A practical guide to Rock Microstructure

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Preface

Learning about rocks can give much pleasure to anyone interested in Earth and its development. I hope that readers of this book will share my enthusiasm for examining rocks with the microscope. I planned the book to be an introductory review of the main processes responsible for the microstructures of Earth rocks. However, I soon realized that if I did that, the book would be a collection of half-truths, with little scientific value. Though many rock microstructures are understood fairly well, the interpretation of many others involves considerable controversy, and new ideas are being published all the time. So, I have felt compelled to mention problems of interpretation and to present alternative views, where appropriate. Thus, the book has evolved into (1) a basic explanation of the main processes, (2) an introduction to more complex issues of interpretation and especially to the relevant literature, and (3) an outline of modern approaches and techniques, in order to emphasize the ongoing, dynamic nature of the study of rock microstructure. Because complicated problems cannot be discussed in detail in a book of this kind, I have tried to provide a sufficient number of references to enable the reader to delve more deeply.

I assume that the reader has a basic knowledge of geology, rock types and microscopic mineral identification. Thus, the book is aimed mainly at senior geoscience undergraduates and above. Emphasis is placed on higher-temperature processes, i.e. those that occur under igneous and metamorphic conditions, although the book begins with a brief discussion of sedimentary microstructures, as background for some of the metamorphic microstructures. The mineral abbreviations used follow those suggested by Kretz (1983), as extended by Bucher & Frey (1994), and are listed at the start of the book. There is an extensive glossary of microstructural terms at the end of the book.

I also hope that materials scientists may also gain some benefit and interest from the microstructures discussed and illustrated, because rocks are the ‘materials’ of Planet Earth, in the sense of ‘materials science’: the branch of science that links all solid materials, such as metals, ceramics, glass, organic polymers and, of course, rocks.

I took all the photographs, except where otherwise acknowledged. I am also responsible for most of the line drawings, with the assistance of Dean Oliver.
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(Figs. 5.11, 5.37) and Daleth Foster (Fig. 5.93). I thank David Durney, Dick Flood, Scott Johnson and Scott Paterson for critically reading parts of the typescript, Judy Davis for assistance with computer techniques, Geoff Clarke for access to specimens at the University of Sydney, Ross Both, John Fitz Gerald and Neil Mancktelow for providing images, and John Lusk, John Ridley, David Durney and Pat Conaghan for providing specimens of opaque minerals, deformed rocks/veins and sedimentary rocks at Macquarie University. People who kindly provided other samples or thin sections are acknowledged in the figure captions.
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Chapter 1
Background

1.1 Introduction

I wrote this book to help you to interpret what you see when you look at thin and polished sections of rocks with the microscope. I say ‘help’, rather than ‘teach’, because I don’t want to give the impression that every microstructure you see can be easily and unambiguously interpreted in terms of processes that produced the rock. Many can, but in many other instances, conventional interpretations are ambiguous or poorly understood. So I intend the book to be only a guide, and I present alternative ideas where appropriate. A healthy scepticism should be maintained when interpreting rock microstructures yourself and also when reading the interpretations of others.

1.2 History of the examination of rocks with the microscope

Rocks in natural outcrops, in samples knocked off these outcrops and in drill cores, are beautiful and instructive. We can see different minerals, and identify many of them with the aid of a hand lens. We can also see some of the more obvious structures in the rocks. However, cutting a slice (section) though a rock with a diamond-impregnated circular saw and polishing the sawn surface shows us the various minerals alongside each other, rather than piled confusingly all around each other. This reveals the structure even more clearly, as can be seen in the polished facing slabs on many buildings and bench tops.

But we always want to see more. So, when D. Brewster in 1817 and William Nicol about 1830 showed how to make a slice of crystalline material thin enough to transmit light (0.03 mm is the standard thickness), stuck to a glass microscope slide (Shand, 1950, p. 6; Loewinson-Lessing, 1954), it wasn’t surprising that a curious person, such as Henry Sorby, should start looking at these thin sections of rocks (Sorby, 1851, 1853, 1856, 1858, 1870, 1877, 1879, 1880, 1908). Sorby learnt the technique of making thin sections from W. C. Williamson in 1848 (Judd, 1908; Folk, 1965) and made the first rock thin section in 1849 (Judd, 1908). He was the first to look seriously at rock sections with the microscope, beginning with a study of chert, a siliceous sedimentary rock that was a very
appropriate choice for microscopic investigation, in view of its very fine grain-size. He described and suggested a mechanical origin for slaty cleavage (Sorby, 1853, 1856), noticed many of the basic features of igneous and metamorphic rocks, made many important observations on sedimentary rocks, including carbonate rocks (Sorby, 1851, 1879), investigated pressure-solution (using fossil crinoids), described meteorites, and published the first papers on the examination of polished sections of metals with the microscope (Sorby, 1864, 1887). So not only is he the founder of petrography (the description of rocks), but the founder of metallography as well (Smith, 1960). He also investigated fluid inclusions in minerals, heating crystals to watch the gas bubbles disappear, in order to get an estimate of the temperature of crystallization of the mineral (Sorby, 1858; Folk, 1965).

Sorby was followed soon after by many others, as discussed by Johannsen (1939) and Loewinson-Lessing (1954). Among them were Dawson (1859), Zirkel (1863, 1866, 1876), Vogelsang (1867), Fouqué & Michel-Lévy (1879), Allport (1874), Rosenbusch (1873, 1877) and Teall (1885, 1886). Since those days, the light microscope has become the main tool for identifying minerals and examining their microstructures, although it has been augmented by many modern techniques (Section 1.6).

1.3 How relevant is the microscope today?

Many petrologists concentrate on the mineralogical and chemical aspects of rocks, without spending much time looking at rocks with the microscope. In fact, in these days of marvellous techniques for the chemical and isotopic analysis of minerals, some people may feel that simply looking at and measuring the shapes and arrangements of crystals in rocks with the microscope is a little out of date. However, carrying out detailed chemical and isotopic analyses of minerals when you don’t understand the relationships of these minerals to other minerals in the rock makes little sense. It’s a waste of expensive resources, at very least.

On the other hand, many structural geologists look at the physical or structural aspects of minerals and rocks, especially from the viewpoint of deformation processes and preferred orientations of grains, without being concerned about the chemical aspects of these processes. Both approaches are valuable, of course, but their interrelationships can be particularly illuminating. Fortunately, many researchers are attempting to integrate the chemical and physical approaches, and the study of rocks with the microscope provides a link between them. In fact, the detailed study of processes in rocks at the microscopic scale is now a major area of research, especially among younger people, in many universities and other research institutions. Moreover, new observational techniques are being developed and used, as discussed in Section 1.6.

Research microscopes commonly have both transmitted and reflected light facilities. An excellent example of the simultaneous use of transmitted and
reflected light microscopy is the study of Columbia River basalts by Long & Wood (1986), in which reflected and transmitted light photographs are arranged side by side, clearly revealing the dendritic shapes of the opaque Fe–Ti oxide minerals and their relationships to the transparent and translucent silicate minerals. Some leading books and review articles on minerals in reflected light, with emphasis on microstructures, are those of Bastin (1950), Edwards (1947, 1952), Cameron (1961), Ramdohr (1969), Stanton (1972), Craig & Vaughan (1994) and Craig (1990a,b).

1.4 Mineral identification

Learning to identify minerals takes time and practice, and is outside the scope of this book. Close teaching in a laboratory situation is the best way to learn about the optical properties of minerals, using textbooks specifically written for the purpose (e.g. Fleischer et al., 1984; Shelley, 1985a; Nesse, 1991; Gribble & Hall, 1992; Deer et al., 1992). Ideally, this should go hand-in-hand with learning about microstructures.

1.5 The concept of a section

Thin and polished sections are two-dimensional sections through three-dimensional objects, and this must always be kept in mind, as explained in some detail by Hibbard (1995). Mineral grains can have unexpectedly complex three-dimensional shapes (see, for example, Rigsby, 1968; Byron et al., 1994, 1995, 1996). Two or even three orthogonal sections may be necessary to reveal the structure of structurally anisotropic rocks, and some recent detailed microstructural studies have used: (1) serial sectioning (see, for example, Byron et al., 1994, 1995, 1996; Johnson & Moore, 1996), coupled with image analysis by computer, to construct a three-dimensional image of the microstructure, and (2) computed X-ray tomography (Section 1.6), to reveal the three-dimensional distribution of large crystals (porphyroblasts) in metamorphic rocks (Carlson & Denison, 1992; Denison & Carlson, 1997), plagioclase chains in basalts (Philpotts et al., 1999), and former melted rock (leucosome) in migmatites (Brown et al., 2002).

1.6 Newer techniques

This book deals mainly with microstructures visible in the optical (light) microscope, in standard thin or polished sections, with the use of polarized light. However, some newer techniques are very useful for revealing features not apparent or less clearly shown in polarized light, as outlined below. Several examples of photographs taken with these techniques will be presented in the book. The new techniques underline the fact that the study of rock microstructure is a dynamic, progressive field of research.
1 Background

(1) **Cathodoluminescence** (CL) is a technique that can reveal internal microstructures of grains of some minerals, for example, compositional zoning, overgrowths, microcracking and replacement veining in quartz, calcite, dolomite, magnesite, zircon, plagioclase, K-feldspar, diamond, fluorite, sphalerite, kyanite, pyrope garnet, corundum, cassiterite, anhydrite and apatite (Sippel & Glover, 1965; Smith & Stenstrom, 1965; Zinkerngels, 1978; Matter & Ramseyer, 1985; Owen & Carozzi, 1986; Sprunt, 1978, 1981; Sprunt & Nur, 1979; Field, 1979; Reeder & Proskey, 1986; Padovani et al., 1982; Hanchar & Miller, 1993; Marshall, 1988; Ramseyer et al., 1988; Morrison & Valley, 1988; Mora & Valley, 1991; Yardley & Lloyd, 1989; Hopson & Ramseyer, 1990; Barker & Kopp, 1991; Shimamoto et al., 1991; Williams et al., 1996; Mora & Ramseyer, 1992; D’Lemos et al., 1997; Watt et al., 1997, 2002; Hayward, 1998; Müller et al., 2000; Pagel et al., 2000; Janousek et al., 2000; Ahn & Cho, 2000; Götze, 2000; Rubatto & Gebauer, 2000; Penniston-Dorland, 2001; Rubatto et al., 2001; Hermann et al., 2001; Barbarand & Pagel, 2001; Peppard et al., 2001; Rusk & Reed, 2002; Viljoen, 2002; Rougvie & Sorensen, 2002). CL is especially useful for revealing microstructural details in minerals that are colourless in the light microscope, for example calcite, quartz and feldspar. For example, quartz, untwinned K-feldspar and untwinned plagioclase may be distinguished by their CL colours in fine-grained aggregates, and CL can assist in provenance and diagenetic studies in sedimentary rocks. The technique is not suitable for iron-rich minerals. Some applications of CL are discussed in Sections 3.11.7, 3.11.9 and 5.9.3.

It can be used with the light microscope (producing true CL colours with a spatial resolution of 1–2 μm), or the scanning electron microscope (producing grey-scale variations with a spatial resolution of less than 1 μm), and may be combined with CL spectroscopy to enable spectral spot analysis of revealed features. CL is combined with X-ray tomography to reveal the internal structure of diamonds (see, for example, Field, 1979).

CL is caused by defect structures in the crystal lattice, such as impurity atoms (transition metals, rare earth elements, lead, titanium, actinides), vacancies and dislocations produced during formation and/or deformation of the mineral, which therefore reflect conditions of crystallization, deformation and alteration. The technique involves coating a polished thin section with carbon and bombarding it with electrons in a vacuum. This bombardment produces light from substitutional atoms in an excited state.

(2) **Laser-interference microscopy** is a relatively new optical technique that detects small differences in refractive index, and so can reveal, in great detail, subtle compositional differences (on which refractive index depends), for example in zoned plagioclase (Chao, 1976; Pearce, 1984a,b; Pearce et al., 1987a,b).

(3) **Scanning electron microscopy** (see, for example, Lloyd, 1987) is capable of revealing sharp microstructural details in shades of grey, although arbitrary colours may also be assigned to form a false-colour image. It involves
backscattered and forescattered imaging in the scanning electron microscope (SEM). This is particularly useful for (a) revealing the detailed microstructure of small grains and fine-grained aggregates and intergrowths (see, for example, Vernon & Pooley, 1981; Wirth & Voll, 1987; Cashman, 1988; Simpson & Wintsch, 1989; Swanson et al., 1989; Johnson & Carlson, 1990; van der Voo et al., 1993; Brodie, 1995; Lloyd & Prior, 1999; Drüppel et al., 2001; Blundy & Cashman, 2001; Rickers et al., 2001; de Haas et al., 2002; Schieber, 2002), (b) identifying very fine-grained minerals (see, for example, Prior et al., 1999), (c) revealing fine-scale compositional zoning in minerals (see, for example, Yardley et al., 1991; Müller et al., 2000; Piccoli et al., 2000; Kuritani, 2001; Alexandrov, 2001; Rubatto et al., 2001; Hermann et al., 2001; Ginibre et al., 2002a,b; Lentz, 2002), (d) measuring orientation differences between grains and subgrains as small as 1 μm across (Prior et al., 1996, 1999; Lloyd et al., 1997; Trimby et al., 1998; Wheeler et al., 2001; Bestmann & Prior, 2003), and (e) revealing domains of different orientation in optically isotropic minerals, such as garnet (Spiess et al., 2001; Prior et al., 2000, 2002) and pyrite (Boyle et al., 1998).

(4) **Transmission electron microscopy** (TEM) assists in the interpretation of some microstructures that can be optically ambiguous, such as some recovery features in deformed quartz (Section 5.4) and fine exsolution lamellae (Section 4.11). The principles and some applications have been reviewed by Champness (1977), Putnis & McConnell (1980), McLaren (1991) and Putnis (1992). TEM resolves very small objects, such as very fine to submicroscopic intergrowths, exsolution features, inclusions and twins and can reveal the arrangement of defects (including dislocations, discussed in Section 5.3.2) in the atomic structure of individual grains of both optically transparent and opaque minerals (see, for example, McLaren et al., 1967; McLaren, 1974, 1991; McLaren & Retchford, 1969; Phakey et al., 1972; Green & Radcliffe, 1972; McLaren & Hobbs, 1972; Champness & Lorimer, 1976; Champness, 1977; McLaren & Etheridge, 1976; Zeuch & Green, 1984; Doukhan et al., 1985; Allen et al., 1987; Cox, 1987a; Couderc & Hennig-Michaeli, 1989; Hennig-Michaeli & Couderc, 1989; Green, 1992; Ando et al., 1993; Doukhan et al., 1994; Vogelé et al., 1998).

(5) **X-ray tomography** (in full: high-resolution computed X-ray tomography) is a more recent development in the study of rock microstructure (see, for example, Mees et al., 2003). This technique maps the variation of X-ray attenuation within solid objects, the attenuation varying with each mineral present. A source of X-rays and a set of detectors revolves around the rock sample, producing images in layers or cross-sections. The series of two-dimensional images can be computed into a three-dimensional representation of the grains and aggregates in the rock (see below), which gives a clearer picture of spatial relationships and crystal size distributions (see, for example, Carlson & Denison, 1992; Carlson et al., 1995, 1999; Denison et al., 1997; Brown et al., 1999; Philpotts et al., 1999).
6 Background

Computer-aided construction of three-dimensional images is a technique whereby serial two-dimensional optical or X-ray tomographic images can be scanned and imported into suitable computer graphics programs to provide three-dimensional constructions (Johnson & Moore, 1993, 1996; Carlson et al., 1995, 1999; Pugliese & Petford, 2001; Castro et al., 2003). This approach is used in confocal laser scanning microscopy (CLSM), which enables objects (such as inclusions in minerals) to be viewed in focus through a thickness of sample in a single image (Petford & Miller, 1992; Petford et al., 1995; Sheppard & Shotton, 1997; Bozhilov et al., 2003), as described in Section 4.4.5. Readily available computer software can also be used to animate images, producing a more complete visualization of features such as grain shapes, grain distributions and vein networks (see, for example, Johnson & Moore, 1996; Carlson et al., 1999; Pugliese & Petford 2001).

X-ray compositional mapping produces maps of compositional zoning in crystals (see Sections 3.11, 4.12). Such maps are produced by multiple stage-scan chemical analyses made with wavelength-dispersive spectrometers on an electron microprobe, different colours being assigned to different concentrations of the analysed element. Examples are shown in Section 4.12. The technique can also be used for more clearly revealing mineral or compositional domains in fine-grained aggregates (see, for example, Lang & Gilotti, 2001; Williams et al., 2001; Clarke et al., 2001; Daczko et al., 2002a, b). Raw X-ray intensity maps may be converted to maps of oxide mass percent by appropriate matrix corrections (Clarke et al., 2001).

1.7 Quantitative approaches

Although most work on rock microstructures is qualitative, involving description and interpretation, quantitative methods are also used. For example, grain measurement is important in the classification and interpretation of clastic sedimentary rocks in terms of transport and depositional environments (Section 2.2.3). Grain size is also used in the classification of igneous rocks, though less precisely, and crystal size distributions (CSD) are being increasingly investigated in igneous and metamorphic rocks (Sections 3.4, 4.3.1). Various statistical techniques are used to determine spatial distribution patterns (SDP) of grains and crystals in rocks (see, for example, Flinn, 1969; Kretz, 1966b, 1969; Jerram et al., 1996; Denison & Carlson, 1997; Daniel & Spear, 1999; Jerram & Cheadle, 2000). Numerical modelling has been used to convert two-dimensional measurements of grain shapes and sizes in thin section to three-dimensional grain shapes and true crystal size distributions (Higgins, 1994, 2000; Peterson, 1996). Moreover, computer software is readily available to do this and to make animated images, as mentioned in the previous section. Interfacial angles have been measured in many metamorphic rocks, sulphide rocks and igneous cumulates, as indicators of mutual solid-state growth of minerals (Section 4.2). In addition, the orientations
of inclusion trails in porphyroblasts have been used as indicators of tectonic processes (Section 5.10). Numerical simulation of the development of metamorphic and deformation microstructures is also well under way (Jessell, 1988a,b; Jessell et al., 2001).

1.8 Some terms

Although no hard and fast rule exists, it is probably best to use crystal for a volume of crystalline mineral with well-formed, planar faces (called crystal faces or facets), and grain for any other volume of crystalline mineral. For me, the shapes, arrangements and orientation of the minerals constitute a rock’s fabric. At the microscope scale, the fabric (microfabric) consists of the grain shapes and arrangement (the microstructure) and the spatial orientation of the minerals (the preferred orientation). However, many people equate ‘fabric’ with ‘preferred orientation’, as recommended by the IUGS Subcommission on the Systematics of Metamorphic Rocks (Brodie et al., 2002).

It would be good to make materials scientists more interested in rocks, as they are the great class of natural solid materials. Therefore, because ‘texture’ means ‘preferred orientation’ to most materials scientists and an increasing number of structural geologists, it would be best not to use it instead of ‘microstructure’ as many petrologists do. However, although ‘microstructure’ is gaining in popular usage, ‘texture’ is common, and no ambiguity is caused among petrologists by using it. Actually, ‘microstructure’ appears to have priority, because the first publications on the microscopic examination of rocks referred to ‘microscopical structure’ or ‘microscopic structure’ (see, for example, Sorby, 1851, 1858; Dawson, 1859; Allport, 1874). Moreover, the IUGS Subcommission on the Systematics of Metamorphic Rocks has recommended that the term ‘texture’ be replaced by ‘microstructure’, which is defined as ‘structure on the thin section or smaller scale’ (Brodie et al., 2002). The term ‘microtexture’, which unfortunately is starting to enter the literature, is a tautology, because ‘texture’ refers to the microscopic scale.

Of course, every gradation in scale exists between the microscopic and macroscopic (outcrop) scales, and so I have not been able to confine the discussion to the microscopic, although this is by far the main scale discussed.

1.9 Traditional rock groupings

Many rock-forming processes apply to more than one of the traditional igneous, sedimentary and metamorphic rock groups. For instance, similar basic principles governing the nucleation and growth of crystals apply to all rocks, and grain growth in the solid state occurs not only in metamorphic rocks (in which it is a universal process), but also in the late stages of formation of some rocks conventionally regarded as igneous. In addition, growth of new minerals in the solid state
(neocrystallization) occurs not only in metamorphic rocks, but also in the late-stage alteration (deuteric alteration) of igneous rocks, and in the low-temperature alteration (diagenesis or burial metamorphism) of rocks that many people would consider to be still sedimentary. Moreover, metamorphic rocks begin to melt at high temperatures, producing rocks with both igneous and metamorphic features. In addition, radiating crystal aggregates (spherulites) commonly grow in glass, which, though technically solid, is liquid-like with regard to its atomic structure. Furthermore, exsolution, which is a solid-state process, occurs in both igneous and metamorphic minerals. As if that isn’t enough, fragmental material thrown out of explosive volcanoes produces rocks that are technically sedimentary, but consist entirely of igneous material, and may also show evidence of solid-state flow of glass. The result of this cross-linking of processes is that, though this book adheres roughly to the traditional sedimentary–igneous–metamorphic subdivision, processes discussed under one of these headings may also be relevant to another of these groups. These instances are cross-referenced.

1.10 Importance of evidence

Science relies on evidence. An assertion made without evidence isn’t worth very much. Yet I often read statements such as: ‘the microstructural (textural) evidence indicates . . . ’ This implies that the writers are asserting that their interpretations are so obviously right that they don’t have to go to the bother of describing what they saw and evaluating the evidence.

Of course, recognizing evidence takes practice. As noted by Chalmers (1999):

It is necessary to learn how to see expertly through a telescope or microscope, and the unstructured array of bright and dark patches that the beginner observes is different from the detailed specimen or scene that the skilled viewer can discern.

Whenever you make interpretations based on microscopic examination of rocks, you should: (1) describe clearly what you see and (2) evaluate the possible interpretations. If one or more interpretations are valid, you should not arbitrarily favour one of them, unless other evidence (e.g. field or chemical evidence) clearly points in that direction. This is the ‘method of multiple working hypotheses’ advocated by Chamberlain (1890). In many instances, the microstructural evidence may not be at all clear, in which case you shouldn’t use it to support a hypothesis. Maybe you will have to suggest equally valid alternative interpretations and leave it at that.

The paramount importance of evidence in making scientific inferences is emphasized in the following quotation.

On so important a question, the evidence must be airtight. The more we want it to be true, the more careful we have to be. No witness’s say-so is good enough.
People make mistakes. People play practical jokes. People stretch the truth for money or attention or fame. People occasionally misunderstand what they’re seeing. People even sometimes see things that aren’t there.

Carl Sagan (1996) was referring to UFOs, but at least some of these statements could refer to petrologists interpreting rock microstructures. People do make mistakes and even see things that aren’t there, and though practical jokes may be uncommon in such a serious pursuit as petrology(!), people certainly do occasionally misunderstand what they’re seeing. We all do, in fact. Most important, we often want something to be true so much that we may be tempted to gloss over the evidence, whereas we should be doubly careful, in order to save ourselves falling into the trap of a woefully wrong interpretation, no matter how attractive it may seem.

It doesn’t matter how many times an assertion is repeated or how loudly it is trumpeted in conversation, in the scientific literature, in textbooks or even on the Internet, it is only as good as the evidence for it. Another point to remember is that an interpretation presented by a great authority on the subject, though worthy of respect perhaps, is also only as good as the evidence for it. Such ‘arguments by authority’ can subdue interpretations based on careful accumulation of evidence (Vernon, 1996b).

Too often we see examples of interpretations based on inadequate evidence used to support a preferred model. Even some well-accepted interpretations may be wrong. A good example is the common belief that an ‘order of crystallization’ in igneous rocks can be inferred by looking at the microstructure. Generally this is impossible, as explained in Section 3.6. If the microstructure cannot give you the evidence, please don’t try to extract it anyway!

In fact, the more I examine and read about rock microstructures, the more cautious I become about interpreting them, and that will be a constant theme in this book. As mentioned below (Section 1.11), recent work on the direct microscopic observation of developing microstructures in organic compounds used as mineral analogues has revealed many unexpected processes, and has shown that similar microstructures may have very different histories. They remind us of the necessity for caution in the interpretation of natural rocks.

So I will try to give explanations that are sufficiently general to be regarded by most people as ‘reasonable’ on the available information, and that students can infer largely from the optical microstructure. Where alternatives need to be discussed, the relevant publications will be mentioned. Moreover, where pitfalls exist, they will be pointed out, and it must be re-emphasized that the book is only the most general of guides. It indicates what to look for and how to start (not stop) thinking about what is observed. Finally, it is as well to keep in mind the wise words of John Spong (Sydney Morning Herald, June 9, 2001): ‘Explanations are always compromised by the levels of knowledge available to the explainer.’ That applies to me just as much as to any other explainer!
1.11 Kinds of evidence used

What evidence is useful in interpreting rock structures? Imagine you had never seen a rock section, either a thin section or a slab cut through a hand sample. How could you begin to interpret the crystal shapes and arrangements you see? You must have some guides. These are field relationships and experimental evidence on rocks and minerals, assisted by some general inferences from experiments on other materials, such as metals, ceramics, organic polymers and synthetic ice. For example, when Sorby first looked at thin sections of slates with the microscope, he would have already known that slates are formed by strong deformation and that the deformation is in some way responsible for their characteristic strong foliation (slaty cleavage). Furthermore, once Sorby had observed and described the features shown by the microstructure of the slate, others were in a position to recognize similar cleavages in thin sections of rocks from other areas. In this way, general guides to the interpretation of rock microstructures have been established.

If we can observe rocks forming, as with sedimentary and volcanic rocks, we are on firm ground for making inferences about how the microstructures were formed. We are on much shakier ground when it comes to intrusive igneous and metamorphic rocks. However, we can learn much from careful interpretation of field relationships, although strong differences of interpretation are common. In addition, experiments on the cooling of melted rocks and the melting of solid rocks are valuable guides to the interpretation of rocks involving melts, and many recent experimental advances have been made in the interpretation of igneous microstructures (see, for example, Lofgren, 1971b, 1973, 1974, 1976, 1980; Lofgren et al., 1974; Fenn, 1974, 1977, 1986; Donaldson, 1976, 1977, 1979; Swanson, 1977; Swanson & Fenn, 1986; London, 1992). However, we should keep in mind possible problems caused by the short duration of experiments.

Experimentally determined stability fields of mineral assemblages in different bulk chemical compositions reveal the conditions of pressure, temperature and fluid composition that occur during metamorphism. However, it is not as easy to conduct successful experiments on the development of microstructures in metamorphic rocks, because of the high temperatures and pressures involved in the experiments and the generally small size of the samples used.

Many important experiments on mineral and rock deformation have been carried out (Chapter 5), but again we can only observe the finished product, not the stages along the way. Fortunately, experiments on ice deformation have helped our understanding of progressive microstructure development during deformation (see, for example, Wilson, 1984, 1986; Wilson et al., 1986). Moreover, a new experimental technique based on transparent and translucent organic compounds that behave somewhat similarly to minerals has been developed, and is being applied with great effect to the interpretation of microstructures, especially deformation features (see, for example, Means, 1977, 1981, 1983, 1989; Means & Jessell, 1986; Means & Park, 1994; Means & Ree, 1988; Park & Means, 1996;
Because these compounds deform, melt and crystallize rapidly at room temperatures, the processes can be observed and photographed in progress in the microscope (‘see-through’ experiments). Of course, we must keep in mind that these materials are generally not minerals, but they have provided some startling insights into possible grain-scale processes that may occur in natural rocks.

In the absence of reliable experimental evidence, it is necessary to fall back on ‘common-sense’ interpretations, based on accumulated experience of the type outlined previously. This applies especially to metamorphic rocks. Unfortunately, common sense isn’t so common, and what makes perfectly good sense to one person may make no sense at all to somebody else. The most important thing is to be as honest and logical as possible, and to evaluate (and if necessary retain as possibilities) every interpretation that can reasonably explain the observations. If the end result is the unsatisfying conclusion that you cannot make an unambiguous interpretation on the available evidence, leave it at that. No harm will be done. On the contrary, many a doubtful interpretation, presented as being reliable, has been accepted at face value and used in later work, thereby misleading subsequent researchers.

1.12 Complexity

A rock’s microstructure is the product of a complicated sequence of events and processes. So is a rock’s chemical analysis. Both may tell us something about the rock’s history, but neither can fully reveal all the historical complexities. This is a problem that petrologists have to accept. We do our best with the evidence available, without taking it too far, and we must acknowledge that our interpretations are often incomplete.

Another point to add to the complexity is that superficially similar microstructures may be formed in different ways, as with exsolution and epitactic (epitaxial) replacement producing similar intergrowths (see, for example, Craig, 1990). For example, hematite lamellae in magnetite, usually inferred to be of replacement origin, may be due to exsolution in some rocks (Edwards, 1949). Another complication is the optical similarity between subgrains formed by recovery and similar features formed by fracture, as discussed in Section 5.4. Other complexities of rock microstructure will become apparent in the following chapters.
2.1 Introduction

Although this book is mainly concerned with igneous, metamorphic and deformation processes, in this chapter I briefly review the main sedimentary microstructures, partly because they need to be understood in order to interpret residual sedimentary microstructures in some metamorphic rocks.

The basic sedimentary microstructures are relatively straightforward, but variations (reflecting variable sedimentary environments) can be very complex, and many complicated classification schemes to deal with this complexity have been suggested. Classification schemes, microstructural details and discussions of sedimentary environments can be found in many excellent books (e.g. Pettijohn, 1949; Williams et al., 1954; Carozzi, 1960; Milner, 1962; Folk, 1968; Selley, 1970; Blatt et al., 1972; Bathurst, 1975; Friedman & Sanders, 1978; Adams et al., 1984, Greensmith, 1989; Boggs, 1992; McPhie et al., 1993; McLane, 1995).

Sediments are loose, unconsolidated fragments, and sedimentary rocks are the consolidated or lithified equivalents. Residual (pedogenic) sediments are essentially in situ deposits of the products of rock weathering, for example weathering crusts, soils and regoliths. Epiclastic (terrigenous) sediments are accumulations of solid fragments formed by erosion of existing rocks. Pyroclastic sediments are deposits of fragmented igneous material (e.g. volcanic glass and crystal fragments) ejected from volcanoes in explosive eruptions directly onto Earth’s surface. Bioclastic sediments are accumulations of organic skeleton or shell fossils that have been at least slightly transported. Chemical sediments are precipitated directly from aqueous solution or by replacement of existing sediment. Polygenetic sediments consist of mixtures of the foregoing types, and are named according to the dominant kind of sediment present (e.g. fossiliferous limestone, tuffaceous sandstone). Moreover, epiclastic rocks commonly contain chemical components in the form of a cement (Section 2.2.5).
2.2 Epiclastic (‘terrigenous’) sedimentary rocks

2.2.1 Detrital (clastic, fragmental) minerals

Quartz and the clay minerals constitute up to about 70–80% of the epiclastic sedimentary rocks, with less abundant feldspar, mica and carbonate. The clay minerals belong to a group of very fine-grained, water-rich, complex aluminosilicates with various other elements, especially potassium, magnesium and iron. Generally they are too fine-grained to be identified with the optical microscope, and so other techniques (especially X-ray diffraction) are used for their accurate determination. Other chemically and physically resistant minerals, such as zircon, tourmaline, ilmenite, magnetite, monazite, rutile, topaz and garnet, are also commonly present in very small quantities: generally no more than 1%, except in local ‘black sand’ concentrations.

The detrital (clastic, fragmental) minerals in epiclastic sediments depend on (1) their presence in the source rocks, (2) their resistance to mechanical abrasion, and (3) their chemical stability in the surface environment. Common fragmental minerals are those that are most stable in Earth’s atmosphere, especially quartz and the clay minerals. Quartz is the most common mineral that is stable in Earth’s atmosphere, and is released from rocks undergoing weathering as individual grains or fragments. In contrast, clay minerals are formed by the chemical breakdown of minerals that are not as stable in the atmosphere, especially feldspars and the ferromagnesian minerals (pyroxene, olivine, biotite and amphibole). However, these less stable minerals can occur as fragments in sediment that has been transported relatively short distances and/or deposited rapidly.

2.2.2 Fragment size and sorting

Transport at normal conditions in water tends to sort the fragments (detritus) into different sizes, namely: pebbles (> 2 mm in diameter), sand (2 – \(\frac{1}{16}\) mm in diameter), silt (\(\frac{1}{16} – \frac{1}{256}\) mm in diameter) and clay (< \(\frac{1}{256}\) mm in diameter). Prolonged washing of sediment, for example in waves in relatively shallow water (involving traction currents), leads to a well-sorted sedimentary rock with a relatively even grainsize (Figs. 2.1, 2.2), whereas if the sediment is deposited rapidly (for example, in turbidity currents produced by submarine slumps on continental slopes), the fragments tend to have very different sizes (said to be poorly sorted or unsorted), as shown in Figs. 2.3–2.7.

Generally, well-sorted sedimentary rocks tend to have rounded fragments (Section 2.2.4) and *vice versa* (Figs. 2.1, 2.5), but not necessarily (Figs. 2.2–2.4). Sedimentary rocks composed mainly of pebbles, sand and clay are called conglomerates or rudites (Figs. 2.7, 2.8), sandstones or arenites (Figs. 2.1–2.6) and shales, claystones, mudstones or pelites (Fig. 2.9), respectively.