Interrelationship between Mg/Fe Ratio and Octahedral Al Content in Biotite

CHARLES V. GUIDOTTI, JOHN T. CHENEY,
Department of Geology and Geophysics, University of Wisconsin,
Madison, Wisconsin 53706

AND PAUL D. CONATORE
Department of Geology, Colorado School of Mines,
Golden, Colorado 80401

Abstract

The trend of a plot by Foster (1960) of Mg vs (Fe$^{2+}$ + Mn) vs R$^{3+}$ (Al, Ti, Fe$^{3+}$) in the octahedral sheet of biotite has led some workers to infer an inverse correlation between Al$^{VI}$ and Mg/Fe$^{2+}$ ratio. This paper presents arguments and data which indicate that such a correlation does not exist over most of the biotite composition range and that the trend mentioned above is in part an accident of sample availability.

Introduction

Foster (1960) made a very useful and comprehensive survey of the compositional range of the trioctahedral micas that are broadly designated as biotites. Included were compositions encompassed by the end members phlogopite, K(Mg$_8$Al$_2$O$_{10}$)(OH)$_2$; eastonite, K(Mg$_{8.5}$Al$_{0.5}$)(Al$_1.5$Si$_2$O$_{10}$)(OH)$_2$; annite, KFe$_7$(Al$_{1.5}$Si$_2$O$_{10}$)(OH)$_2$; and siderophyllite, K(Fe$_{2.5}$Al$_{0.5}$)(Al$_1.5$Si$_2$O$_{10}$)(OH)$_2$. In this note “biotite” refers to micas plotting within the field bounded by these four end members.

Based upon a compilation of a large number of analyses from many different parageneses, Foster (1960, Fig. 11; Fig. 1, this paper) made an interesting triangular plot of Mg vs (Fe$^{2+}$ + Mn) vs R$^{3+}$ (includes Al, Ti$^{IV}$, Fe$^{3+}$) in the octahedral sheet of biotite. The points lie in a distinct band (Fig. 1) in which Al$^{VI}$ appears to decrease with increase of Mg/(Fe$^{2+}$ + Mn). However, in discussing this band, Foster (1960, p. 25) states, “The trioctahedral micas may, therefore, be thought of as members of a system characterized by concurrent but independent replacement of Mg by Fe$^{2+}$ and R$^{3+}$ ions.” Nonetheless, because of the shape and orientation of the band shown in Figure 1, several workers have inferred that changes in the Mg vs (Fe$^{2+}$ + Mn) content of the octahedral sheet control the amount of Al$^{VI}$. For example, such a relationship has been tacitly implied by Hazen and Wones (1972, p. 124) when they state, “As the Fe$^{3+}$ content of these micas increases, so does the octahedral aluminum content.”

Gower (1957) constructed a diagram quite similar to that of Foster (1960) using data from specimens of diverse origin. He also inferred (p. 155) “A greater tolerance for octahedral aluminum is indicated in iron-rich varieties.”

The goal of this paper is to demonstrate that, with the possible exception of biotites very near the annite-siderophyllite join, no clear relationship exists between the Mg/(Fe$^{2+}$ + Mn) ratio and octahedral Al content. Hence, any inferences of a direct relationship between these two factors should be reconsidered. Moreover, it will be shown that the specific, detailed pattern shown on Figure 1 (or on Gower’s Fig. 5) is in part an accident.

Our approach is largely empirical, inasmuch as little attention is given to the substitutional models applicable to the octahedral and tetrahedral sheets of trioctahedral micas. We have followed the common practice of assigning Mg, Fe$^{2+}$, Ti, and Al in excess of that needed to fill tetrahedral sites to the octahedral sheet.

Discussion of the Band of Points on Figure 1

Several factors appear to explain the orientation of the band of points shown on Figure 1.

The tapering of the biotite field toward the
Fig. 1. Plot of octahedral $R^+3(=Al + Fe^{3+} + Ti)$ vs $Mg$ vs $(Fe^{2+} + Mn)$ in more than 200 natural biotites from various parageneses. Diagram based on Foster (1960, Fig. 11).

Fig. 2. Plot of octahedral $Al$ vs $Mg$ vs $(Fe^{3+} + Mn)$ in biotites from northwestern Maine. Biotites A, B, and C from sillimanite-bearing adamellite; biotites D, E, F, and G from two-mica adamellite (see Table 1E). All other biotites from pelitic schists with sillimanite, staurolite, or cordierite (see Tables 1-4). Best fit line is visually located.

Fig. 3. Plot of octahedral $(Al + Ti)$ vs $Mg$ vs $(Fe^{2+} + Mn)$ in the same biotites as on Figure 2. Best fit line is visually located.
phlogopite end-member appears to be due in part to the plotting of Mg-rich biotites from Al-poor rocks such as marbles, calc-silicate granulites, and ultramafics. The vast majority of analyzed, Mg-rich biotites reported in the literature are from such rocks. In these rocks, biotite usually does not coexist with aluminous minerals. Hence, it tends to plot toward the phlogopite end of the phlogopite-estonelite series. Moreover, Figure 1 and Gower's diagram show considerable scatter in $R^3+/ (Mg + Mn + Fe^2+).$ This scatter results from plotting biotite analyses from specimens from a multitude of different parageneses and bulk compositions. It will become evident below that this has also contributed to obscuring the true relationship between $Al^VVI$ content and the $Mg/(Fe^{2+} + Mn)$ ratio.

Figure 1 includes $Fe^{3+}$ and $Ti^{4+}$ as part of the $R^3+$ term. As a result, $Fe$ biotites, siderophyllites, and epidotolaminates plot noticeably nearer the $R^4+$ corner (cf Figs. 2 and 3). Moreover, analyses discussed below show that Mg-rich biotites are in fact low in Ti. Hence, the trend shown by the band of points in Figure 1 is in part an accident due to the availability of samples, and in part a function of what has been included with $R^4+.$ The trend resulting from the latter aspect is of course real, but it should not be interpreted as implying a marked interrelationship between $Al^VVI$ content and the $Mg/(Fe^{2+} + Mn)$ ratio.1

1 It must be reemphasized that in biotites near the pure Fe end members such a relationship may exist. This is evident from the discussion of Hazen and Wones (1972). Our data do not bear directly on this composition range.
New Data from Pelitic Schists and Adamellites of Northwestern Maine

Numerous biotites from the pelitic schists of northwestern Maine have been analyzed during the course of petrologic studies by the authors (Guidotti, 1974; Guidotti, Cheney, and Conatore, 1975; and previously unpublished data). These data are presented in Tables 1-5 and were obtained by electron microprobe analysis using the general technique of Bence and Albee (1968). Simple anhydrous silicates and oxides were used as standards. The analytical error is about two percent for elements constituting more than two percent of a given sample.

A rather wide range of biotite Mg/Fe ratios has been found because specimens from pyrrhotite-rich rocks as well as more ordinary pelitic schists have been analyzed. Some of the biotites from the former group of specimens plot well within the phlogopite field of Foster (Fig. 1). All of the biotites from pelitic schists under consideration coexist with minerals such as sillimanite, staurolite, or cordierite. They would plot along the high-Al boundary of the biotite field on an ARM projection, and are clearly Al-saturated for the prevailing P,T conditions. As a result, there is little variation of AlIV or AlII due to differences in bulk composition. Moreover, because virtually all of the specimens from northwestern Maine have graphite present, one can assume that

Table 3. Analyses of Biotites from Sulfide-Rich Small's Falls Formation, Southwest Rumford Quadrangle, Maine. Upper Sillimanite to K-Feldspar + Sillimanite Zone*

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<th>Ru-P14-73</th>
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* Data from Guidotti et al (in preparation). Values cited for H₂O by difference from 100 percent. All specimens contain sillimanite.

Table 4. Analyses of Biotite from the Lower Sillimanite Zone of the Southeast Portion of the Old Speck Mountain Quadrangle, Maine

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Table 5. Analyses of Biotites from Two-Mica Adamellites of Northwestern Maine

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* Data from Cheney (in preparation). Values cited for H₂O by difference from 100 percent. All specimens contain sillimanite.
Fe$^{3+}$ is probably present in low and uniform amounts. The above considerations will now enable us to make a concise evaluation of any relationship between Al$^{VI}$ and the Mg/(Fe$^{2+}$ + Mn) ratio.

Figure 2 shows the relative amounts of (Fe$^{2+}$ + Mn), Mg, and Al in the octahedral sheets of biotites from pelitic schists of northwestern Maine. Also shown are analyses of biotites from several igneous parageneses. With respect to the biotites from pelitic schists, it is evident that no marked relationship exists between Al$^{VI}$ and the Mg/(Fe$^{2+}$ + Mn) ratio. Biotites A, B, and C come from sillimanite-bearing adamellites which intrude the pelitic schists of northwestern Maine. Hence, they can be legitimately compared with the biotites discussed above and thereby extend the range of Mg/Fe ratios to still lower values (to 0.466).

Biotites D, E, F, and G are from adamellites which contain no sillimanite. The absence of sillimanite suggests a lower activity for Al$_2$O$_3$ in these specimens. Of note is the fact that on Figure 2 these biotites plot at lower Al values than do biotites A, B, and C.

Figure 3 is similar to Figure 2, but includes Ti$^{4+}$ in with R$^{3+}$. Comparison of the trends on the two diagrams shows that in Figure 3 there is a distinct suggestion of a decrease of R$^{3+}$ as Mg increases. Moreover, our data do show that Ti$^{4+}$ decreases as Mg increases in biotite (see Tables 1-5) and that this inverse trend must be controlled by crystallochemical factors, because all specimens coexist with a Ti-saturating phase$^2$ (rutile in the pyrrhotite-rich specimens and ilmenite in the others).

It is thus clear that at least part of the trend shown on Figure 1 can be explained by inclusion of Ti$^{4+}$ with R$^{3+}$. The remainder of the trend can be attributed to inclusion of Fe$^{3+}$ in R$^{3+}$ and to the availability of samples as exemplified by previous analyses of low-Al phlogopites only from marbles, calc-silicates, and ultramafic rocks. Moreover, our use of biotites from only a few similar parageneses has greatly reduced the amount of scatter in terms of the R$^{3+}/$(Mg + Fe$^{2+}$ + Mn) ratio on both Figure 2 and Figure 3 relative to that seen on Figure 1.

**Conclusions**

1. The present data demonstrate no clear relation between Mg/(Fe$^{2+}$ + Mn) and Al$^{VI}$ content of biotite over most of the biotite composition range. However, our data do not preclude such a relationship in biotites approaching the annite-siderophyllite join. For these compositions, structural arguments suggest that a pure Fe octahedral sheet may not be stable (Hazen and Wones, 1972).

2. Our data demonstrate the dangers of trying to infer crystallochemical relationships in rock-forming solid solutions using data based on specimens from many different parageneses.

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**References**


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