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EXPERIMENTAL ANALYSIS OF SOFT-TISSUE FOSSILIZATION: OPENING THE BLACK BOX

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Abstract: Taphonomic experiments provide important insights into fossils that preserve the remains of decay-prone soft tissues, tissues that are usually degraded and lost prior to fossilization. These fossils are among the most scientifically valuable evidence of ancient life on Earth, giving us a view into the past that is much less biased and incomplete than the picture provided by skeletal remains alone. Although the value of taphonomic experiments is beyond doubt, a lack of clarity regarding their purpose and limitations, and ambiguity in the use of terminology, are hampering progress. Here we distinguish between processes that promote information retention and those that promote information loss, in order to clarify the distinction between fossilization and preservation. Recognizing distinct processes of decay, mineralization and maturation, the sequence in which they act, and the potential for interactions, has important consequences for analysis of fossils, and for the

design of taphonomic experiments. The purpose of well-designed taphonomic experiments is generally to understand decay, maturation and preservation individually, thus limiting the number of variables involved. Much work remains to be done, but these methodologically reductionist foundations will allow researchers to build towards more complex taphonomic experiments and a more holistic understanding and analysis of the interactions between decay, maturation and preservation in the fossilization of non-biomineralized remains. Our focus must remain on the key issue of understanding what exceptionally preserved fossils reveal about the history of biodiversity and evolution, rather than on debating the scope and value of an experimental approach.

Key words: decay, fossilization, taphonomy, experiment, exceptional preservation.

ARGUABLY the most scientifically valuable evidence of ancient life on Earth comes from fossils preserving remains of the decay-prone soft tissues (e.g. integument and muscle) that are usually degraded and lost prior to fossilization. These examples of ‘exceptional preservation’, and the fossil biotas from which they are recovered, represent invaluable fossil archives, giving us a view into the past that is much less biased and incomplete than the partial picture provided by skeletal remains alone. Recent technological and methodological advances have allowed the acquisition of progressively more detailed anatomical and chemical data on fossil soft tissues and, as a result, reports of high fidelity preservation of anatomy (at micro- and macro-scales) and biomolecules are expanding the known limits of morphological and chemical fossil preservation. One component of current

advances in the field is a reinvigoration of experimental investigations into the taphonomy of non-biomineralized organisms and tissues: laboratory-based analyses of post-mortem decay, maturation and mineralization, and the implications for processes of fossilization of soft tissue remains and biomolecules (e.g. Raff *et al.* 2008; Sansom *et al.* 2010; Sansom *et al.* 2011; Cunningham *et al.* 2012a, b; McNamara *et al.* 2013; Murdock *et al.* 2014; Coleary *et al.* 2015; Naimark *et al.* 2016). This type of taphonomic experiment, focused on non-biomineralized remains, is the subject of this contribution; throughout, all references to ‘taphonomic experiments’ exclude those designed to address questions of skeletal taphonomy. We use the terms ‘soft tissues’ and ‘non-biomineralized tissues’ interchangeably, and to include sclerotized tissues.

The application of experimental taphonomy to exceptional preservation has developed over several decades (see Briggs & McMahon 2016 for a recent review), and has made major contributions to our understanding of how non-biomineralized tissues become fossilized. Experiments have provided significant insights, for example, into preservation of soft tissues through maturation and stabilization of organic compounds (e.g. Gupta *et al.* 2006; Gupta *et al.* 2009), processes of microbially mediated authigenic mineralization (e.g. Sagemann *et al.* 1999), microbial pseudomorphing of soft tissues (e.g. Raff *et al.* 2008, 2013), and how non-random patterns of anatomical decay can introduce systematic biases into the interpretation of exceptionally preserved fossils (e.g. Sansom *et al.* 2010; Murdock *et al.* 2014).

The value of experimental and analytical approaches applied to the study of exceptionally preserved fossils is beyond doubt, but we identify two issues that are hampering further progress. First, a lack of clarity regarding the purpose and limits of experimental approaches, and second, ambiguity in the use of terminology. More precise use of language and greater clarity regarding the rationale for conducting taphonomic experiments will allow researchers to focus on the key issue of what exceptionally preserved fossils reveal about the history of biodiversity and evolution, rather than the scope and value of an experimental approach.

DECAY, MATURATION, PRESERVATION AND FOSSILIZATION

Clarity regarding ‘fossilization’ and ‘preservation’ are clearly crucial to taphonomic analysis, otherwise we risk confusing processes with results, yet the terms are used interchangeably by some authors and to mean distinct and different things by others.

Fossilization is one outcome of the range of processes that affect an organism after death (Fig. 1). These processes cumulatively result in both loss and retention of information, and can balance out in different ways, with the most obvious alternative outcome to fossilization being non-fossilization: the loss of features or complete absence from the fossil record. Every part of every organism ends up somewhere on a spectrum from partial to complete non-fossilization. We focus here on decay, maturation and mineralization as distinct processes resulting in information retention (preservation) and information loss.

Decay is the post-mortem process by which original biomolecules, tissues and structures are degraded and lost through abiotic processes (such as chemical thermodynamics) and biotic processes, such as autolysis and microbially mediated decomposition. For many

researchers, the antithesis of decay is preservation, but preservation is not the same thing as fossilization (see below).

Preservation refers to the processes that directly result in retention of information (Fig. 1); processes by which inorganic and/or organic chemical activity replicates the form or converts the remains of non-biomineralized tissues into minerals (mineralization) and organic compounds that are stable over geological timescales (maturation). This conversion can be via replacement or replication by minerals (mineralization), chemical transformation through maturation of organic compounds, or a combination. Different types of preservation may occur at different stages of decay and maturation. This definition of preservation is neither new nor out of step with its widespread use in the taphonomic literature; it serves merely to clarify the distinction between preservation and fossilization.

Mineralization is replacement or replication of non-biomineralized tissues by minerals. It can occur at any stage post-mortem, pre- or post-burial, early or late, and different modes of mineralization can act at different times (and under different conditions) in the taphonomic history of a fossil. Different modes of mineralization can occur in different parts within the same carcass, linked to differences in microenvironments (McNamara *et al.* 2009). Although many exceptionally preserved fossils have been mineralized, mineralization is not a requirement for exceptional fossilization.

Maturation of organic remains occurs post-mortem, mostly post-burial, primarily in the diagenetic realm, and can involve processes resulting in degradation and loss of biological information and/or processes resulting in stabilization of organic compounds such that they can survive over geological timescales. Maturation thus includes elements of both loss and retention of information, with preservation of organic remains through maturation involving *in situ* polymerization of more labile compounds (e.g. Gupta *et al.* 2006, 2009; McNamara *et al.* 2016).

Secondary transformations might be considered as an additional stage in the post-mortem history of a fossil. Like maturation, they can involve either loss or retention of information. Mineral replacements from early stages of preservation can be lost or altered by subsequent chemical activity under different conditions of diagenesis (iron sulphides converted to iron oxides, for example). And organic remains stabilized through low temperature polymerization might be oxidized, depolymerized or otherwise mobilized and lost during later, higher grades of maturation. Information loss through weathering will also be a factor, but we do not consider this further here.

Focusing on these processes of decay, mineralization and maturation clarifies the distinction between fossilization and preservation: fossilization of an organism’s remains is one outcome of the balance and interactions

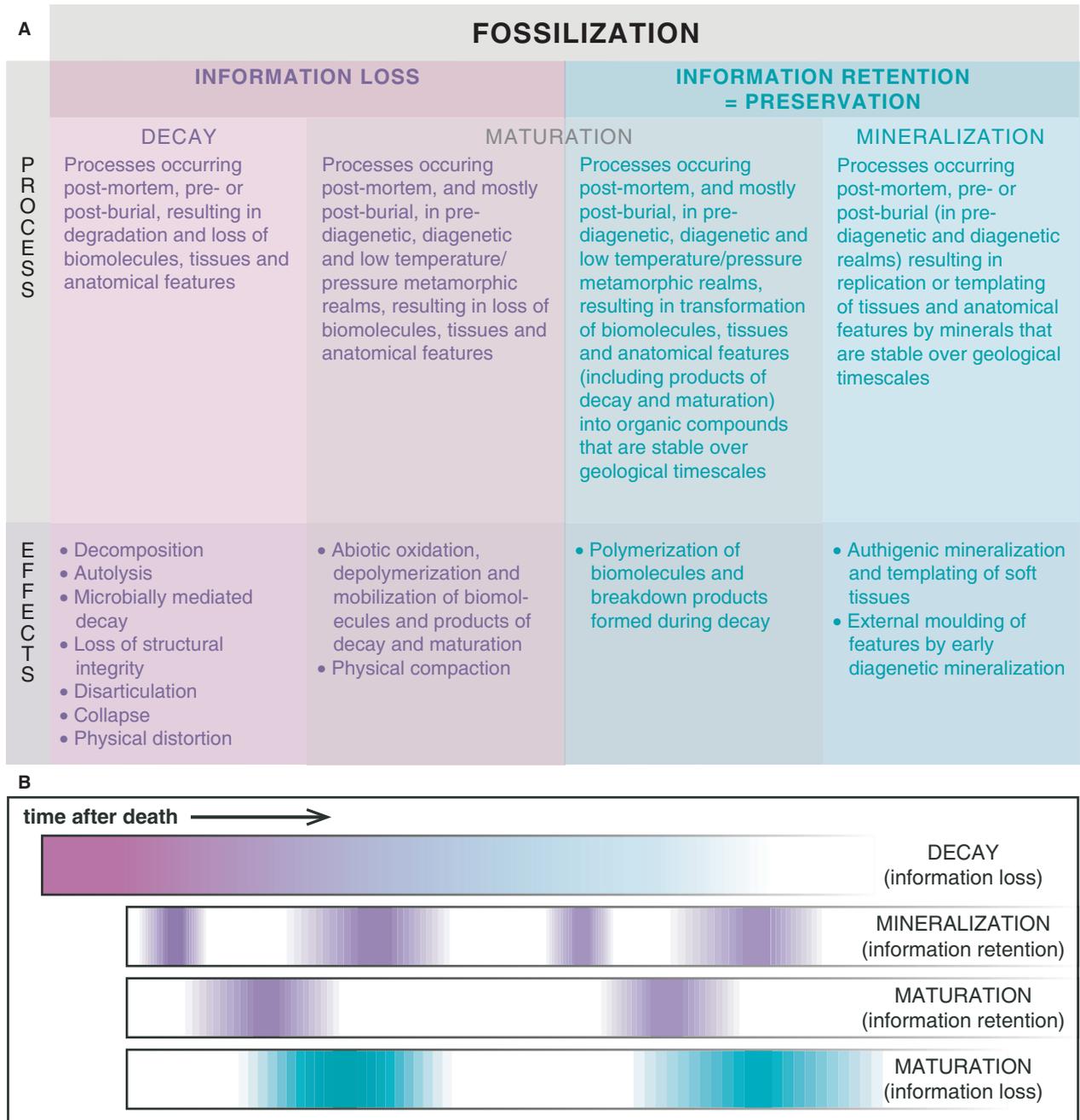


FIG. 1. A, terminology for processes involved in fossilization. B, sequence of action, effects and potential timespan over which processes act. Processes are not continuous; intervals when processes are operating more intensely are shown by more intense colours (B); the relative timing and duration of periods of more intense loss and retention of information (via mineralization and maturation) are schematic. Decay starts before other processes but the potential timespan over which it operates is shorter. It is irreversible, and the rate of decay is not constant. Mineralization can start before maturation, but not before decay has commenced, and can occur at any subsequent point, although not continuously (multiple phases of mineralization are possible). Mineralization that post-dates decay can only preserve information previously retained through either mineralization or maturation. Maturation, both information loss and retention, can start before or after mineralization and can occur at any subsequent point. That maturation is a process that promotes both information loss and information retention does not imply separation of these processes in fossilization. We do not attempt to show interactions between processes; e.g. information loss and information retention through maturation do not necessarily occur simultaneously (although maturation can affect different tissues differently). Late stage information loss through weathering processes is not shown. ‘Information’, in this context, refers to primary anatomical, microanatomical and biochemical data, not information regarding the geological processes of fossilization and palaeoenvironment.

between processes of information loss (decay and maturation) and preservation (information retention via maturation and mineralization). This balance can be expressed as two very simple equations:

1. processes of information loss > processes of information retention = no fossil
2. processes of information loss < processes of information retention = fossil

The balance and outcome reflected in equation 1 are far more prevalent than those of equation 2.

Original biological tissues, or components thereof, are either lost or transformed into materials that are stable over geological timescales, and the existence of a fossil, even the most exquisitely preserved, does not imply survival of its biological information without alteration and loss. In these terms, 'exceptional preservation' is shorthand for *part of the process* that results in exceptional fossilization; we should not ignore the unexceptional aspects of information loss.

THE RATIONALE FOR CONDUCTING TAPHONOMIC EXPERIMENTS

The goal of the vast majority of taphonomic experimental analyses is not 'experimental fossilization': their aim is not to *replicate* the fossilization process in the laboratory or field. This is because fossilization involves many variables, including both known and unknown unknowns, and multiple confounding variables mean that (if our focus is on understanding the processes of fossilization) the results of such fossil replication experiments, whether in general or for specific Lagerstätten, are unlikely to tell us much of interest. While their success could be evaluated in terms of whether they produce something that might look more or less like a fossil, the experiments can reveal little if anything about the various processes involved in fossilization. This is not to say that observations of what happens when organisms decay under natural conditions have no place; these observations can provide useful constraints on the design of taphonomic experiments, or guidance on what a fossil jellyfish might look like, for example. But crude experiments that attempt to replicate fossilization without controlling variables are, in effect, treating fossilization as a black box (Fig. 2). To understand the processes that control information loss and information retention, the processes which ultimately produce the outcome of fossilization, we need to see inside the box.

This is the focus of robust taphonomic experiments: experimental decay, experimental maturation, and experimental preservation. The purpose of well-designed taphonomic experiments is generally to understand these processes individually (thus limiting the number of

variables involved and making experiments tractable). In particular, robust taphonomic experiments investigate how specific variables (such as environmental pH, availability of ions, diffusion rate, burial temperature, etc.) have potentially affected the loss and retention of anatomical information and biased exceptionally preserved biotas.

A simplifying initial assumption of this reductionist approach is that there are no direct causal links between processes of decay, maturation and preservation. However, it is important to note that this assumption applies only to the *design* of taphonomic experiments and does not preclude finding evidence of links between the various processes of decay, preservation and maturation, either in fossil data or in experimental results (a good example being taphonomic experiments demonstrating that certain forms of mineralization require decay to establish the geochemical gradients across which they operate; Sagemann *et al.* 1999). In general, analysing and understanding each of these processes is a prerequisite for clear understanding of potential interactions between them.

Recognizing the distinction between processes, the sequence in which they act (Fig. 1), and the potential for interactions, has important consequences for analysis of fossils; missing this point has led to some unjustified criticism of experimental approaches. There is no reason to assume, for example, that sequences of decay should provide a guide to sequences of preservation: decay commences first (Fig. 1) and provides a timeline and pattern of morphological modification and loss; the products of decay, in the form of incomplete carcasses and decay-modified characters, are the substrate upon which the processes of maturation and mineralization act. It is the timing and interplay between processes of loss and retention that govern what anatomical structures, tissues and biomolecules are ultimately fossilized, and this can be unravelled only if we understand the taphonomic patterns resulting from decay. Similarly, claims that decay experiments are not applicable to the fossil record because either the environment or the experimental taxon used is not a good analogue for what occurred in deep time are missing the point of controlled taphonomic experiments. The same is true of criticisms that taphonomic experimental models are not informative because fossilization in each deposit, taxon and even specimen reflects a suite of unique conditions. Were these criticisms to be levelled at experiments designed to transform carcasses into fossils, we would agree, but they carry little weight as arguments against the validity of taphonomic experiments as an approach to modelling the general parameters that control decay, maturation and mineralization and their respective roles in fossilization across environments and taxa.

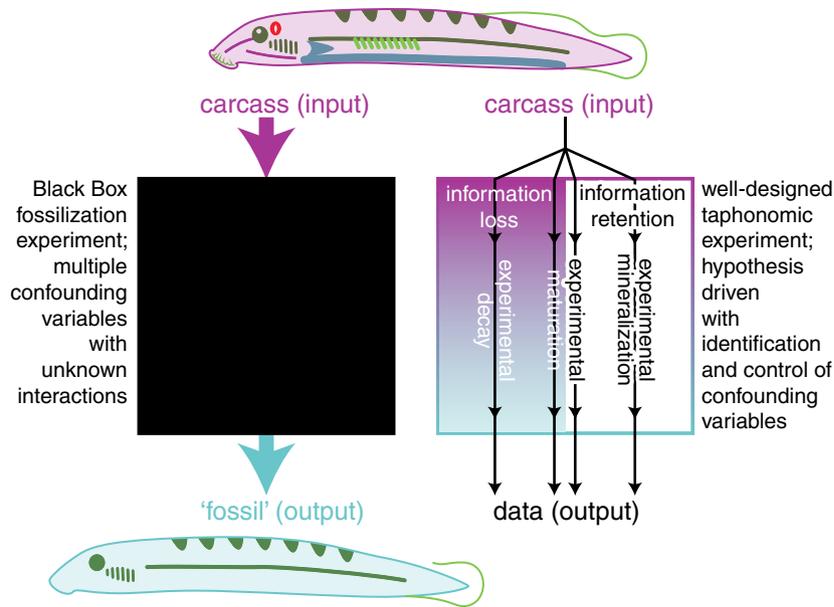


FIG. 2. Cartoon illustrating the difference between experiments that attempt to replicate fossilization, treating the process as a Black Box, and those that focus on the processes of decay, maturation or mineralization. The Black Box approach reveals little about the processes of information loss and information retention, the cumulative effects and interactions of which ultimately results in a fossil (or, more often, not). Well-designed taphonomic experiments do not attempt to replicate fossilization and do not result in a fossil, but provide reproducible data concerning the processes of fossilization. An ideal taphonomic experiment, with all variables identified and controlled for, is rarely attainable in practice.

TAPHONOMIC EXPERIMENTS AND COMPARATIVE ANATOMY OF EXCEPTIONALLY PRESERVED FOSSILS

It is important to point out here that taphonomic experiments can be designed to address a range of different questions. For some, the goal is to understand the environmental conditions in which organisms decayed and were ultimately preserved (e.g. Plotnick 1986; Kidwell & Baumiller 1990; Hellawell & Orr 2012), or the degree to which a fossil biota is diminished in diversity by the loss of soft bodied organisms. In such cases taphonomic signatures are a proxy for palaeoenvironment or for faunal completeness, and the loss of anatomical information translates into gains in geological data.

For much recent work, however, the purpose of taphonomic experiments is to provide better data upon which to base interpretations of the anatomy of fossil organisms and, in turn, more accurate reconstructions of phylogenetic relationships and evolutionary patterns. These endeavours rely on comparative anatomical analysis, and when applied to exceptionally preserved non-biomineralized fossils, this analysis is predicated on the assumption that differences between taxa are not simply the result of random taphonomic processes. This is a crucial point that is often overlooked, particularly in the most fundamental elements of comparative anatomy: the individuation of body parts and characters, and determination of the suite of characters

present in a taxon (see Rieppel & Kearney 2002 for discussion of individuation). Comparative analysis is possible only if the similarities and differences in characters reflect original anatomy and, critically, if taphonomic factors can be detected and taken into account. Furthermore, because comparative analysis ultimately requires comparison with extant organisms, investigators must distinguish differences that arise because of evolutionary history from those that simply reflect the incompleteness of the fossil (Donoghue & Purnell 2009). This decision can be based either on intuitions and assumptions, or on the crucial evidence generated by taphonomic experiments. We advocate the latter approach.

TAPHONOMY, EXCEPTIONAL PRESERVATION, AND EXPERIMENTS: A WAY FORWARD

The processes of decay, maturation and mineralization are controlled by diverse factors that vary both spatially and temporally in how they act. A focus of future studies must be on deconvolving the relative impact of these processes on our reading of the fossil record of exceptionally preserved organisms and particular Lagerstätten. In many exceptionally preserved fossils, the suite of characters present does not correspond to what might be predicted from a simplistic application of experimental decay results

(i.e. the fossils are not simply a collection of the most decay resistant body parts). This is because the effects of preservational processes, sometimes highly selective with regard to tissue types, are superimposed upon the results of decay; it is false reasoning to suggest that fossilization of more than just decay resistant remains in itself indicates that patterns and sequences of character transformation and loss observed in experimental decay deviate from those that occurred in particular fossils. To use a specific example, invertebrate nervous tissues decay rapidly under controlled experimental conditions (e.g. Murdock *et al.* 2014; Sansom *et al.* 2016), yet mounting evidence supports the interpretation that some exceptionally preserved biotas include specimens with fossilized nervous tissues (e.g. Ma *et al.* 2012; Strausfeld *et al.* 2016). There is no conflict here: the fossils do not falsify the experiments, and the experiments do not falsify the anatomical interpretations of the fossils. Together they highlight a gap in our understanding of how nervous tissues become fossilized (Murdock *et al.* 2014). We argue that decay experiments are the best starting point for understanding the biases and filters of fossilization because it is upon decayed remains that the processes of maturation and preservation must operate.

Further work is also required to better understand the processes of mineralization and authigenic mineral replication of original tissue. Despite experimental studies of the processes of replication of soft tissues in calcium phosphate (Briggs & Kear 1993*a, b*; Briggs & Wilby 1996) and in pyrite (Grimes *et al.* 2001, 2002) other authigenic minerals have received little attention (but see Martin *et al.* 2004; McCoy *et al.* 2015). Soft tissues may be preserved via multiple pathways in a single fossil (e.g. Butterfield 2002; McNamara *et al.* 2009) but the controlling factors have yet to be elucidated experimentally. Future studies focusing on these preservational processes will be especially critical for attempts to extract tissue-specific and taxonomic signatures from fossil specimens that preserve different tissues in different minerals. Similarly, greater understanding of how microbial communities are mediating both decay and mineralization is likely to yield significant new insights into preservational biases.

We have attempted here to clarify the essential elements and the terminology that form the conceptual framework of taphonomic experiments, and the value of considering decay, maturation, mineralization, and preservation as distinct but interacting processes. Greater emphasis on the rationale for conducting particular experiments will enhance experimental design, allowing taphonomists to construct more tightly constrained models of taphonomic processes. This reductionist approach allows us to see beyond the Black Box view of fossilization, and from these foundations we can build towards a

more holistic understanding of the roles of, and interactions between, decay, maturation and preservation in the fossilization of non-biomineralized remains.

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