Using biotite composition of the Devonian Lake George granodiorite, New Brunswick, as a case study for W-Mo-Au-Sb mineralized magmatic hydrothermal systems

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Abstract: The granodioritic Lake George polymetallic deposit is located approximately 35 km southwest of Fredericton, New Brunswick. This intrusion is a metaluminous to weakly peraluminous, calc-alkaline body that shows an evolved I-type, volcanic arc affinity. Based on an Early Devonian age determination, (412 +5/-4 Ma, zircon U-Pb) it is related to the Hakshaw granite phase of the Pokiok Batholith. The Lake George stock is cut by Hibbard stibnite -quartz veins and quartz-scheelite-molybdenite veinlets that contain significant amounts of gold. Fresh biotites from this intrusion were analyzed at both the core and rim by electron microprobe, and along rim to rim transects by laser ablation ICP-MS at the University of New Brunswick to build an understanding of the halogens responsible for mineralization and trace element distribution within this deposit.

Biotites of this intrusion are reddish brown in colour (indicative of reduced I-type source) and mostly altered to chlorite. They usually contain apatite, zircon, titanite, ilmenite, rutile, and sulphide minerals as mineral inclusions. Temperature was calculated with the Ti-In-biotite geothermometer, in which the results showed a variation between 583 and 745°C.

Two depths of emplacement were determined based on the Al in hornblende and biotite geobarometries confirming the porphyritic texture of this intrusion (4.3 and 1.5 km, respectively). Forming amphiboles at that depth clearly indicates a high water content of the source magma; in addition, hydroxyl is the most dominant component of the hydroxyl site (Average 1.89 wt%) in biotites. The limited range of IV(F/Cl) values of the Lake George biotites suggested that they all equilibrated with one fluid.

Even though there is no noticeable difference in major elements from core to rims, evidence of magma evolution is recorded by biotite grains by their trace elements. For instance, Cu, Rb, Cr, K, Mo, Sn, Cs and W increase from core to rims, whereas Ba, Ni, Mn, and Li act inversely. Sb has a negligible variation from core to rim. Interestingly, the partition coefficient (biotite/whole-rock) is significantly small for Sb, W, and Mo (main associated mineralization) with 0.06, 0.28 and 0.13 in pure magmatic biotites and increase to 0.77, 1.93 and 0.20 in more altered biotites reflecting enrichment of these elements towards the late stage fluid.

Based on these observations, the concept of using mica composition to help identify fertile Acadian magma systems was proved; this method may be a useful tool to indicate the difference between barren and mineralized granophileelement rich systems.

Originally presented Fredericton 2014: Geological Association of Canada - Mineralogical Association of Canada Joint Annual Meeting. May 25, 2014.

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Azadbakht, Z., Lentz, D.R., and McFarlane, C.R.M., 2015. Using biotite composition of the Devonian Lake George granodiorite, New Brunswick, as a case study for W-Mo-Au-Sb mineralized magmatic hydrothermal systems; *in* TGI 4 – Intrusion Related Mineralisation Project: New Vectors to Buried Porphyry-Style Mineralisation, (ed.) N. Rogers; Geological Survey of Canada, Open File 7843, p. 459-474.



Project Goals

- Document detailed chemistry of coexisting phyllosilicate mineral phases to reveal distinctive element signatures that may be specific to fertile granitoids.
- Provide a geochemical fingerprint for morespecific mineralization styles and prospectivity in the Acadian Plutonic Complex.

























type; I-SCR, strongly contaminated and reduced I-type; I-WC, weakly contaminated I-type; after Ague and Brimhill (1987).













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		Tr	ace	Ele	mei	nts	
(ppm)	80-33A	1291	1351	1647	1741	1886	
u	257.42	265.23	275.35	321.72	281.47	392.44	
Р	2.93	23.86	14.04	72.06	19.43	2174.51	
Са	292.38	452.92	347.68	266.09	732.63	6141.64	
Ті	24872.94	31702.44	27927.93	27990.00	28495.14	27782.75	
v	385.32	640.43	454.60	523.67	486.95	544.28	Yellow: Highe
Cr	47.51	130.86	60.52	69.05	70.14	63.88	concentration
Mn	6634.01	2263.85	4287.13	4652.31	3940.02	4649.05	
Со	44.53	31.43	58.46	49.38	62.84	45.65	Orange: lowe
Ni	44.84	46.29	61.08	48.67	62.42	40.49	concentration
Cu	3.66	4.99	1.15	0.55	3.81	12.50	
Zn	531.73	391.19	455.21	476.70	449.32	484.71	
Ga	50.72	63.22	46.02	49.68	45.97	50.80	
Rb	802.79	843.97	733.01	763.71	755.35	746.56	
Sr	1.73	5.65	1.59	1.70	2.32	5.69	
Mu	0.97	0.34	0.74	0.35	0.62	0.32	
Sn	63.94	18.77	8.02	10.57	8.21	12.11	
Sb	0.77	2.36	0.27	0.07	0.33	-0.09	
Cs	90.91	58.39	17.94	26.41	20.55	42.73	
Ва	1304.40	1439.19	1885.19	2053.30	1466.99	2468.81	
w	1.07	1.29	0.78	0.76	0.59	0.84	



Trace Elements

	Мо	Sn	Sb	w	
1351	6	<1	4.8	<1	
1741	3	<1	4.3	8	
1647	7	<1	6	6	
1291	6	1	6.5	<1	
1886	2	<1	5.6	5	

Whole rock trace element Yang et al. (2002)

Averages of Mo, Sn, Sb and W in biotites of the Lake George granodiorite (Right)

	Mo	Sn	Sb	W
1351-6	0.64	9.91	0.26	0.64
1351-7	0.75	9.31	0.22	0.67
1351-8-1	0.71	6.89	0.33	0.92
1741-2	0.64	8.66	0.47	0.64
1741-4	0.54	8.05	0.75	0.69
1741-5	0.65	8.01	0.4	0.59
80-33-A-2	1.00	66.18	0.76	1.30
80-33-A-3	1.12	62.56	0.66	1.06
80-33-A-5	0.74	62.82	1.16	0.92
1647-3	0.33	10.82	0.06	0.56
1647-4	0.44	10.62	0.01	0.48
1647-5	0.29	10.14	0.16	1.54
1291-2	0.29	20.78	0.32	0.64
1291-7	0.39	16.37	3.59	2.24
1291-8	0.34	17.18	11.14	2.91
1886-4	0.33	11.70	-0.2	0.77
1886-6	0.29	12.56	0.11	0.95







Conclusion

- Biotite chemistry reflects whole rock geochemistry
- *f*H₂O is higher than *f*HF and *f*HCl
- Disequilibrium partitioning of halogens and high metal concentrations in biotite, good reflection of contamination?
- Internal zoning of trace element good record of physio-chemical changes during biotite crystallization
- Proof of concept that mica chemistry may aid in identification of fertile Acadian magma systems