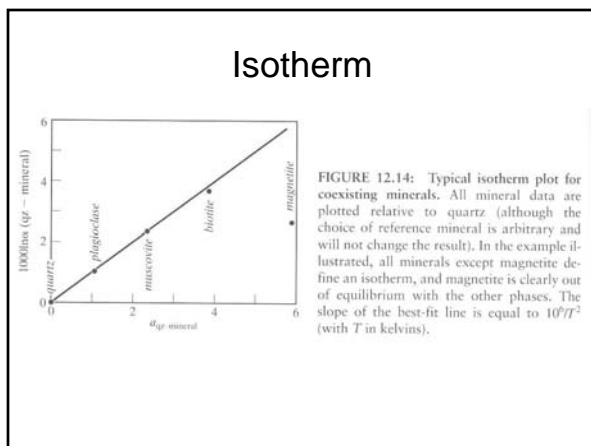


TABLE 12.P1: Some commonly used coefficients of oxygen isotope fractionation relative to quartz. In form of equation 12.11 (after Javoy, 1977).

Mineral	a	b
Plagioclase	$0.97 + 1.04\beta$	0
Olivine	2.75	0
Muscovite	2.2	-0.6
Biotite	3.69	-0.6
Amphibole	3.15	-0.3
Ilmenite	5.29	0
Magnetite	5.57	0
Garnet	2.88	0
Chlorite	5.44	-1.63

β is the fraction of anorthite in plagioclase.



Oxygen isotope ratios in coexisting minerals of regionally metamorphosed rocks

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Increasing
"isotopic
stability" ↑ Garnet
Quartz
Ilmenite, muscovite, biotite

Geospeedometry

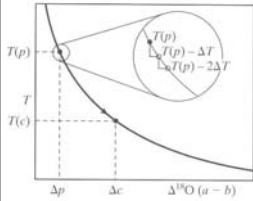


FIGURE 12.16: Schematic plot of $\Delta^{18}\text{O}$ value for phases a and b vs. temperature. The initial, peak temperature is $T(p)$. If the sample is cooled rapidly to room temperature, it will preserve fractionations Δp , corresponding to equilibrium at $T(p)$. If the sample is cooled slowly, it will continue to reequilibrate during cooling, following the equilibrium fractionation-vs.-temperature path (e.g., $T(p)$ to $T(p) - \Delta T$, etc.). The sample will continue to reequilibrate until the closure temperature $T(c)$ is reached, at which point intracrystalline diffusion effectively ceases.

*Diffusional exchange in this case is limited by the intracrystalline diffusion rate, or self-diffusion rate. This is the diffusion rate of oxygen within a crystal. The intercrystalline, or grain-boundary, diffusion rate, which allows oxygen to migrate between phases, is considered to be orders of magnitude more rapid than the intracrystalline diffusion rate. Limited experimental studies support this assumption (e.g., Farver and Yund, 1991a).
