1):41–54 DOI 10.1007/s10853-011-5914-9.

Curtis N, Jones MEH, Evans SE, O’Higgins P, Fagan MJ. 2013. Cranial sutures work collectively to distribute strain throughout the reptile skull. *Journal of the Royal Society Interface* **10**(86):1–9 DOI 10.1098/rsif.2013.0442.

De Margerie E, Sanchez S, Cubo J, Castanet J. 2005. Torsional resistance as a principal component of the structural design of long bones: comparative multivariate evidence in birds. *Anatomical Record – Part A Discoveries in Molecular, Cellular, and Evolutionary Biology* **282**:49–66 DOI 10.1002/ar.a.20141.

Di Vincenzo F, Profico A, Bernardini F, Cerroni V, Dreossi D, Schlager S, Zao P, Benazzi S, Biddittu I, Rubini M, Tuniz C, Manzi G. 2017. Digital reconstruction of the Ceprano
calvarium (Italy), and implications for its interpretation. *Scientific Reports* 7(1):13974 DOI 10.1038/s41598-017-14437-2.

Dutel H, Gröning F, Sharp AC, Watson PJ, Herrel A, Ross CF, Jones MEH, Evans SE, Fagan MJ. 2021. Comparative cranial biomechanics in two lizard species: impact of variation in cranial design. *Journal of Experimental Biology* 224(5):717 DOI 10.1242/jeb.234831.

Dzialo C, Wood SA, Berthaume MA, Smith AL, Dumont ER, Benazzi S, Weber GW, Strait DS, Grosse IR. 2014. Functional implications of squamosal suture size in *Paranthropus boisei*. *American Journal of Physical Anthropology* 153(2):260–268 DOI 10.1002/ajpa.22427.

Eraghi SH, Toofani A, Khaheshi A, Khorsandi M, Darvizeh A, Gorb S, Rajabi H. 2021. Wing coupling in bees and wasps: from the underlying science to bioinspired engineering. *Advanced Science* 2004383(16):2004383 DOI 10.1002/advs.202004383.

Esteve J, Marcé-Nogué J, Pérez-Peris F, Rayfield EJ. 2021. Cephalic biomechanics underpins the evolutionary success of trilobites. *Palaeontology* 64(4):519–530 DOI 10.1111/pala.12541.

Etienne C, Mallet C, Cornette R, Houssaye A. 2020. Influence of mass on tarsus shape variation: a morphometrical investigation among Rhinocerotidae (*Mammalia: Perissodactyla*). *Biological Journal of the Linnean Society* 129(4):950–974 DOI 10.1093/biolinnean/blaa005.

Fitzpatrick CK, Rullkoetter PJ. 2012. Influence of patellofemoral articular geometry and material on mechanics of the unresurfaced patella. *Journal of Biomechanics* 45(11):1909–1915 DOI 10.1016/j.jbiomech.2012.05.028.

Fleps I, Enns-Bray WS, Guy P, Ferguson SJ, Cripton PA, Helgason B. 2018. Correction: on the internal reaction forces, energy absorption, and fracture in the hip during simulated sideways fall impact (PLoS ONE (2018) 13:8 (e0200952) DOI: 10.1371/journal.pone.0200952). *PLOS ONE* 13(11):e0208286 DOI 10.1371/journal.pone.0208286.

Fortuny J, Marcé-Nogué J, Gil L, Galobart À. 2012. Skull mechanics and the evolutionary patterns of the otic notch closure in capitosaurs (*Amphibia: Temnospondylia*). *The Anatomical Record: Advances in Integrative Anatomy and Evolutionary Biology* 295(7):1134–1146 DOI 10.1002/ar.22486.

Gasser TC, Ogden RW, Holzapfel GA. 2006. Hyperelastic modelling of arterial layers with distributed collagen fibre orientations. *Journal of The Royal Society Interface* 3(6):15–35 DOI 10.1098/rsif.2005.0073.

Guan W, Sun Y, Qi X, Hu Y, Duan C, Tao H, Yang X. 2019. Spinal biomechanics modeling and finite element analysis of surgical instrument interaction. *Computer Assisted Surgery* 24(sup1):151–159 DOI 10.1080/24699322.2018.1560086.

Gupta A, Bayraktar HH, Fox JC, Keaveny TM, Papadopoulos P. 2007. Constitutive modeling and algorithmic implementation of a plasticity-like model for trabecular bone structures. *Computational Mechanics* 40(1):61–72 DOI 10.1007/s00466-006-0082-5.

Gislason MK, Foster E, Bransby-Zachary M, Nash DH. 2017. Biomechanical analysis of the Universal 2 implant in total wrist arthroplasty: a finite element study. *Computer Methods in Biomechanics and Biomedical Engineering* 20(10):1113–1121 DOI 10.1080/10255842.2017.1336548.

Habegger L, Motta P, Huber D, Pulaski D, Grosse I, Dumont ER. 2020. Feeding biomechanics in billfishes: investigating the role of the rostrum through finite element analysis. *The Anatomical Record* 303(1):44–52 DOI 10.1002/ar.24059.

Hedenstierna S, Halldin P, Brolin K. 2008. Evaluation of a combination of continuum and truss finite elements in a model of passive and active muscle tissue. *Computer Methods in Biomechanics and Biomedical Engineering* 11(6):627–639 DOI 10.1080/17474230802312516.
Huang H, Tang W, Tan Q, Yan B. 2017. Development and parameter identification of a visco-hyperelastic model for the periodontal ligament. *Journal of the Mechanical Behavior of Biomedical Materials* 68:210–215 DOI 10.1016/j.jmbbm.2017.01.035.

Hutchinson JR. 2021. The evolutionary biomechanics of locomotor function in giant land animals. *Journal of Experimental Biology* 224(11):125 DOI 10.1242/jeb.217463.

Ito K, Nakamura T, Suzuki R, Negishi T, Oishi M, Nagura T, Jinzaki M, Ogihara N. 2022. Comparative functional morphology of human and chimpanzee feet based on three-dimensional finite element analysis. *Frontiers in Bioengineering and Biotechnology* 9:1–13 DOI 10.3389/fbioe.2021.760486.

Jasinoski SC, Rayfield EJ, Chinsamy A. 2009. Comparative feeding biomechanics of Lystrosaurus and the generalized dicynodont Oudenodon. *Anatomical Record* 292(6):862–874 DOI 10.1002/ar.20906.

Kim N. 2015. *Introduction to nonlinear finite element analysis.* New York, NY, USA: Springer.

Klunk CL, Argenta MA, Casadei-Ferreira A, Economo EP, Pie MR. 2021. Mandibular morphology, task specialization and bite mechanics in Pheidole ants (*Hymenoptera: Formicidae*). *Journal of The Royal Society Interface* 18(179):20210318 DOI 10.1098/rsif.2021.0318.

Krings W, Marcé-Nogué J, Gorb SN. 2021. Finite element analysis relating shape, material properties, and dimensions of taeniolissus radular teeth with trophic specialisations in Paludomidae (*Gastropoda*). *Scientific Reports* 11(1):22775 DOI 10.1038/s41598-021-02102-8.

Kulkarni SG, Gao XL, Horner SE, Mortlock RF, Zheng JQ. 2016. A transversely isotropic visco-hyperelastic constitutive model for soft tissues. *Mathematics and Mechanics of Solids* 21(6):747–770 DOI 10.1177/1081286514536921.

Lautenschlager S. 2016. Reconstructing the past: methods and techniques for the digital restoration of fossils. *Royal Society Open Science* 3(10):160342 DOI 10.1098/rsos.160342.

Lautenschlager S, Figueirido B, Cashmore DD, Bendel E-M, Stubbs TL. 2020. Morphological convergence obscures functional diversity in sabre-toothed carnivores. *Proceedings of the Royal Society B: Biological Sciences* 287(1935):20201818 DOI 10.1098/rspb.2020.1818.

Lee T, Choi JB, Schafer BW, Segars WP, Eckstein F, Kuhn V, Beck TJ. 2009. Assessing the susceptibility to local buckling at the femoral neck cortex to age-related bone loss. *Annals of Biomedical Engineering* 37(9):1910–1920 DOI 10.1007/s10439-009-9751-9.

Ma W, Pittman M, Butler RJ, Lautenschlager S. 2021. Macroevolutionary trends in theropod dinosaur feeding mechanics. *Current Biology* 32(3):1–10 DOI 10.1016/j.cub.2021.11.060.

Marcé-Nogué J, Fortuny J, Gil L, Sánchez M. 2015. Improving mesh generation in finite element analysis for functional morphology approaches. *Spanish Journal of Palaeontology* 31:117–132 DOI 10.7203/sjp.30.1.17227.

Marcé-Nogué J, Liu J. 2020. Evaluating fidelity of CT based 3D models for Zebrafish conductive hearing system. *Micron* 135:102874 DOI 10.1016/j.micron.2020.102874.

Marcé-Nogué J, Püschel TA, Daasch A, Kaiser TM. 2020. Broad-scale morpho-functional traits of the mandible suggest no hard food adaptation in the hominin lineage. *Scientific Reports* 10(1):6793 DOI 10.1038/s41598-020-63739-5.

Matsumura Y, Jafarpour M, Reut M, Shams Moattar B, Darvizeh A, Gorb SN, Rajabi H. 2021. Excavation mechanics of the elongated female rostrum of the acorn weevil Curculio glandium (*Coleoptera; Curculionidae*). *Applied Physics A: Materials Science and Processing* 127(5):1–11 DOI 10.1007/s00339-021-04353-8.

McCabe K, Henderson K, Pantinople J, Richards HL, Milne N. 2017. Curvature reduces bending strains in the quokka femur. *PeerJ* 5(3):e3100 DOI 10.7717/peerj.3100.
Miller CV, Pittman M, Kaye TG, Wang X, Bright JA, Zheng X. 2020. Disassociated rhamphotheca of fossil bird Confuciusornis informs early beak reconstruction, stress regime, and developmental patterns. *Communications Biology* 3(1):519 DOI 10.1038/s42003-020-01252-1.

Nagel-Myers J, Mastorakos I, Yuya P, Reeder G. 2019. Modelling crushing crab predation on bivalve prey using finite element analysis. *Historical Biology* 33:1373–1382 DOI 10.1080/08912963.2019.1699555.

Neenan JM, Ruta M, Clack JA, Rayfield EJ. 2014. Feeding biomechanics in Acanthostega and across the fish–tetrapod transition. *Proceedings of the Royal Society B: Biological Sciences* 281(1781):20132689 DOI 10.1098 rspb.2013.2689.

Noailly J, Planell JA, Lacroix D. 2011. On the collagen criss-cross angles in the annuli fibrosi of lumbar spine finite element models. *Biomechanics and Modeling in Mechanobiology* 10(2):203–219 DOI 10.1007/s10237-010-0227-5.

Ogden RW. 1984. *Non-linear elastic deformations*. Mineola: Dover.

Palmer C, Dyke G. 2012. Constraints on the wing morphology of pterosaurs. *Proceedings of the Royal Society B: Biological Sciences* 279(1731):1218–1224 DOI 10.1098 rspb.2011.1529.

Pataky TC, Koseki M, Cox PG. 2016. Probabilistic biomechanical finite element simulations: whole-model classical hypothesis testing based on upcrossing geometry. *PeerJ Computer Science* 2:e96 DOI 10.7717/peerj-cs.96.

Patiño S, Pérez Zerpa J, Fariña RA. 2019. Finite element and morphological analysis in extant mammals’ claws and quaternary sloths’ ungual phalanges. *Historical Biology* 33:1–11 DOI 10.1080/08912963.2019.1664504.

Pierce SE, Angielczyk KD, Rayfield EJ. 2009. Shape and mechanics in thalattosuchian (Crocodylomorpha) skulls: implications for feeding behaviour and niche partitioning. *Journal of Anatomy* 215(5):555–576 DOI 10.1111/j.1469-7580.2009.01137.x.

Preuschoft H, Witzel U. 2002. Biomechanical investigations on the skulls of reptiles and mammals. *Senckenbergiana Lethaea* 82(1):207–222 DOI 10.1007/BF03043785.

Püschel TA, Marcé-Nogué J, Chamberlain AT, Yoxall A, Sellers WI. 2020a. The biomechanical importance of the scaphoid-centrale fusion during simulated knuckle-walking and its implications for human locomotor evolution. *Scientific Reports* 10(1):3526 DOI 10.1038/s41598-020-60590-6.

Püschel TA, Marcé-Nogué J, Gladman J, Patel BA, Alméciya S, Sellers WI. 2020b. Getting its feet on the ground: elucidating paralouatta’s semi-terrestriality using the virtual morpho-functional toolbox. *Frontiers in Earth Science* 8:79 DOI 10.3389/feart.2020.00079.

Püschel TA, Sellers WI. 2016. Standing on the shoulders of apes: analyzing the form and function of the hominoid scapula using geometric morphometrics and finite element analysis. *American Journal of Physical Anthropology* 159(2):325–341 DOI 10.1002/ajpa.22882.

Rajabi H, Rezasefat M, Darvizeh A, Dirks J-H, Eshghi S, Shafiei A, Mostofi TM, Gorb SN. 2016a. A comparative study of the effects of constructional elements on the mechanical behaviour of dragonfly wings. *Applied Physics A* 122(1):19 DOI 10.1007/s00339-015-9557-6.

Rajabi H, Shafiei A, Darvizeh A, Dirks J-H, Appel E, Gorb SN. 2016b. Effect of microstructure on the mechanical and damping behaviour of dragonfly wing veins. *Royal Society Open Science* 3(2):160006 DOI 10.1098/rsos.160006.

Rayfield EJ. 2007. Finite element analysis and understanding the biomechanics and evolution of living and fossil organisms. *Annual Review of Earth and Planetary Sciences* 35(1):541–576 DOI 10.1146/annurev.earth.35.031306.140104.
Rayfield EJ. 2019. What does musculoskeletal mechanics tell us about evolution of form and function in vertebrates? The Science and Regulations of Naturally Derived Complex Drugs. Berlin: Springer International Publishing, 45–70.

Richmond BG, Wright BW, Grosse I, Dechow PC, Ross CF, Spencer MA, Strait DS. 2005. Finite element analysis in functional morphology. The Anatomical Record Part A: Discoveries in Molecular, Cellular, and Evolutionary Biology 283A:259–274 DOI 10.1002/ar.a.20169.

Sagl B, Schmid-Schwamp M, Piehslinger E, Kundi M, Stavness I. 2019. A dynamic jaw model with a finite-element temporomandibular joint. Frontiers in Physiology 10:1–12 DOI 10.3389/fphys.2019.01156.

Sellers WI, Pond SB, Brassey CA, Manning PL, Bates KT. 2017. Investigating the running abilities of Tyrannosaurus rex using stress-constrained multibody dynamic analysis. PeerJ 5(79):e3420 DOI 10.7717/peerj.3420.

Sellés AG, Marcé-Nogué J, Vila B, Pérez MA, Gil L, Galobart À, Fortuny J. 2019. Computational approach to evaluating the strength of eggs: implications for laying in organic egg production. Biosystems Engineering 186(4):146–155 DOI 10.1016/jbiosystemseng.2019.06.017.

Shan G, Visentin P, Elsheikh A, Ballo A, Moritz N, Zhao J, Liao D. 2010. Biomechanics: principles, trend and applications. New York: Nova Science Publishers, Inc.

Shearer T. 2015. A new strain energy function for the hyperelastic modelling of ligaments and tendons based on fascicle microstructure. Journal of Biomechanics 48(2):290–297 DOI 10.1016/j.jbiomech.2014.11.031.

Skamniotis CG, Elliott M, Charalambides MN. 2019. Computer simulations of food oral processing to engineer teeth cleaning. Nature Communications 10(1):3571 DOI 10.1038/s41467-019-11288-5.

Stansfield E, Kumar K, Mitteroecker P, Grunstra NDS. 2021. Biomechanical trade-offs in the pelvic floor constrain the evolution of the human birth canal. Proceedings of The National Academy of Sciences of The United States of America 118(16):e2022159118 DOI 10.1073/pnas.2022159118.

Tanaka E, Miyawaki Y, Del Pozo R, Tanne K. 2000. Changes in the biomechanical properties of the rat interparietal suture incident to continuous tensile force application. Archives of Oral Biology 45(12):1059–1064 DOI 10.1016/S0003-9969(00)00082-0.

Timoshenko S. 1955. Strength of materials. New York: Van Nostrand.

Timoshenko SP, Gere JM. 1961. Theory of elastic stability. New York: McGraw-Hill.

Tseng ZJ. 2021. Rethinking the use of finite element simulations in comparative biomechanics research. PeerJ 9(5):e11178 DOI 10.7717/peerj.11178.

Tuset L, Fortuny G, Herrero J, Puigjaner D, López JM. 2019. Implementation of a new constitutive model for abdominal muscles. Computer Methods and Programs in Biomedicine 179(9395):104988 DOI 10.1016/j.cmpb.2019.104988.

van Dongen B, Manning PL, Warwood S, Buckley M, Kitchener AC. 2017. A fossil protein chimera; difficulties in discriminating dinosaur peptide sequences from modern cross-contamination. Proceedings of the Royal Society B: Biological Sciences 284(1855):20170544 DOI 10.1098/rspb.2017.0544.

Vorp DA. 2007. Biomechanics of abdominal aortic aneurysm. Journal of Biomechanics 40(9):1887–1902 DOI 10.1016/j.jbiomech.2006.09.003.

Williams CJ, Pani M, Bucchi A, Smith RE, Kao A, Keeble W, Ibrahim N, Martill DM. 2021. Helically arranged cross struts in azhdarchid pterosaur cervical vertebrae and their biomechanical implications. iScience 24(4):102338 DOI 10.1016/j.isci.2021.102338.
Witmer LM. 1995. The extant phylogenetic bracket and the importance of reconstructing soft tissues in fossils. In: Thomason J, ed. Functional Morphology in Vertebrate Paleontology. Cambridge: Cambridge University Press.

Zhou Z, Fortuny J, Marcé-Nogué J, Skutschas PP. 2017. Cranial biomechanics in basal urodèles: the Siberian salamander (Salamandrella keyserlingii) and its evolutionary and developmental implications. Scientific Reports 7(1):10174 DOI 10.1038/s41598-017-10553-1.

Zhou Z, Winkler DE, Fortuny J, Kaiser TM, Marcé-Nogué J. 2019. Why ruminating ungulates chew sloppily: biomechanics discern a phylogenetic pattern. PLOS ONE 14(4):e0214510 DOI 10.1371/journal.pone.0214510.

Zienkiewicz OC, Taylor RL. 1981. Finite element method – the basis (Volume 1). New York: McGraw-Hill.