

## **Coupled dichotomies of apatite and fluid composition during contact metamorphism of siliceous carbonate rocks**

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### **ABSTRACT**

The  $X_{\text{CO}_2}$  recorded by mineral-fluid equilibria in contact metamorphosed siliceous carbonates commonly defines two groups of rocks in the same aureole. One group records relatively low  $X_{\text{CO}_2}$  that results from infiltration of chemically reactive  $\text{H}_2\text{O}$ -rich fluid. The other records relatively high  $X_{\text{CO}_2}$ , up to 0.99, that results from decarbonation reactions with little or no infiltration. A complementary dichotomy in apatite compositions exists in five contact aureoles in Italy, Scotland, and the U.S.A. Apatite in the low- $X_{\text{CO}_2}$  group is close to an F-OH solid solution. Apatite in the high- $X_{\text{CO}_2}$  group is a relatively Cl-rich Cl-F-OH solution. The halogen content of fluid coexisting with analyzed apatite was characterized in two aureoles to determine the origin and significance of the dichotomy in apatite composition. Calculated  $a_{\text{HF}}/a_{\text{H}_2\text{O}}$ ,  $a_{\text{HF}}/a_{\text{HCl}}$ ,  $a_{\text{HF}}$ , and  $m_{\text{F}_T}$  (the total F molality of fluid) are systematically higher in fluid coexisting with the low- $X_{\text{CO}_2}$  group. In contrast,  $a_{\text{HCl}}/a_{\text{H}_2\text{O}}$  in the high- $X_{\text{CO}_2}$  group may be higher than or overlap with  $a_{\text{HCl}}/a_{\text{H}_2\text{O}}$  in the low- $X_{\text{CO}_2}$  group. Calculated  $a_{\text{HCl}}$  and  $m_{\text{Cl}_T}$  in the high- $X_{\text{CO}_2}$  group are lower than or overlap with  $a_{\text{HCl}}$  and  $m_{\text{Cl}_T}$  in the low- $X_{\text{CO}_2}$  group. The Cl-rich apatites in the high- $X_{\text{CO}_2}$  group are explained by crystallization at relatively low  $a_{\text{H}_2\text{O}}$ ,  $a_{\text{HF}}$ , and  $m_{\text{F}_T}$  rather than at high  $a_{\text{HCl}}$  or  $m_{\text{Cl}_T}$ . In comparison, the F-OH apatites in the low- $X_{\text{CO}_2}$  group formed by infiltration of rock by and equilibration with relatively  $\text{H}_2\text{O}$ -rich, high  $m_{\text{F}_T}/m_{\text{Cl}_T}$  fluid, reflecting the same metasomatic process responsible for F-rich humite-group minerals and skarns in many contact aureoles. Calculated halogen contents indicate that the non- $\text{CO}_2$  fraction of fluid in equilibrium with both groups had modest, seawater-like salinity, and that the reactive  $\text{H}_2\text{O}$ - and F-rich fluid that infiltrated the low- $X_{\text{CO}_2}$  group had a plutonic source.