

Table 3
Electron microprobe analyses of amphiboles from selected wet runs on the high-alumina quartz tholeiite composition.

Conditions of run	10 kb	10 kb	9 kb	9 kb	10 kb	10 kb	10 kb	10 kb
	920°C	960°C	1040°C	1040°C	960°C	920°C	940°C	920°C
	7½ hr WET	4 hr WET	4 hr WET	4 hr WET	4 hr WET	7½ hr WET	6 hr WET	5½ hr WET
Runs conducted in platinum capsules					Analyses adjusted for iron loss			Runs conducted in graphite capsules
Co-existing phases	cpx *, opx, plag *	cpx *, opx	cpx *, opx *	cpx *, opx *	cpx *, opx	cpx *, opx, plag *	cpx *, opx, mt	cpx, opx, mt
SiO ₂	40.5	40.8	39.8	39.3	40.2	39.9	40.0	40.2
TiO ₂	2.9	2.8	3.9	3.8	2.8	2.9	3.0	3.1
Al ₂ O ₃	15.8	14.4	15.8	15.6	14.2	15.6	14.8	15.4
FeO	10.0	8.2	7.6	9.8	10.6	12.6	9.4	9.8
MgO	13.1	14.9	14.9	13.7	13.6	11.6	12.9	12.1
CaO	12.4	11.7	12.0	11.8	11.5	12.2	11.6	11.8
Na ₂ O	2.5	2.7	2.9	2.9	2.7	2.5	2.0	1.8
K ₂ O	0.4	0.4	0.2	0.2	0.4	0.4	0.3	0.3
	97.6	95.9	97.1	91.7	96.0	97.7	94.0	94.5
$\frac{100 \text{ Mg}}{\text{Mg} + \text{Fe}}$	70.3	76.4	77.8	71.4	69.6	62.1	71.0	68.8
Structural formulae — calculated on 'dry' basis of 23 [0] since H ₂ O content could not be determined								
$z \left\{ \begin{array}{l} \text{Si} \\ \text{Al} \end{array} \right.$	5.932	6.025	5.802	5.790	6.002	5.909	6.288	6.039
	2.068	1.975	2.198	2.210	1.998	2.091	1.712	1.961
$y \left\{ \begin{array}{l} \text{Al} \\ \text{Ti} \\ \text{Fe} \\ \text{Mg} \end{array} \right.$	0.659	0.531	0.516	0.498	0.500	0.631	1.029	0.766
	0.320	0.311	0.427	0.422	0.314	0.323	0.354	0.350
	1.209	1.013	0.927	1.208	1.324	1.561	1.236	1.231
	2.862	3.281	3.239	3.011	3.029	2.561	3.024	2.710
$z \left\{ \begin{array}{l} \text{Ca} \\ \text{Na} \\ \text{K} \end{array} \right.$	1.945	1.851	1.874	1.862	1.840	1.936	1.954	1.899
	0.709	0.861	0.820	0.829	0.781	0.717	0.610	0.524
	0.074	0.075	0.037	0.037	0.075	0.075	0.060	0.058
z	8.00	8.00	8.00	8.00	8.00	8.00	8.00	8.00
y	5.05	5.14	5.11	5.14	5.17	5.08	5.64	5.06
z	2.73	2.79	2.73	2.73	2.70	2.79	2.62	2.48

* Denotes co-existing phase analyzed.

enrichment of iron relative to magnesium would have been decreased if the experiments had not been carried out under such reducing conditions, since under more oxidising conditions a significant amount of iron in the trivalent state may be taken into the amphibole and pyroxene structures. For example amphiboles and pyroxenes from plutonic and effusive members of the calc-alkaline series contain a significant amount of ferric iron ($(100 \text{ Fe}_2\text{O}_3) / (\text{FeO} + \text{Fe}_2\text{O}_3)$) ranges from 15.7–48.6 for amphiboles and from 19.1–32.8 for clinopyroxenes [13-16]).

* Denotes co-existing phase analyzed.

The most significant feature of this experimental investigation at 9–10 kb under hydrous conditions where $P_{\text{H}_2\text{O}} < P_{\text{load}}$, is that the amphiboles and pyroxenes are subsilicic, so that extraction of these phases provides a very efficient means of enriching liquid fractionates in silica. Thus the residuum in equilibrium with a liquid of basaltic andesite composition will consist of approximately equal proportions of pyroxene and amphibole. However, amphibole and subordinate pyroxene will form the residuum in equilibrium with a liquid of andesite composition (the