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Lazar, O. R., K. M. Bohacs, J. Schieber, J. H. S. Macquaker, and T. M. Demko, 2022, Tools and techniques for studying mudstones, in K. M. Bohacs and O. R. Lazar, eds., Sequence stratigraphy: Applications to fine-grained rocks: AAPG Memoir 126. p. 35–88.

Tools and Techniques for Studying Mudstones

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What Tools and Techniques Should Be Used? Carefully define the questions and equally carefully select the tools and techniques that will give the best answers within a time—cost framework.

—Potter et al., 1980, p. 102

ABSTRACT

Mudstone properties vary widely, but systematically, both vertically and laterally at millimeter-to-kilometer scales. This variability can be detected by applying a range of physical, petrophysical, chemical, and paleontological methods to characterize the rocks at different scales and interpreting the resulting data using the sequence-stratigraphic approach.

This chapter concentrates on introducing the tools and techniques that provide data about texture, bedding, composition, and grain origin, which enable the characterization of mudstone strata at millimeter-to-kilometer scales. Specifically, it addresses the application of such tools and techniques to decipher depositional conditions and construct sequence-stratigraphic frameworks. This chapter provides key references for the tools and techniques commonly used to generate the texture, bedding, composition, and grain-origin data for further consideration. It then outlines our approach toward making detailed and systematic observations of the key attributes of mudstones in outcrops, cores, and thin sections. It concludes with an overview of the key sequence-stratigraphic concepts that we find useful for studying mudstones and a summary of the methods we use to construct and test a sequence-stratigraphic framework for mudstones. This approach is elaborated and illustrated in the following chapters.

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INTRODUCTION

Mudstones are important sources, reservoirs, and seals of hydrocarbons and dominate the sedimentary record (e.g., Sorby, 1908; Picard, 1971; Potter et al., 1980; Stow, 1981; Blatt, 1982; Klemme and Ulmishek, 1991; Curtis, 2002, 2009). Mudstone properties vary systematically—both vertically and laterally at millimeter-to-kilometer scales. This variability is controlled by the changes in the depositional environment and burial history, and is recorded in the stratal stacking (e.g., Bohacs and Schwalbach, 1992; Bohacs, 1998; Schieber, 1998a, b, 1999; Macquaker et al., 1998, 2007; Bohacs et al., 2005, 2014; Lazar, 2007; Lazar et al., 2010, 2015a, b). Sequence stratigraphy provides a genetic framework within which the varying source, reservoir, and seal attributes of mudstones can be predicted. The attributes of mudstones that need to be recorded to utilize this method are their (1) texture (grain size), (2) bedding, (3) composition, and (4) grain origin (see Lazar et al., 2022a, Chapter 2 this Memoir). Information regarding these attributes reveals essential aspects of the processes that form, transport, deposit, and then modify the component grains and their associated pore spaces in all depositional and burial environments.

TOOLS

This section covers visual, geochemical, well-log, and seismic tools and techniques. Visual tools include optical inspection, digital imaging, and optical and electron microscopy. Geochemical tools comprise bulk, elemental, and molecular techniques. Well-log tools include techniques for estimating lithology, porosity, pressure, hydrocarbon presence and type, and structural characteristics (fractures, faults, folds, in situ stress). Seismic tools cover seismic stratigraphy and facies analyses and mapping.

Visual Tools

You can observe a lot by just looking.
—Berra, 2010

Careful visual observation at the macro-to-microscale is the essential first step in characterizing mudstones, selecting appropriate samples for additional analyses, and deciphering the depositional conditions. We recommend that mudstone be examined visually at a millimeter-to-centimeter scale in outcrops, cores, and thin sections to determine texture, bedding, and composition, as well as physical sedimentary structures;

type, diversity, and abundance of body and trace fossils; bioturbation index; taphonomy of macrofossils; type, habit, and distribution of diagenetic products; fractures; and color (Lazar et al., 2015a, b; Lazar et al., 2022a, Chapter 2 this Memoir).

Visual observations should be made on surfaces of fresh rock in outcrops and cores (Figure 1), and then supplemented with observations made using tools that include high-intensity white and ultraviolet light, and optical, electronic, and digital imaging of hand specimens and thin sections. Observations should include the abundance of each physical, biological, and chemical attribute to assist in the identification of facies, facies associations, and associated stacking patterns (e.g., Bohacs et al., 2014; Lazar et al., 2015a, b). Visual observations can be facilitated and enhanced by using the tools and approaches that are discussed next.

Preparation of Hand Specimens and Core Slabs

A clean, dry rock surface is an essential prerequisite to any form of visual examination. The time spent preparing fine-grained rocks is amply rewarded by the many details revealed (Figure 1). A list of tools and supplies we recommend for preparing fresh rock surfaces and describing mudstones in outcrops and cores is given in Table 1. We recommend the following tools and approaches for preparing mudstone surfaces for visual examination:

- Sharp-rock saw: Prepare a flat, relatively polished surface. A portable concrete saw can be used in the field to acquire continuous, perpendicular-tobedding slot samples; samples should be stabilized with fiberglass matting and epoxy resin prior to removal (e.g., Schieber, 1998b).
- Clean: Scrub and polish using a paint brush, scrub brush, abrasive paper or sponge (polished using a series of sandpaper of grades from no. 80 to no. 800 grit); scrubbing pad (softer—white and harder—green); or pencil eraser (~1 in. wide, pink or white). Pencil erasers can be quite effective for removing dust, precipitated minerals, or oil-based mud from slabbed-core pieces. Beware of rocks that are sensitive to water—minimize exposure time and dry each piece immediately, using compressed air, if possible, or a clean absorbent material (e.g., paper towel). Use a dilute degreasing solution on cores taken with oil-based mud.
- Contrast enhancement techniques: Several approaches are available that are fast, simple, and useful for visualizing sedimentary details if further geochemical analyses of the studied rocks are not planned. Many of these approaches are

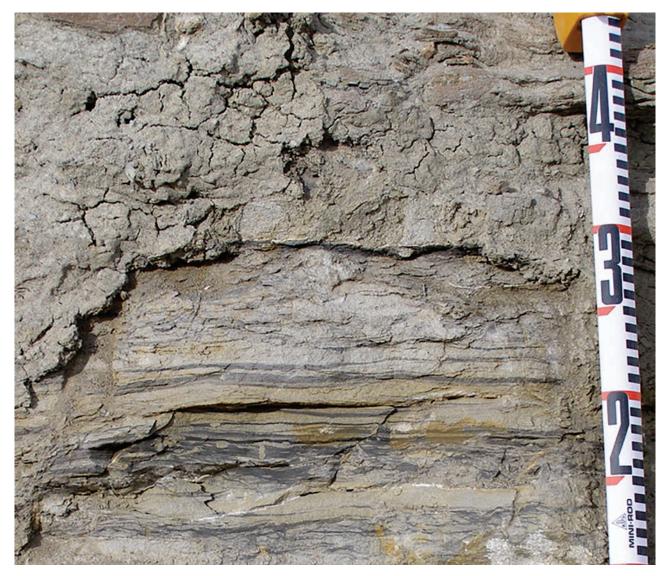


Figure 1. Example of the Cretaceous Mowry Shale in outcrop, Wyoming, contrasting weathered and fresh surfaces. A variety of sedimentary features become apparent in mudstones once we obtain a fresh surface of an outcrop or core.

summarized in the literature (e.g., Ekdale et al., 1984; Miller, 1988; Scholle and Ulmer-Scholle, 2003; Flügel, 2010); some of these approaches are:

- Etching in dilute hydrochloric acid (Flügel, 2010): excellent for enhancing texture.
- Wetting with water (Frey, 1970): excellent for emphasizing bioturbation.
- Staining with an ink: Alizarin Red S (e.g., Hamblin, 1962; Farrow, 1966), a mixture of India ink and carborundum powder (West, 1965), and periodic acid-Schiff (Risk and Szczuczko, 1977); excellent for identifying composition and porosity (e.g., Friedman, 1959; Evamy, 1963; Yanguas and Dravis, 1985; Flügel, 2010).

Modified Bushinsky oil technique (Bushinsky, 1947; Bromley, 1981): excellent for enhancing the visibility of internal structures and details of fabric in chalk. Samples are slabbed, polished, and then painted with lightweight oil.

Light Sources

A strong light source is an essential tool for making visual observations in mudstones. The source, intensity, and angle of illumination are all important factors. Useful light sources include the following:

White light: Sunlight is best, but it is not always available. Remember that the intensity of light

Table 1. Suggested List of Equipment and Supplies for Describing Mudstones (after Lazar et al., 2015b).

Backpack

Geologic hammer

Compass

Jacob staff

Magnifiers:

Hand lens: 10×, 20×

Pocket microscope: 25×, 50×

Binocular microscope, reflected light*

Portable lamps (white and UV light)*

Pocket scratcher (sharp steel probe)

Acid bottle (10% HCl)

Water spray bottles*

Grain-size chart

Towel paper, bucket, sponges*

Brushes, chisels

Penknife/multitool

Tape, ruler

Sample bags

Pencils, markers

Eraser*

Field notebooks

Clipboard or clipboard case (letter size)

Digital camera

Laptop

White multiuse ¾" × 1" labels to mark the depth of samples and relevant details to be photographed (include core name and depth)*

Color Post-it flags or notes to mark significant candidate sequence-stratigraphic surfaces (red-sequence boundaries; green-maximum flooding surfaces; blue—transgressive surfaces and parasequence boundaries)*

Whistle, flashlight, small first aid kit

*supplemental gear for core description UV = ultraviolet

changes during the day. We recommend a broadspectrum, artificial light source that provides an even intensity across the examined sample (at least 10–15 cm across). Several models of batteryoperated or AC-power-supplied lights with multiple light-emitting diodes (LEDs) in linear or circular arrays are available.

- Ultraviolet (UV) light: We recommend using several wavelengths of ultraviolet (UV) light to enhance different aspects:
 - ◆ Longer wavelengths (~365 nm/UV-A) enhance detection of hydrocarbons and various types of organic matter (e.g., spores, cutin, resins, waxes, fats, algal cysts, and bacterial degradation products; e.g., Taylor et al., 1998; Lazar, 2007).
 - Shorter wavelengths (~254 nm) enhance visualization of mineral species.
 - ◆ It is helpful to construct an illumination and viewing bench suitable for examining slabbedcore samples (e.g., Lazar, 2007).
 - Eye and skin protection are essential when using UV light sources.

Optical Imaging

Handheld magnifiers are useful for determining the grain-size distribution and details of bedding, body, and trace fossils. These tools include the following:

- **Hand lens:** Typically, the resolution of the naked eye is no more than 100 µm. The most useful magnifications are 10× and 20×; a 10× hand lens can resolve down to 10 µm, and a 20× hand lens can resolve down to 5 µm under optimal conditions. A good magnifier has a moderately large field of view and flat image, with minimal spherical aberration.
- **Pocket microscope:** A 50× pocket microscope, with attached light, is also useful, especially for a quick identification and description of microfossils and small nodules.
- Digital handheld microscope: These units come with a variety of magnifications and focal distances and are able to capture images with scale bars. Some can broadcast images via Wi-Fi to tablet computers and smartphones, which are very useful when working with large groups.

Digital Imaging

Digital images are useful for both revealing subtle aspects of the rocks that might not be readily apparent to the naked eye and documenting the observations made. These images can be acquired in several ways:

Camera system: A camera system includes the camera body, lens, and lighting. All the three aspects need to be optimized for capturing useful images of mudstones:

- **Camera body:** Of especial importance are the sensor type and configuration, the megapixel range, and the construction (rugged, durable, dust and weather resistant, compact but easy to use).
- Lens: A macro lens is very useful for closeup photography. It is specifically designed to provide maximum resolution and sharpness with minimum distortion at its closest focus. Macro lenses come in various fixed focal lengths (typically 50–200 mm). Shorter focal lengths have a slightly greater depth of field but closer working distances that can make lighting a challenge. Longer focal lengths allow more flexibility in lighting but with lesser depth of field. We recommend a lens at the long end of standard macro lenses (between 100 and 105 mm) as it provides a comfortable working distance (9–12 in. [20–30 cm]) with an acceptable depth of field. Several compact digital cameras have quite good macro capability (part of this capability is inherent in the small lens and sensor size), but it is essential to test the actual model on typical subjects to determine which one is most appropriate for your intended uses. Lens filters might be necessary to allow artificial colors or change in light intensity.
- **Lighting:** Achieving even illumination of the subject at close working distances is a challenge in photographing mudstones. Sufficient intensity is also necessary to allow short exposure times at small f-stops—this leads to maximal sharpness and depth of field. For shorter focal length lenses, we recommend using a ring flash or ring light that mounts directly around the lens and provides illumination that can be varied in intensity and ratio. A normal flash unit that connects to the top of the camera can be used, but the camera must be at least a meter or more away from the subject—this necessitates a longer focal length lens to fill the frame with a small subject.
- **Accessories:** Depending on the type of sample and the amount of photography to be done, a copy stand or a tripod setup, large memory capability for photo storage, and wireless remote triggers are helpful.
- Digital scanner: A flatbed scanner is a fast and easy-to-use imaging tool for an object that has a flat surface and is relatively small (e.g., a piece of slabbed core or a thin section).
 - Flatbed scanner: We recommend a flatbed scanner with a range of optical resolution

- capabilities (300-6400 dpi), high maximum density (4.0 D_{max}), 48-bit color and 16-bit grayscale output, dust/scratch removal, and sharpness capabilities.
- Fluid-immersion scanning can be helpful for certain types of mudstones:
 - As with optical microscopy, fluid immersion increases the resolution of the lens, and that enables the lens to achieve higher magnifications. (The numerical aperture of the lens is proportional to the refractive index of the fluid.)
 - For fluid-immersion scanning, the slabbed sample is immersed using a glass-bottomed tray specially designed for the scanner used, with fluid between the glass and the slabbed face. The benefits of this method include reducing losses through reflection and scattering (making the image appear brighter and with greater color saturation) and increasing transmission integrity (providing better definition of shadowed areas and greater dynamic range).
 - The usual considerations and precautions must be taken with selecting and using a fluid that does not adversely affect the rocks being scanned.

Image Processing

Image manipulation of hand specimens, cores, and thin sections using Adobe PhotoshopTM or similar software can reveal the presence of subtle sedimentary features and enable estimation of the degree of compaction of the studied mudstone (e.g., Lobza and Schieber, 1999; Schieber, 2003). For example, image enhancement with image processing software (following manipulation of autolevels, brightness, contrast, color balance, and saturation) can reveal subtle discontinuities and disruptions of laminae and subtle burrows in mudstones with very little variation in color, texture, and composition (e.g., Schieber, 2003). These software applications can also be used to produce an image that approximates the rock's precompaction state. The steps we recommend for achieving this "virtual decompaction" are as follows:

- Select perpendicular cuts of horizontal burrows.
- Resize image by holding the horizontal dimension constant and changing the vertical dimension stepwise until the selected burrows show circular cross sections (Lobza and Schieber, 1999).

Microscopy

By the help of Microscopes, there is nothing so small as to escape our inquiry; hence there is a new visible World discovered to the understanding.

—Robert Hooke, 1665

Examination of rock samples and properly prepared thin sections is an essential early step in characterizing mudstones, specifically providing information about grain size and composition, as well as insights into the depositional and diagenetic processes. Because mudstones are generally rather soft, preparation of good thin sections can pose a challenge. Various methods of thin-section preparation, using special epoxy resins, multiple epoxy impregnation, diamond laps instead of loose grit, double polish, and so on, have been developed (e.g., Catt and Robinson, 1961; Altemüller, 1974; Murphy, 1986; Lindholm, 1987; Miller, 1988). Critical to remember is that mudstones are highly variable in composition and cementation history, and thus, the preparation methodology has to be fine-tuned to the specific features of a given rock. A set of tips for thin-section preparation are given later in this section.

For a good number of years, only a relatively small number of individuals made a sustained effort to communicate the benefits of thin-section petrography in mudstones (e.g., O'Brien, 1981; Zimmerle, 1982, 1994; Schieber, 1989, 1998a, b, 1999, 2002, 2009, 2011; O'Brien and Slatt, 1990; Macquaker and Gawthorpe, 1993; Macquaker and Taylor, 1996; Macquaker et al., 1998). The recent boom in hydrocarbon production from these rocks, however, has brought the general attention to the complexity of mudstones and the significance of observations made under the microscope. As a consequence, studying thin sections of mudstones has become significant as shown by a growing number of related recent publications (e.g., Milliken et al., 2007; Virtasalo et al., 2010; Leonowicz, 2011; Trabucho-Alexandre et al., 2012; Milliken and Day-Stirrat, 2013; Könitzer et al., 2014; Plint, 2014). Examination of thin sections under the microscope is, in our opinion, a crucial source of information on sedimentary features, mineralogy, diagenesis, and microfossil content, and should be performed early on in any investigation of a mudstone succession.

Thin Sections—Evaluations of primary rock attributes described directly in outcrops and cores can be enhanced by integration with observations made in thin sections. Examination of polished thin sections can be performed under a petrographic microscope

and, if available, a scanning electron microscope (SEM). Examination of thin sections is particularly useful for identifying the presence of composite grains (e.g., floccules, faecal pellets, and intraclasts) and recognizing the effects of cementation, dissolution, and bioturbation on grain size (e.g., Lazar et al., 2015a, b). Examination of large, polished thin sections allows for imaging and investigating of multiple genetic beds (e.g., Lazar et al., 2015a, b). Additionally, examination of polished thin sections is particularly useful in distinguishing the composition of individual grains and cement, particularly using an SEM equipped with combined energy-dispersive, cathodoluminescence, and backscattered electron detectors, as discussed in the following sections (e.g., Macquaker and Gawthorpe, 1993; Milliken, 1994, 2013; Macquaker et al., 1998; Schieber, 1999, 2011; Schieber et al., 2000; Schieber and Baird, 2001; Schieber and Riciputi, 2004; Milliken et al., 2012b; Milliken and Day-Stirrat, 2013).

We recommend this workflow for the process from mudstone sampling to preparation and examination of thin sections:

- Obtain representative samples of all facies identified in the core or outcrop (see "Sample Selection—General Considerations" section in this chapter and Lazar et al., 2015b for guidelines for mudstone sampling). Wrap outcrop samples in aluminum foil and tape to prevent rock breakage and excessive development of mold during sample transportation.
- We recommend that thin sections be prepared early in the investigation. If there are no sample-size limitations, it is particularly valuable to make large, polished thin sections (76 mm × 48 mm [~3 in. × 2 in.]).
- We recommend that, after a quick examination under the microscope, and depending on rock texture and composition, "ultra-thin" sections (<20 μm) be prepared to maximize the textural information that can be obtained from the examined mudstones (e.g., Bowles, 1968; Macquaker and Gawthorpe, 1993; Schieber and Zimmerle, 1998; Lazar et al., 2015b). Many constituent particles in mudstones tend to be smaller than the 30 um thickness of a standard thin section, and thus, larger grains (coarse silt and larger) are more conspicuous under the microscope. In contrast, particles that measure in the 5–30 µm size range become "invisible" within the thin "slab" of material that has been sectioned, especially when encased in a matrix that does not provide much contrast. To counteract this problem, one can

resort to the manufacture of "ultra-thin" sections or simply take advantage of the fact that many thin sections develop "wedged" margins while thinned to the final thickness. Simply examining the wedged part allows one to view particles that are much smaller than the thickness of the standard thin section. Double-polished thin sections can also be helpful because grain relations are more readily observed when surface roughness on both sides of the "slab" has been minimized. (Grain size is a fundamental attribute of sedimentary rocks. See, for example, McCave and Syvitski [1991] for a review of principles and other methods of particle-size analysis. Interpretation of results obtained by such methods should be made with caution, bearing in mind that mudstone samples may be disaggregated to particle sizes that are actually not present in the rock—and reveals more about the disaggregation process rather than the original grain-size distribution.)

- Polished thin sections have multiple benefits: They do not only improve the visibility of features for transmitted light microscopy but also enable reflected light microscopy, fluorescence microscopy, cathodoluminescence (CL) microscopy, scanning electron microscopy, and electron microprobe analysis.
- Preparing polished thin sections of mudstones is labor intensive because constituent minerals have a wide range of hardness (clay minerals are soft, and cementation may be absent or variably developed), and it is difficult to inject stabilizing epoxy into the tiny pores of these rocks. Samples can also be damaged during the preparation of thin sections because mudstones commonly contain relatively high proportions of hydratable clay minerals. To minimize sample damage, argillaceous mudstones should be exposed to as little water as possible during the preparation process (Lazar et al., 2015b). Thin-section petrography of soils deals with similar aspects as thin-section petrography of mudstones, and thus, the handbook for soil thin-section description by Bullock et al. (1985) is a handy and compact supplement to other texts on thin-section preparation and description.
- Scan the thin sections on a flatbed, high-resolution scanner to produce high-quality, low-magnification images that enable a detailed study of the physical, biological, and chemical sedimentary features
- Capture the relevant features with a digital camera attached to a petrographic microscope.
- Use image processing software (e.g., Adobe PhotoshopTM) to bring out seemingly obscure features

(image enhancements: levels, brightness, contrast, color balance, sharpening, etc.).

We recommend documenting thin-section observations of texture, bedding, and composition using a fit-for-purpose template (see Lazar et al., 2015b for template examples). Thin-section templates should enable one to record observations made on key attributes of the mudstones and should be adapted to the objectives of and the time available for a specific project.

Optical Microscope—Optical microscopy can, for example, inform about the not-always-so-obvious origin of quartz grains (e.g., Schieber, 1996; Milliken et al., 2012a; Milliken, 2013), the formation history of small spots of cherty-looking material (e.g., Milliken et al., 2007), depositional parameters (e.g., Macquaker and Taylor, 1996; Schieber, 1999; Wilson and Schieber, 2014; Lazar et al., 2015a, b), and sequence-stratigraphic packaging and parasequence stacking patterns (e.g., Schieber and Lazar, 2004; Lazar, 2007; Lazar et al., 2015a, b). For example, Schieber (1996) was able to show that diagenetic infilling of algal cysts produced sand-size diagenetic quartz grains that are easily mistaken for detrital grains, which can lead to erroneous interpretations of mudstone-associated sandstone beds. This avenue of research was extended further when in distal mudstones, large proportions of siltsize (and presumably detrital) quartz grains were linked to early diagenetic processes as well (Schieber et al., 2000). Chertlike grains may initially suggest a diagenetic origin, however, these grains can also be detrital with some help from grain-concentrating benthic agglutinated foraminifera (e.g., Milliken et al., 2007; Schieber, 2009).

When examined closely, many mudstone successions also show a wide variety of primary sedimentary structures and bioturbation features at thin-section scale and can provide excellent clues to sedimentary conditions, such as the presence of bottom currents (e.g., Schieber, 1999; Schieber et al., 2007), event deposition (e.g., Schieber, 1989, 1999; Loucks and Ruppel, 2007; Macquaker et al., 2010), and microbial mats (e.g., Schieber, 1989, 1999), as well as to substrate consistency (e.g., Wetzel and Uchman, 1998; Lobza and Schieber, 1999) and more subtle forms of animal-sediment interaction (e.g., Schieber, 2003; Pemberton et al., 2008). Lazar et al. (2022b, Chapter 4 this Memoir), as well as all of the case study chapters (Bohacs and Ferrin, 2022; Bohacs and Grabowski, 2022; Bohacs and Guthrie, 2022; Bohacs et al., 2022d, e; Campo et al., 2022; Lazar and Schieber, 2022; Potma et al., 2022; Chapters 8–15 this Memoir) include many representative examples.

Another microscopic approach to mudstone petrography is the examination of silt- and sand-size constituents in grain mounts after separation from the mudstone matrix. This furnishes additional information on the provenance of the fine-grained sediment and the diagenetic alteration of grains. Separation of this grain population by means of heavy liquids or magnetic separators or both allows a further split into light and heavy minerals, and facilitates the study of rare constituents that are not manifest in conventional thin sections. Auxiliary methods for analysis are an SEM with an attached energy-dispersive x-ray analysis system (EDS) or an electron microprobe (discussed in "Elemental Mapping" paragraphs in the "Scanning Electron Microscope" section).

Thin-section examination under either the optical microscope or the SEM has also been used to detect grain-size trends in stratigraphic successions (e.g., Macquaker and Taylor, 1996), but some caution is advisable in that regard. Grains smaller than section thickness tend to be less readily observed, and grains larger than section thickness will not only be more conspicuous, but their observed average size will be smaller than the actual size because few will be sectioned through the widest part of the grain. Thus, trends from sand to silt can probably be determined with reasonable assurance, whereas trends within, say, the silt range are difficult to reproduce consistently. One could, of course, also try to measure grain-size trends via settling-tube analysis from the disaggregated samples, but there the problem is that it is nearly impossible to disaggregate a lithified mudstone into its component grains. As such, results of this kind of analysis need to be evaluated critically, unless the studied rock is indeed very poorly consolidated.

Scanning Electron Microscope—The scanning electron microscope (SEM) has long been a powerful tool in its various applications to sedimentary rocks (e.g., O'Brien, 1981; Pye and Krinsley, 1984; Trewin, 1988) and has proven indispensable for the investigation of mineral relations and textures in fine-grained rocks (e.g., Camp et al., 2013). Examination under an electron microscope provides useful insights into grain composition and origin as well as depositional processes and diagenetic history (e.g., O'Brien and Slatt, 1990; Schieber, 1996; Schieber et al., 2000; Milliken and Day-Stirrat, 2013; Schieber, 2013). Composition-content SEM studies have revealed decameter-scale stratigraphic patterns that assisted sequence-stratigraphic interpretations (e.g., Macquaker and Taylor, 1996; Williams et al., 2001).

Fundamentally, *SEM imaging* relies on secondary electron emission that is sensitive to the surface relief of the sample as well as to compositional differences

when operated in the backscattered electron mode. Several chapters in AAPG Memoir 102 (Camp et al., 2013) introduce up-to-date operation principles and sample preparation methods (Erdman and Drenzek, 2013; Huang et al., 2013).

Although SEM magnifications can range from single digits to more than 100,000×, it is the 1000×–10,000× range that is commonly used for textural studies, whereas the 20,000×–120,000× range is commonly used for studies of porosity and detailed mineral interrelations. Because of its great depth of focus (compared with petrographic microscopes), the SEM allows detailed imaging of rough and broken surfaces (e.g., drill cuttings), but its power as a petrographic tool really comes to the forefront when flat surfaces are examined.

SEM analyses require high-quality polished surfaces for reliable results. Although such flat surfaces have traditionally been prepared by mechanical grinding with successively finer polishing media (polished thin sections), the associated mechanical damage manifests itself via surface "smear" and obscured details of mineral relations and pore spaces. The more recent application of argon ion milling, a sample preparation technique developed by material scientists (Bollinger and Fink, 1980), overcame this limitation and enabled imaging of pore spaces and mineral interrelations at the magnifications needed for mudstone studies (Jiang et al., 1990; Hover et al., 1996; Rask et al., 1997; Schieber, 1998c, 2002, 2004). Pores are readily observed in transmission electron microscopy (TEM) images of argon-ion-milled samples (e.g., Hover et al., 1996), but today the preferred technique for imaging pores in mudstones is the SEM examination of argon-ion-milled surfaces (e.g., Tomutsa et al., 2007; Loucks et al., 2009; Schieber, 2010, 2011, 2013; Driskill et al., 2013; Jennings and Antia, 2013; Schieber et al., 2013). The potential for generating artificial pores in organic-material-rich areas resulting from ion beam heating has been a serious concern. See Schieber et al. (2013) for a discussion about a procedure to eliminate ion-milling artifacts.

SEM analyses can be used for *elemental mapping*—One of the very useful by-products of electron beam imaging is the emission of x-rays from the surface of a sample. Because each chemical element has a unique atomic structure, it produces a unique set of characteristic x-rays that can be differentiated with an x-ray spectrometer. In SEM applications, energy-dispersive x-ray spectroscopy (commonly abbreviated as EDS or EDX) is the most widely employed method, although wavelength-dispersive spectroscopy (WDS; used in electron microprobes) is superior for quantitative analysis. Within the resolution limits of modern EDS detectors (about 0.2–1 μ), each image spot that is scanned with an SEM can also be associated with an EDS spectrum that carries information about the

chemical composition of that spot (Goldstein, 2003). Software manipulation of spectral datasets can be used to extract maps for chemical elements of interest and also to map the object classes (e.g., minerals) within the field of view. These element maps are very useful for the identification of minerals and for estimates of rock composition.

Another useful by-product of electron beam imaging is *cathodoluminescence*. Cathodoluminescence (CL) is the photoelectric effect in reverse. Electrons interact with a luminescent material (usually nonconductive or semiconductive) and cause emission of photons in the wavelengths of the visible light spectrum (as well as beyond: infrared and ultraviolet). CL is observable by petrographic microscope (floodgun) or by SEM where other detectors can support mineral identification (e.g., Long and Agrell, 1965; Boggs and Krinsley, 2006). Rock-forming minerals (e.g., quartz, feldspar, carbonate, and apatite) can be identified on the basis of their CL colors (e.g., Long and Agrell, 1965; Smith and Stenstrom, 1965; Zinkernagel, 1978). A drawback of the "floodgun" approach is that high densities of energetic electrons can induce undesirable high specimen temperatures, cause reduced CL emissions, and even cause permanent damage to the CL emission of the specimen. In contrast, scanning for CL in the electron microscope (SEM-CL) gives much higher spatial resolution, and because the beam "rests" on a given spot for only a few microseconds, heating effects are greatly reduced. State-of-the-art detector designs, such as the Gatan ChromaCL, enable live digital color imaging at a range of pixel resolutions. All CL information is acquired in a single imaging pass, and potential beam damage to CL-related features is greatly reduced.

Cathodoluminescence emissions are dependent on several factors, including but not limited to crystal structure, lattice defects, and distribution of trace or minor elements. As a tool for detection of chemical differences, CL is, in most cases, much more sensitive than EDS and can allow visualization of differences in the ppm concentration range. By showing the distribution of trace and minor elements, as well as crystal growth patterns, SEM-CL can be a very valuable petrographic technique that is able to show features that are undetectable with other techniques.

Another application where CL imaging has the potential to add to the understanding of sedimentary successions is in the determination of sediment provenance (e.g., Seyedolali et al., 1997). The CL color (or spectrum) of detrital quartz grains reflects the rocks from which the quartz originated. This was first comprehensively documented in an early study by Zinkernagel (1978) and has been confirmed by subsequent studies. Traditionally, provenance assessments of mudstones used bulk rock geochemical

parameters, which are prone to bias by weathering and diagenesis. Thus, for a long time, mudstones have been underused for provenance studies, although they constitute the bulk of most sedimentary basin fills. With time, increasingly more studies of CL textures for quartz sourced from a variety of igneous, metamorphic, and sedimentary rocks have been conducted (e.g., Milliken, 1994). The new generation of color CL detectors (like ChromaCL2TM) makes it possible to obtain good-quality provenance information from large quartz grains down to those in the silt-size range by identifying populations of quartz silt grains of similar provenance and diagenetic history through the combined use of the CL color and grain internal textures (e.g., Schieber and Wintsch, 2005; Krinsley et al., 2009; Thyberg et al., 2010). Applying SEM-CL to quartz grains, the chemically and mechanically most robust major component of these rocks, will in the future enable us to greatly improve our understanding of provenance and uplift history of the source area of sedimentary basin fills.

X-Ray Computed Tomography Scan

X-ray CT (computed tomography) examination is most useful for 3-D visualization of bedding and (subtly expressed) bioturbation; it can also provide useful insights into grain-size distribution and composition (organic-rich or cemented strata). Information on depositional conditions comes from interpreting the lamina geometries and bedding styles, which can be challenging to resolve with normal optical means. For more information on this method, see Ketchum and Carlson (2001), Spaw et al. (2011), Ketchum and Hildebrandt (2014), and Spaw (2015).

Sample Preparation and Analytical Considerations

- X-ray CT scans are rapid and nondestructive; they reveal the 3-D internal bedding and properties of mudstones through directing x-rays at a core from multiple angles and measuring the energy transmitted through the core. The images obtained are compiled into video formats and record variations in x-ray attenuation by the core material, which is a function of the density and composition (atomic number) of the rocks (as well as the incident x-ray energy and path length of the x-ray beam).
- This analysis is best suited for cylindrical samples. The only length limitation is the capacity of the machine.
- It can be done on slabbed or whole core (in or out of the liner/sleeve).

CT Data Interpretation

- CT data are captured in a succession of image files suitable for visualization and analysis using various 2-D and 3-D image processing tools.
- Standard modes of 3-D visualization include volume rendering and isosurfacing. Volume rendering is commonly the most useful for sequence-stratigraphic analysis.

Tips and Traps

- CT scans are best acquired when the core is intact because artifacts from broken core pieces and imaging procedures can complicate interpretations.
- CT scans work best on mudstones with relatively high contrast among their components (e.g., clay minerals, carbonate, or pyrite). It would, however, be difficult to differentiate carbonate shells in a fine-grained carbonate matrix.
- Resolution is typically limited to 1000–2000 times the object's cross-sectional diameter (~50–100 μ for a typical 4 in. core).
- Each CT scan produces multiple gigabytes of data and requires significant computing resources for manipulation, visualization, and analysis.

Procedures to Capture Mudstone Description in Cores and Outcrops

Millimeter-to-centimeter-scale observations of mudstone strata and their stacking patterns are recorded keeping track of the abundance of each physical, biological, and chemical attribute. One can then recognize laminae, laminasets, beds, and bedsets, and aggregate these observations into facies, facies associations, and facies-association successions (e.g., Bohacs et al., 2014; Lazar et al., 2015a, b). An example of a description form and symbols we use to capture lamina-to-bedset-scale observations in mudstone successions is given in Figure 2. Such a form clearly separates observations from interpretations, along with differentiating the various levels of interpretations; it captures observations on the left side; first to the right are lower level interpretations (e.g., sedimentary structures) and then higher level interpretations (e.g., benthic-energy and oxygen levels, depositional environments; Figure 2).

Observations—Key attributes to capture include the following (see also Table 2 and Lazar et al., 2022a, b, Chapters 2 and 4 this Memoir):

- Depth or elevation
- Sample or photograph unique identifier

- Grain size: minimum, mode, maximum
- Lamina geometry: continuity, shape, geometry (figure 1B of Lazar et al., 2022a, Chapter 2 this Memoir)
- Bedding surfaces: location, shape, spacing
- Composition: argillaceous, siliceous, and calcareous (figure 1C of Lazar et al., 2022a, Chapter 2 this Memoir)
- Physical sedimentary structures: primary, secondary
- Biogenic material: type, abundance, diversity, and taphonomy (figure 1F, G of Lazar et al., 2022a, Chapter 2 this Memoir)
- Ichnofauna: type, diversity, and abundance
- Diagenetic products: nodules, cement
- Bioturbation index (figure 1E of Lazar et al., 2022a, Chapter 2 this Memoir)
- Fractures: type, abundance, orientation, mineralization

Interpretations—It is important to comment on the following (see also Table 2; and Lazar et al., 2022b; Bohacs et al., 2022a; Chapters 4 and 5 this Memoir):

- Origin of grains
- Transportation, depositional, and reworking processes
- Diagenetic processes
- Environment of deposition
- Sequence stratigraphy: surfaces, stratal units

Comparison Charts

Consistent capture of hand specimen and core observations is facilitated by using various charts for estimating grain size, percentage of a particular grain type, degree of bioturbation, or color. A brief introduction to these charts follows:

Grain-Size Charts—These charts classify sediments and sedimentary rocks based on the range of the diameter of individual grains (e.g., Udden, 1914; Wentworth, 1922; Krumbein and Sloss, 1963). See Lazar et al. (2022a, Chapter 2 this Memoir) for a discussion and definitions of mud grain-size boundaries. As a reminder, before examination of mudstone strata, it is important to obtain and examine fresh surfaces of rocks from each potential stratigraphic package. Then, estimate the percentage of sand-size grains with visual comparison to percentage charts. If less than 50 % are sand-sized grains, the rock is a mudstone and next, a scratch test can be performed to determine the dominant grain size (Lazar et al., 2022a, Chapter 2 this Memoir).

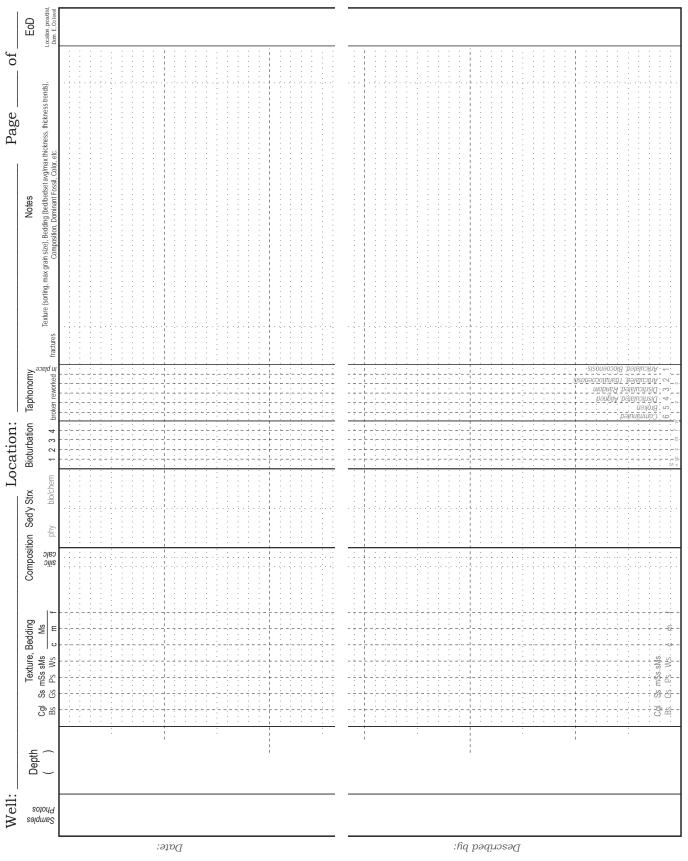


Figure 2. (A) Example of a form that we recommend using to capture the key attributes of mudstones observed in cores.

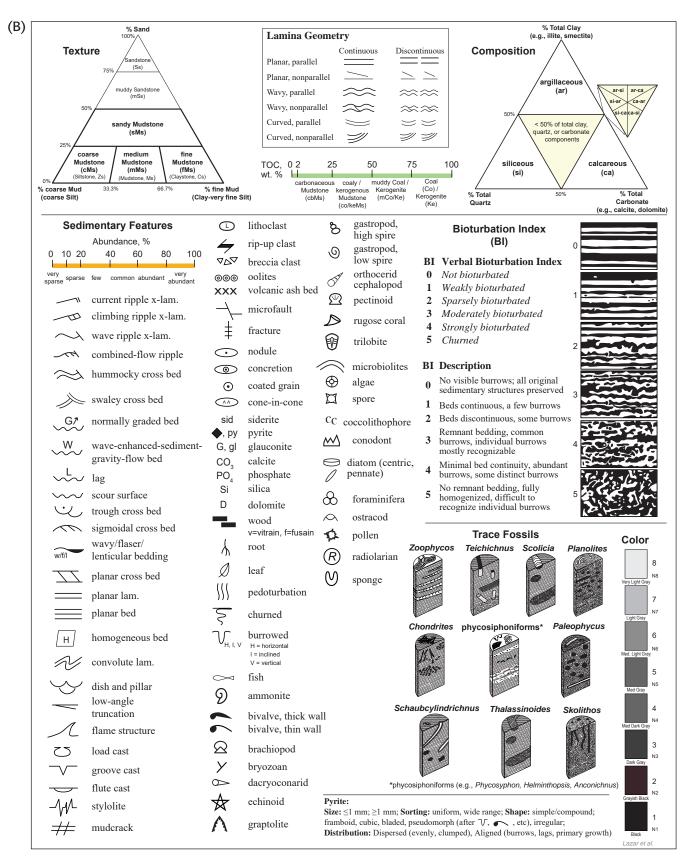


Figure 2. (*Continued*) (B) Example of symbols useful for capturing mudstone observations in cores and hand specimens (after Lazar et al., 2015b, used with permission).

Table 2. Mudstone Attributes Most Useful for Sequence-stratigraphic Analyses.

Mudstone character	•	
	Texture	Grain size, sorting
	Bedding	Geometry, shape, continuity
	Composition	Detrital, biogenic, diagenetic
Aspects of deposition	onal environment	
	Origin of grains	Physical, biogenic, chemical (see figures 1C and 2 of Chapter 2 for common origins of mudstone grains)
	Transport and	Benthic energy levels
	deposition	Mechanisms: current, wave, gravity, aeolian
		Instantaneous sedimentation rate
		O ₂ /Eh levels: persistence/history and recurrence
		pH levels: persistence/history and recurrence
		Salinity levels: persistence/history and recurrence
	Reworking and	Completeness of record (hiatuses)
	diagenesis	Types and successions of diagenetic changes: early (microbial), late (thermal)
		Early: depositional environment and rock character (reductants/oxidants)
		Late: decipher original character

Percentage-Estimation Charts—These charts can be used for visual comparison of percentage estimation of angular grains (Terry and Chilingar, 1955), small and large angular grains (Baccelle and Bosellini, 1965), spherical grains (Schafer, 1969), irregularly shaped grains of various shapes (Flügel, 1982), or nearly spherical (crinoidal) or blady (pelecypod) skeletal grains (Schafer, 1969).

Bioturbation-Index Estimator—The degree of bioturbation can be assessed using a 0-5 scale (figure 1E of Lazar et al., 2022a, Chapter 2 this Memoir; Lazar et al., 2015a, b). Bioturbation tends to be subtle in mudstone strata; do not expect too many textbook-style ichnofossils because burrows tend to be obscure where matrix and fill do not differ in composition and rheology (Lazar et al., 2015b). Prior to mud compaction, benthos do not burrow but swim through muddy sediment with high water content (70%–90%) and disturb the sediment fabric, producing deformational structures (e.g., Lobza and Schieber, 1999; Schieber, 2003). For in-depth discussions and reviews of trace fossils and trends in depth, and extent of bioturbation in fine-grained rocks, see, for example, Bromley and Ekdale (1984), Bromley (1990),

Pemberton et al. (2001, 2008), Droser and Bottjer (1986, 1988, 1989, 1991), Bottjer et al. (1988), Bottjer and Droser (1992), Schieber (2003), Hasiotis (2004), and Gerard and Bromley (2008).

Consistency of Marine Substrate—Organisms bioturbating fine-grained sediments may produce different burrows depending on substrate consistency (e.g., Wetzel, 1991; Bromley, 1996; Brett and Allison, 1998; Wetzel and Uchmann, 1998; Lobza and Schieber, 1999; Schieber, 2003). Five levels of substrate consistency can be inferred from ichnofossil analysis—from "soupground" to "hardground" (Table 3).

Rock-Color Charts—A rock-color chart is useful in describing the colors of mudstones. The colors of mudstones can be compared to the Geological Society of America or Munsell color charts. The color of mudstones varies significantly and may be in various shades of black, gray, green, or brown.

It is important to record the time of day that a mudstone description was made, as well as the light type and intensity, and the wetness state (dry or wet).

Inferred Substrate Consistency	Inferred Water	Burrow Architecture								
	Content	Tunels Vertical Horizontal	Galleries Vertical Horizontal	Branching	Pelleted Walls	Sculpted Lined Unlined	Ornated Lined Unlined	Mantle and Swirl	Spreite	Meniscate
1. Soupy	>75%									
2. Soft	50%–75%	^ ^	^	1	1				1	
3. Stiff	25%–50%								\downarrow	1
4. Firm	10%–25%			.		1	1		·	\downarrow
5. Hard	<10%									
6. Wood										

Table 3. Inferred Substrate Consistency Based on Ichnofossils Analysis.

After Wetzel (1991), Bromley (1996), Brett and Allison (1998), Wetzel and Uchmann (1998), Lobza and Schieber (1999), and Schieber (2003).

- Composition, especially the amount of iron (and its oxidation state) and organic-carbon content, is the main factor that determines mudstone color (e.g., Potter et al., 1980, 2005; Schieber and Zimmerle, 1998; Lazar et al., 2015b). Organic-carbon-rich rocks are typically dark chocolate brown (5YR3/4–5YR2/1). Gray or green mudstones are typically organiccarbon lean. Mudstones with a high content of disseminated pyrite are typically grayish black to black (N2–N1; Lazar et al., 2015b).
- Weathering and diagenetic processes can significantly alter mudstone color.

Geochemical Tools

Chemistry: that most excellent child of intellect and art.

> —Sir Cyril Norman Hinshelwood (quoted in Nye, M. J., 1993)

Numerous molecular, elemental, and isotopic tools that are useful for characterizing mudstones are available. These geochemical tools range in use from a quick screening (e.g., reaction to hydrochloric acid) to extremely refined estimates of microbial respiratory pathways (e.g., compound-specific isotopic analysis). Listed in this section are tools that are either commonly used by a geologist during initial mudstone investigations or available later to augment and expand the understanding of mudstone character and deciphering depositional conditions under which these rocks formed. Many other analytical tools are used for characterizing mudstones and inferring depositional conditions. For example, such analytical tools can be used to determine total uranium concentrations (delayed neutron counting) or base metal content, and trace and rare earth elements (inductively coupled plasma mass spectrometer or inductively coupled plasma atomic emission spectroscopy).

Sample Selection—General Considerations

As with all methods that analyze a very small volume of rock, it is essential to select representative samples within a geologic-stratigraphic context that enables interpretation and application (upscaling) of analytical results appropriately. Spot-sample analyses provide insights into mudstone character and depositional conditions. Profiles of analytical results for successions of samples are required for sequence-stratigraphic analyses. In either case, a sampling strategy that considers geological context and project goals is essential. Special care must be exercised in selecting samples for vertical profiles to allow robust conclusions to be drawn—samples need to be taken from equivalent facies or locations with respect to significant stratigraphic surfaces (e.g., parasequence boundaries) to discern secular trends accurately. Some of these concerns are mitigated with analytical tools that provide a near-continuous record (mm-to-cm scale), such as an x-ray fluorescence (XRF) scanner. See Lazar et al. (2015b) for more detail on sampling strategies.

Reaction to Hydrochloric Acid

Carbonate minerals effervesce to various degrees when exposed to dilute hydrochloric (HCl) acid. The onset, extent, and rate of effervescence are influenced by mineralogy, amount, and particle size, along with the temperature of the sample and the temperature and concentration of HCl (Soil Survey Division Staff, 2014). Carbonate minerals are an important component of mudstones that attest to rock character and depositional conditions. Carbonate minerals can be of biogenic, detrital, or diagenetic origin. The acid test is useful for making a first approximation of carbonate content and mineralogy, but further work is required to reveal its origin. The composition of early-formed cements and nodules in mudstones is a function of depositional conditions, and certain compositions are associated with specific types of sequence-stratigraphic surfaces. For example, calcite or calcite and pyrite are common below parasequence boundaries, dolomite with maximum flooding surfaces, and siderite with sequence boundaries or other bypass surfaces (see discussions in Bohacs et al., 2022a, c, Chapters 5 and 6 this Memoir). For safety notes and more information on the acid test, see Soil Survey Division Staff (2014).

Standard Procedure

- Obtain a fresh surface of a mudstone sample.
- Scratch the fresh surface of the mudstone sample with a sharp steel probe.
- Apply a drop of dilute HCl onto the fresh surface and powder.
- Observe and rate reaction:
 - Instant, vigorous effervescence of fresh surface indicates calcite.
 - Rapid, strong effervescence of powder (and slow, weak effervescence of fresh surface) indicates dolomite or siderite.
 - Siderite can sometimes be differentiated from dolomite as it tends to weather bright orange to red-brown.

Tips and Traps

- Standard concentration of HCl for this method
- The degree of effervescence can be classified using a table such as the one found in Soil Survey Division Staff (2014).
- The degree of effervescence is influenced by the temperature of the sample and the temperature and concentration of HCl.
- Hydrochloric acid is corrosive. Always protect the eyes and skin when working with concentrated acids.

Staining for Mineral Identification

Rapid identification of some carbonate minerals can be done using stains (e.g., Friedman, 1959; Reid, 1969; Friedman et al., 2007). Stains are inexpensive, easy to use, rapid to apply, and quite versatile as they can be used in the field, on rock slabs or thin sections. Stains

work by reactions whose occurrence and extent are influenced by mineral composition and crystallography. Of especial interest for sequence-stratigraphic analyses and insights into depositional conditions is the potential to differentiate aragonite, calcite, ferroan dolomite, and siderite. For more information, see Friedman (1959), Evamy (1963, 1969), Dickson (1965, 1966), and Friedman et al. (2007).

Common Procedures

- The most widely used stain is a mixture of Alizarin Red S (ARS) and potassium ferrocyanide (PF) in dilute HCl for 1-2 min. It differentiates calcite and aragonite from dolomite and siderite.
- For samples with low iron concentrations, a dual staining procedure can be used: PF + HCl, followed by ARS + PF stains as above.
- Aragonite can sometimes be distinguished from calcite using Feigl's stain. See Friedman (1959) and Schneidermann and Sandberg (1971) for more information.

Tips and Traps

- The occurrence and extent of staining is a function of the reactivity of carbonate mineral species with the acid, so the crystal size and temperature of the sample, and the temperature and concentration of the acid influence the outcome.
- Hydrochloric acid is corrosive. Always protect the eyes and skin when working with concentrated acids.

X-Ray Fluorescence

X-ray fluorescence (XRF) is a relatively nondestructive and rapid method for detecting the occurrence and concentration of major and trace elements (Na through U) in bulk samples. Element concentration can be used to infer such depositional conditions as paleo-redox levels, terrigenous influence, and detrital or biogenic content through geochemical proxies (e.g., Pratt, 1984; Jones and Manning, 1994; Bertrand et al., 1996; Dean and Arthur, 1998; Murphy et al., 2000a, b; Werne et al., 2002; Lazar and Schieber, 2003; Lyons et al., 2003; Sageman et al., 2003; Algeo, 2004; Rimmer, 2004; Rimmer et al., 2004; Bohacs et al., 2005; Algeo and Lyons, 2006; Tribovillard et al., 2006; Lazar, 2007; Algeo and Maynard, 2008; Rowe et al., 2012). Interpretation of depositional conditions of fine-grained sediments is always enhanced when geochemical proxies are integrated with physical, biological, and chemical observations from outcrops, cores, and thin sections.

Vertical profiles taken through appropriately spaced samples or with an XRF scanner are helpful supplementary information for sequence-stratigraphic analysis, especially using proxies that estimate such key factors as detrital or biogenic content, redox levels, and terrigenous influence. For more information on this technique, see, for example, Norrish and Chappell (1977), Tertian and Claisse (1982), Fairchild et al. (1988), Rollinson (1993), and Fitton (1997).

XRF is well suited for bulk chemical analysis of major and minor elements (i.e. Al, Ca, Fe, K, Mg, Mn, Mo, Na, P, S, Si, Ti) and trace elements (>1 ppm; i.e. Ba, Ce, Cd, Co, Cr, Cu, Ga, La, Mo, Nb, Ni, Rb, Rh, Sc, Sr, U, V, Zn, Zr).

Sample Preparation for Bulk Analysis

- XRF is a bulk analysis, so sample size must be several times larger than the largest grain size or particle in the rock. For most mudstones, this means a representative sample of about 8 g (approximately 2 cm³ cube) for quantitative XRF. Some laboratories can use as little as 0.05 g for a "semiquantitative" XRF analysis.
- The sample is crushed to an average grain size of a few millimeters, and then a representative subsample is selected and powdered. A powdered aliquot is typically about 100 ml (~250 mg).
- The powdered sample can then be analyzed directly (appropriate for trace elements) or mixed with a chemical flux, melted to make a homogeneous glass, and analyzed (useful for samples with a wide range of elemental abundances, especially Fe).
- If sample availability is an issue, powdered mudstone samples that have been extracted for molecular geochemical analysis can be used for XRF analyses. Alternatively, if XRF was conducted on samples that were powdered-only, those samples can be used for other analytical techniques.

XRF Scanning Spectrometer

- Samples must have a relatively planar face (e.g., slabbed core). The main limitation is the length of sample that can be scanned at one time.
- Spatial resolution can be selected, typically from 0.1 to 10 mm between locations (width from 2 to 15 mm); an analysis takes from 1.5 to 30 s per location, with approximately 0.5 s between scans (depending on the machine used).
- Output is a profile with typically the same suite of major and trace elements as for single sample outputs (Na through U).
- See Fitton (1997), Jansen et al. (1998), Rothwell et al. (2006), Banerjee (2011), and Liu et al. (2013) for details on applications.

Portable XRF Spectrometer

- The power of this approach is that it allows taking the instrument to the sample instead of the sample to the instrument. It is potentially useful for relatively rapid acquisition of vertical profiles in cores or outcrops.
- Two categories of portable XRF spectrometers, both of which use an x-ray tube source, are used:
 - ◆ The first category consists of handheld, radar-gunlike units that are applied to a sample ("point and shoot") to analyze a spot typically 3–10 mm in diameter.
 - ★ The second category consists of portable, boxlike units with a compartment into which a relatively small sample is placed. This type tends to be higher powered and more flexible in use.
- All portable units represent some trade-off of range and number of elements, flexibility, and accuracy for portability and speed. In general, they can detect more than 25 elements but are more reliable for heavier elements.
- Key factors in choosing an appropriate unit include a range of elements detected, excitation range, detector resolution and total count rate, ability to modify x-ray beam, and software and display options.
- Serious consideration must be given to evaluating whether the time in the field is best spent examining the rocks and collecting representative samples for later analyses instead of analyzing on the spot. Optimum use in the field requires at least one person dedicated to the instrument and a second person locating and preparing sample locations.

Paleo-Redox Proxies

- It is not a simple matter to decipher paleoredox conditions during the formation of organic-carbon-rich fine-grained rocks. Hence, a series of geochemical proxies are used to interpret paleo-redox conditions.
- These proxies are ratios of a redox-sensitive element to an element that does not fluctuate with redox conditions: the degree of pyritization (DOP = Fe_{pyritic}/[Fe_{pyritic} + Fe_{acid soluble}]), V/Cr, V/(V+Ni), Ni/Co, U/Th, and Mo/Al (e.g., Adams and Weaver, 1958; Berner, 1969, 1970, 1980, 1981, 1982; Ernst, 1970; Lewan and Maynard, 1982; Dypvik, 1984; Lewan, 1984; Raiswell and Berner, 1985; Dill, 1986; Calvert, 1987; Raiswell et al., 1988; Canfield, 1989; Dean and Arthur, 1989; Calvert and Karlin, 1991; Calvert and Pedersen, 1992, 1993; Arthur and

Sageman, 1994; Jones and Manning, 1994; Wignall, 1994; Werne et al., 2002; Lyons et al., 2003; Rimmer et al., 2004; Lazar, 2007; Lazar et al., 2015b).

Clastic Dilution Proxies

- Geochemical proxies commonly used for inferring the amount of clastic dilution include the Ti/Al, Si/Al, and K/(Fe + Mg) ratios (e.g., Pratt, 1984; Bertrand et al., 1996; Dean and Arthur, 1998; Murphy et al., 2000a, b; Werne et al., 2002; Sageman et al., 2003; Rimmer et al., 2004; Bohacs et al., 2005; Lazar, 2007; Lazar et al., 2015b).
- Detritus has been also predicted from the concentration of potassium using the following equation (e.g., Isaacs, 1981; Schwalbach and Bohacs, 1992):

Detritus (weight %) = $36.3 \times K$ (%) + 0.9.

Paleo-Productivity Proxies

Examples of geochemical proxies used to infer past rates of primary production include TOC, TOC/Al, and TOC/P_{tot} (e.g., Hay, 1995; Rühlemann et al., 1999; Wortmann et al., 1999; Murphy et al., 2000a, b; Werne et al., 2002; Sageman et al., 2003; Rimmer et al., 2004; Bohacs et al., 2005; Tribovillard et al., 2005; Lazar, 2007; Lazar et al., 2015b).

Tips and Traps

- Do not apply strictly or in isolation from other data (proxies can be misleading if sediment has been reworked, thus altering the original concentrations).
- Examine a variety of proxies; integrate with physical, biological, and chemical observations from outcrops, cores, and thin sections; and then make interpretations based on preponderance of evidence, not on any single proxy.

X-Ray Powder Diffraction

X-ray powder diffraction (XRD) is a relatively rapid and nonconsumptive method for detecting the occurrence and concentration of mineral species and phases in bulk samples. It is a basic tool for determining mudstone composition and especially useful in differentiating clay-mineral types that are difficult to determine optically. Its output should be used to infer depositional conditions only when used in conjunction with optical microscopy, electron microscopy, or both (because XRD by itself cannot differentiate among detrital, biogenic, or diagenetic origin of any mineral species). For more information on XRD, see Hardy

and Tucker (1988), Bish and Post (1989), Mumme et al. (1996), Moore and Reynolds (1997), or other standard references.

Sample Selection and Analytical Considerations

- Typically, 4-8 g of a mudstone sample is ground finely (<10 µm) and homogenized, and a 10–30 mg aliquot selected. The sample is spread on a homogenous holder and air-dried.
- The sample is inserted into an x-ray diffractometer and scanned through a wide range of incidence angles.
- Each analysis takes 20 minutes or less.

Tips and Traps

- Because of the small sample size, it is, as always, essential to understand the geological context of the samples analyzed.
- Detection is limited to minerals constituting as low as 0.5% of the volume of a sample.
- Differentiating specific types and mixtures of clay minerals typically requires more advanced treatment and multiple analyses (e.g., ion saturation, glycolation, heating to various temperatures). The degree and amount of change in crystal structure, especially of swelling clays, provides supplementary information essential to identification.
- XRD analysis is quite straightforward; interpretation of XRD output, however, requires care and experience, especially for complex mixtures of clay minerals.
- Inferring depositional conditions requires optical microscopy or electron microscopy or both, as discussed in the Microscopy section.

Total Organic Carbon Content

Vertical and lateral profiles of total organic carbon (TOC) content have been used for rapid screening of the quality and distribution of mudstones as source, reservoirs, and seals of hydrocarbons (Table 4). The TOC content of a rock is the result of the nonlinear interaction of the rates of organic-matter production, destruction, and dilution (e.g., Tyson, 1995, 2001, 2005; Bohacs et al., 2000, 2005; Sageman et al., 2003; Rimmer et al., 2004; Harris, 2005; Katz, 2005; Lazar, 2007). High TOC contents record the optimized combinations of moderately high production, low destruction, and low dilution rates. Low TOC contents can be the result of low production, high dilution, or high destruction rates, as well as high thermal maturity. As referenced in the section on Well-Log Tools (and discussed in Bohacs et al., 2022b, Chapter 7 this Memoir),

Table 4. Essential Attributes of Mudstones as Hydrocarbon Sources, Reservoirs, and Seals.

Source	Quantity of hydrocarbons	TOC, HI
	Types of hydrocarbons	HI, OMT
	Kinetics of generation	OMT
Reservoir	Storage	Porosity (proportional to TOC, biogenic content, clay-mineral content)
	Transmissivity	Permeability, Young's modulus, Poisson's ratio
Seal	Transmissivity	Clay-mineral content, early cements
	Ductility/ strength	Clay-mineral content, early cements, Porosity, TOC

HI = hydrogen index; OMT = organic-matter type; TOC = total organic carbon.

the vertical profile of TOC has been related to physiographic setting and sequence-stratigraphic units (i.e., parasequence, parasequence set, sequence, and sequence-set scale; see discussions in Creaney and Passey (1993), Bohacs (1998), and Bohacs et al. (2005)).

TOC content can be measured through direct combustion, modified direct combustion, indirect (by difference), and pyrolysis plus combustion products. Each method has advantages and disadvantages; see Peters and Cassa (1994) for a detailed discussion. We recommend the direct combustion method for most mudstones.

Sample Selection and Analytical Considerations—A few considerations regarding sample selection and analysis include the following:

- Sample size is typically approximately 250 mg; richer rocks require smaller samples. (See discussion in the previous sections about selecting representative samples for sequence-stratigraphic analysis.)
- TOC is expressed as the weight percent of organic carbon in a sedimentary rock.
- TOC analyses are relatively rapid and inexpensive.

Tips and Traps

 This measure is useful for the identification of mudstones enriched in organic matter as well

- as for generating vertical profiles of TOC content in mudstone successions.
- Samples for reliable analyses should be at least several centimeters in thickness, free of modern-day plant roots, and relatively unweathered (as indicated by the lack of oxidation of such reactive minerals as pyrite).
- The measured values of mature samples need to be corrected to estimate the original values at the time of deposition because as thermal maturity increases, TOC decreases.

Rock-Eval® Pyrolysis: Hydrogen and Oxygen Indices

Rock-Eval® pyrolysis is a rapid screening method for estimating the hydrogen and oxygen content of organic matter and provides insights into (1) the character of the original organic material (algal versus land plant), (2) the preservational conditions in the depositional environment (e.g., Peters et al., 2005), and (3) the quality and distribution of hydrocarbon sources and reservoirs (Table 4). Vertical profiles of hydrogen indices (HIs) have been related to physiographic setting and sequence-stratigraphic units and surfaces (e.g., Creaney and Passey, 1993; Bohacs, 1998; Bohacs et al., 2013). For more information on the Rock-Eval method and parameters, see Espitalié et al. (1977, 1985), Lafargue et al., 1998, Peters (1986), and Tissot and Welte (1984).

Sample Selection and Analytical Considerations

- Sample size can range from 100 to 500 mg, depending on the TOC content; richer rocks require smaller samples. (See discussion in the previous sections about selecting representative samples for sequence-stratigraphic analysis.)
- The Rock-Eval analysis is relatively rapid and inexpensive, making it quite useful for screening many samples and generating vertical profiles in mudstone successions.
- It requires the measurement of TOC content to calculate HI and oxygen index (OI).

Rock-Eval® Parameters

- Measurements include three peaks, S1, S2, and S3, as well as T_{max}.
- **Hydrogen Index (HI):** HI = [S2/TOC] × 100 (mg "hydrocarbons"/g organic carbon). It estimates the quantity of pyrolyzable organic compounds or "hydrocarbons".
- Oxygen Index (OI): OI = $[S3/TOC] \times 100$ (mg CO_2/g TOC). It is a function of the type and

preservational state of the original organic matter. It tends to be higher in terrigenous organic matter and in oxidized aquatic organic matter. It is used together with HI to make inferences about the organic-matter type.

- **Production Index (PI):** PI = S1/[S1+S2]. It estimates the amount of free hydrocarbons in the rock, which is a function of porosity and thermal maturity.
- T_{max} : It is the temperature at which the maximum amount of S2 is generated. It is related to thermal maturity (as well as the organic-matter type).

Tips and Traps

- Insights into the original organic-matter type and depositional conditions are best provided by cross-plots such as S2 versus TOC and HI versus OI. See Peters and Cassa (1994) and Peters et al. (2005) for more information.
- Insights into sequence stratigraphy are best provided by vertical profiles of HI and TOC plotted together with well-log and paleontological data.
- Rock-Eval results are significantly affected by the level of thermal maturation: HI and OI decrease with increasing maturity.
- Rock-Eval results are also affected by outcrop weathering (decreases HI and increases OI) and by staining by migrated hydrocarbons or oil-based drilling muds (increases S2 and HI).
- Rock-Eval analysis is not useful for rocks with less than 0.5 wt.% TOC.

Visual Kerogen Description and Palynology

The definition of the term "kerogen" has evolved through time. See the discussions in Hunt (1979), Durand (1980), Tissot and Welte (1984), Tyson (1995), and Vandenbroucke and Largeau (2007). Here, we define kerogen as the sedimentary organic matter that is not soluble in common organic solvents. Following TOC and Rock-Eval analyses, selected samples may undergo visual kerogen description to determine the type of organic material present in a mudstone. Selected samples may also be analyzed for palynomorph (e.g., spores) and phytoclast identification.

Organic material includes aquatic algae, bacterial debris, land plant parts (wood, cuticle, etc.), charcoal, spores, pollen, and bitumen. Commonly, the structure of the original material is preserved, and a variety of macerals can be identified under the microscope. Both visual kerogen and palynology analyses provide direct insights into the origin of organic-material particles and depositional conditions (including redox levels, proximity to terrigenous provenance, and paleoclimate from land-plant spores and pollen), biostratigraphy, and maturity. Vertical trends of appropriately spaced samples add an almost invaluable dimension to sequencestratigraphic analysis. Sequence-stratigraphic surfaces are commonly marked by distinct changes in organic matter across them (see discussions and examples in Bohacs et al., 2022a, b, c, Chapters 5–7). Assemblages of palynomorphs in lacustrine mudstones are helpful for discerning lake-basin type (sensu Carroll and Bohacs, 1999), which then provides specific hypotheses about the associated sequence-stratigraphic expression (e.g., Bohacs et al., 2000; Carroll and Bohacs, 2001).

During these analyses, organic material is separated from crushed samples of whole rock using a variety of physical and chemical techniques in succession (including ultrasonic disaggregation, heating, oxidation, surfactants, flotation, and centrifugation). The resulting concentrated organic material is examined by optical (including reflected and UV light) or electron microscopy or both, and the occurrence, character, abundance, and taphonomy of organic material separated from whole-rock samples are described. Sample size is generally about 1–5 g (0.5–2 cm³) as a function of the organic-matter content of the whole rock (estimated by TOC). For more information, see Durand (1980), Tyson (1995), Taylor et al. (1998), and Vandenbroucke and Largeau (2007).

Sample Selection and Analytical Considerations

- Sample suites should span the full range of lithofacies present in an interval, with similar facies collected at multiple stratigraphic levels.
- One can sometimes make good use of small pieces that have fallen off into the core box if there is not too much vertical shifting.
- Outcrop samples with some weathering can still be quite useful, as the first step in many processing protocols is oxidation.
- The presence of solid bitumen in rocks with low permeability suggests that some oil generation has occurred locally.

Tips and Traps

Identification of particular types of organic matter is quite challenging, especially in thermally mature samples. This can cause varying interpretations by different analysts as well as complicate your own. Terminology also varies

- widely, and some translation between different analysts is commonly needed.
- Cutting samples can be used, but they are prone to contamination from cavings and the presence of drilling additives (such as lignite and walnut shells).

Molecular Geochemistry

Molecular geochemistry has also been used to characterize the organic matter present in mudstones and to make inferences about past depositional environments. Biomarkers are molecular-compound fossils that are structurally related to their precursors in once-living organisms (e.g., Peters et al., 2005). The character of a biomarker is a function of the original organism as modified during deposition and later thermal maturation and biodegradation. Hence, biomarkers can provide insights into the age and composition of the original mix of organic matter and such depositional conditions as redox levels, salinity, and water column stratification (Peters et al., 2005, and references therein). Vertical profiles of the type and abundance of molecular markers are needed for sequence-stratigraphic analyses—these profiles must be specifically acquired from representative samples to reveal long-term secular trends (e.g., Mello et al., 1993; Isaksen and Bohacs, 1995; Bohacs et al., 2000). Samples need to be taken from equivalent facies or locations with respect to significant surfaces (e.g., parasequence boundaries) to discern secular trends accurately. In lacustrine mudstones, small sample suites can be quite useful in determining the most likely lake-basin type (sensu Carroll and Bohacs, 1999), which then provides specific hypotheses about the sequence-stratigraphic expression in the strata (e.g., Bohacs et al., 2000; Carroll and Bohacs, 2001). For more information on analytical techniques and applications, see Philp and Gilbert (1986), Philp (1993), Schouten et al. (2001), and Peters et al. (2005).

Sample Selection and Analytical Considerations

- The size of rock samples is typically approximately 50 g (~20 cm³ = 0.25 cm thick piece of a 4 in. slabbed core), although samples as small as 1–2 g can be analyzed successfully if they are sufficiently rich in organic matter.
- Screen the potential samples first for TOC content and Rock-Eval parameters. Molecular analyses are not recommended for samples with TOC <1 wt.% (e.g., Rowland and Maxwell, 1984; Farrimond et al., 1989).
- The output of the molecular analysis represents the mixture of the many compounds of organic matter present in the tens of grams of a sample.

Tips and Traps

- Examine a variety of molecular markers; make interpretations based on preponderance of evidence and not on any single compound or ratio.
- Calculated molecular ratios may differ from laboratory to laboratory as a function of the instrument(s) and techniques used.
- Caution must be exercised in interpretation as biomarker character is the result of a long chain of transformations from photosynthesis to thermal maturation. For best results, use rock samples of low thermal maturity and, in the case of oil, of minimal biodegradation.
- Biomarkers represent only a tiny fraction of the organic matter, which itself is only a small part of the whole rock, so obtaining representative samples is essential, as is restraint in extrapolation (upscaling) of analytical results.
- All analyses are more meaningful when conducted on representative samples and interpreted within the geological context.
- Avoid contamination of samples: Enclose them in an aluminum foil (baked at ~500°C) before wrapping or bagging in plastic. Do not label rock samples directly as marker pen ink is particularly problematic.
- Contamination with oil- and water-based drilling muds and mud additives may impact the results of molecular analyses.

Stable Isotopic Analysis

Isotopes are variants of a chemical element with different numbers of neutrons (e.g., Faure, 1986; Hoefs, 1997). The two types of isotopes are unstable (radioactive) and stable (nonradioactive). Radioactive isotopes have provided the information needed for establishing the geological timescale. Stable isotopes have provided valuable insights into the origins of the component grains in mudstones, timing of cementation, and temperatures of precipitation (e.g., Hoefs, 1997; Brownlow, 1996; Klein et al., 1999). In addition, stable isotopes have been used to link regional stratigraphy to events and settings characterized by distinct isotopic excursions from typical values present in adjacent sediments (e.g., Veizer et al., 1980; Joachimski and Buggisch, 1993; Jenkyns et al., 1994; Menegatti et al., 1998; Jenkyns and Wilson, 1999; Kump and Arthur, 1999; Danelian et al., 2004; Buggisch and Joachimski, 2006; Dumitrescu and Brassell, 2006). Stable isotopes of five elements, carbon (C), oxygen (O), hydrogen (H), sulfur (S), and nitrogen (N), have been used extensively for the study of mudstones. See discussion and references in Potter et al. (2005) and Schieber (1998c). A short introduction to C and O stable isotopes follows next.

The isotopic signature of a chemical element is expressed by the "delta" (δ) notation. The relative abundance of isotopes is recorded and reported as the ratio of the heavier-to-lighter isotopes in the sample compared to the same ratio in a chosen standard (e.g., the Vienna Peedee Belemnite for C and the Vienna Peedee Belemnite or the Vienna Standard Mean Ocean Water for O). Ratios of stable isotopes are expressed as parts per thousand ("per mil," %). The varying stable-isotope ratios of elements such as C and O are present in mudstones in minerals such as calcite, quartz, and clay (see Potter et al., 2005, appendix A.7, for examples of C and O isotopic variations in rocks and minerals). Isotopic values reflect varying contributions of detrital minerals, weathering products (e.g., clay minerals), cements, and organic materials (e.g., Lonstaffe, 2003; Potter et al., 2005), and are influenced by diagenetic processes and thermal maturity (e.g., Hudson, 1977; Joachimski et al., 2003; Machent et al., 2007, 2012).

In sequence-stratigraphic studies, stable-isotope analyses can be used to refine the understanding of the origin of specific components, thereby aiding the interpretation of the environments of deposition (e.g., Schidlowski et al., 1983; Machent et al., 2007, 2012). Such data are particularly useful where they can show the changing balance of reservoirs that contributed solutes to specific reaction products. For instance, isotopic analyses of C in carbonate cements can be used to identify whether or not cements precipitated from (1) waters in equilibrium with the sea-water marine reservoir (δ^{13} C typically on the order of 0‰), (2) the anoxic pore waters associated with sulfate reduction ($-25\% \delta^{13}C$), or (3) the anoxic conditions associated with methanogenesis (+15% δ^{13} C; e.g., Irwin et al., 1977). As part of these analyses, it is common to compare the δ^{13} C data with data derived from δ¹⁸O (having made assumptions concerning starting isotopic composition of the pore waters), so that estimates of the temperature of cement precipitation can also be made.

Variations in the sources of organic carbon can also be tracked by isotopic analyses in mudstones as stable isotopic signatures of C vary depending upon the precise mechanisms of organic-carbon production and the subsequent degree of oxic degradations, and are thus very useful when relating variations in source-rock quality in sequence-stratigraphic context. For instance, because of the existence of different photosynthetic pathways and differences in the stable isotopic composition of C in the marine carbonate

reservoir (0% δ^{13} C) and CO₂ in the atmosphere (-8% δ^{13} C), marine phytoplankton typically exhibits stable isotopic values on the order of -10% to -31% δ^{13} C, whereas land plants are typically in the range of -33% to -23% for C₃ plants and -16% to -9% δ^{13} C for C₄ plants (e.g., Smith and Epstein, 1971; Stuiver, 1978; Farquhar, 1980, 1983; O'Leary, 1981; Schidlowski, 1988; Fogel and Cifuentes, 1993; Killops and Killops, 2005).

Sample Selection and Analytical Considerations

- Representative samples of each facies within the geological context are collected to enable robust interpretation and extrapolation.
- Generally, samples are combusted or pyrolyzed, and then the target gas species is purified by some combination of filters, traps, catalysts, and gas chromatography (GC). Following separation by GC, isotope ratios can be obtained using an isotope-ratio mass spectrometer.
- For inorganic carbon and oxygen of diagenetic cements, 50-100 mg of rock are used in the analysis. These samples are treated with acid to remove all organic matter before analysis.
- For organic carbon, 5-20 mg of rock extract is used in the analysis. Alternatively, 50–100 mg of whole rock is crushed and treated with acid to remove all carbonate carbon. Caution must be used, as the acid treatment can dissolve some labile carbon compounds (typically enriched in ¹³C) and thus change the isotopic value of the bulk carbon.
- For more information on the technique, see, for example, Faure (1986), Hoefs (1997), Coplen (1996), Werner and Brand (2001), and Lonstaffe (2003).

Tips and Traps

- Samples should be as clean as possible to not introduce contaminants.
- Reliable interpretation of stable isotopic data requires that the mudstones being analyzed have not undergone significant alteration since they accumulated (e.g., recrystallization or some other form of isotopic exchange), and the range of materials that are actually being sampled and analyzed is broadly known, particularly where sampling constraints mean that multiple grains or cement generations are being sampled.
- As for all analytical data, the most effective interpretations should always be made within a geological context.

Well-Log Tools

From bathtub experiments to modern nuclear physics, formation logging has been the backbone of oil and gas development, letting geologists, geophysicists and engineers see more and more of the subsurface.

—Pike and Duey, 2002, p. 52

Well-log analysis is the most complete and least expensive method of characterizing mudstones downhole. This method provides robust characterization of the stratigraphic distribution of potential source, reservoir, and seal rocks. It is particularly valuable where sample data are not available or for lake strata where thin-bedded source rocks are common. Few well-log types, however, directly measure the rock properties sought to evaluate source, reservoir, or seal potential—it is essential to understand the assumptions made, and caution must be used in interpretation.

The three main categories of well logs are electric, radioactive, and structural; most well logs were originally designed and mainly used for estimating lithology, porosity, rock strength, pressure, hydrocarbon presence and type, and structural characteristics (fractures, faults, folds, in situ stress). Table 5 summarizes the principles of well-logs commonly used for sequence-stratigraphic analysis and provides helpful tips and traps of log applications to mudstone successions. Many excellent textbooks provide more detail on well-logs and their interpretation (e.g., Serra, 1984; Schlumberger Limited, 1991; Emery and Myers, 1996; Rider, 2002; Evenick, 2008). A few general tips and traps to have in mind when using well logs are as follows:

- Direct measurement of rock properties from samples is always the most accurate information you can obtain and is invaluable for calibrating well-log estimates. Sample coverage, however, tends to be sparse and unevenly distributed; hence, calibrated well-logs are essential for evaluating the entire section under examination.
- Well-log estimates should be calibrated against values measured from appropriate samples whenever possible. Note, however, that the difference in sample volume analyzed versus that measured by a well-log tool (cm³ vs m³) makes caution essential in any calibration. See Passey et al. (1990) for a discussion of this issue for organic-matter-rich mudstones.
- Many unusual lithologies can occur in mudstone successions, especially in those that accumulated in lake systems—and mudstones are not always identifiable by a high gamma-ray signature. Lacustrine rocks commonly pose special challenges

- to well-log analysis because of their thin beds and varied lithologies. Their well-log responses can differ significantly from marine rocks, especially with gamma-ray and resistivity tools (e.g., Bohacs and Miskell-Gerhardt, 1998; Bohacs et al., 2001; Martel, 2013).
- Well-log analyses of mudstones require special attention to the standard procedures for depth alignment in combined log techniques, tool calibration, and tool baseline. Pay particular attention to the caliper log, for some mudstones tend to wash out, whereas others drill "gunbarrel straight."

For source evaluation, TOC depth profiles estimated from well-log data can be used to calculate TOC thickness, organic-matter volume, and the amount of hydrocarbon generated from a source rock over a particular area of a basin. The TOC profile also helps to interpret mudstone depositional environments and to correlate stratigraphic sequences (e.g., Creaney and Passey, 1993; Bessereau and Guillocheau, 1995; Bohacs, 1998). Mudstone reservoir and seal character can be estimated through analogous analyses using combinations of various well-logs (e.g., Spears and Jackson, 2009; see Table 5); the vertical profiles generated also provide useful insights for constructing sequence-stratigraphic frameworks (e.g., Bohacs, 1998).

Seismic Tools

The unconformity-bounded sequences and their content comprise the most meaningful segments of the present stratigraphic record, without which valid interpretation of geologic history is patently impossible.

—Wheeler, 1964, p. 600

This section is intended as an introduction and overview to this vast area of investigation. For more detail, we heartily recommend the reader to consult Payton (1977), Sheriff (1980), Abreu et al. (2010), and Veeken and van Moerkerken (2013).

Seismic data have been used over a large range from single wells to large basins such as across the Gulf of Mexico to provide continuous images of the subsurface to delineate stratal patterns and infer depositional history, environments of deposition, lithotypes, and fluid content (e.g., Sheriff, 1980; Veeken and van Moerkerken, 2013).

Seismic stratigraphy is an essential component of sequence stratigraphy. Seismic stratigraphy involves both the (1) seismic-sequence analysis and (2) seismic-facies analysis. The seismic-sequence analysis

Table 5. Well-logs: Applications, Tips, and Traps. See, for Example, Serra (1984), Schlumberger (1991), Emery and Myers (1996), Rider (2002), and Evenick (2008) for More Detail on Well-log Interpretations, Applications, and Caveats.

Log Name/Applications	Tips	Traps
Total gamma ray (GR) • It records the naturally occurring radioactivity of rocks. GR logs provide good estimates of grain size (and subsequently inferred depositional energy). Mudstones commonly have a relatively high GR activity (because of clay-mineral, heavy-mineral, and organic-matter content). GR values are measured in American Petroleum Institute (API) units and range from very few units (in anhydrite) to over 200 API units (in mudstones).	 Marine mudstones have a variety of GR signatures proportional to the amounts of both radioactive clays and organic-matter content. Commonly, a general increase in radioactivity is related to organic matter, but this relation is not always consistent. The GR curve is useful for correlation on a basin to prospect scale. GR logs are particularly useful for identifying parasequence boundaries (flooding surfaces and their correlative conformities; e.g., Bohacs, 1998; Bohacs et al., 2001; Lazar, 2007). 	 Increased GR has in the past been linked to "shale" volume, but increased radioactivity can result from clay, organic matter, heavy minerals, or even increased feldspar. A decrease in GR can be because of relative coarsening or decreased amounts of radioactive clays within a biogenic-rich mudstone. As a general rule of thumb, the GR is quite useful for stratigraphic correlation but before interpreting organic and/or clay content, the spectral GR should be examined and sample calibration should be completed.
• It measures the radiation emitted during the breakdown of uranium (U), thorium (Th), and potassium (K)—key components of mudstones. SGR logs typically provide the concentrations in percentage (K) and in parts per million (ppm) (U and Th). • Various ratios of U, Th, and K have been used to make estimates of the rock character and depositional conditions (e.g., Zelt, 1985; Myers and Wignall, 1987; Isaksen and Bohacs, 1995).	 GR spectra can also be obtained for outcrops or cores using a portable GR scintillometer (e.g., Meyers and Wignall, 1985; Zelt, 1985; Meyers, 1987; Schwalbach and Bohacs, 1992; Lazar et al., 2010, 2015b). The geometry of the outcrop relative to the detector influences the volume sampled—operator must strive for constant geometry (Schwalbach and Bohacs, 1992). 	• See the following for U, Th, and K.
Uranium (U) It is a good indication of relative total organic carbon (TOC) content (in marine organic-matter-rich rocks with significant hydrogen content) and phosphates. It can also relate to TOC content in balanced-filled lakes with crystalline rocks in drainage catchment.	The relation of U to TOC is an indirect one: The presence of U is actually because of the reducing conditions that can arise from the presence of elevated TOC—this is only the case when the TOC is associated with sufficient hydrogen index (HI). Thus, only rocks with elevated TOC and HI will have significant enrichment in U. Rocks with lower HI (because of oxidation or primary input) will not necessarily have a consistent relation of TOC with U.	 Apparent variations in this relation through geologic time are more likely because of different HI of the organic material instead of biological evolution. Caution must be exercised when applying U as a proxy for TOC, and local calibration is required. It is generally not correlative to TOC content in lacustrine strata (especially freshwater or overfilled lake basins with sedimentary rocks in drainage catchment).
Thorium (Th) • It is commonly a good indicator of relative clay content, volcanic ash, and heavy minerals such as monazite and zircon.	The relation of Th to clay appears to be consistent through geologic time. Elevated Th is commonly found in volcanic ash beds.	Quantitative relation of clay to Th might change from basin to basin, but more definitely, it changes with geological age. If no calibration exists, Th is still a good indicator of relative increase in clay, but Th can be elevated also because of volcanic ash.

 Table 5. (Continued)

Log Name/Applications	Tips	Traps
Potassium (K) • K is commonly a good indicator of illite, K-feldspar, micas, and sylvite.	It is a general indicator of clay content because most mudstones are dominated by a subequal mixture of illite and smectite.	The K curve can be severely influenced by K-based muds, which would give the impression of a higher total GR or higher K content.
Density (RHOB) • It measures the formation's bulk density (mineral matrix and pore fluids) and is most commonly used to estimate porosity. Different lithologies can also be determined based on the measured density value. It is commonly used in conjunction with neutron logs to determine the lithology of formation (density-neutron suites such as Schlumberger FDC-CNL suite). Organic matter, because of its low grain density, can have a significant impact on the bulk density measured by well-log tools (e.g., Meyer and Nederlof, 1982; Passey et al., 1990).	 In mudstone reservoirs, the RHOB curve typically provides the most accurate estimate of total porosity. Lower RHOB can give a fair indication of elevated TOC content (if there are no large local variations in other components that affect bulk density), but this complicates the calculation of porosity in organic-matter-rich rocks. This log's relatively fine vertical resolution (~30 cm [1 ft]) can be useful for differentiating small and closely spaced vertical variations in TOC. It can also be used in combination with a resistivity log to estimate the TOC content of mudstones (e.g., Passey et al., 1990). 	 Lower bulk density can indicate higher TOC; care must be taken to explicitly account for the density of all grains (including kerogen) when developing a mineralogy-based porosity from RHOB. It is likely affected by borehole conditions (need to examine the caliper log).
Neutron (NPHI) • It measures the formation's porosity through detecting the quantity of hydrogen present (assumed to be in the water or hydrocarbons in the pores). The log is calibrated to a limestone matrix. The linear limestone porosity units are calibrated using the API neutron test pit wherein the 19% porosity, water-filled limestone is defined as 1000 API units. This log is also useful in estimating rock composition.	 Increased NPHI can give a good indication of increased clay content resulting from the presence of hydroxyl (OH⁻) in clay minerals. Similar to conventional reservoirs, the NPHI-RHOB separation can show relative clay content, and the NPHI-RHOB crossover can be a good indication of light hydrocarbons (i.e., gas). Generally, the NPHI log by itself is a poor indicator of organic-matter content as it is affected not only by the hydrogen in the organic matter but also by the hydrogen in clay minerals (OH⁻), formation water, and any liquid or gaseous hydrocarbons present. 	• Note that the clay effect on the NPHI log is opposite to the gas effect, so in clay-mineral–rich intervals, gas crossover may not be apparent. To confirm this relation, all the available data (e.g., Th, K, and RHOB measurements) should be used.
Sonic (acoustic; DT) • It measures the capacity of a formation to transmit sound waves and is related to both the porosity and the lithology of the rocks being measured. Thus, if the lithology of a formation is known, this log can be used to determine its porosity. Organic-matter content has a significant influence on compressional (P-wave) velocity, which can be calibrated to TOC content (assuming no significant local variations in porosity or bulk mineralogy). • It assists in seismic interpretation (gives interval velocities and the acoustic impedance log).	 Mudstones have a lower velocity (higher transit time) than sandstones with the same porosity, making this log a good indicator of grain size. It can be useful for estimating porosity but is not as accurate in low-porosity rocks as the RHOB curve. TOC content is likely to impact the shear (S) wave response (e.g., Zhu et al., 2010). The DT log is best used in combination with a resistivity log to estimate the TOC content of mudstones (e.g., Passey et al., 1990). 	DT logs can provide the most accurate estimate of porosity when pyrite and other heavy minerals are present.

 Table 5. (Continued)

Log Name/Applications	Tips	Traps
Spontaneous Potential (SP) • It measures the electrical current that occurs naturally in boreholes as a result of salinity differences between the formation water and the borehole mud filtrate (formation and surface). SP logs are used as indicators of permeable beds or for locating bed boundaries.	 The SP log was one of the first tools to be used to distinguish mudstone from sandstone in clastic sequences (zero matches pure mudstone, and high SP values match sandstone; Schlumberger et al., 1932). It can be useful in conventional and unconventional reservoirs as it gives an indication of possible permeable intervals (potential reservoirs). 	It can be useful, but the interpreter must be aware of the water type (saline or fresh).
 Resistivity-lateral (R-L) It measures the bulk resistivity of a formation (the reciprocal of conductivity). This log is very useful for determining the type of fluids in formations and is commonly used as an indicator of formation lithology. Resistivity is, in general, a function of porosity and pore and bound fluid in a rock. 	 It is the preferred log for rocks with resistivity values greater than 100–200 Ω m²/m. Resistivity typically decreases when rocks are saturated with saline water and increases with increasing hydrocarbon saturation. Clay content, pyrite content (only a factor when very concentrated), and other conductive components present at high thermal maturities in mudstones can influence resistivity (see discussion in Passey et al., 2010). Resistivity tends to be lower in argillaceous fine mudstones and higher in oil-bearing porous lithologies. 	• In mudstone reservoirs, other lithologic effects can affect the log response even if present in small amounts. Intervals with pyrite content exceeding 5–10 wt. % can be conductive and can lower the resistivity response, masking a hydrocarbon response. Moreover, small amounts of bound water on clay films can lower resistivity resulting in a similar outcome (e.g., a shaly sand). In contrast, slightly elevated resistivity can be measured where carbonate cements are present. It is recommended to use other logs and geological context to build a case for presence of hydrocarbons.
Resistivity-induction (R-I)	• It is the preferred log for rocks with resistivity values smaller than 100–200 Ω m ² /m. Resistivity typically decreases when rocks are saturated with saline water and increases with increasing hydrocarbon saturation.	• As for Resistivity-lateral (R-L), above.
It measures borehole size, shape, and orientation.	Borehole shape may ultimately indicate borehole stability and local stress state.	 One-arm caliper does not always give an accurate representation of the complete borehole. Look for the multiple-arm caliper tool, if present, for density correction.
 Borehole-image log It provides a relatively high-resolution image of the face of the borehole, most commonly using resistivity, but also through DT, RHOB, GR, or video, or combinations thereof. It also serves as a high-resolution dipmeter. This log can be used to measure the occurrence, orientation, and spacing of bedsets, bedding surfaces (including unconformities), folds, fractures, and faults. These data can be used to interpret vertical trends in bedset thickness and bedding orientation, sediment transport direction, and fold axes. It can also 	• A resistivity-image log can resolve features of less than a centimeter (~10× that of an acoustic image log). It can be run in aqueous or nonaqueous drilling fluids and on wireline or logging while drilling. An overall resistivity curve calculated from this log has been used to conduct very-high-resolution Delta Log R analysis (e.g., Passey et al., 2010). From this analysis, individual parasequences (as thin as 0.9 m) can be identified readily; parasequences have high TOC and are more resistive (bright) at the base, and have low TOC and are more conductive (darker) upward.	• The relatively high resolution does result in large datasets that must be processed, analyzed, and displayed—it is very helpful to use the variety of displays (raw, static, dynamic range) and try different color schemes to highlight various features in the rocks. Vertical resolution is likely to be coarser in nonaqueous fluids.

 Table 5. (Continued)

Log Name/Applications	Tips	Traps
help distinguish natural and induced fractures and provide insights into the relation of fractures to bedding.		
Elemental/mineralogic/geochemical logs It estimates rock composition through the use of a suite of well logs, typically including GR, resistivity, DT, RHOB, and NPHI. Some methods use active source nuclear spectroscopy (e.g., Radtke et al., 2012), although their generally very slow logging speeds tend to result in sparse usage. Log measurements, tool uncertainties, and component response parameters are used to compute statistically the volume of minerals and fluids in the formation (Herron and Herron, 1990).	 Various approaches are employed for these computations: simultaneous sets of response equations, multivariate cluster analysis, and error minimization. These logs can be reasonably accurate for total clay, carbonate, and quartz content (at least within ±5–10 wt. %). Vertical profiles of rock compositions can be used to augment the recognition of sequence-stratigraphic surfaces and units by providing more dimensions of variation. Compositional parameters that can be related to sedimentary provenance, depositional conditions (especially redox), and organic-matter character could be of particular utility. These logs appear to be useful in identifying pyrite and/or siderite-rich intervals—intervals that are commonly associated with sequence-stratigraphic surfaces in distal environments where clastic input is minimized (see discussion in Passey et al., 2010, their figure 4). 	Log estimates of rock composition can provide some insights into grain origin and depositional conditions, subject to the caveats discussed in Chapter 2 and "Geochemical Tools" section (e.g., carbonate content can have multiple origins: detrital, biogenic, or diagenetic).
Derived logs		
• It employs the overlaying of a porosity log (DT or RHOB) on a resistivity curve. The curve separation is termed "Delta Log R" (Passey et al., 1990). A TOC vertical profile can be calculated knowing the value of the separation and the level of thermal maturity of the rocks.	 It is useful for displaying vertical trends in organic-carbon content and identifying source-rock settings and sequence-stratigraphic units and surfaces (Creaney and Passey, 1993; Bohacs, 1998). Both types of porosity tools (sonic, density) should be tried, and the results for each particular unit should be compared. 	 Logs should be depth-aligned; be aware of washed-out wellbore segments. Note that TOC calculations are only calibrated up to an R_o of 1.0; the vertical trends in log separation (Delta Log R) are still useful at higher maturities for identifying organically enriched intervals and trends. Local calibration with samples is essential, especially in higher maturity zones.
Poisson's ratio It is a measure of rock compressibility perpendicular to the applied stress. It is based on Poisson's effect—rocks compressed or stretched in one direction tend to expand or contract in the other directions.	 It is useful for showing variations in rock ductility. It is essentially a Vp/Vs curve and so requires a dipole sonic log for measuring both P and S waves. Different cutoffs should be used in different plays to show which rocks are more suitable for hydraulic fracturing. 	The dipole can be affected (which can affect the calculation of Poisson's ratio) if there is borehole washout.
Young's modulus • It is a measure of rock stiffness, that is, the ratio of stress to strain.	It is useful for showing which rocks will stay propped open more readily when hydraulically fractured.	• It is derived from the RHOB and DT logs, so the caliper character in the zones of interest is noted.

subdivides the seismic section into geometrically defined intervals that are the seismic expression of depositional sequences—stratigraphic units of relatively conformable, genetically related strata bounded by unconformities and their correlative conformities. The seismic-facies analysis examines the configurations of reflections interpreted as strata within depositional sequences to determine the environmental setting and to estimate lithology (Mitchum and Vail, 1977). Although there are numerous software tools for assisting these analyses, it is essential that the user understands the fundamentals to appropriately apply these tools.

Seismic Sequence Analysis

Seismic stratigraphy is the study of stratigraphy and depositional facies as interpreted from reflection seismic data. Seismic reflection terminations and configurations are interpreted as stratification patterns and are then used for recognition and correlation of depositional sequences, interpretation of depositional environment, and estimation of lithofacies (e.g., Mitchum et al., 1977). It is basically a geological approach to interpreting geophysical data that aims to extract the maximum amount of information regarding paleogeomorphology, depositional environments, and facies through adding geological context and sedimentological insights.

Seismic reflections approximate bedding surfaces that record depositional profiles and therefore chronostratigraphic surfaces. This enables one to use the geometry of seismic reflection patterns to interpret

- chronostratigraphic correlations,
- genetically related depositional units,
- environments of deposition from depositional topography,
- paleobathymetry,
- topography and relief on unconformities,
- post-depositional structural deformation and syndepositional thickness changes,
- burial history, and
- paleogeography and geological history.

Rock type cannot be determined directly from reflection patterns but can be estimated by the two-step process of interpreting environments of deposition and then using insights from models and observations of the depositional processes and facies.

Seismic-sequence analysis subdivides seismic sections into packages of concordant reflections (seismic sequences, interpreted as depositional sequences) that

are separated by surfaces of discontinuity identified by systematic reflection terminations (e.g., Mitchum et al., 1977; Ramsayer, 1979). The two fundamental types of terminations are lapout and truncation. Lapout is the lateral termination of a stratum at its original depositional limit; truncation is the lateral termination of a stratum as a result of being cut off from its original depositional limit by subsequent erosion or deformation (e.g., Mitchum et al., 1977). Reflection terminations interpreted as stratal terminations include onlap, downlap, toplap, and erosional truncation (e.g., Mitchum et al., 1977).

Seismic-Stratigraphy Procedure

The first step in seismic stratigraphy is to identify seismic terminations, and then use those terminations to recognize surfaces and delineate them. Next, key surfaces such as sequence boundaries and maximum flooding surfaces (maximum transgressive surfaces) are traced on the intersecting lines in a grid of seismic lines or propagated through a seismic-data volume until the boundaries have been correlated and tied within the entire dataset. This is intended to verify the regional extent of major discontinuity surfaces and to identify which other potentially significant surfaces are local in nature. Table 6 outlines the workflow for seismic stratigraphy. Table 7 details the criteria for identifying key stratigraphic surfaces.

Seismic Facies Analysis

Seismic-facies analysis is the description, mapping, and geologic interpretation of seismic-reflection parameters within a chronostratigraphic framework

Table 6. Seismic-stratigraphy Workflow.

Step	Action
1	Investigate the background of the dataset. a. Geophysical—acquisition, processing, display b. Geological—age, tectonic setting, basin phase
2	Identify large-scale packages (tectonostratigraphic units).
3	Make detailed interpretation of sequences: surfaces, grid/volume ties, maps.
4	Map seismic facies within genetically related packages (sequences or systems tracts).
5	Interpret seismic facies in terms of depositional environments and potential lithofacies.
6	Integrate seismic stratigraphy with other datasets: well logs, core, paleontology, geochemistry, etc.

Table 7. Surface Definitions With Translation Terms, and Primary and Secondary Recognition Criteria.

Surface	Translation Terms	Primary Recognition Criteria	Secondary Recognition Criteria (based on limited available data)
Maximum Flooding Surface (MFS)	Maximum Transgressive Surface (MTS)	Atop maximum landward position of the shoreline within a sequence	Downlaps ^s ; Turn around in stacking pattern from retrogradation to aggradation or progradation ^{w,c,o}
Transgressive Surface (TS)	Maximum Regressive Surface* (MRS)	Atop maximum basinward position of the shoreline within a sequence	Surface beneath first backstep (landward step) of shelf-slope break ^s ; Turn around in stacking pattern from progradation or aggradation to retrogradation ^{w,c,o}
Sequence Boundary (SB)	Sequence Boundary (SB)	Below abrupt basinward shift in shoreline position at base of sequence	Surface beneath first increase in accommodation after progradation or degradation; Break in shoreline trajectory "S"s; Truncation and/or toplap below, onlap above ^s ; Abrupt occurrence of proximal facies over distal facies ^{w,c,o}

^{*}sensu Embry and Catuneanu (2002).

Data for recognition criteria marked with superscript letters: s = seismic; w = well; c = core; o = outcrop (after Abreu et al., 2014).

of sequence boundaries and downlap surfaces (after Mitchum et al., 1977). The interpreter delineates the external form, internal reflection characteristics, and 3-D associations of seismic-facies units—this enables interpretation of the units in terms of environmental setting, depositional processes, and estimates of lithotype. This interpretation is always done within a stratigraphic framework of depositional sequences to insure the analysis of genetically related strata. The genetically related stratal intervals are usually at the sequence or systems-tract scale but may range from sequence sets in very distal areas on 2-D regional lines to parasequences in sculpted subvolumes of 3-D data.

Basic seismic-facies information includes

- the relation of reflections to their upper and lower bounding surfaces (segments of onlap, downlap, toplap, truncation, and concordance; and direction of onlap and downlap), and
- the dominant types of reflection configuration between the upper and lower bounding surfaces (parallel, divergent, sigmoid, oblique, etc.).

In many instances, one aspect of the seismic facies proves more significant than others, and a separate map is prepared to portray that single aspect (Mitchum et al., 1977). For example, dip direction of downlapping reflections is important in interpreting the sediment transport direction and possible provenance areas. In other cases, displaying some combination of characteristics, such as thickness and seismic amplitude, can be revealing.

Seismic-Facies Parameters: Internal Form—Seismicfacies units are mappable, 3-D seismic units composed of groups of reflections whose parameters differ from those of adjacent facies units (Mitchum et al., 1977). These parameters include the configuration or geometry, continuity, amplitude, frequency, and interval velocity (Table 8). Each parameter provides considerable information on the geology of the subsurface (Table 8). Reflection configuration reveals gross stratification patterns from which depositional processes, erosion, and paleotopography can be interpreted. In addition, fluid contact reflections (flat spots) are commonly identifiable. Reflection continuity is closely associated with continuity of strata; continuous reflections suggest widespread, uniformly stratified deposits. Reflection amplitude contains information on the velocity and density contrasts of individual bedding interfaces and their spacing. It is used to predict lateral bedding changes and hydrocarbon occurrences. "Frequency" (reflection spacing), although a characteristic of the seismic pulse, is also related to such geologic factors as the spacing of reflectors or lateral changes in interval velocity (because of organic-matter or hydrocarbon content or lithofacies changes).

Grouping these seismic parameters into mappable seismic-facies units facilitates their interpretation in terms of depositional environment, sediment

Table 8. Seismic-reflection Characteristics of Seismically Definable Rock Bodies.

Parameter	Geologic Information
Amplitude	 Impedance contrast (significant stratal surfaces) Bed spacing/tuning Lithofacies Fluid content
Continuity	 Lateral stratal continuity Depositional processes
Frequency	Bed thicknessFluid content
Geometry	• Depositional processes

provenance, and geologic setting. Seismic-reflection configuration or geometry is the most obvious and directly analyzable seismic parameter. Stratal configuration or geometry is interpreted from seismic-reflection configuration and refers to the geometric patterns and relations of strata within a stratigraphic unit. These commonly indicate depositional setting and processes as well as later structural movement.

Seismic-Facies Mapping—Seismic-facies mapping involves analysis, on a qualitative or quantitative basis, of seismic character to infer areal trends in paleoenvironment and lithofacies. This section introduces a variety of techniques for mapping seismicfacies that can be used on the range of data from 2-D paper to 3-D volumes. The overall approach remains the same, but a particular technique is chosen based on dataset, resources, business stage, and objectives of study. Even with high-resolution 3-D datasets, it is generally useful to plot representative regional strike and dip lines on paper as a first step in defining the range of seismic facies and their rough distribution. Once the range of facies is defined, along with the dip and strike of the basin, then further interpretation and mapping is commonly conducted on a workstation.

Seismic-analysis computer programs can provide quantitative measures of frequency, continuity, and amplitude to support mapping. Seismic-amplitude mapping is particularly well developed in industry. Seismic-volume interpretation allows viewing of seismic-amplitude 3-D objects and their projected polygons in proper spatial and temporal relations.

As a quick reminder, some seismic parameters relate to lithofacies or the processes responsible for deposition. Other parameters relate to acoustic-impedance contrast, tuning, and the interaction of the seismic wave front with the strata, and thus, seismic frequency and resolution are factors in their patterns of occurrence. In the following discussion, stratal continuity is assumed to be detectable (i.e., continuity exceeds the Fresnel-zone width) for a given seismic frequency.

To extract the maximum geological information, one needs to map the smallest unit of genetically related strata, as a practical function of the objective of the interpretation project. For example, individual systems tract commonly have different seismic facies, and each needs its own map. Stacking all the systems tracts of one depositional sequence onto one map, by contrast, makes for a very complicated map that is difficult to interpret in terms of depositional environment or lithofacies (e.g., Armentrout et al., 1993).

External Form and Internal Geometry: A-B/C Mapping—Seismic-facies mapping was definitively explained by Ramsayer (1979), based on 2-D seismic data. It is referred to as the "A-B/C" mapping approach, as observations are made upon the upper boundary (A), the lower boundary (B), and the internal reflection character (C). The A, B, and C categories of Ramsayer's (1979) seismic facies codes each included five types initially, thus providing 15 different variations for a given seismic interval of interest (Table 9). For example, a prograding seismic package with oblique clinoforms, toplap at its upper surface, and downlap at its base would be noted as Tp-Dn/Ob. Subsequent work and incorporation of Campbell's (1967) approach to stratal description expanded the range of descriptors for "C" (internal character) to span continuity, reflection shape, reflection inter-relations, and amplitude (Table 9, the lower part). We highly recommend using all these aspects of reflection character to describe the internal reflection configuration.

The A-B/C mapping is still necessary even with the most advanced and high-resolution 3-D datasets and visualization tools because seismic facies are most meaningful within their stratigraphic context—that is, when applied to genetically related strata (which is the whole point of sequence stratigraphy). Bounding surfaces provide important geological and stratigraphic information. Seismic facies are nonuniquely related to depositional environment and lithofacies (e.g., clinoform reflections can occur in deltaic sandstones, carbonate platforms, and mudstone-dominated shelves); hence, one needs more information from the stratigraphic context of the seismic facies—their relation to lower and upper bounding surfaces and position along the depositional profile.

Table 9. Seismic Facies A, B, and C (Expanded after Mitchum et al., 1977; Ramsayer, 1979).

External Bounding Surfaces—A-B

Тор—А	Con or C	Concordant	Commonly associated with parallel internal reflections
	Top or Tp	Toplap	Nonerosional
	Dwn or Dn	Downlap	Uncommon at the top surface
	Div or Di	Divergent	
	Tr	Truncation	Associated with erosional unconformities
Base—B	Con or C	Concordant	
	Top or Tp	Toplap	Uncommon at the basal surface
	Dwn or Dn	Downlap	Double downlap commonly associated with mounding
	Div or Di	Divergent	
	Tr	Truncation	Uncommon at the basal surface

Internal Reflection Configuration—C*

	Configuration			
		Inter-reflection		
Continuity	Shape	Relations	Reflection Form	Amplitude
Continuous (C)	Planar (Pl)	Parallel (P)	Divergent (Div)	Low (La)
Semicontinuous (sC)	Curved (Cv)	Subparallel (sP)	Convergent (Cnv)	Moderate (Ma)
Discontinuous (D)	Wavy (Wy)	Nonparallel (nP)	Chaotic (Cha)	High (Ha)
		•	Shingled (Sh)	Variable (Va)
			Oblique (Ob)	
			Oblique-parallel	
			Oblique-tangential	
			Sigmoidal (Sig)	
			Complex Sigmoidal-oblique	
			Hummocky (Hum)	

^{*} Seismic facies are defined within genetically related strata bounded by sequence boundaries or downlap surfaces (the A and B surfaces). Evaluation of relative continuity, configuration, and amplitude changes needs to be conducted over equivalent length scales.

Although the A-B/C technique was developed largely using 2-D seismic data, it can be used on 2-D and 3-D sections displayed on conventional industry workstations. The details vary according to the software tools used but generally involve defining a stratigraphic "pseudo-surface" for each depositional sequence or systems tract to represent its internal seismic facies and assigning a distinct time or depth value to represent each seismic facies. A 3-D dataset offers the advantages of allowing determination of the accurate direction of depositional dip and strike before selecting the orientation of the 2-D grid. Such datasets also allow the use of additional nonorthogonal lines to illustrate key stratigraphic relations and to aid interpolation. One can propagate and extend the initial grid of seismic facies interpretation using a variety of 3-D interpretation and visualization tools, some of which are discussed in the following section.

Seismic Facies: Emphasis on Amplitude Continuity—Since Ramsayer's seminal paper in 1979, seismic facies techniques have evolved to include additional information on internal amplitude characteristics within individual systems tracts. Robust seismic facies information related to reflection amplitude, continuity, and frequency can be described in qualitative terms or quantified using various analysis techniques and software products (e.g., Gao, 2011). This approach has been particularly well developed for deep-water paleoen-vironments where amplitude commonly provides significant information on lithofacies and depositional environments (e.g., channel axis versus margin). This technique is most accurate where local well control is

available to calibrate the amplitude-based seismic facies (e.g., Garfield, 2000). Calibrated observations of internal and external seismic character provide a means of interpreting depositional systems directly from seismic data in adjacent areas with little or no well control.

Seismic Facies by Trace Classification—Another innovation in seismic facies involves discrimination and classification of seismic wavelet trace shapes (using various computer programs). The approach must be used within an individual sequence or systems tract to extend what is essentially a one-dimensional seismic-facies analysis to a volume of genetically related strata.

Numerous other methods, both automated and semiautomated, for classifying the seismic data based on detailed geophysical parameters (frequency, wavelet duration, etc.) and on image analysis are available. Techniques using image analysis are particularly suitable for use in seismic facies mapping (e.g., Vinther et al., 1995; DeGroot, 1999; West and May, 2003; Marroquin et al., 2009; Gao, 2011). In one way, they represent just another way to describe and classify seismic facies and can be slotted into the general workflow at step 4 (Table 6). Remember that no matter how high-powered these techniques appear, they still must be applied to a genetically related volume of strata to provide the maximum amount of information on the depositional setting.

TECHNIQUES

The more technique you have, the less you have to worry about it.

> —Pablo Picasso (in Parmelin, 1963, p. 42)

This section details workflows for examining and interpreting mudstones across a broad range of scales, from µm to km, using the broad array of tools discussed previously in this chapter. We start with a workflow for sample-based analyses at the thinsection to outcrop scale. We then specify a workflow for sequence-stratigraphic analysis that summarizes key concepts for incorporating both sample and remote-sensing (well-log and seismic) data into a comprehensive stratigraphic framework. These workflows emphasize the appropriate and efficient use of the tools discussed in this chapter as well as the integration of observations of physical, chemical, and biogenic attributes of the rocks.

Integrated Workflow for Description of Mudstones in Outcrops, Cores, and Thin Sections

This section provides an introduction to the workflow we recommend for examining mudstones in outcrops, cores, and thin sections. Please see Lazar et al. (2015b) for more details on the specific steps that we take and the tools we recommend for describing these finegrained rocks. Our approach is based on the fact that the appearance of mudstones can change significantly as a result of weathering in outcrops or cores (the relatively large content of reactive minerals such as clay and pyrite is very responsive to air and water and mudstones change readily). Describing these rocks in outcrops and cores is therefore a challenge, and, at first sight, one may argue that not much more than color variations can be observed. After preparing a fresh surface (Figure 1), however, detailed and systematic examination of mudstones reveals much about their texture, bedding, composition, vertical and lateral variation, and modes of accumulation. It also enables acquisition of appropriate and representative samples within their sedimentologic and stratigraphic context, and then further analyses under the microscope and in the laboratory.

The steps of the workflow we recommend for examining mudstones in outcrops, cores, and thin sections are:

I. Make observations

A. Establish stratigraphic context

- 1. Step back, examine, and photograph the entire exposure, walking the section several times.
- 2. Check core depths and core box order, clean the core, and photograph the entire core. Step back, examine, and walk the section several times with well logs in hand.
- 3. Look for changes in texture, bedding, composition, thickness, continuity, and stacking; and presence of erosional surfaces. Tentatively identify stratigraphic packages.
- 4. Make note of weathering characteristics and their vertical and lateral distribution.
- B. Start at the base of the section or core, identify, examine, and describe stratigraphic packages and surfaces as you proceed up-section. Integrate outcrop or core observations with thin-section observations. Observe and take pictures of stratigraphic features at all scales (laminae, laminasets, beds, bedsets, parasequences, parasequence sets, and sequences).
 - 1. Examine a fresh face of rocks from each potential stratigraphic package; sample

- each significant facies; mudstones tend to weather deeply and may require much digging to remove the weathered rock and to obtain fresh exposure and samples (Figure 1).
- 2. Describe the mudstones: texture, bedding, and composition. Figure 1A–D in Lazar et al. (2022a, Chapter 2 this Memoir) summarize the terms and definitions we recommend for texture, bedding, and composition. A practical, proxy method, the "scratch test," we recommend to use to determine the dominant grain size at hand specimen scale is detailed in Table 10. Fine-tune outcrop and core estimations of texture, bedding, and composition by integrating observations under optical and electronic microscopes and analytical data obtained on samples taken in sedimentologic and stratigraphic context.
- 3. Describe biogenic sedimentary structures and characterize the degree of bioturbation using a 0–5 scale (figure 1E of Lazar et al., 2022a, Chapter 2 this Memoir).
- 4. Describe the type, size, diversity, abundance (figure 1F of Lazar et al., 2022a, Chapter 2 this Memoir), preservation, and taphonomy (figure 1G of Lazar et al, 2022a, Chapter 2 this Memoir) of body fossils. Similarly, describe trace fossils.
- 5. Describe the type, size, composition, and abundance of diagenetic products

- (figures 1C, F of Lazar et al., 2022a, Chapter 2 this Memoir).
- 6. Organize recurring, representative, and diagnostic facies attributes into facies associations.
- 7. Identify and describe stratal packages and key surfaces (sequence boundaries and flooding surfaces; Figure 3; Tables 7, 11). Obtain a spectral gamma-ray profile to further characterize stratigraphic units (Figure 4).
- 8. Record all information consistently in an appropriate format, designed for your particular setting or unit.

II. Make interpretations

- A. Integrate outcrop, core, or thin-section observations with analytical, well-log, and seismic data as available. Make a hierarchical interpretation of stratal units (laminae, laminasets, beds, bedsets, parasequences, parasequence sets, sequences; significant stratal boundaries).
- B. Make interpretations of dominant sediment provenance, input mode, physical reworking, sediment accumulation rate, completeness of sedimentary record, bottom-water redox conditions, and environment of deposition (e.g., Bohacs et al., 2005; Bohacs et al., 2014; Lazar et al., 2015a, b; Table 2; see also Bohacs et al., 2022a, c; Lazar et al., 2022b; Chapters 4–6 this Memoir)

Table 10. Scratch Test (After Lazar et al., 2015a, b).

Step	Action			
1	Obtain and examine fresh surfaces of rocks from each potential stratigraphic package.			
2	Estimate the percentage of sand-size grains with visual comparison to percentage charts. If less than 50% are sand-size grains, the rock is a mudstone.			
3	Perform a scratch test to determine the dominant grain size. The scratch test is performed by scratching the fresh face of outcrop rocks or the clean back of the core with a sharp steel probe and then observing the luster of the scratch and the color of the powder generated: a Distinctly waxy luster and dark color generally indicates more than two-thirds fine-mud-size material (termed a "fine mudstone"). b Distinctly dull luster and light color indicates more than two-thirds coarse-mud-size material (termed a "coarse mudstone"). c Mid-lustrous scratch and intermediate colored powder indicates medium-mud-size material between two-thirds and one-third (termed a "medium mudstone").			

[—]Interpret results from the scratch test critically because cement (which can lead to apparent coarser interpretations of grain size), aggregate grains (which can lead to underestimation of grain size especially when argillaceous aggregate grains are present), and microcrystalline quartz (which can lead to overestimates of grain size; e.g., Milliken et al., 2012; Milliken, 2013) might be present.

[—]Grain-size assessment can be further enhanced if thin sections are available (e.g., recognize the effects of diagenesis and bioturbation on grain size.

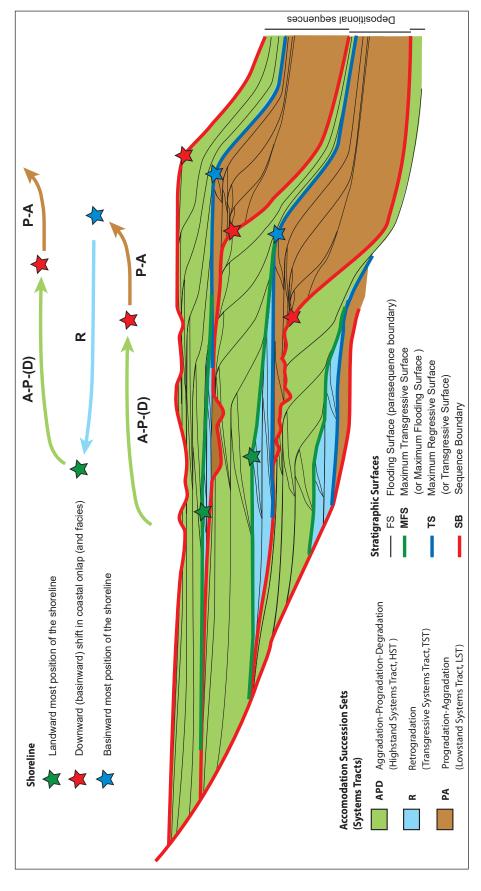


Figure 3. Accommodation succession showing key stratigraphic surfaces, stacking patterns, and depositional sequence expression using definitions outlined in Tables 7 and 11 (after Neal and Abreu, 2009; Abreu et al., 2010, 2014).

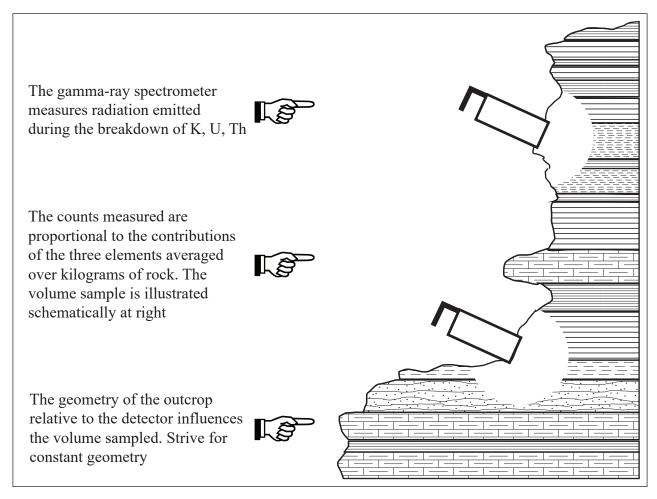


Figure 4. Schematic diagram of using a gamma-ray spectrometer in the field (after Schwalbach and Bohacs, 1992, 1995).

Introduction to Sequence Stratigraphy

The most perfect technique is that which is not noticed at all.

—Pablo Casals (in Kahn, 1970, p. 224)

The current concepts of sequence stratigraphy emerged from investigations of the relations among sedimentation, unconformities, and changes in base level (e.g., Grabau, 1913; Barrell, 1917; Sloss et al., 1949; Wheeler and Murray, 1957; Wheeler, 1958, 1959, 1964; Sloss, 1962, 1963; Curray, 1964; Frazier, 1974) as well as from the development of seismic stratigraphy (e.g., Vail, 1975; Mitchum and Vail, 1977; Vail et al., 1977a, b, c; Vail et al., 1991). Concepts of sequence stratigraphy were put into practice by using the large-scale perspective of seismic cross sections and analysis of outcrop and well-log data within a chronostratigraphic framework (e.g., Bohacs, 1990, 1993, 1998; Van Wagoner et al., 1990; Posamentier et al., 1992; Schwalbach and Bohacs, 1992; Van Wagoner and Bertram, 1995; Macquaker and Taylor, 1996; Schieber, 1998a; Posamentier and Allen, 1999; Adams et al., 2006; Lazar, 2007; Abreu et al., 2010).

Sequence stratigraphy was first applied to continental margin settings and has since expanded to address deep-water, fluvial, aeolian, mire, and lacustrine systems. The sequence-stratigraphic approach was explicitly extended to fine-grained sedimentary rocks beginning with the work on condensed sections (summarized in Loutit et al., 1988) and with fully integrated studies of source- and reservoir-prone mudstones from slope and basinal settings (Bohacs, 1990; Bohacs and Schwalbach, 1992). Subsequent work extended from the Proterozoic to the Holocene and shoreward through epicontinental seaways to lake systems (e.g., Miskell-Gerhardt, 1989; Schieber, 1989, 1990, 1998a, b, 1999; Wignall and Maynard, 1993; Macquaker and Taylor, 1996; Witzke et al., 1996; Bohacs and Suter, 1997; Bohacs, 1998; Macquaker et al., 1998; Bohacs et al., 2000, 2001, 2003, 2005, 2007; Carroll and Bohacs, 2001; Macquaker and Jones, 2002; Brett et al., 2004; Lazar, 2007; Macquaker et al., 2007; Lazar et al., 2015a, b). Each new environment sharpened our focus on the essentials and reinforced the validity of the basic approach. Although depositional processes and basin-filling mechanisms differ significantly, the basic approach of identifying key surfaces and recognizing repeated stratal geometries and stacking patterns works in all depositional settings. This approach is particularly essential in understanding the occurrence, character, and distribution of mudstones, and therefore, it is quite essential to locate and extract natural resources effectively and efficiently.

Key Concepts of Sequence Stratigraphy

Sequence stratigraphy is the study of rocks within a framework in which the vertical succession of rocks is subdivided into genetically related 3-D units that have characteristic stacking patterns and physical, biogenic, and chemical properties, and are bounded by surfaces, including unconformities and their correlative conformities (e.g., Vail, 1975; Mitchum, 1977; Mitchum and Vail, 1977; Vail et al., 1977a, b, c; Posamentier et al., 1988; Van Wagoner et al., 1988; Vail et al., 1991; Bohacs and Schwalbach, 1992; Bohacs, 1998; Abreu et al., 2010). Sequence stratigraphy has five major components (Bohacs and Schwalbach, 1992; Bohacs, 1998; Abreu et al., 2010):

- **Method** that recognizes a hierarchy of various types of stratal surfaces, rock packages, and stacking patterns.
- Observations of physical, biological, and chemical aspects of surfaces and rocks within a 3-D framework.
- Models that summarize and generalize detailed observations.
- **Mechanisms** that seek to explain the origin of stratal patterns and facies distributions in terms of small-to-large-scale processes and to provide predictive capabilities.
- Prediction and Testing that uses further observations to test predictive capabilities, refine models, and enhance the understanding of key mechanisms.

Sequence stratigraphy allows the construction of a comprehensive stratigraphic framework based on a single criterion—the physical relations of the strata themselves—that reveals genetically related rocks. Additionally, a comprehensive sequence-stratigraphic framework is ideal for integrating all types of data, from reflection seismic, through well-log, core, outcrop, thin section, to organic and inorganic geochemistry. This integrated response allows recognition,

correlation, and prediction of sequence-stratigraphic units and surfaces in all these datasets across all scales.

The main building blocks of the stratigraphic record are depositional sequences that contain parasequences composed of beds and bedsets. All of these stratal building blocks, although discontinuous at some scale, comprise relatively conformable successions of smaller scale units bounded by surfaces of erosion, nondeposition, or correlative conformity.

Sequence stratigraphy is an informal method that combines various chronostratigraphic approaches that have evolved over the past 400 years. It organizes and systematically applies stratigraphic insights from Steno's (1667, in Maar, 1910) principles of superposition, lateral continuity, and original horizontality, through Smith's (1815) and Lyell's (1830, 1835) rock systems, Walther's (1894) law of facies succession, Chamberlain's (1898, 1909) and Sloss's (1962, 1963) unconformity-bounded cratonic sequences, and Van Siclen's (1958) depositional topography, to the Exxon group's lamina, laminaset, bed, and bedset, and pattern correlation (Campbell, 1967), seismic stratigraphy (Vail et al., 1977a, b, c), parasequences (Van Wagoner et al., 1990; Van Wagoner, 1995; Bohacs et al., 2014), to today's fully integrated sequence stratigraphy (e.g., Bohacs et al., 2005, 2007; Lazar, 2007; Guthrie and Bohacs, 2009; see also the case studies in Chapters 9 to 15).

Throughout the following discussion it is essential to remember that the rock record is fundamentally discontinuous, with breaks in sediment accumulation at all scales, from kilometers to submillimeters. Hence, the terms "isochronous" or "diachronous" and "conformable" or "unconformable" are to be understood in relative terms that are a function of the resolution of data being used to make the interpretation. Practically, all surfaces result from a change in the sediment accumulation rate at some level of significance—a supercritical decrease or increase in the rate of sediment supply relative to the accommodation rate that produces a change in system behavior. (Level of significance refers to the spatial (and inferred temporal) distribution of the surface and the rocks it bounds—how much area it covers and how much time it records.) This change in system behavior results in erosion, nondeposition, or different rock characteristics (texture, bedding, or composition). Our observations indicate that most surfaces record a pause in sediment accumulation.

Stratal Units and Surfaces

This section presents an overview of all scales and components of the stratal hierarchy used in sequence stratigraphy. They are all elaborated in the following chapters.

Stratal Units—Sequence-stratigraphic stratal units are defined using geometric criteria, with the supporting evidence of other physical, biogenic, and chemical attributes. Although thickness, areal extent, and time for formation are neither essential attributes nor part of the definition of sequence-stratigraphic units, these units do tend to have characteristic spatial and temporal scales as well as common modes of formation. Note that characteristic thicknesses tend to be a function of grain size and are typically thinner in mudstones. Characteristic timescales tend to be strongly related to depositional setting and basin size, with relatively short intervals in small lacustrine basins and relatively long intervals in large marine basins (according to the response time of the basin, which scales to the second power of its characteristic length scale; see Paola et al., 1992).

The **depositional sequence** is the fundamental unit of sequence stratigraphy; it is a relatively conformable succession of strata bounded at base and top by laterally extensive (regional scale) unconformities and their correlative conformities (Figure 3; Mitchum, 1977; Neal and Abreu, 2009; Abreu et al., 2010). Depositional sequences are meters to hundreds of meters thick and extend over many thousands of square kilometers. They are inferred to represent multiple episodes of shoreline progradation with significant shifts in coastal onlap and base level over tens to thousands of millennia. A complete depositional sequence can be subdivided into systems tracts defined by their position within the sequence and by the stacking patterns of the parasequence sets within each systems tract. Parasequence sets are bounded by parasequence set boundaries that are flooding surfaces and their equivalents. Systems tracts include lowstand, transgressive, and highstand (see Table 11).

A parasequence, the main building block of the depositional sequence, is a relatively conformable succession of beds or bedsets bounded below and above by parasequence boundaries (surfaces that record a pause in sediment accumulation, formed by nondeposition, local erosion, or very slow sedimentation and include

flooding, abandonment, or reactivation surfaces and their correlative surfaces; after Van Wagoner et al., 1988, 1990; Bohacs, 1998; Bohacs et al., 2014). Parasequences range from tens of centimeters to tens of meters in thickness and extend over significant parts of a basin, on the order of hundreds to thousands of square kilometers. In shelf or lacustrine settings, they typically represent one episode of shoreline or mudbelt progradation—the dominant depositional "motif" or building block. Equivalent-scale units in other settings include fan "lobes" in submarine-fan settings and channel-belt sets in fluvial or submarine-slope settings. Parasequences are interpreted to form in centuries to millennia.

A **bed** is a relatively conformable succession of genetically related laminae or laminasets bounded at base and top by bedding surfaces—surfaces of erosion, nondeposition, or correlative conformity (Campbell, 1967).

A lamina is the smallest megascopic layer in a sedimentary succession without internal layers (typically ≥0.1 mm or so; Campbell, 1967). See Lazar et al. (2022a, b, Chapters 2 and 4 this Memoir) for detailed discussions of laminae, beds, and parasequences.

Stratal Surfaces—The two distinct types of widespread and mappable surfaces are parasequence boundaries and sequence boundaries (Figure 3; Vail, 1975; Mitchum and Vail, 1977; Vail et al., 1977a, 1991; Posamentier et al., 1988; Van Wagoner et al., 1988; Bohacs and Schwalbach, 1992; Bohacs, 1998; Bohacs et al., 2004). Identification of these surfaces in a stratal succession relies on both their local character and their lateral extent (Van Wagoner et al., 1990; Bohacs and Schwalbach, 1992; Bohacs, 1998; Lazar, 2007; Bohacs and Lazar, 2008; Abreu et al., 2010; Bohacs et al., 2014; Lazar et al., 2015a, b).

A parasequence boundary (flooding surface and correlative surfaces) records a supercritical increase in accommodation relative to sediment supply that significantly changes system behavior (Bohacs, 1998;

Table 11. Definitions of Systems Tracts With Stacking Patterns and Recognition Criteria (after Abreu et al., 2014).

Systems Tract	Observed Stacking Pattern	Bounding Surfaces	Accommodation/Sediment Supply Trend
Highstand Systems Tract (HST)	Aggradation to Progradation to (possible) Degradation [A-P-(D)]	Above: SB Below: MFS (MTS)	Decreasing at an increasing rate
Transgressive Systems Tract (TST)	Retrogradation (R)	Above: MFS (MTS) Below: TS (MRS)	Rapidly increasing to a maximum
Lowstand Systems Tract (LST)	Progradation to Aggradation (P-A)	Above: TS (MRS) Below: SB	Increasing at an increasing rate

MFS = maximum flooding surface; MRS = maximum regressive surface; MTS = maximum transgressive surface; SB = sequence boundary; TS = transgressive surface.

Bohacs et al., 2004). Commonly, strata above a parasequence boundary are deposited in deeper water, and less energetic and more distal environments, whereas strata below a flooding surface are deposited in shallower water, and more energetic and proximal environments (Bohacs, 1998; Bohacs et al., 2004, 2014; Lazar et al., 2015a, b). Parasequence boundaries are marked by a sharp decrease in coarse sediment supply, increased and laterally extensive accumulation of pelagic and authigenic components (e.g., organic matter, remains of plankton and nekton, volcanic ash, dropstones; cements, nodules, and concretions), early lithification or cementation, and increased continuity of laminae, beds, and bedsets (Bohacs, 1990, 1998; Bohacs and Schwalbach, 1992; Macquaker and Taylor, 1996; Bohacs et al., 2004, 2014). See Bohacs et al. (2022a, Chapter 5 this Memoir) for a full discussion.

Within each depositional sequence, two specific parasequence boundaries are interpreted as lower order surfaces, one as the transgressive surface (TS) and another as the maximum flooding surface (MFS), based on their position and geometric relations (Figure 3; Bohacs et al., 2022c, Chapter 6 this Memoir).

The transgressive surface is the parasequence boundary atop the most basinward position of the shoreline of the progradational-aggradational (PA) or lowstand systems tract. It defines the top of the lowstand systems tract and separates progradationally to aggradationally (stepping basinward) stacked parasequences below from retrogradationally (stepping landward) stacked parasequences above (after Bohacs and Schwalbach, 1992; Bohacs, 1998). It is also known as the maximum regressive surface (MRS; Abreu et al., 2014). The surface is interpreted beneath the first landward shift (backstep) of the shelf-slope break; in vertical successions, at the turnaround in parasequence stacking pattern from progradation or aggradation to retrogradation.

A maximum flooding surface is the one particular parasequence boundary representing the maximum landward extent of basinal facies within a sequence. It defines the top of the transgressive systems tract and separates retrogradationally (stepping landward) stacked parasequences below from aggradationally to progradationally (stepping basinward) stacked parasequences above (after Bohacs and Schwalbach, 1992; Bohacs, 1998). It is also known as the maximum transgressive surface (MTS; Abreu et al., 2014). The presence of prograding strata above identifies the maximum flooding surface as a downlap surface on reflection seismic profiles. It represents the greatest landward extent of the sea or lake within a depositional sequence (e.g., Loutit et al., 1988; Posamentier and Vail, 1988; Bohacs and Schwalbach, 1992).

Sequence boundaries are the laterally extensive (regional scale) unconformities and correlative

conformities that bound a depositional sequence (Mitchum, 1977); they are fundamentally different from flooding surfaces. Sequence boundaries (in contrast to flooding surfaces) record a supercritical decrease in accommodation relative to sediment supply, commonly accompanied by an increase in depositional energy or a significant change in sediment supply (e.g., erosional bypass in marine environments), over hundreds to thousands of square kilometers (e.g., Bohacs, 1998; Bohacs et al., 2004; Lazar, 2007; Bohacs and Lazar, 2008, 2010). They are easiest to recognize in medial reaches of the shelf. Common attributes of sequence boundaries are summarized in Table 7 and discussed further in Bohacs et al. (2022c, Chapter 6 this Memoir). Sequence boundaries are surfaces across which there is a basinward shift in coastal onlap, marked by laterally extensive erosional truncation of underlying strata (with evidence of exposure and presence of reworked clastics in lag deposits) and toplap below and onlap and downlap above (e.g., Mitchum, 1977; Schwalbach and Bohacs, 1992; Bohacs, 1998; Schieber, 1998a; Bohacs et al., 2004; Lazar, 2007, Bohacs and Lazar, 2008). It occurs below the abrupt basinward shift in shoreline position at the base of a depositional sequence. It is placed at the surface beneath the first increase in accommodation above progradationally or degradationally stacked parasequences, at the break in shoreline sandstone trajectory (Figure 3).

Constructing and Testing a Sequence-Stratigraphic Framework for Mudstones

As discussed in Bohacs and Lazar (2022, Chapter 1 this Memoir), method, observations, models, mechanisms, and prediction and testing are the five major components of sequence stratigraphy (Bohacs and Schwalbach, 1992; Bohacs, 1998; Abreu et al., 2010). An optimal workflow for establishing a sequence-stratigraphic framework follows this fivefold organization and includes the following (Bohacs and Schwalbach, 1992; Bohacs, 1998):

- Establishing geological context and geophysical background of the region under investigation
- Identifying large-scale stratal units by locating terminations and interpreting surfaces
- Tying surfaces and stratal units around a grid
- Making maps of terminations, stratal units, and their internal geometry
- Interpreting depositional environments and estimating facies character from those maps and sections
- Summarizing and generalizing observations and first-order interpretations into a model for the 3-D distribution of surfaces, strata, and rock properties
- Inferring the mechanisms that led to the formation of the surfaces and strata, nesting small-scale

Table 12. Construction of a Sequence-stratigraphic Framework.

- I. Establish general geological context and history of the area of interest.
 - A. **Reconstruct plate-tectonic setting**, establish the location of continental and oceanic crust, transition zones, and major tectonic and structural elements.
 - B. Determine the location, size, shape, and evolution of the "container"—basins and sub-basins (using seismic, well logs, outcrop, core, biofacies data).
 - C. Establish general locations of depocenters, sediment provenance areas, and general depositional environments.
- II. Examine lithologic and stratal characteristics of the rocks in vertical section (outcrops, cores, well logs—see Chapter 1).
 - A. **Physical attributes:** bed thickness, bed-surface and lamina geometry, primary and secondary sedimentary structures, gamma-ray spectra.
 - B. **Biogenic attributes:** abundance and diversity of paleofauna from macro, micro, nanno, and trace fossils; abundance and diversity of paleoflora from macrofossils, phytoliths, palynomorphs, and kerogen.
 - C. Chemical attributes: major and accessory lithotypes; nodules; major, minor, and trace elements; organic carbon; kerogen type; isotopic signatures of inorganic and organic components; molecular geochemistry.
- III. Group the lithotypes and stratal characteristics at the bed and bedset scale into facies packages based on recurring associations of attributes.
- IV. Relate the stacking patterns of facies packages to the depositional environment.
 - A. Facies stack in distinctive manners and proportions according to their depositional environment. Stacking patterns can be recognized in outcrops or cores from bedding-surface arrangements, laminae geometries, and lithofacies and biofacies changes, and in the subsurface from various well logs. The interpretation of a depositional environment for any one map point is strengthened by a three-dimensional perspective gained from mapping paleotopography, regional paleogeography, and isopach thickness of the stratigraphic interval on a palinspastic base map, as well as from estimates of open-ocean conditions (surface currents, undercurrents, upwelling zones, etc.). The interpretation of a depositional environment, therefore, includes input at all scales of observation—from cores, outcrops, well logs, and seismic data to plate-tectonic setting.
 - B. Calibrate well-log response of each lithofacies association/depositional environment. Beware of assuming what the well-log signature of a particular lithotype or environment will be: volcaniclastic sandstone can have a high gamma-ray response (a "shaley" signature); fine-grained cherts and porcelanites have very low gamma-ray activity and somewhat "blocky" log patterns that mimic a siliciclastic braided-stream signature (among others...). Calibrate the well logs locally.
- V. Use the stacking patterns and lateral relations of rock packages from the different depositional environments and their bounding surfaces to interpret the sequence stratigraphy. It is generally best to start with sections that represent medial positions in the depositional systems: These are the locations that have the largest contrasts among lithofacies and depositional environments.
 - A. Divide the stratigraphic section into large-scale packages bounded by significant surfaces: key discontinuities and major floodings (Figure 5). These packages are an initial approximation of the systems tracts. In proximal environments, the major flooding surfaces (maximum flooding surface; transgressive surface) are most easily recognized; in moderately distal environments, the sequence boundary (unconformity) is commonly fairly obvious.
 - B. Group bedsets of depositional environments into parasequences. Group parasequences into parasequence sets according to their stacking patterns (progradational, aggradational, retrogradational), location relative to preexisting shelf break, and relations to bounding surfaces. Note the relation of the parasequence sets to the significant surfaces recognized in step V.A, and refine your estimation of systems tracts.
 - C. Group parasequence sets into depositional sequences and assign tentative sequence stratigraphic surfaces and systems tracts to each vertical section or location.
 - D. Correlate vertical sections in a three-dimensional grid, carrying the major flooding and discontinuity surfaces first, then filling in with smaller scale stratal packages; use seismic and biostratigraphic data as well as the regional geology. This three-dimensional correlation is essential as, of course, all sorts of random events occur during deposition—storms, floods, earthquakes, slope failures—but the observations that went into the development of the sequence-stratigraphic approach indicate that there is a nonrandom, repeated pattern in how the record of these events are stacked and packaged. At any one vertical section, rocks are passive recorders of local conditions (bottom energy, pH, Eh, etc.)—hence, the geologist must correlate over a reasonably large area and look at the three-dimensional array of surfaces and rock packages to distinguish the significant time-stratigraphic units from random local noise.
 - E. As the correlation progresses around the grid ("tying loops"), make maps of particular time slices on plate-tectonic reconstruction bases. This insures that the regional geological information is factored into the interpretations. Ask: Do the patterns of depositional environments on the map make sense, in terms of both general principles and local geological history? Map patterns that are obviously unrealistic point out miscorrelations and can indicate that significant surfaces have been misidentified or not identified at all.
 - F. **Iterate to a final result by fine-tuning interpretations of vertical sections, cross sections, and maps** until a consistent, geologically reasonable picture emerges that accounts for the full range of data (physical, biogenic, chemical).

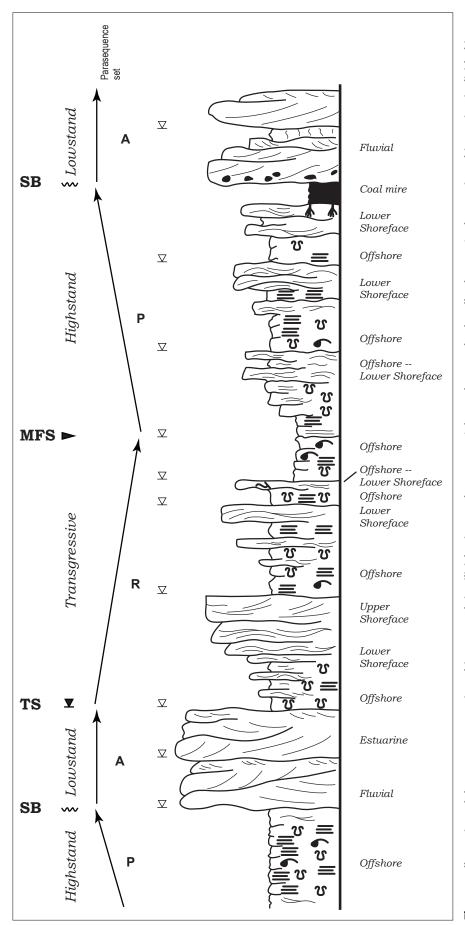


Figure 5. Illustration of the sequence-stratigraphic approach in siliciclastic nearshore strata (after Bohacs and Schwalbach, 1992). The stratigraphic section is divided into large-scale stratal packages bounded by significant surfaces that are identified by their characteristics and stratigraphic context (place in stacking patterns, etc.). See Table 12 for more details. MFS = maximum flooding surface; SB = sequence boundary; TS = transgressive surface.

processes of grain formation, erosion, transport, and deposition within larger scale processes of climate, tectonics, and biological evolution and their forcing functions

Details of this workflow are given in Table 12 and elaborated throughout the following chapters.

Now, we can proceed to apply these tools and techniques at successively larger scales, starting with the smallest scales, laminae, laminasets, beds, and bedsets, in Lazar, et al. (2022b, Chapter 4 this Memoir).

CONCLUSIONS

Mudstone vertical and lateral variability at millimeter to kilometer scales can be detected by applying a range of physical, petrophysical, chemical, and paleontological methods to characterize and interpret the rocks within an integrated sequence-stratigraphic approach.

This chapter introduced key tools and techniques that provide data about texture, bedding, composition, and grain origin. Such data enable characterization of mudstone strata at lamina to sequence-set scales. The application of such tools and techniques to decipher depositional conditions and construct sequence-stratigraphic frameworks was specifically addressed. Outlines of our approach to making detailed and systematic observations of key attributes of mudstones in outcrops, cores, and thin sections as well as an introduction to key sequence-stratigraphic concepts that we find useful for studying mudstones were included for quick reference. Our approach is elaborated and illustrated in all of the following chapters.

ACKNOWLEDGMENTS

We thank ExxonMobil for support and permission to release this information. We acknowledge Quinn Passey, Matt Boyce, and Russell Spears for their feedback on Table 5; John Snedden for his contribution on the seismic interpretation section; and Fred Schroeder for his input on the development of sequence stratigraphy. We thank Jon Kaufman for his timely feedback on the entire chapter. We are most grateful to our reviewers, Dr. Jennifer Eoff and Prof. Paul Potter.

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