

It can readily be seen that this geothermometer is relatively insensitive for Fe/Fe + Mg ratios greater than about 0.7, because the garnet produced is very close to pure almandine (Fig. 2). For ratios lower than this it can be computed that an error in $\ln K$ of 0.2, is equivalent to a temperature error of 50°. For a typical K of 6, such an error would correspond to an error of roughly 1.1 in determination of K . If this error in turn is assigned entirely to analytical error, it would correspond roughly to an error of 10% in the Fe/Mg + Fe ratio of both garnet and cordierite. Determination of this accuracy does not appear to present any problems in the present state of the art, either by electron probe methods, or by concentration of minerals. It follows that for favourable compositions the method should give results accurate within 50°, and probably considerably better.

Eq. 12(a) has been applied to the data in Table 1. These data indicate that the temperature reached about 600° in the southeastern part of the area, and almost 700° in the northwestern part of the area, at pressures of 5.1 and 5.8 kilobars respectively.

We have also applied the geothermometer to the data of Reinhardt (1968), and Wynne-Edwards and Hay (1963) (Table 1). For the simplest mineral assemblage garnet-cordierite-sillimanite quartz (Reinhardt's specimen D-102) the computed conditions are 675°C and 6.2 kilobars. For his other specimens containing biotite, excepting specimen D-56, the garnet-cordierite pair, indicates a temperature of $675 \pm 10^\circ$, and pressure of 6.1 ± 0.3 kbars, in agreement with the conditions indicated by specimen D-102. This agreement strongly suggests that the geothermometer is applicable to nearly all cordierite-garnet-sillimanite-quartz gneiss. It is possible that D-56 illustrates the partial breakdown of one of the assumptions, but it is also possible that an analytical error is involved. The other specimens which show significant variations in Fe/Mg ratios in biotite, demonstrate the independence of the thermometer from other Fe and Mg bearing phases.

Throughout this analysis the pressure of H_2O has been ignored. In Fig. 3 we have combined data for the garnet-cordierite equilibrium with data for the aluminosilicate triple-point (Gilbert *et al.*, 1968), the biotite breakdown curve (Eugster and Wones, 1964), and the muscovite breakdown curve (Evans, 1965). The following conclusions, although based on the Gilbert *et al* triple point, are also applicable to most of the other suggested triple points discussed by Zen (1969). The occurrence of andalusite in equilibrium with sillimanite, as observed in the southeastern part of the region, and the total absence of muscovite cannot be explained if $P_{H_2O} = P_{total}$. However if P_{H_2O} is much less than total pressure, the field data may be reconciled with experimental data. The data of Reinhardt (1968) require very low water pressures, approaching 0.2 total pressure. According to Gilbert *et al.* (1968), for the deduced P - T conditions these rocks would contain sillimanite as the stable polymorph. Under these conditions, garnet and cordierite can coexist with biotite, in the absence of muscovite, and in the presence of either andalusite and sillimanite. These latter polymorphs can both be present in equilibrium with compositions having Fe/Fe + Mg greater than 0.8. In order for the observed coexistence of cordierite with Fe/Fe + Mg less than 0.3 with garnet (Fe/Fe + Mg < 0.7) to take place, as it does north of Loughborough Lake, both temperature and total pressure must increase to roughly 700° and 6.5 kilobars

