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Blandine Gourcerol, D J Kontak, J A Petrus, P C Thurston. Application of LA ICP-MS analysis of arsenopyrite to gold metallogeny of the Meguma Terrane, Nova Scotia, Canada. *Gondwana Research*, 2020, 81, pp.265-290. 10.1016/j.gr.2019.11.011 . hal-02442027

HAL Id: hal-02442027

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Application of LA ICP-MS analysis of arsenopyrite to gold metallogeny of the Meguma Terrane, Nova Scotia, Canada

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ARTICLE INFO

Article history:

Received 1 July 2019

Received in revised form 7 November 2019

Accepted 14 November 2019

Available online xxx

Handling Editor: F. Pirajno

Keywords:

LA-ICP-MS

Metallogeny

Gold

Meguma Terrane

Orogenic deposit

ABSTRACT

Investigation of gold metallogeny in the Paleozoic Meguma terrane (Canada) is conducted through LA-ICP-MS analysis of arsenopyrite collected from eight slate-belt style vein gold deposits using a novel approach integrating elemental distribution maps and their derived elemental paragenesis with multi-element binary plots. The data reveal two distinct gold events: 1) an early event characterized by a Co-Ni-Mo-Sb-Se elemental association related to initial growth of arsenopyrite that reflects the presence of invisible gold (>10 ppm); and 2) a second event, spatially associated with late fracture sets, that is characterized by an Al-Ti-V-Mn element association and reflects either remobilization or upgrading of primary invisible gold and is manifest as visible gold.

The results of this study indicate a complex and protracted history of gold mineralization which has important ramifications for the Meguma gold deposits, as well as other orogenic gold districts globally. In the case of the Meguma Terrane, it involves an initial gold event that is followed by element mobilization and, in the case of precious metals (Au, Ag), an upgrading through a zone refining process. In addition, the variable coupling and decoupling of elements is only revealed using in-situ derived LA-ICP-MS data.

1. Introduction

The Meguma Terrane of Nova Scotia (Canada) part of the larger Appalachian Orogen of eastern North America, hosts numerous orogenic gold deposits that equate to the slate-belt-hosted lode gold deposit type (e.g., Goldfarb et al., 1986; Poulsen et al., 2000; Bierlein and Crowe, 2000). These quartz-vein hosted deposits were exploited from the late 19th to the middle of the 20th century with a total recorded production of ca. 1.3 Moz Au from about 60 deposits (Sangster and Smith, 2007). More recent mining and exploration activity indicate, however, that the true endowment of these deposits is a multiple of this, as evidenced by the current and forecast production of 200,000 oz./year by Atlantic Gold Corp. from four of these historical deposit areas (<http://www.atlanticgoldcorporation.com>).

The nature of these deposits has been well documented by various studies which, in addition to regional- and deposit-scale studies, includes structural analysis, litho-geochemistry, geochronology, stable (C, O, H) and radiogenic (Sr, Pb) isotopes, and fluid inclusions, as summarized by Ryan and Smith (1998) and Kontak and Horne (2010). The results of these studies indicate that similar overall geological settings, veins types, fluid chemistry and wall-rock alteration occurs in all the deposits, which strongly suggests a similar origin for gold mineralization across the entire terrane. However, despite having similar geological and geochemical features, at least two dis-

tinct vein-forming events are defined based on the aforementioned integrated studies: 1) an early event related to the regional Neocadian orogenic deformation at ca. 410–390 Ma; and 2) a later event associated with the widespread emplacement of meta-to peraluminous granitoids and lesser mafic bodies at ca. 370–380 Ma. Importantly, gold mineralization in both cases has been attributed to the circulation of fluids carrying a metamorphic signature which coincide with regional deformation (ca. 408 Ma) and intrusion of large granitic batholiths (ca. 380 Ma) (Kontak and Horne, 2010; Kontak et al., 2011). Thus, despite the exhaustive amount of data available for these deposits (see further discussion below) there remain many outstanding questions about the elemental associations in these deposits, and the nature of both refractory and non-refractory gold in these systems. It is these aspects which are addressed in this paper.

A considerable advance in our understanding of the complex evolution and paragenesis of many gold deposit settings, such as multi-stage mineralizing events and/or remobilization, has been revealed through integration of classic petrographic and imaging methods (e.g., EMPA, SEM-EDS) with quantitative laser ablation inductively coupled plasma-mass spectrometry (LA-ICP-MS) element distribution analyses (i.e., maps, traverses, spots) of Fe—As sulfide phases (e.g., Large et al., 2007, 2009; Zhao et al., 2011; Lawley et al., 2015; Neyedley et al., 2017; Gourcerol et al., 2018a; Kerr et al., 2018; Dubosq et al., 2018; Wu et al., 2018, 2019). As a means to extract further information from such map and traverse data, Gourcerol et al. (2018a, 2018b) introduced the novel concept of time slice domains (TSD) which essentially converts the LA traverse data used to construct maps into point analyses. When the latter approach is com-

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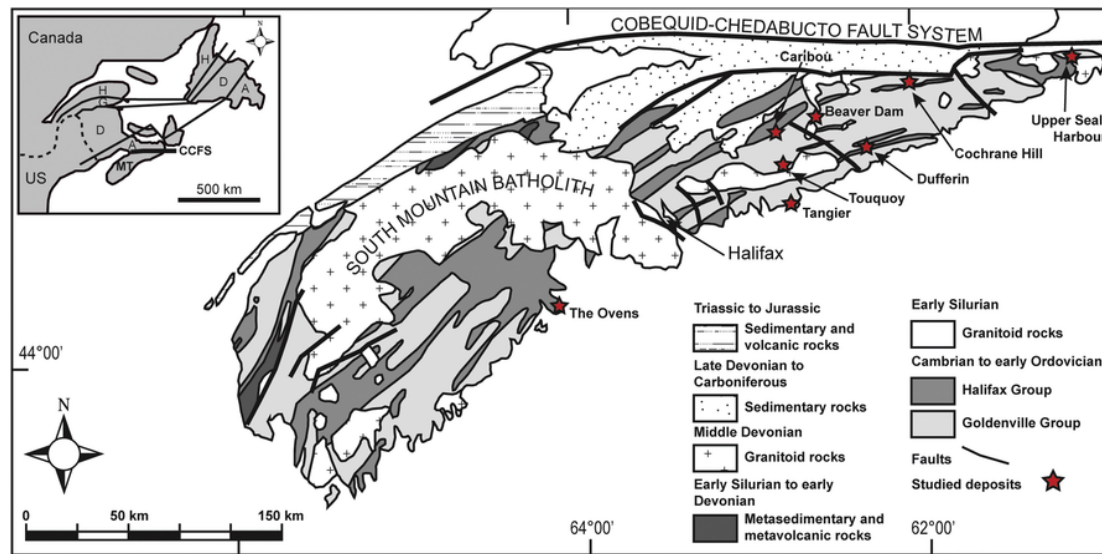


Fig. 1. Simplified geological map of the Meguma terrane (southern Nova Scotia, Canada) showing the distribution of selected gold deposits used in this study (red stars) as well as other Meguma gold deposits (black circles). The inset box outlines the location of the various terranes in the Canadian Appalachians in addition to the terrane bounding (i.e., Meguma and Avalon terranes) Cobequid-Chedabucto Fault System in Nova Scotia. Abbreviations: A: Avalon terrane; D: Dunnage terrane; G: Gander terrane; H: Humber terrane; MT: Meguma terrane. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

combined with other concepts of data treatment, such as elemental paragenesis, multi-element plots and a variety of statistical approaches, it provides the means to assess the complexities of gold paragenesis, that is primary versus subsequent mobilization and upgrading or addition of new gold (e.g., Lawley et al., 2015; Augustin et al., 2017; Masurel et al., 2019). This latter approach has been successfully applied to better constrain the nature of gold (i.e., visible to invisible) in several deposit settings in Canada, including Archean BIF-hosted mineralization (e.g., the Meadowbank and Musselwhite deposits, the Meliadine gold district; Gourcerol, 2018a), the Archean orogenic Bisset Lake deposit, (Neyedley et al., 2017), the Archean intrusion-related Renabie deposit (McDivitt, 2016), and the hybrid orogenic-intrusion related Archean Madrid deposit, Hope Bay (Kerr et al., 2018).

In order to further assess the gold metallogeny of the Meguma terrane, that is the potential source(s) of gold and mineralizing fluid(s) plus the nature of the gold events, we present the results of LA-ICP-MS mapping of grains and traverses of arsenopyrite combined with treatment of these data using the TSD approach for several gold deposits (see Fig. 1). Our goal is to establish elemental associations and paragenesis for each deposit, assess the potential upgrading of Au within the sulfide host, constrain potential reservoirs with which the mineralizing fluids interacted, and assess the apparent similarity of many of these gold deposits. As this approach is both cost and time intensive, we acknowledge the dataset provided is albeit limited compared to more extensive litho-geochemical studies that are traditionally used, such as for the Meguma gold deposits themselves where Kerswill (1988, 1992) used 490 samples to show a strong correlation between both As and S versus Au. The advantage of our approach, as shown below and discussed by Gourcerol et al. (2018b), is that it provides a means to relate elements paragenetically versus spatially as in whole-rock analyses. Thus, this study is intended to be more of a proof of concept.

2. Regional geological setting

The Meguma terrane of southern Nova Scotia (Canada) defines the most easterly allochthonous part of the Canadian Appalachian orogenic belt (Fig. 1; Williams, 1995) with its emplacement coinci-

dent with the Neoproterozoic (van Staal, 2007; White and Barr, 2010). This terrane, around 480 km by 120 km, consists of: 1) late Neoproterozoic (i.e., Ediacaran) to Ordovician metaturbiditic rocks (i.e., former Meguma Group), which includes the basal sandstone-dominant Goldenville Group (ca. 11 km) and conformably overlying shale-dominant Halifax Group (ca. 5 km) (White et al., 2012); 2) the disconformably overlying Silurian-Devonian metasedimentary and bimodal metavolcanic rocks of the Annapolis Supergroup (White et al., 2012); 3) voluminous ca. 380 to 370 Ma granitic batholiths (e.g., South Mountain Batholith) and smaller plutons with lesser gabbroic intrusions and mafic dike rocks (e.g., Kontak and Reynolds, 1994; Clarke et al., 1997;); and 4) rare ca. 360 Ma granitoid bodies restricted to the southwest Meguma terrane (see Kontak et al. (2013) for summary) (Table 1). The metasedimentary and metavolcanic units were intensely deformed and metamorphosed (i.e., greenschist-amphibolite facies) in the Late Devonian (e.g., Kontak et al., 1998; Hicks et al., 1999) during the collision of the Meguma and Avalon terranes (ca. 410–390 Ma) along the east-west trending Cobequid-Chedabucto Fault System (CCFZ; Fig. 1; Murphy et al., 2012). The overlying clastic rocks of the Late Devonian-Early Carboniferous Horton Group, which on-lap the CCFZ, record the uplift and erosion of the aforementioned deformed rocks.

The basement of the Meguma terrane (i.e., substrate to the Meguma metasedimentary rocks) is not well documented and remains equivocal. At the supracrustal level, some workers (Clarke et al., 1993; Dostal et al., 2006) interpret the intensely folded, high-grade rocks represented by ortho- and paragneisses exposed in the so-called Liscomb Complex to be analogous to a gneiss dome, whereas others (White et al., 2009; Scallion et al., 2011) consider these rocks to be part of a contact metamorphic aureole related to proximal granites. The presence, however, of granulite facies xenoliths in ca. 374 Ma lamprophyric dyke rocks of the eastern Meguma terrane (Owen et al., 1988; Ebertz et al., 1991; Greenough et al., 1999) and as fragments in megabreccias in the CCFZ (Gibbons et al., 1996) indicate the unequivocal presence of a high-grade basement infrastructure. The aforementioned lamprophyre-hosted xenoliths, termed the Tangier xenoliths, include sapphirine granulites, mafic gneisses

Table 1

Summary of the major events and their respective ages affecting the Meguma and Avalon terranes (Nova Scotia, Canada).

Events	Age	Methods	Comments	References
Ordovician granitic and gabbroic magmatism	460 Ma	U/Pb (zr)	Located in the Avalon terrane, refers to A-type granites	Escarraga et al., 2012; Murphy et al., 2012
Early Silurian igneous activity	440 Ma	U/Pb (zr)	Reported in the western part of the Meguma terrane, from volcanic rocks within the White Rock Formation and the Brenton pluton	Keppie and Krogh, 2000; MacDonald et al., 2002
Acadian orogeny	410–390 Ma	⁴⁰ Ar/ ³⁹ Ar (wr)	Corresponds to regional metamorphism and deformation	Keppie and Dallmeyer, 1987; Muecke et al., 1988; Kontak et al., 1998, 2002; Hicks et al., 1999
Peraluminous granitoid emplacement	370–380 Ma	⁴⁰ Ar/ ³⁹ Ar (ms, bt) U/Pb (zr, mz)	Described as S-type granites; U/Pb reflects age of crystallization; Ar/Ar reflects cooling of plutons;	Hill, 1988; Reynolds et al., 1981; Bradley, 1983; Keppie and Dallmeyer, 1987, 1995; Krogh and Keppie, 1988; Clarke et al., 1993; Keppie et al., 1993; Kontak et al., 2004; Kontak and Reynolds, 1994; Tate et al., 1997; Keppie and Krogh, 1999; Murphy and Keppie, 2005; Chen, 2015
Emplacement of gneisses of the Liscomb Complex	374–377 Ma	⁴⁰ Ar/ ³⁹ Ar (wr) U/Pb (zr, mz)	Contemporaneous with peraluminous granites, closely associated with amphibolite-granulite facies metamorphism	Kontak and Reynolds, 1994; Dostal et al., 2006
Emplacement of mafic intrusions and dikes	370–374 Ma	⁴⁰ Ar/ ³⁹ Ar (hbl, bt)	Dating performed on hbl and bt; Rapid cooling of dikes and gabbroic bodies (e.g., Liscomb area)	Kempster et al., 1989; Kontak and Reynolds, 1994
Granitic magmatism	360 Ma	⁴⁰ Ar/ ³⁹ Ar (ms) Re/Os (mb)	Described as A- and I-type granites; Reported in the Southwestern part of the Meguma deposit	Kontak et al., 2013

and garnetiferous quartzo-feldspathic gneisses (Owen et al., 1988; Owen and Greenough, 1991).

2.1. Geological and geochemical parameters of Meguma gold deposits

The Meguma lode-gold deposits discussed below are present throughout the metasandstone dominant Goldenville Group with a few also at the base of the overlying, mudstone dominant Halifax Group (Malcolm, 1929; Ryan and Smith, 1998; Horne and Culshaw, 2001). The deposits are well constrained in terms of their structural settings, as it has long been known that they are localized in the hinge areas of anticlines or their southern, often overturned limbs (Malcolm, 1929; Keppie, 1976; Horne and Culshaw, 2001). The deposits consist of a variety of quartz vein types, including bedding concordant, discordant, en echelon, and saddle reef (Malcolm, 1929; Keppie, 1976; Graves and Zentilli, 1982; Horne and Culshaw, 2001; Kontak and Horne, 2010). Despite their various geometrical relationships to their host rocks, most of these veins are now considered to represent emplacement during a late-stage fold tightening and, based on their mutual cross-cutting relationships, to represent in a specific deposit a single vein-forming event (Horne and Culshaw, 2001; Kontak and Horne, 2010). It has also been shown that individual deposits may be dominated by one vein type, thus, for instance, saddle veins at The Ovens, Upper Seal Harbour, and Dufferin deposits, whereas bedding-concordant veins at Beaver Dam, Cochrane Hill, and Moose River deposits, and both discordant and concordant vein types at Caribou and Tangier deposits. These latter named settings representing the focus of this study (Fig. 1).

The auriferous veins are generally dominated by a simple quartz ± carbonate ± sulfide mineral assemblage (Malcolm, 1929; Newhouse, 1936; Kontak and Horne, 2010), hence similar to most orogenic vein systems (Goldfarb et al., 2005; Dubé and Gosselin, 2007), and often have ribbon textures due to inclusion of thin wall-rock slivers or septae. The sulfides include arsenopyrite, which is locally the dominant sulfide phase, in addition to pyrite, pyrrhotite, galena, sphalerite and chalcopyrite. Gold is present in all the vein types mentioned and occurs in a variety of settings (e.g., Kontak and Smith 1993; Kontak et al., 1996; Ryan and Smith, 1998), which includes discrete grains in vein quartz, along wall-rock contacts or adhering to ribbons of wall-rock, attached to sulfides (arsenopyrite ± galena ± pyrrhotite ± pyrite ± sphalerite), or as fracture-filling/inclusions within sulfides, in particular arsenopyrite. Although low-grade (i.e., <1–2 g/t Au) disseminated-type (i.e., vein-poor packages) gold mineralization is present in the metasedimentary host rocks, it has only been recognized as a significant component of the mineralization in a few settings (e.g., Touquoy, Fifteen Mile Stream, Beaver Dam), with the Touquoy deposit of the Moose River district the only site where this has been studied in detail (Bierlein and Smith, 2003). In the latter case, a global resource of a 550,000 oz. @ 1.55 g/t was indicated prior to commencement of mining in 2018 (http://www.atlanticgoldcorporation.com/projects/touquoy_gold_project). As in other metasedimentary rock-hosted gold settings (Bierlein and Crowe, 2000), wall-rock alteration varies from cryptic to intensely developed with zones of sericite, silica, carbonate (e.g., ankerite and calcite), and Fe-sulfides (i.e., arsenopyrite) the most abundant; in addition are rare biotite and tourmaline alteration (e.g., Kontak and Smith, 1989, 1993; Bierlein and Smith, 2003; Kontak and Horne, 2010).

Given the similar settings, vein paragenesis, and fluid chemistry, as constrained by isotopes (S, O, D, C), fluid inclusions (i.e., aqueous carbonic, X_{CO2} = 0.10–0.2, 5–10 wt% equiv. NaCl), and wall-rock al-

Table 2
Summary of the relevant geology of the studied gold deposits from the Meguma terrane (Nova Scotia, Canada).

Deposits	Localization	Stratigraphy	Deposit-type	Metamorphism	Dating	Comments	References
Beaver Dam	Southern limb of the Beaver Dam northeast-trending anticline; close to River Lake granite intrusion	Metasiltstones (Goldenville Formation)	Bedding-parallel and discordant narrow qtz veins	Greenschist	460 Ma (Re/Os), 440 Ma (Re/Os), 370–380 Ma (Ar/Ar)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on vein-fill minerals (hbl, ms, bt) and whole rock; Re/Os performed on Aspy (Ar/Ar)	Kontak et al., 1990b, 1993; Kontak and Smith, 1993; Kontak and Kerrich, 1995; Chen, 2015
The Ovens	Hinge of the Ovens Anticline	Metasiltstones (Goldenville Formation)	Saddle-reef concordant and discordant qtz veins	Greenschist	406 Ma (Re/Os), 380 Ma (Re/Os; Ar/Ar)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on ms from pressure shadows around Aspy and bt alteration; Re/Os performed on Aspy	Hicks et al., 1999; Kontak et al., 1998; Kontak; Horne and Culshaw, 2001; Morelli et al., 2005; Chen, 2015
Dufferin	Hinge of the Crown Reserve Anticline	Metasiltstones (Goldenville Formation)	Saddle-reef concordant qtz veins	Greenschist	408 Ma (Ar/Ar), 380 Ma (Re/Os)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on wall rock adjacent to discordant vein; Re/Os performed on Aspy	Morelli et al., 2005; Horne and Culshaw, 2001; Kontak et al., 1998;
Touquoy	Centred on the Moose River Anticline	Metasiltstones (Goldenville Formation)	Bedding-parallel and discordant narrow qtz veins	Greenschist	440 Ma (Re/Os), 410–400 Ma (Ar/Ar; Re/Os), 380 Ma (Ar/Ar)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on vein-micas and whole rock	Kontak et al., 1990b, 1993; Bierlein and Smith, 2003; Chen, 2015
Cochrane Hill	Eastward extension of the Cochrane Hill Anticline	Metasiltstones (Goldenville Formation)	Bedding-parallel	Amphibolite			Smith, 1983a, b; Smith, 1984
Caribou	Northeast-trending Caribou Anticline	Metasiltstones (Goldenville Formation)	Bedding-parallel and discordant narrow qtz veins	Greenschist	373 Ma (Ar/Ar)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on ms	Kontak et al., 1990b
Upper Seal Harbour	Hinge of an anticlinal	Metasiltstones (Goldenville Formation)	Saddle-reef concordant veins	Greenschist	366 Ma (Ar/Ar)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on ms	Kontak et al., 1990b
Tangier	Southern limb of the Tangier-Harrigan Cove northeast-trending anticline; close to the Musquodobit batholith	Metasiltstones (Goldenville Formation)	Bedding concordant and discordant	Greenschist	374 Ma (Ar/Ar)	$^{40}\text{Ar}/^{39}\text{Ar}$ performed on bt	Corey and Mills, 1992; Smith, 2000; Kontak and Archibald, 2002

Abbreviations: Aspy=arsenopyrite; bt=biotite, hbl=hornblende, ms=muscovite.

teration, these deposits have been suggested to share a metamorphic origin (e.g., Graves and Zentilli, 1982; Kontak et al., 1990a, 2011; Kontak and Kerrich, 1997; Ryan and Smith, 1998; Kontak and Horne, 2010). However, direct dating (Re—Os, $^{40}\text{Ar}/^{39}\text{Ar}$) of some deposits indicate at least two distinct gold mineralizing events (see summary in Table 2): 1) an early one, coincident with regional deformation and recognized locally at Beaver Dam, The Ovens, and Touquoy deposits based on Re/Os dating of arsenopyrite grains (e.g., Morelli et al., 2005; Chen et al., 2014; Chen, 2015); and 2) a later event, which overlaps the felsic(–mafic) magmatism at ca. 380–370 Ma, as constrained by $^{40}\text{Ar}/^{39}\text{Ar}$ (whole rock, amphibole, muscovite, biotite; see summary in Kontak and Archibald (2002) and Kontak and Horne (2010) and Re—Os on arsenopyrite grains (Morelli et al., 2005) dating. This gold event is represented at Beaver Dam, The Ovens, Dufferin, Touquoy, Upper Seal Harbour, Tangier and Caribou deposits. Thus, there is some evidence for possibly multiple hydrothermal events in these deposits (e.g., Beaver Dam), the degree and significance of which remains unquantified, as has also been recorded in other orogenic gold deposit settings (e.g., Wagner et al., 2007; Lawley et al., 2015; Kerr et al., 2018).

2.2. Occurrence of arsenopyrite

As noted above, arsenopyrite is a common sulfide phase in the Meguma deposits in both mineralized and barren veins and in adja-

cent wall rock. Representative examples from among the various deposit sites are shown in Fig. 2 (see Fig. 1 for locations) and includes Beaver Dam (Fig. 2A, B, C, D), Touquoy (Fig. 2E, F, G), The Ovens (Fig. 2H, I, J), Dufferin (Fig. 2K), Mooseland (Fig. 2L, M, N), and Fifteen Mile Stream (Fig. 2O). In general, arsenopyrite is seen to occur as: 1) layers partially to wholly replacing wall rock where it is concentrated as thin layers along bedding-plane horizons (Fig. 2A, B, C, N, O) or replacing beds (Fig. 2I, J); 2) disseminations in the various metasedimentary host rocks (Fig. 2F); 3) coarse porphyroblasts in the host rocks (Fig. 2C, D, E, G); 4) coarse euhedral in quartz-dominant veins (Fig. 2H, M, L) or the dominant phase of small quartz veins (Fig. 2E); and 5) replacing wall-rock ribbons having stylolitic texture, as seen in a mineralized quartz vein (Fig. 2K).

The timing of arsenopyrite in relation to vein formation is poorly constrained, in particular when present as disseminations (Fig. 2F), porphyroblasts (Fig. 2C, G), or replacing beds (Fig. 2J). However, where fabric development is observed, the formation of pressure shadows (Fig. 2E) or boudinaging of veins (Fig. 2H) about arsenopyrite indicates at least some of these grains are pre- to syn-deformation. Given that the emplacement of the majority of the quartz veins have been considered to reflect a late-stage folding tightening event of the Meguma stratigraphy (Horne and Culshaw, 2001), it follows therefore that the timing of arsenopyrite growth would also be considered to be synchronous with vein formation (i.e., fluid flux).

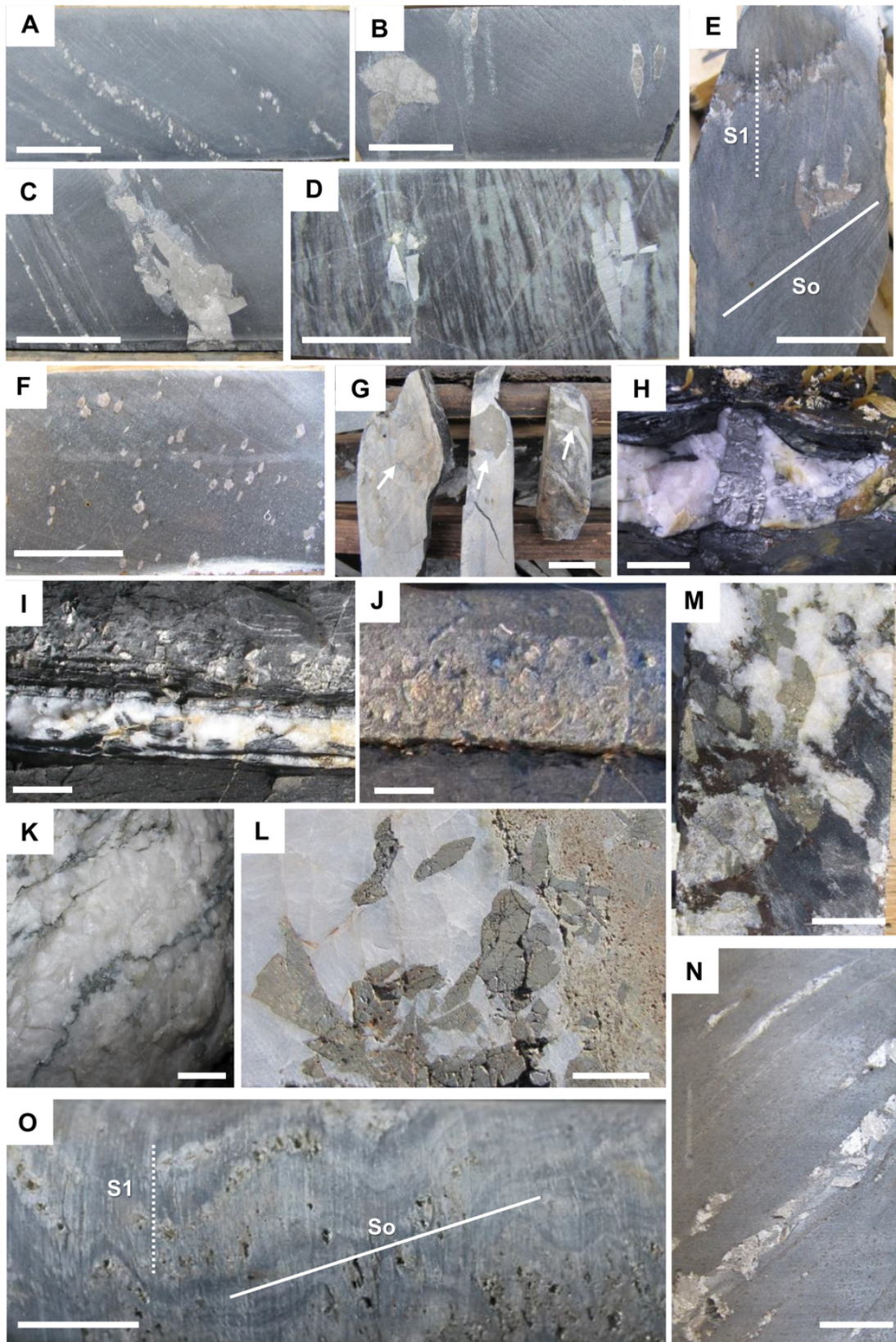


Fig. 2. Representative images of outcrop (H, I, J, K) and drill core (all others) showing the style of arsenopyrite (Aspy) found in the Meguma gold deposits with samples from the following locations (see Fig. 1). Note that the white line on each picture represents 2 cm. A, B, C, D) Beaver Dam; E, F, G) Touquoy; H, I, J) The Ovens; K) Dufferin; L, M, N) Mooseland; O) Fifteen Mile Stream. A) Aspy along bedding-plane horizons and lacking quartz. B, C) Coarse porphyroblastic aspy, in part as aggregates (see image C) lacking pres-

sure shadows, and as finer grains along bedding-plane horizons. D) Bleached metasilstone unit with aspy coarse aspy euhedra growing along bedding-plane horizons. E) Aspy in folded bedding concordant vein at the contact of mudrock and metasilstone and also as randomly oriented porphyroblasts in the metasilstone. Note the development of pressure shadows filled with quartz and carbonate in the latter. The traces of bedding (So) and fabric (S1) are highlighted. F) Disseminated aspy in metasilstone showing incipient development of pressure shadows. G) Coarse aspy porphyroblasts in vein-free metasilstones. H) Coarse euhedral pyrite in boudinaged quartz vein cutting black mudstone. I) Crack-seal or ribbon -textured quartz vein in mudstone with abundant euhedral aspy in the adjacent bed of metasilstone unit. J) Metasilstone bed replaced by massive aspy. K) Aspy replacing thin septae of wall rock material having stylonitic texture in mineralized quartz vein. L) Coarse aspy euhedra at the vein-wall rock contact; the porous texture at the contact is due to dissolution of carbonate. M) Vein sample showing aspy intergrown with quartz and biotite replacing wall-rock inclusions of mudstone. N) Aspy along bedding-plane horizons with some development of quartz \pm carbonate. O) Aspy disseminated in mudstone unit from a fold hinge. Note the presence of both bedding (so) and S1 fabric. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Table 3
LA-ICP-MS Instrumentation and acquisition parameters.

Laser ablation system	
Instrument	Resonetics RESOLUTION M-50
Laser type	ArF excimer
Wavelength	193 nm
Pulse duration	20 ns
Repetition rate	5 Hz (LC-12-16; D4-A; TQ-1C); 6 Hz (OV-6); 7 Hz (everything else)
Spot diameter	48 μ m (LC-12-16; D4-A), 90 μ m (OV-6), 66 μ m (everything else)
Speed	18 μ m/s (LC-12-16; D4-A), 55 μ m/s (OV-6), 33 μ m/s (everything else)
Energy density	5 J/cm ²
Carrier gas (He)	650 ml/min
Additional gases	6 ml/min N ₂
Primary reference material	NIST610
Secondary reference material(s)	NIST612, BHVO2g, Po725
Mass spectrometer	
Instrument	Thermo X-Series II
Plasma RF power	1450–1480 W
Plasma gas flow	780–820 ml/min
Isotopes analyzed for LC-12-16, OV-6, D4-A, and TQ-1C	27Al, 47Ti, 51V, 52Cr, 55Mn, 57Fe, 59Co, 60Ni, 65Cu, 66Zn, 75As, 82Se, 95Mo, 107Ag, 111Cd, 113In, 118Sn, 121Sb, 182W, 197Au, 208Pb, 209Bi 10 ms dwell time per analyte total duty cycle of 276 ms
Isotopes analyzed for CH-K275 and the DM-91-12, Tangier, and CAR-88-7C traverses	27Al, 33S, 47Ti, 51V, 52Cr, 55Mn, 56Fe, 57Fe, 59Co, 60Ni, 65Cu, 66Zn, 75As, 82Se, 95Mo, 107Ag, 111Cd, 113In, 115In, 118Sn, 121Sb, 125Te, 182W, 197Au, 208Pb, 209Bi 10 ms dwell time per analyte total duty cycle of 316 ms
ThO ⁺ /Th ⁺	< 0.6%
U ⁺ /Th ⁺	0.9–1

ThO⁺/Th⁺ and U⁺/Th⁺ were determined on NIST612 during instrument tuning.

3. Analytical methods

3.1. LA-ICP-MS sulfide trace element chemistry

Arsenopyrite grains mounted in pucks or as polished thin sections were analyzed by LA-ICP-MS in Laurentian University's Chemical Fingerprinting laboratory. Areas of the sulfide grains free of mineral inclusions (i.e., other sulfides, oxides, silicates, visible gold) and with minimal amounts of fractures were selected for analysis in either raster or traverse mode, with the former providing the element maps referred to below. The selected areas were ablated with a Resonetics RESOLUTION M-50, 193 nm wavelength, 20 ns pulse duration ArF excimer laser ablation microprobe employing a Laurin Technic two-volume sample cell (Müller et al., 2009). Ablation took place in ultra-pure He flowing at 650 ml/min. The ablated material and He were combined with Ar (780–820 ml/min) and a small flow of N₂ (6 ml/min; to enhance sensitivity, Hu et al., 2008) just outside the cell

and approximately 2.5 m before entering the torch. The combined aerosol was fed into a Thermo X-Series II quadrupole ICP-MS for elemental analysis. The analyses consisted of traverses and maps (themselves consisting of a series of adjacent, equal length traverses) that were acquired by moving the sample under the laser at speeds of 1/3 to 1/2 the spot diameter per second. Between each traverse, 30 to 45 s of gas blank was collected to be used for baseline subtraction during data reduction. Reference materials (a combination of NIST610 +/- NIST612, BHVO2g and Po725) were analyzed periodically to account for instrument drift. The data were collected in several sessions from 2014 to 2016, with each map taking 4 to 7 h to acquire. The laser and ICP-MS operating conditions are listed in Table 3.

The raw data were quantified using the trace-element data reduction scheme of Iolite 3 (Paton et al., 2011), where NIST610 (Jochum et al., 2011) and Fe acted as the external and internal references for quantification, respectively. A slight modification was made to the trace element data reduction scheme so that the Fe was not forced to 34.3% on a point-by-point or a pixel-by-pixel basis, but to the mean of a representative and clean part of the arsenopyrite grain. This was done to avoid over or under estimating concentrations in fractures and inclusions where the Fe and ablation rate can vary. In fractures and inclusions, the data are considered to be semi-quantitative. Although the arsenopyrites and reference materials used differ significantly in terms of their chemistry and ablation efficiency, Wohlgemuth-Ueberwasser and Jochum (2015) and Gourcerol et al. (2018b) have shown that using the NIST610/Fe calibration strategy to quantify PGEs in a sulfide (Po) yields concentrations acceptably close to what is expected. Additional experiments conducted as part of this study demonstrate that a wide range of elements in Fe-rich reference materials including GSD-1G, GSE-1G, BHVO2G and MASS1 (USGS), and Po725 (Sylvester et al., 2005) are mostly within 20% of their accepted values when calibrated this way (cf., Fig. A). A homogeneous, well characterized reference arsenopyrite does not exist that we are aware of, therefore this combined evidence of a wide range of Fe-rich materials being acceptably calibrated with NIST610/Fe provides reasonable confidence in the arsenopyrite concentrations determined this way. It is also noteworthy that the type of data analysis performed here (TSD-based) is concerned more with orders of magnitude concentration variations rather than subtle differences, thus rendering minor inaccuracies irrelevant.

Element maps were stitched together using Iolite's "Image from Selections" functionality (Woodhead et al., 2007), with additional bilinear interpolation between traverses and 3 \times 3 mean pixel smoothing. This was done for the images (i.e., Figs. 3–7), but not for the TSD (i.e., Figs. 14–18), for which no interpolation or smoothing were performed in order to avoid biasing the data.

The analyses were performed on euhedral arsenopyrite grains with element distribution maps acquired for samples from the Beaver Dam, The Ovens, Dufferin, Touquoy and Cochrane Hill deposits, whereas element distribution traverses were acquired for samples from the Upper Seal Harbour, Tangier and Caribou deposits. Arsenopyrite grains were sampled from bedding parallel quartz veins, except for The Ovens, which were hosted in metasilstone adjacent to a bedding parallel vein. The samples used for LA-ICP-MS analysis were examined in both reflected light and also using the scan-

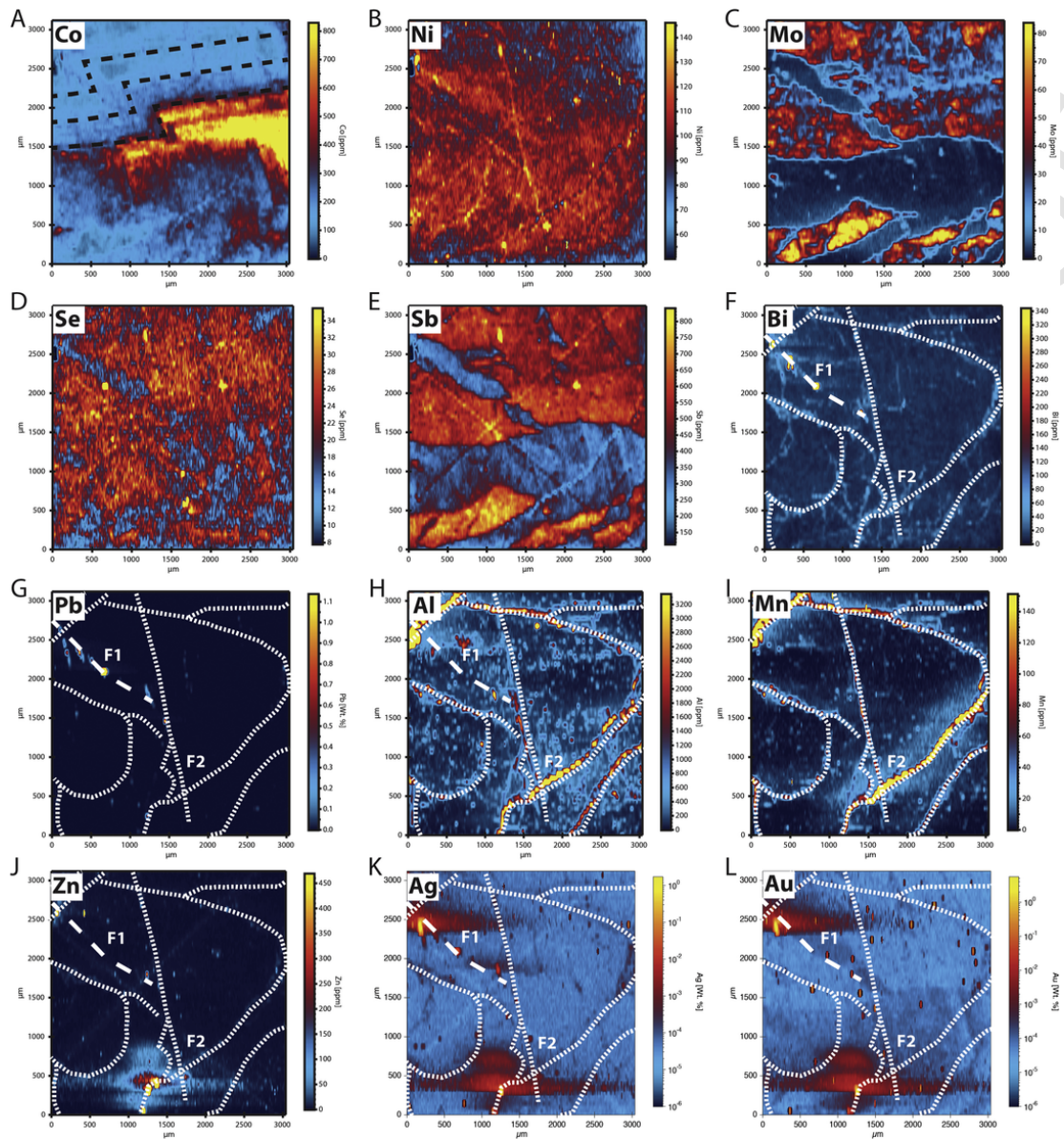


Fig. 3. LA-ICP-MS element maps showing the distribution of selected elements in part of an arsenopyrite grain (sample LC-12-16) from the Beaver Dam gold deposit, Meguma terrane (Nova Scotia). A) to J) Elemental maps plotted as concentrations scaled between the data median ± 3 standard deviations (in ppm). K, L) Element maps plotted as logarithmic concentrations scaled between the minimum and maximum values (in wt%). Note the following features illustrated in the figure: 1) the black dashed lines shown in A represent inferred primary growth zoning; 2) the white dashed lines represent two sets of fractures referred to as F1 (spaced one) or F2 (closer spaced one). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

ning electron microscopy coupled with an energy dispersive X-ray spectrometry (SEM-EDS).

4. Results and interpretation

4.1. Elemental paragenesis

Quantitative LA-ICP-MS element distribution maps and traverses (Table 4) were performed on several arsenopyrite grains from eight different deposits from the central and eastern part of the Meguma terrane (Fig. 1): Beaver Dam (Fig. 3), The Ovens (Fig. 4), Dufferin (Fig. 5), Touquoy (Fig. 6), Cochrane Hill (Fig. 7), Upper Seal Harbour (Fig. 8), Tangier (Fig. 9), and Caribou (Figs. 10, 11). These analyses, in combination with detailed petrographic and/or SEM-

EDS study, provide the basis for establishing an elemental paragenesis for each deposit, which is shown in Fig. 12 and discussed in detail below. This procedure is paramount in determining elemental associations for each of the deposits and is the first step towards understanding the mineralizing event in a deposit. In regards to the paragenesis presented, we note that not all the elements summarized in Fig. 12 are shown in the elemental distribution maps presented.

We first note in regards to the elemental maps and related elemental paragenesis, that despite being located in different stratigraphic settings with respect to both potential facies variation and host-rock relative age in the Meguma stratigraphy, the studied samples and hence deposits reveal a very similar elemental paragenesis. The euhedral arsenopyrite grains studied show an early enrichment in Co, Ni, Te, Se, Mo, and Sb (Fig. 12) which is reflected in both the elemen-

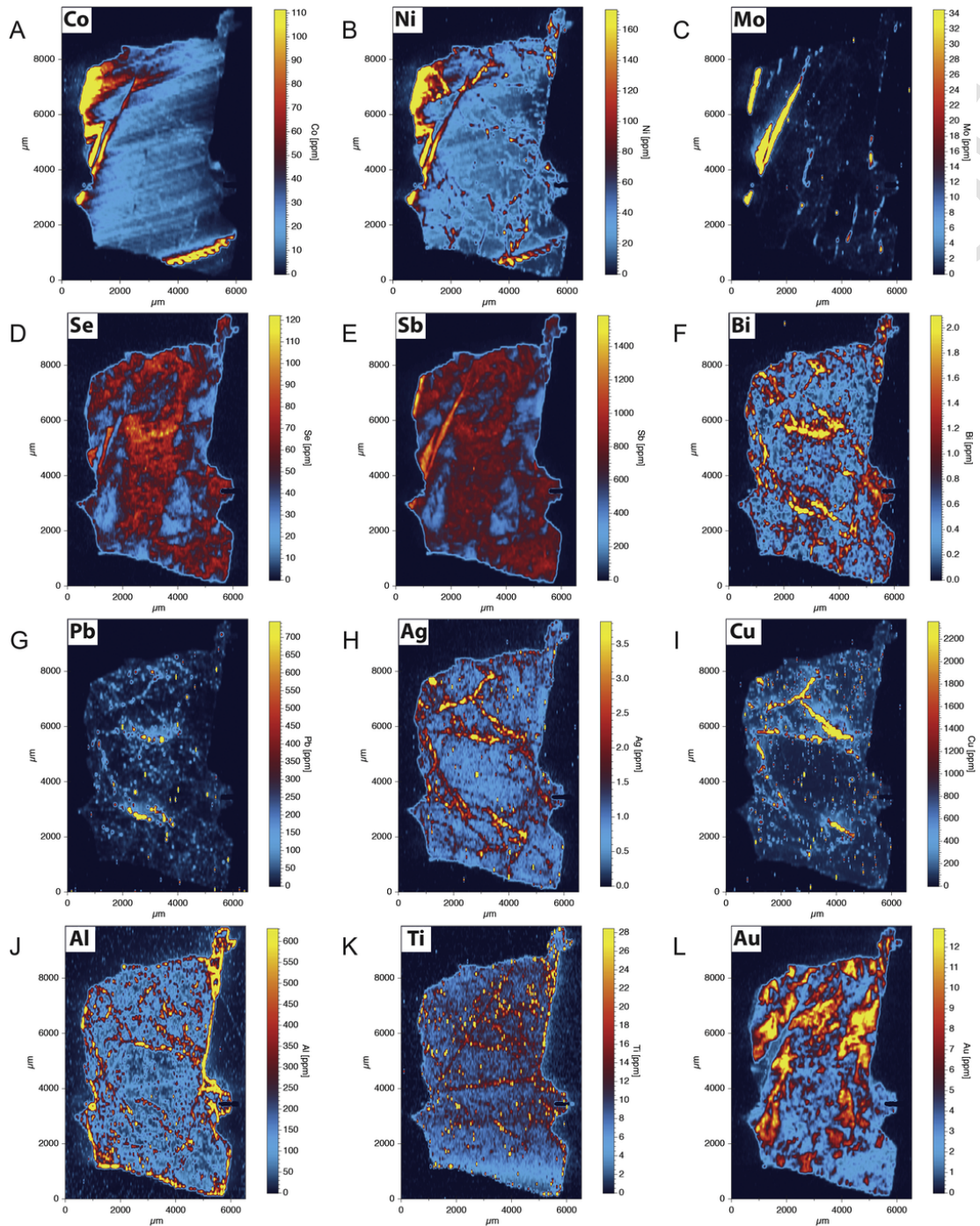


Fig. 4. LA-ICP-MS element maps showing the distribution of selected elements in part of an arsenopyrite grain (sample OV-6) from The Ovens gold deposit, Meguma terrane (Nova Scotia). A) to L) Elemental maps plotted as concentrations scaled between the data median \pm 3 standard deviations (in ppm). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

tal distribution maps (e.g., Dufferin, Touquoy, Cochrane Hill; Figs. 5, 6 and 7, respectively) and traverse patterns (e.g., Cochrane Hill, Upper Seal Harbour, Tangier and Caribou deposits; Figs. 7 to 11, respectively). The maps and traverses suggest this elemental enrichment overlaps the growth of the host phase. Moreover, traverse analyses from the Caribou samples (Figs. 10, 11) suggest that Au, and to a lesser extent Ag, also mimic this growth zoning and were introduced early as nanoparticles or a background enrichment (Figs. 10, 11). Importantly, the concentration of Co and Ni in an “external rim”

in some cases (i.e., Touquoy, Cochrane Hill deposits; Figs. 6 and 7, respectively) are also interpreted to reflect primary growth zoning.

Selenium, Mo, Sb, and to a lesser extent Ni and Te all show similar distribution patterns, which either mimic or locally overprint the zoning of the host arsenopyrite. This elemental distribution also reflects a primary feature. Traverse and some map analyses show that the putative earlier background concentrations of Au appear to be relatively antithetic to Mo zonation (e.g., Beaver Dam, The Ovens, Touquoy, Cochrane Hill, and Caribou deposits; Figs. 3, 4, 6, 7,

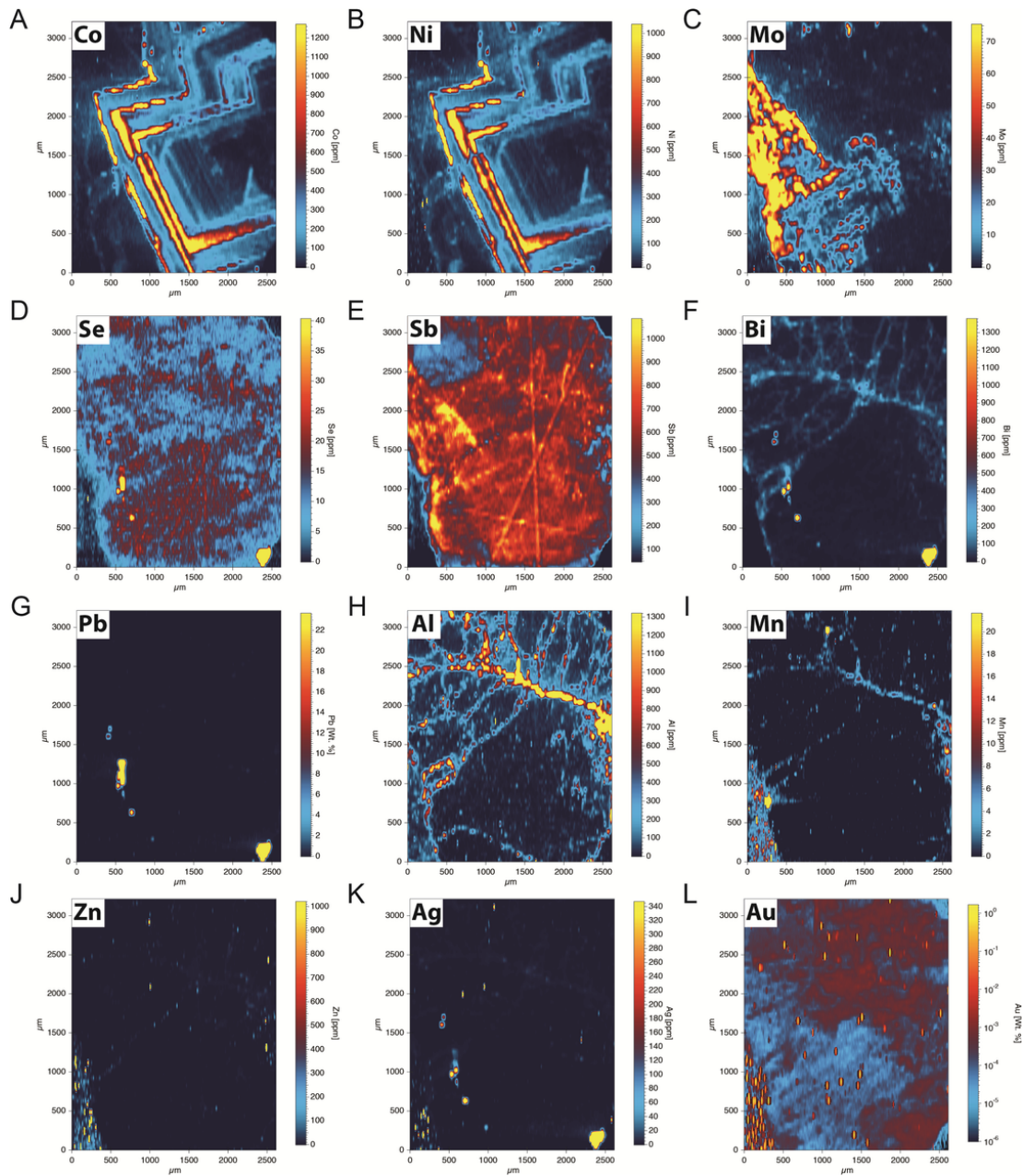


Fig. 5. LA-ICP-MS element maps showing the distribution of selected elements in part of an arsenopyrite grain (sample D4-A) from the Dufferin gold deposit, Meguma terrane (Nova Scotia). A) to K) Elemental maps plotted as concentrations scaled between the data median \pm 3 standard deviations (in ppm). L) Au map plotted as logarithmic concentrations scaled between the minimum and maximum values (in wt%). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

10, 11, with it particular well illustrated for the Dufferin deposit sample (Fig. 5).

Several events that post-date the primary growth of arsenopyrite discussed above are recognized in all the samples, as illustrated by: 1) an early Bi, Pb, Cd, In, Se, Ag, Au and locally Zn (i.e., Beaver Dam, Touquoy and Cochrane Hill, Caribou deposits; Figs. 3, 6, and 7, respectively; but note that In and Cd are lacking in the maps shown) element assemblage associated with an early fracture set (F1 assemblage herein). This assemblage is also associated with formation of galena and likely sphalerite either as micro-inclusions or as fracture-fillings (Fig. 13A, B, C, D). Further support of this was presented by Kontak and Smith (1993) in their study of the Beaver Dam de-

posit; and 2) a later stage of enrichment of a broad suite of elements that includes Al, Ti, V, Mn, Au, Ag, (e.g., Beaver Dam, Dufferin and Touquoy; Figs. 3, 5, 6, respectively; also Cr, W, Sn which are not shown in the maps), Zn (e.g., Dufferin and Touquoy; Figs. 5 and 6, respectively), Cu (e.g., Beaver Dam, Cochrane Hill; Figs. 3 and 7, respectively). This later elemental assemblage is associated either with a later fracture set (F2 assemblage herein), which clearly crosscuts the earlier set (cf., note F1 in Beaver Dam; Fig. 3), or with an external rim on the arsenopyrite (e.g., Touquoy; Fig. 6). This elemental assemblage is reflected mineralogically by the presence of various inclusions or fracture-fillings in arsenopyrite (e.g., rutile, ilmenite, chlorite; Fig. 13D, E, F). Furthermore, examination of the elemen-

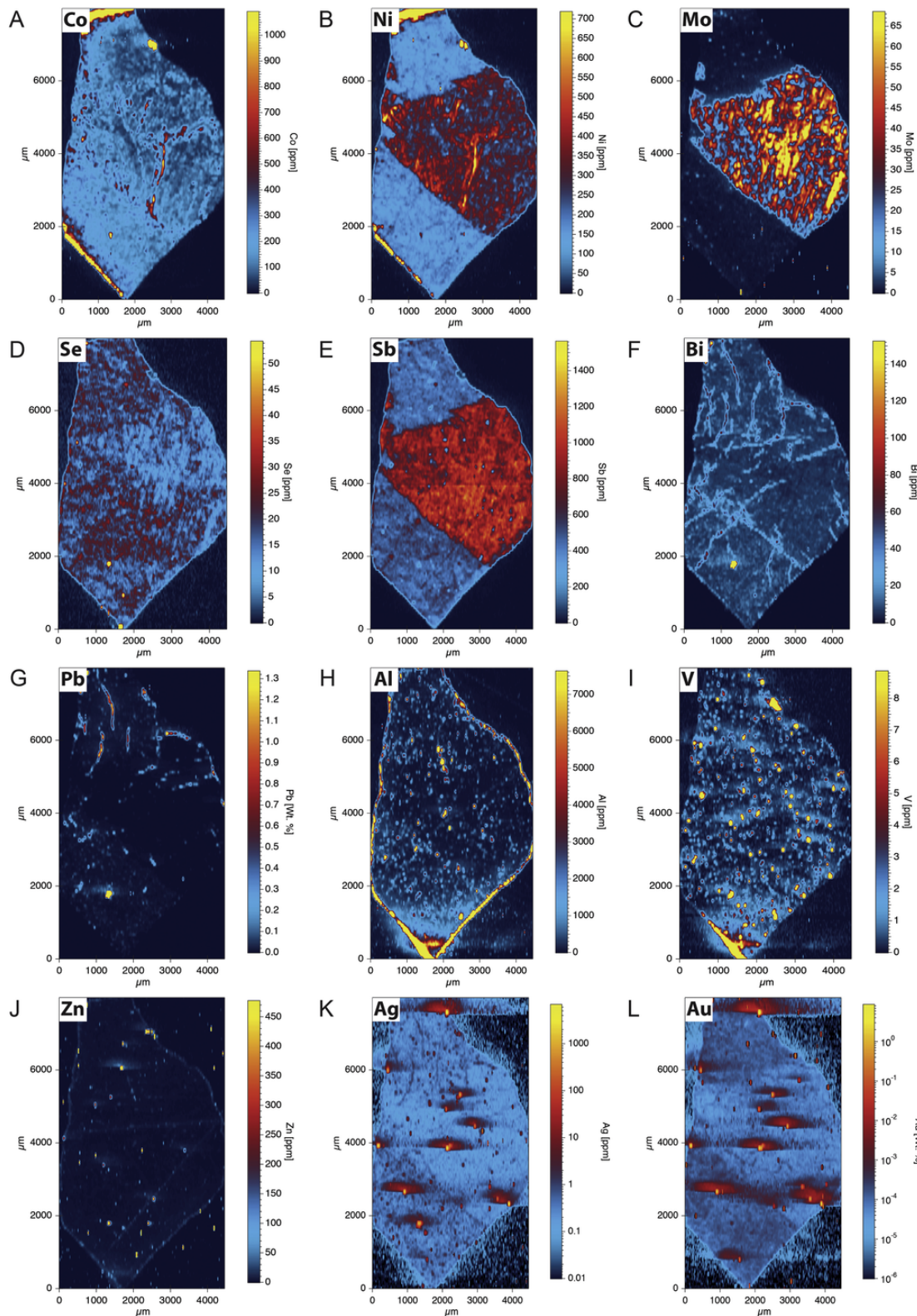


Fig. 6. LA-ICP-MS element maps showing the distribution of selected elements in part of an arsenopyrite grain (sample TQ-1C) from the Touquoy gold deposit, Meguma terrane (Nova Scotia). A) to J) Elemental maps plotted as concentrations scaled between the data median \pm 3 standard deviations (in ppm). K, L) Element maps plotted as logarithmic concentrations scaled between the minimum and maximum values (in ppm and wt%). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

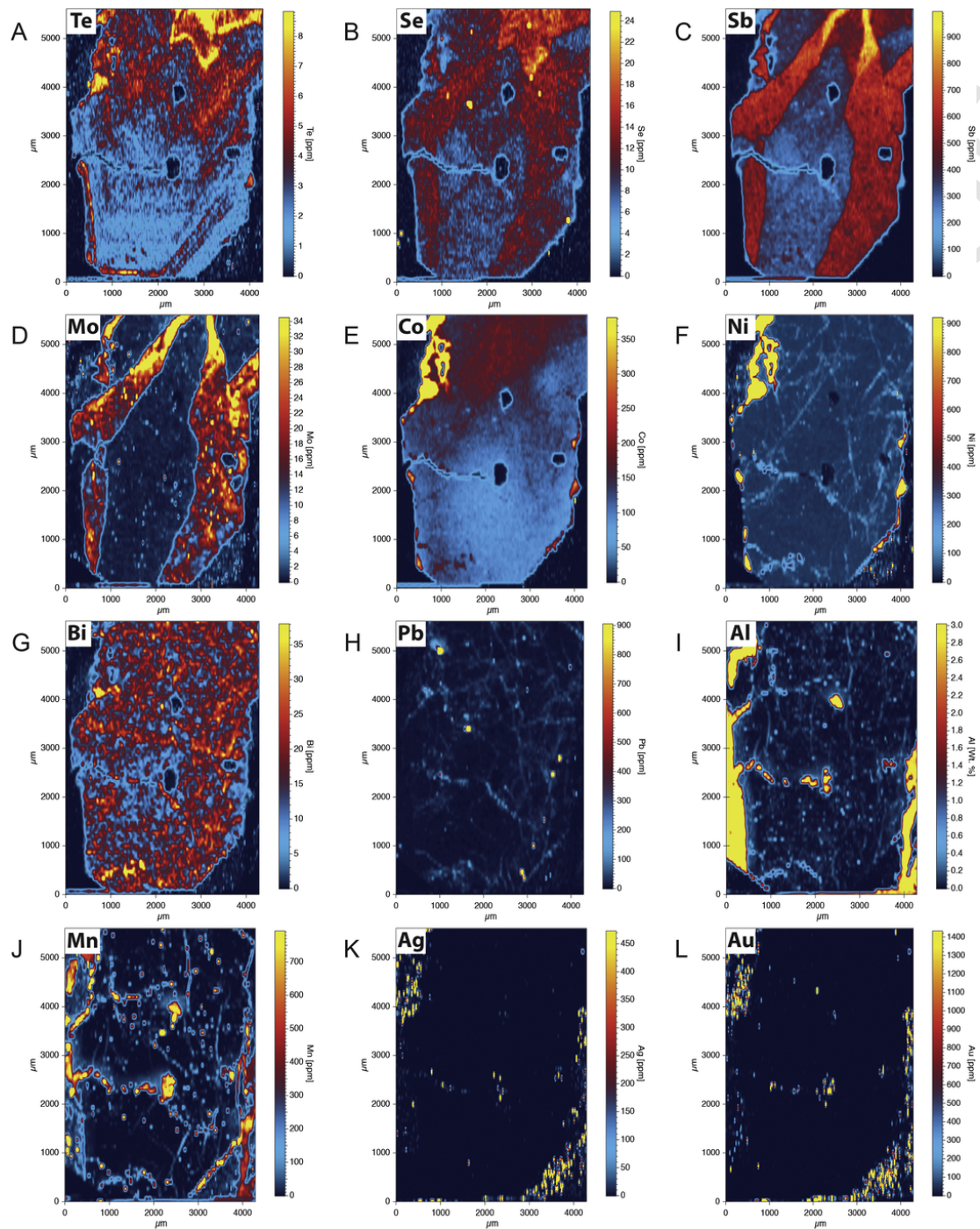


Fig. 7. LA-ICP-MS element maps showing the distribution of selected elements in part of an arsenopyrite grain (sample CH-K275) from the Cochrane Hill gold deposit, Meguma terrane (Nova Scotia). A) to L) Elemental maps plotted as concentrations scaled between the data median \pm 3 standard deviations (in ppm). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

tal distribution analyses, minor concentrations of this latter element association can be observed in the early fracture set, which suggests reactivation and mobilization of these elements out of these earlier fractures during continued or subsequent fluid circulation (e.g., Beaver Dam, Upper Seal Harbour, and Tangier). In addition, Ni may show a later stage of superimposition along the F1 and F2 fractures sets (cf. Beaver Dam, Touquoy; Figs. 3 and 6, respectively), which further suggests a post-crystallization history of elemental mobilization.

Regarding Au and Ag concentrations in arsenopyrite, we note the F1 and F2 fracture sets are mainly associated with micro-inclusions, as demonstrated by SEM observations (e.g., Beaver Dam, Touquoy, Cochrane Hill; Fig. 13), and higher concentrations of these elements than in the arsenopyrite core (>10 ppm), as is clearly evident in traverse analyses. Thus, the element maps and traverses, which offer complementary information, provide evidence for a late-stage enrichment of Au and Ag along both fracture sets: 1) F1 with a Bi-Pb-Cd-

Table 4

A compilation of the LA-ICP-MS maps and traverses produced from the various analyzed arsenopyrite grains and the location of the data presented in this paper.

Deposits	Maps	Traverses	Figures	Appendix
Beaver Dam	LC-12-16		Fig. 2	
The Ovens	OV-6		Fig. 3	
Dufferin	D4-A		Fig. 4	
Touquoy	TQ-1C		Fig. 5	
Cochrane Hill	CH-K275	CH-K092	Fig. 6	Fig. A
Upper Seal Harbour		DM-91-12_1 DM-91-12_2	Fig. 7	Fig. B
Tangier		TG-01	Fig. 8	
Caribou		CAR-88-47_1 CAR-88-47_2 CAR-88-47_1 CAR-88-7C_2 CAR-88-7C_3	Fig. 9	Fig. C Fig. D

In elemental assemblage; and 2) F2 with an Al-Ti-V-Cr-Mn elemental assemblage. This late-stage precious-metal endowment relates to the post-crystallization stage of the arsenopyrite grains and reflects a potential upgrading of Au and to a lesser extent Ag due to either continued and or subsequent fluid circulation. Similar findings have been noted and similarly interpreted by, for example, Wagner et al. (2007) and Lawley et al. (2015).

4.2. Elemental distributions and Au mineralizing events

The evaluation of original Au distribution in arsenopyrite (i.e., the primary gold event) and its potential upgrading can be assessed using binary element plots of Au versus Ag, which are further enhanced using color coding, as a proxy for concentration, for single elements or element associations (Gourcerol et al., 2018b). Distribution of the TSD data reveals the relationships between gold concentration (i.e., its tenor) and the relative timing of its sequestering into the host arsenopyrite. Based on this treatment of the TSD data, three dis-

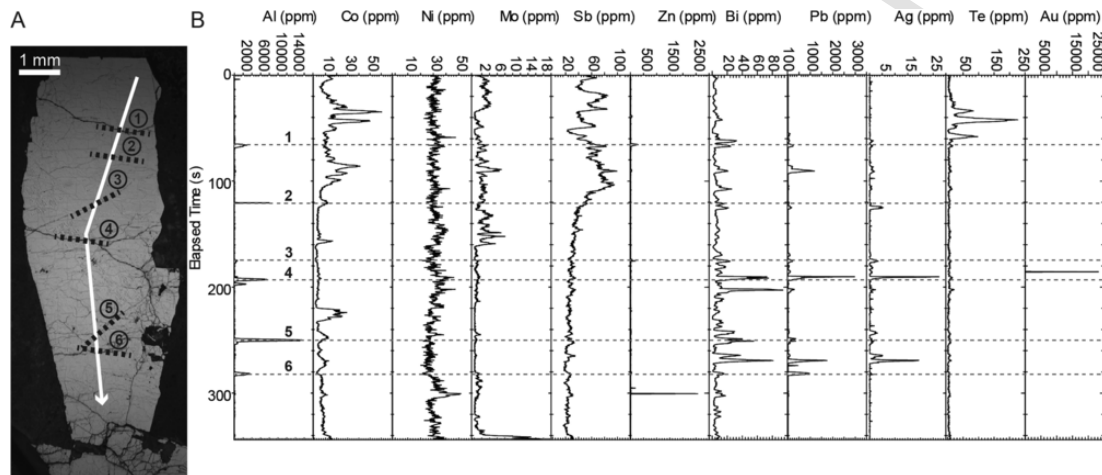


Fig. 8. Quantitative LA-ICP-MS elemental profiles (in ppm) acquired from a traverse across an arsenopyrite grain (sample DM-91-12-2) from the Upper Seal Harbour gold deposit, Meguma terrane (Nova Scotia). A) Reflected light photomicrograph of the analyzed arsenopyrite grain. The solid white line represents the traverse whereas the numbered (1 to 8) dashed black lines represent micro-fractures identified in reflected light. B) Concentration profiles for selected elements along the traverse shown in image A. Note that the dashed black lines refer to the micro-fractures identified in the image. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

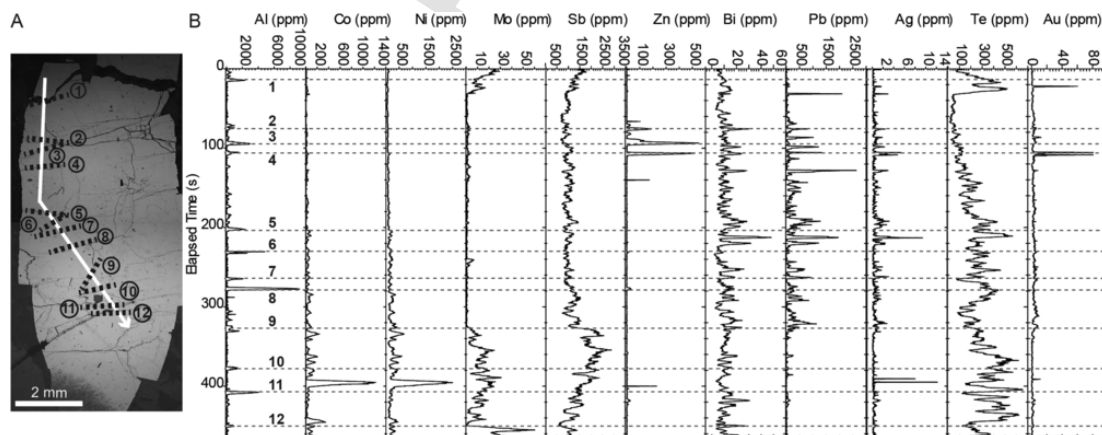


Fig. 9. Quantitative LA-ICP-MS elemental profiles (in ppm) acquired from a traverse across an arsenopyrite grain (sample Tangier) from the Tangier gold deposit, Meguma terrane (Nova Scotia). A) Reflected light photomicrograph of the analyzed arsenopyrite grain. The solid white line represents the traverse whereas the numbered (1 to 12) dashed black lines represent micro-fractures identified in reflected light. B) Concentration profiles for selected elements along the traverse shown in image A. Note that the dashed black lines refer to the micro-fractures identified in image A. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

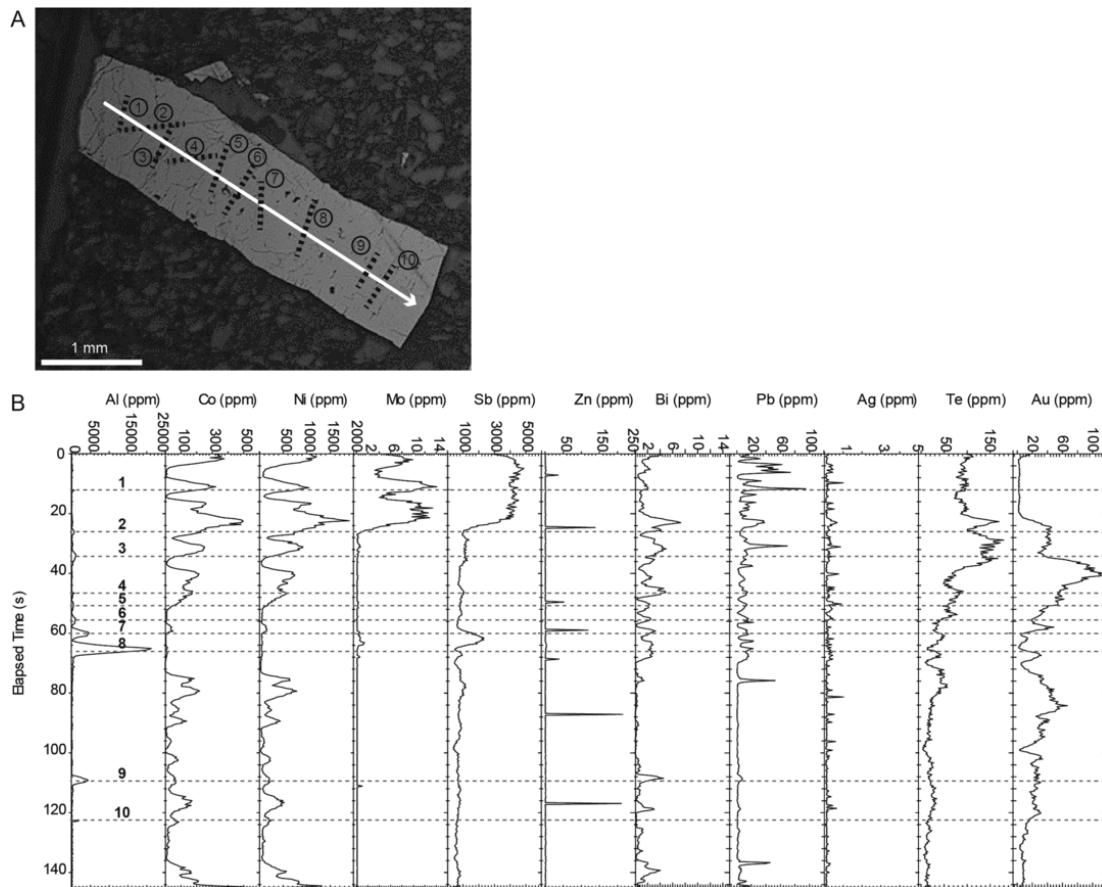


Fig. 10. Quantitative LA-ICP-MS elemental profiles (in ppm) acquired from a traverse across an arsenopyrite grain (sample CAR-88-7C_1) from the Caribou gold deposit, Meguma terrane (Nova Scotia). A) Reflected light photomicrograph of the analyzed arsenopyrite grain. The solid white line represents the traverse whereas the numbered (1 to 10) dashed black lines represent micro-fractures identified in reflected light. B) Concentration profiles for selected elements along the traverse shown in image A. Note that the dashed black lines refer to the micro-fractures identified in image A. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

tinct groupings of deposits are identified based on data distribution and precious-metal tenors.

The first group, illustrated by Beaver Dam and Touquoy deposits (Figs. 14 and 15, respectively), suggests a strong correlation of Au and Ag tenors (with a consistent 10:1 ratio). The data confirm that Au and Ag (both <10 ppm) were initially concentrated as invisible metals, that is, either structurally-bound or present as nanoparticles (Gourcerol et al., 2018b), during the primary growth of the arsenopyrite whereby Co and Ni are used as proxies for its early growth (Figs. 14A, B, 15A). A Mo, Sb and Se element assemblage, which represents a relatively late-stage primary feature in both deposits, shows a slight increase in Au and Ag to about 100 ppm (Figs. 14C, 15B, C). F1 fractures associated with the Bi-Pb-In-Cd element assemblage are associated with an increase of Ag and Se tenors (up to 1000 ppm; Fig. 14E, 15D), whereas an Al-Ti-V-Cr-Mn elemental assemblage (i.e., later F2 fractures for Beaver Dam deposit and the external rim for the Touquoy deposit) is associated with an increase in both Au and Ag tenors (up to 10 wt% for Au and 1 wt% for Ag) along with a 10:1 ratio and probable formation of gold micro-inclusions in the fracture set (Figs. 13E, F, 14F, 15E). Lastly, there is the presence, albeit minor, of Ag-poor micro-inclusions of gold in both samples (Fig. 14A, 15A).

A second group, which includes The Ovens and Dufferin deposits (Figs. 16 and 17, respectively), shows a different elemental distribution with the most notable difference compared to that for Beaver Dam and Touquoy being the absence of a well-defined and uniform Au:Ag relationship (i.e., the 10:1 ratio). There is however, at least

for the Dufferin dataset, the presence of Ag-poor micro-inclusions of gold, which are lacking at The Ovens (Figs. 16A, 17A). Nevertheless, as in the previous plots, common elemental associations and trends are seen in the data for these two deposits such as: 1) presence of primary Au and Ag in the arsenopyrite (both up to 10 ppm; Figs. 16A to C, 17A to C); 2) a Bi-Pb-In-Cd element assemblage, present either as inclusions or in fracture sets, that is associated with an increase of Ag and minor Se tenors (e.g., up to 1 wt% Ag for the Dufferin deposit) along with consistent Au values (up to 10 ppm). Note that for Dufferin there is a higher proportion of data defining this latter trend than for any other data set (Fig. 17D); and 3) a relatively late fracture set or late-stage rim overgrowth with an Al-Ti-V-Cr-Mn element association, which is slightly correlated with a higher Au tenor (up to 100 ppm for the Dufferin deposit (Fig. 17E) versus 10 ppm for The Ovens (Fig. 16F)).

Finally, data from the Cochrane Hill deposit (Fig. 18) defines a third group which differs from the previous datasets in regard to the absence of Ag-rich outliers and a poorly defined Au:Ag ratio, but it does show the presence of Ag-poor micro-inclusions of gold. The primary nature of Au and Ag is however still well defined by the data with up to 100 ppm of each. In addition, the data do not define the Bi-Pb-In-Cd element association with Ag enrichment, as seen in the other deposits, but does show the Al-Ti-V-Cr-Mn element association related to minor (e.g., Beaver Dam and Touquoy deposits) Au and Ag enrichment (Fig. 18D; up to 1000 ppm).

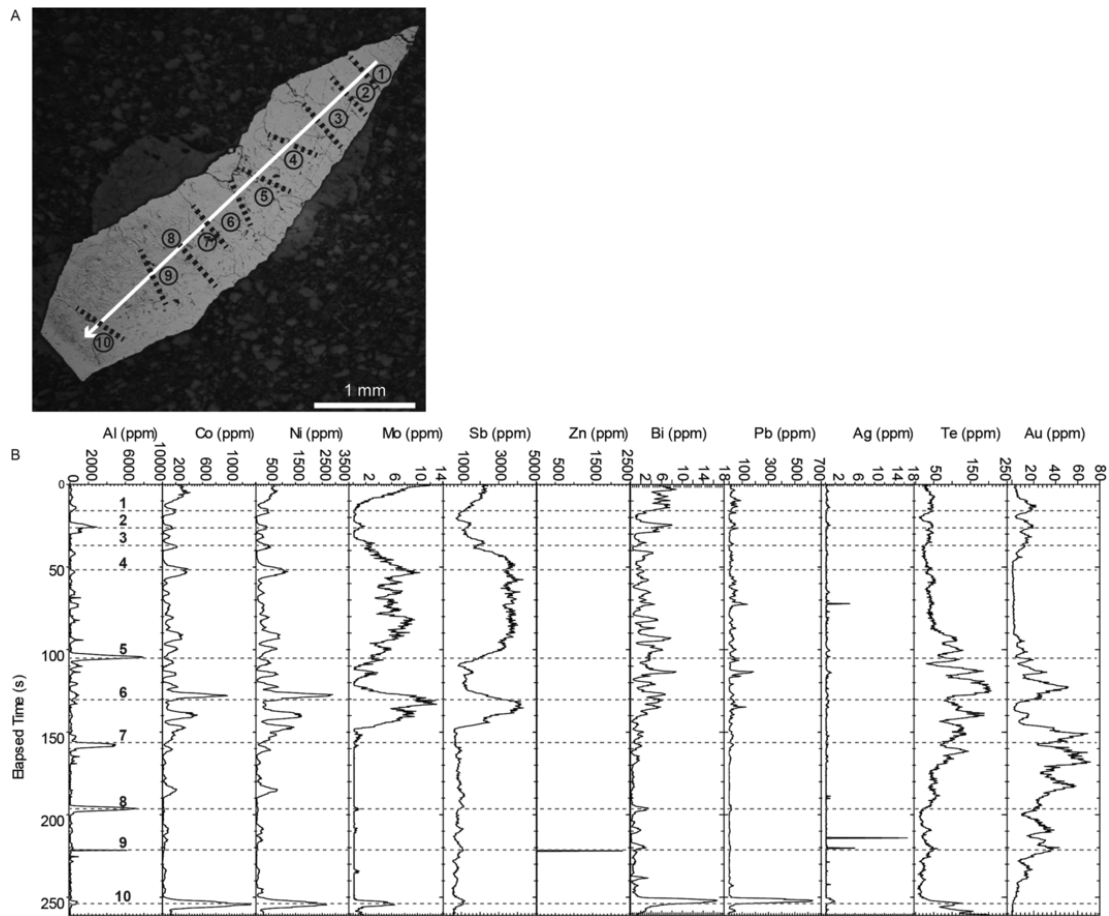


Fig. 11. Quantitative LA-ICP-MS elemental profiles (in ppm) acquired from a traverse across an arsenopyrite grain (sample CAR-88-7C_2) from the Caribou deposit, Meguma terrane (Nova Scotia). A) Reflected light photomicrograph of the analyzed arsenopyrite grain. The solid white line represents the traverse whereas the numbered (1 to 10) dashed black lines represent micro-fractures identified in reflected light. B) Concentration profiles for selected elements along the traverse shown in image A. Note that the dashed black lines refer to the micro-fractures identified in image A.

Based on the observations for the various binary element plots discussed above, at least two distinct gold events are identified: 1) the first is associated with primary sulfide growth and reflects a continued Au input, at times increasing until the latest stage of arsenopyrite growth (i.e., coincident with Mo-Se-Sb endowments); and 2) a second event, which varies in its gold tenor and is similar (Beaver Dam and Touquoy) or dissimilar (other deposits) regarding its Au:Ag ratio and is associated with an Al-Ti-V-Cr-Mn elemental assemblage. This latter association reflects upgrading of the tenor of Au and is seen in all the deposits along with a good correlation between primary and late-stage secondary element associations (e.g., Figs. 14H, 15G, H, 16G, H). Importantly, the presence of the Ag-, Au-Ag- and Au-rich outliers (Figs. 14 to 18), which reflect micro-inclusions, may be related to post-crystallization events and/or a continuation of the primary mineralizing system. Moreover, the change in the Au:Ag ratio of the former population may be attributed to a zone refining process. As for the Bi-Pb-Cd-In-Ag assemblage, which is Au poor, this is attributed to remobilization (Figs. 15H, 16G, 17G, 18F), hence the association with elevated Al-Ti-V-Cr-Mn, or subsequent upgrading (Fig. 14G) of earlier Au tenors as confirmed by Au micro-inclusions along the F1 fracture set in the Beaver Dam deposit.

Importantly, strong to good correlation between Bi-Pb-Cd-In and Al-Ti-V-Cr-Mn element associations with respect to Au (Figs. 15G, H, 16G, 17G) suggests the exploiting of fracture sets, either reactivated or newly formed, during subsequent fluid circulation, as sug-

gested in the previous section. An apparent anomaly here is the apparent lack of Au enrichment (i.e., maximum at about 10ppm) for The Ovens sample compared to the other analyzed samples despite the evidence of this later fluid event.

Quantitative element distribution data from traverses, rather than elemental maps, were acquired for samples from the Upper Seal Harbour, Tangier and Caribou deposits. These data show elemental distributions similar to those seen in the Beaver Dam and Touquoy deposits (Table 4; Figs. 8 to 11, Figs. B to E), such as primary Au enrichment, Au upgrading, the Bi-Pb-Cd-In-Ag association, and Au enrichment correlated with an Al-Ti-V-Cr-Mn. We also note that the presence of spikes in elements such as Te-Bi-Pb, Zn and Te (Upper Seal Harbour; Fig. 8), Pb, Pb—Ag, and Zn (Tangier; Fig. 9), and Pb, Zn, and Ag (Caribou; Figs. 10, 11) likely indicate the presence of micro-inclusions. For these samples, the Au versus Ag plots for arsenopyrite (Fig. 19) reveal similarities to the three types of plots discussed previously in terms of either a well-defined Au:Ag ratio (e.g., Cochrane Hill data; Fig. 19A) versus the lack thereof for Tangier, Upper Seal Harbour and Caribou samples (Fig. 19B, C, and D, respectively). In addition, there is a large range in Au enrichment with the Cochrane Hill samples showing the highest values (10,000ppm, Fig. 19A), whereas the Caribou data show the highest values for inferred primary Au enrichment with up to 100ppm Au (Fig. 19D) versus <10ppm for Tangier and Upper Seal Harbour (Fig. 19B and C, respectively).

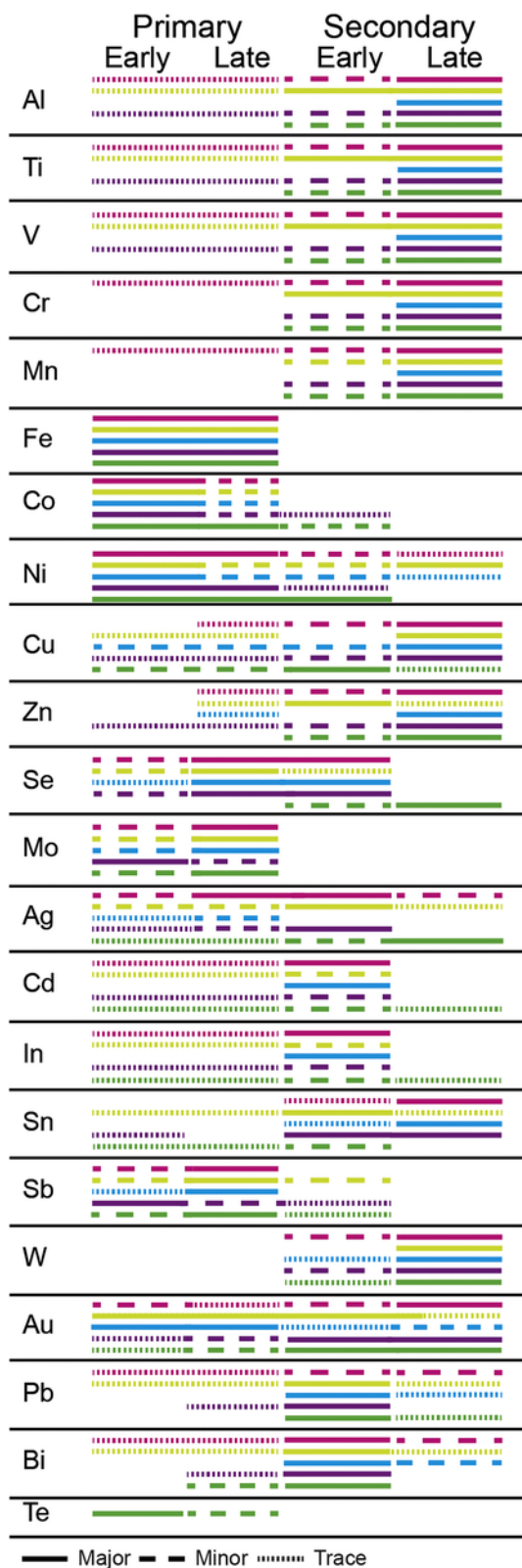


Fig. 12. Compilation of the inferred elemental paragenesis for the Meguma gold deposits studied based on the integration of reflected light petrographic studies and quantitative LA-ICP-MS element distribution maps produced for arsenopyrite grains. The color coding represents the Beaver Dam (pink), The Ovens (yellow), Dufferin (blue), Touquoy (purple) and Cochrane Hill (green) deposits. Note that, in general, there is

a broadly similar elemental paragenesis for each of the deposits studied. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

5. Discussion

In this study, arsenopyrite samples from several gold deposit sites across the Meguma terrane of Nova Scotia were studied in order to constrain several important aspects of gold metallogeny in general and metasedimentary-hosted deposits specifically. The application of data derived from element distribution maps, element profiles based on traverses, and a variety of multi-element plots using TSD data derived from continuous LA-ICP-MS analysis provides the basis for addressing several important issues that pertain to the nature of the gold mineralization: 1) elemental paragenesis and element associations; 2) nature of gold and its possible remobilization or upgrading; 3) significance of the Au:Ag ratio; 4) possible wall-rock influence inferred from the whole-rock, isotopic and mineralogical data bases; and 5) the implications of the data for the source of gold and gold mineralizing processes in a gold metallogenic province.

5.1. Elemental paragenesis and associations in the Meguma deposits

The integration of petrographic observations and SEM-EDS imaging with the elemental maps and the traverses together with the multi-element plots provide the basis for establishing elemental paragenesis. Comparison of the paragenesis of the eight studied deposits indicates very similar features and suggests their common evolution and thus strongly argues for a similar fluid history (Figs. 12, 20).

The arsenopyrite grains studied reveal a protracted and complex growth history with two very distinct early mineralizing episodes (i.e., primary events). An initial stage of primary growth is reflected by oscillatory changes in the Co-Ni-(Te) contents during the initial stage of arsenopyrite nucleation and subsequent development. This type of zonation has been reported in other studies of arsenopyrite (e.g., Cathelineau et al., 1989; Morey et al., 2008; Cook et al., 2013) and has been attributed to the influence of the primary hydrothermal fluid, rather than a later metamorphic fluid (e.g., Shore and Fowler, 1996), with the alternating zones reflecting fluctuations in fluid chemistry coupled with a low diffusion rate in the crystal, and hence preservation of the zonation, and possibly varying temperature. The second mineralizing episode, which has a Mo-Se-Sb element association, is inferred paragenetically to be later in the initial growth history (Fig. 12); it locally follows the growth zoning but mainly shows a non-uniform distribution which may suggest precipitation under near-equilibrium conditions with uniform mixing (Krinov, 2008). Importantly, preservation of this initial oscillatory zoning in the arsenopyrite suggests that diffusional equilibration did not affect these domains. In addition to the above mineralizing events, two later stage fluid events, as illustrated by fracture-controlled element associations (i.e., F1, F2), affected the arsenopyrite grains: 1) early F1 fractures with a Bi-Pb-Cd-In-Ag element association; and 2) later F2 fractures, which crosscut F1, which have an Al-Ti-V-Cr-Mn element association. The early Bi-Pb-Cd-In-Ag bearing fluid is associated with the upgrading of Ag tenors along with Se, and formation of galena micro-inclusions. The later Al-Ti-V-Cr-Mn elemental association is related to Au upgrading, both where Au:Ag is 10:1 and Ag-poor, and involved the local formation of visible gold grains in these fractures along with rutile, chlorite and ilmenite.

Thus, the integration of the elemental maps with ore petrology and SEM-EDS observations provide evidence for ore-system longevity and complexity, in this case across an entire metallo-

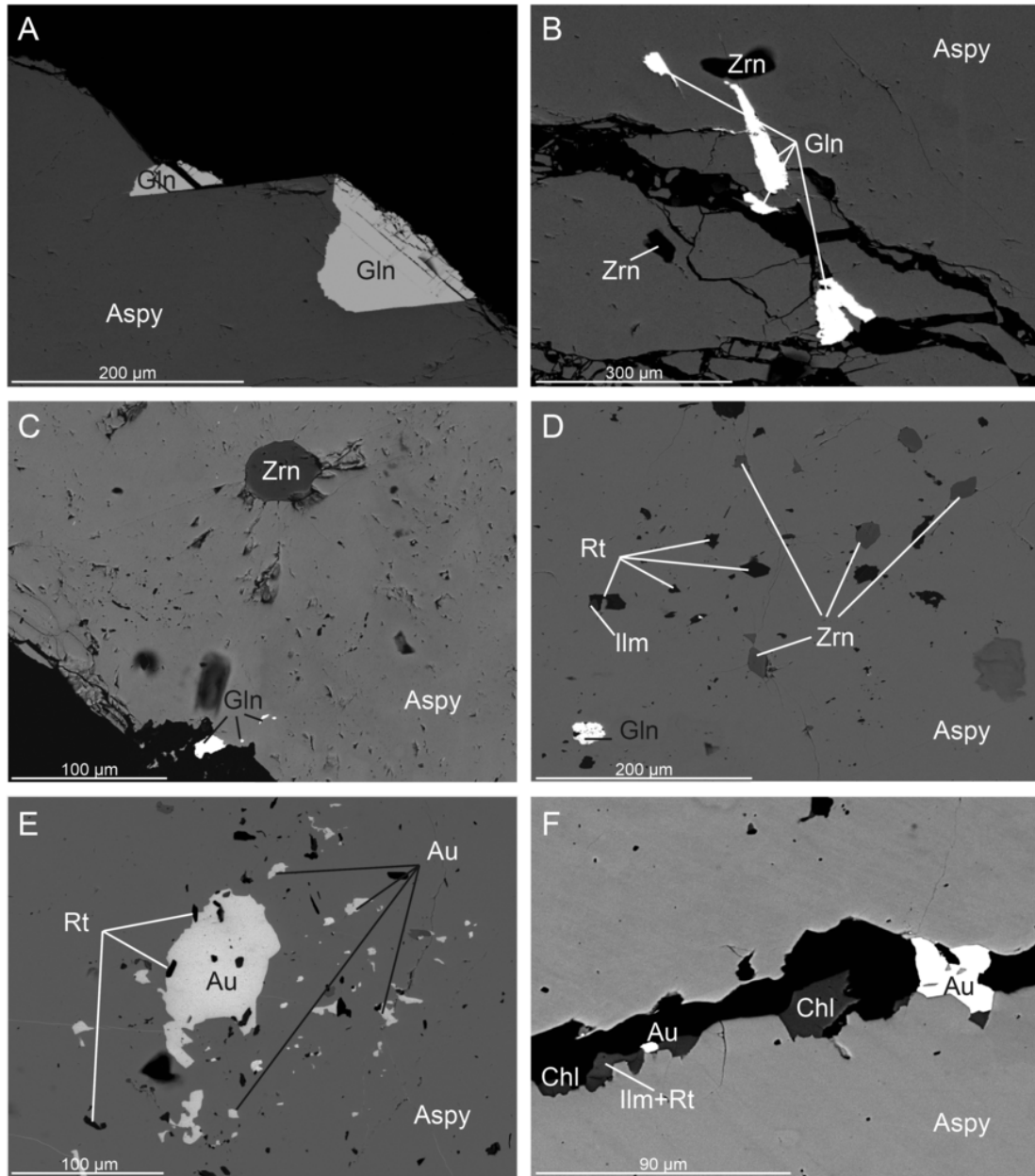


Fig. 13. Scanning electron microscope backscatter images of arsenopyrite grains from selected deposits in the Meguma terrane (Nova Scotia). A) Subhedral galena grains forming an external rim on arsenopyrite (Dufferin deposit). B) Galena filling fractures in addition to zircon inclusions closely associated with a micro-fracture set (Dufferin deposit). C) Small subhedral zircon grain in addition to inclusions of galena in the host arsenopyrite (Touquoy deposit). Also note the extensive pitting of the arsenopyrite grain, a texture which is often associated with coupled dissolution precipitation in silicates. D) Zircon, rutile, ilmenite and galena inclusions in arsenopyrite (Touquoy deposit). E) Gold and rutile inclusions, some of which are closely associated with micro-fracture sets, in arsenopyrite. Note that rutile appears either to overprint gold inclusions suggesting a late crystallization (Touquoy deposit) or reflect hydrothermal processes such as an ilmenite breakdown reaction. F) Gold, chlorite and ilmenite-rutile in a late stage fracture cutting arsenopyrite (Beaver Dam deposit). Abbreviation: Aspy: arsenopyrite, Au: gold, Chl: chlorite, Gln: galena, Ilm: ilmenite, Rt: rutile, Zrn: zircon. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

genic terrane based on our sampling (regarding geochemistry of the arsenopyrite grains). The establishment of such elemental associations and paragenesis has not previously been possible and, as shown here, provides the means to demonstrate a commonality to mineralizing processes among similar ore deposit settings within a given metallogenic terrane. The previous use of bulk-rock chemistry via traditional whole-rock analysis is clearly flawed in terms of establishing elemental associations, as illustrated by the complexities revealed in the elemental maps shown above.

5.2. Nature and implications of gold mineralizing events

Elemental paragenesis established from both the element distribution maps and traverses and the derived Au versus Ag plots using the TSD data suggest two distinct gold events: 1) the first is evident from the presence of invisible gold or inferred nanoparticles; and 2) the second is seen as an association of Au enrichment, sometimes as visible particles, with an Al-Ti-V-Cr-Mn elemental association concentrated along fractures (i.e., the F2 set).

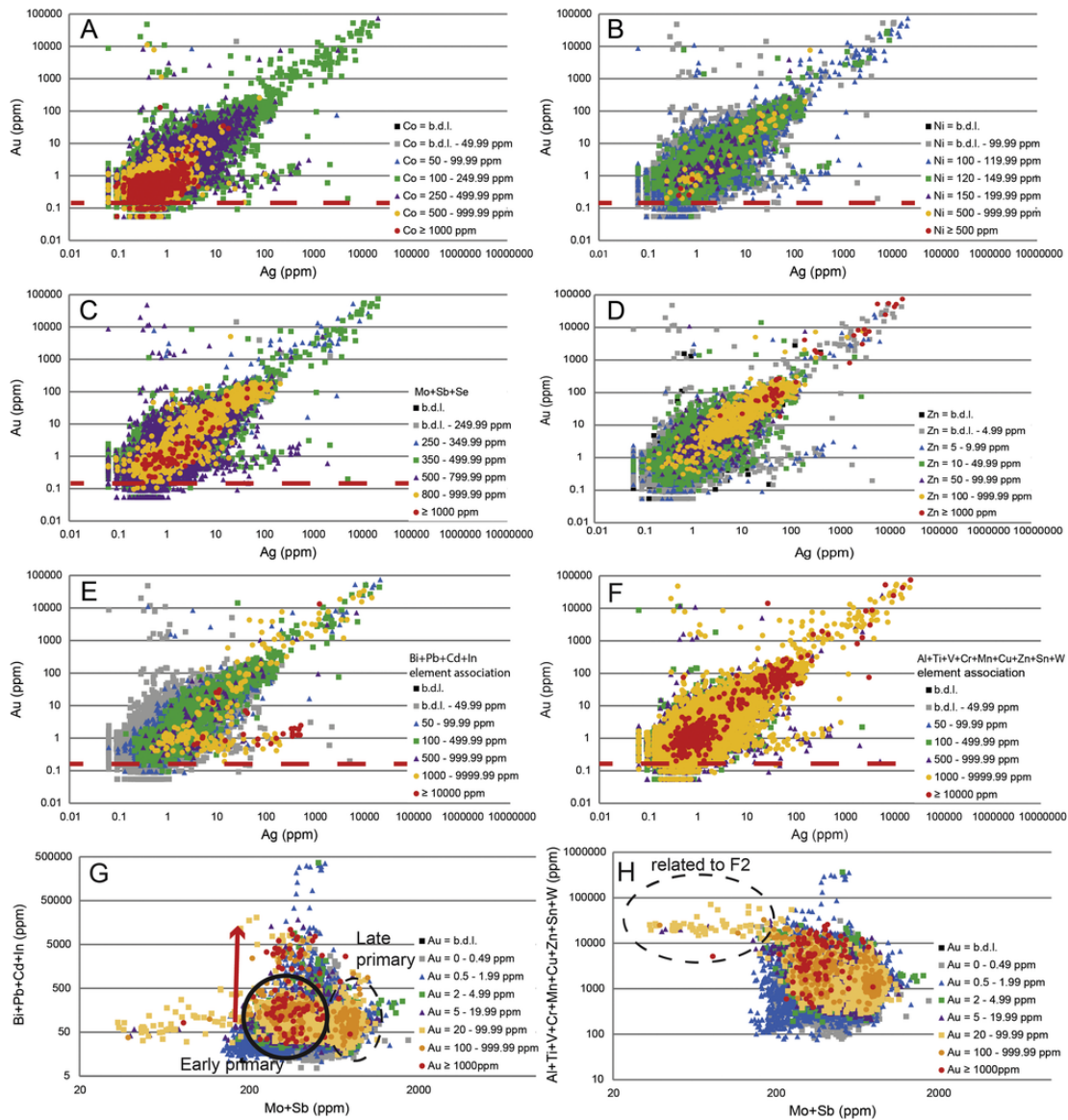


Fig. 14. Binary element plots (in ppm) based on the TSD for the mapped arsenopyrite grain from the Beaver Dam gold deposit (Nova Scotia). Note that 39,800 data points are summarized in each of these plots and the points are color coded (i.e., cold to hot) in order to reflect the enrichment in either an element or elemental association. A to F) Binary plots (in ppm) of Au versus Ag. G) F1 element association (Bi-Pb-Cd-In) versus Mo—Sb concentrations. H) F2 element association (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) versus Mo—Sb concentrations. Note that red arrow is interpreted to suggest a subsequent remobilization and upgrading of Au tenor. The dashed red horizontal line represents the detection limit for Au. Few plots from this deposit are also presented in Gourcerol et al., 2018b. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

The first Au event coincided with the growth of arsenopyrite, which was monitored by using Ni and Co as proxies of primary growth. In this early stage of arsenopyrite, gold is inferred to have been deposited either as invisible gold (e.g., Reich et al., 2005) or as nanoparticles (Fougerouse et al., 2016b) within oscillatory Co—Ni zoned arsenopyrite; the traverse mode analyses provide a good representation of this oscillatory zoning via the coupling of Co and Ni (e.g., Figs. 10 and 11). Gold content in such zones is mostly present at values <10 ppm, but can in some cases (i.e., Caribou, Beaver Dam, Touquoy) extend up to 100 ppm (Figs. 10, 14, and 15, respectively). This early event reflects a continued and uniform proportion of Au, although at times increasing, until the latest stage of sulfide crystallization, as marked paragenetically by enrichment of Mo—Se—Sb and confirmed by the Au versus Ag plots. This type of mineralization is common in a variety of hydrothermal ore systems, such as metasedi-

ment-hosted orogenic (e.g., Bendigo-Ballarut; Thomas et al., 2011), Carlin type (e.g., Large et al., 2009) and even VMS (e.g., Mercier-Langevin et al., 2015) deposits.

The second Au event shows higher gold contents, but is variable in its tenor (Figs. 20, 21). Traverse analyses provide evidence for coupled dissolution-precipitation (CDP) associated with the late fracture set, as is illustrated in fractures 7, 8, 9 of Fig. 11. Thus, the CDP process is related to dissolution of the host and commensurate release of invisible gold from the arsenopyrite structure and its subsequent precipitation as visible gold along fractures. This latter process essentially represents a zone refining processes, as documented by others (e.g., Wagner et al., 2007; Fougerouse et al., 2016a; Wu et al., 2019). In combination, the diffusional process developed by Reich et al. (2006) may have conducted to precipitation of gold nanoparti-

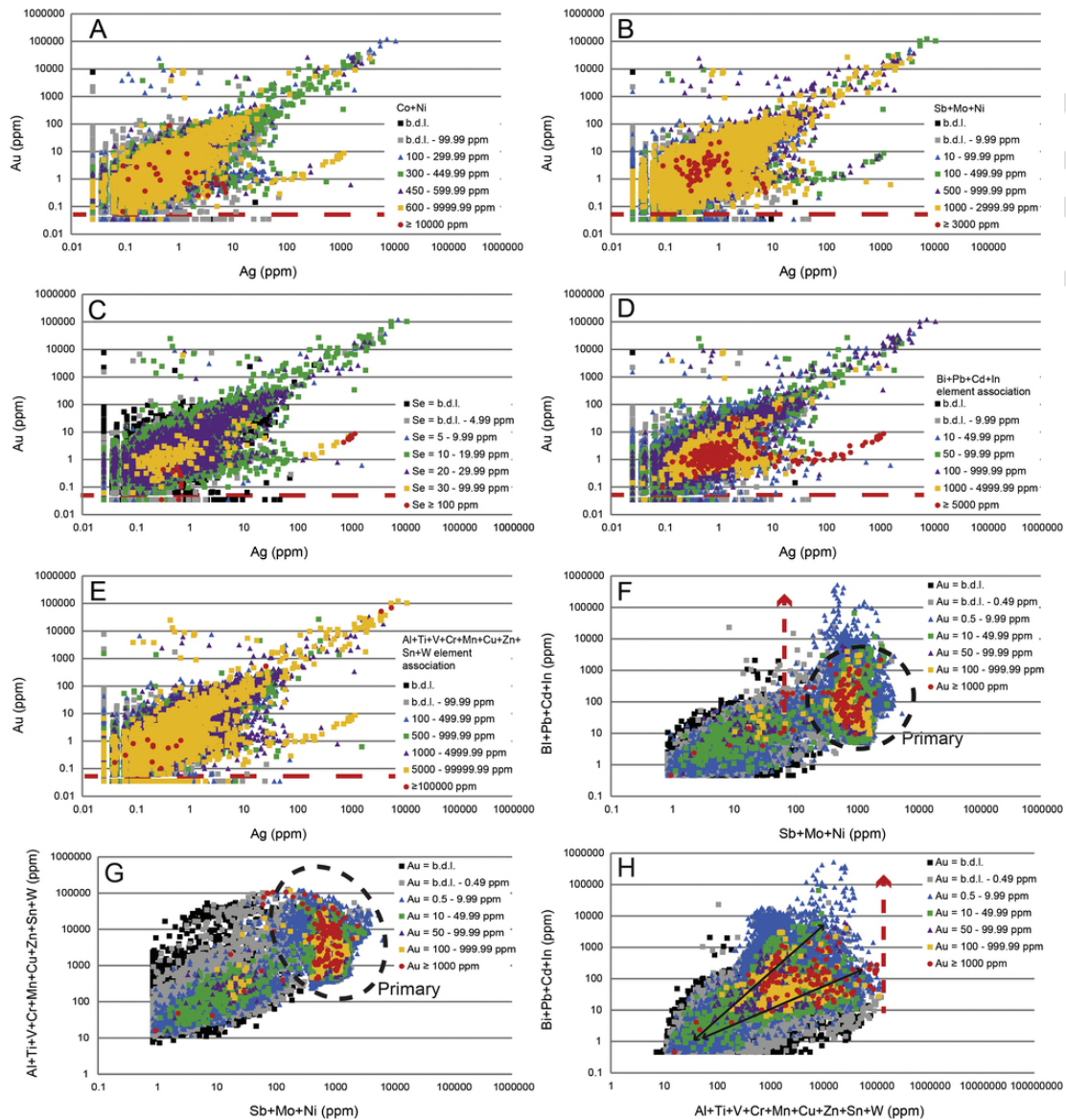


Fig. 15. Binary element plots (in ppm) based on the TSD for the mapped arsenopyrite grain from the Touquoy gold deposit (Nova Scotia). Note that 58,820 data points are summarized in each of these plots and the points are color coded (i.e., cold to hot) in order to reflect the enrichment in either an element or elemental association. A) to E) Binary plots (in ppm) of Au versus Ag. F) F1 element association (Bi-Pb-Cd-In) versus Ni-Mo-Sb concentrations. G) F2 element association (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) versus Sb-Mo-Ni concentrations. H) F1 (Bi-Pb-Cd-In) versus F2 element associations (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) concentrations. Note that dashed red lines (13F and 13H) highlight strong correlation between element associations. The dashed red horizontal line represents the detection limit for Au. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

cles within the arsenopyrite matrix during the early stage of the second metamorphic thermal event prior to remobilization.

Two additional points are noted in the context of these two gold events. Firstly, for both of the events identified, the Au:Ag ratio is uniform at 10:1 in some of the deposits, which is a common value in orogenic gold deposits and must therefore be reflective of a geologically meaningful and relevant process (e.g., Groves et al., 1998; Dubé and Gosselin, 2007; Hastie et al., 2019); this aspect is discussed in more detail below. Secondly, elemental paragenesis and enrichment are very similar in the eight studied deposits (Figs. 12 and 20, respectively) and suggest similar processes and likely similar fluid reservoirs for all of the Meguma gold deposits. We also note in regards to the distribution of Au abundance, based on the tabulation of approximately 300,000 TSD data (Fig. 21), that there are clearly two dis-

tinct concentrations of Au in the arsenopyrite analyzed (i.e., invisible and micro-inclusions).

Independently, the Au versus Ag plots show Au-rich versus Ag-poor values (i.e., Figs. 14, 15, 17, 18, 19) which reflect presence of gold micro-inclusions along fracture sets (Fig. 13) or disseminated. As alluded to before, this gold mineralization represents a zone refining process and accompanies mobilization of low-grade invisible gold to form free gold grains.

In addition to the two gold events noted above, a third event is identified based on the Bi-Pb-Cd-In-Ag elemental association in the Au—Ag plots and is related to fluid migration along the F1 fracture set. This association has been recognized at most deposits, thus indicating it is common across the Meguma terrane. Importantly, this latter event predates the second gold event (i.e., formation of micro-in-

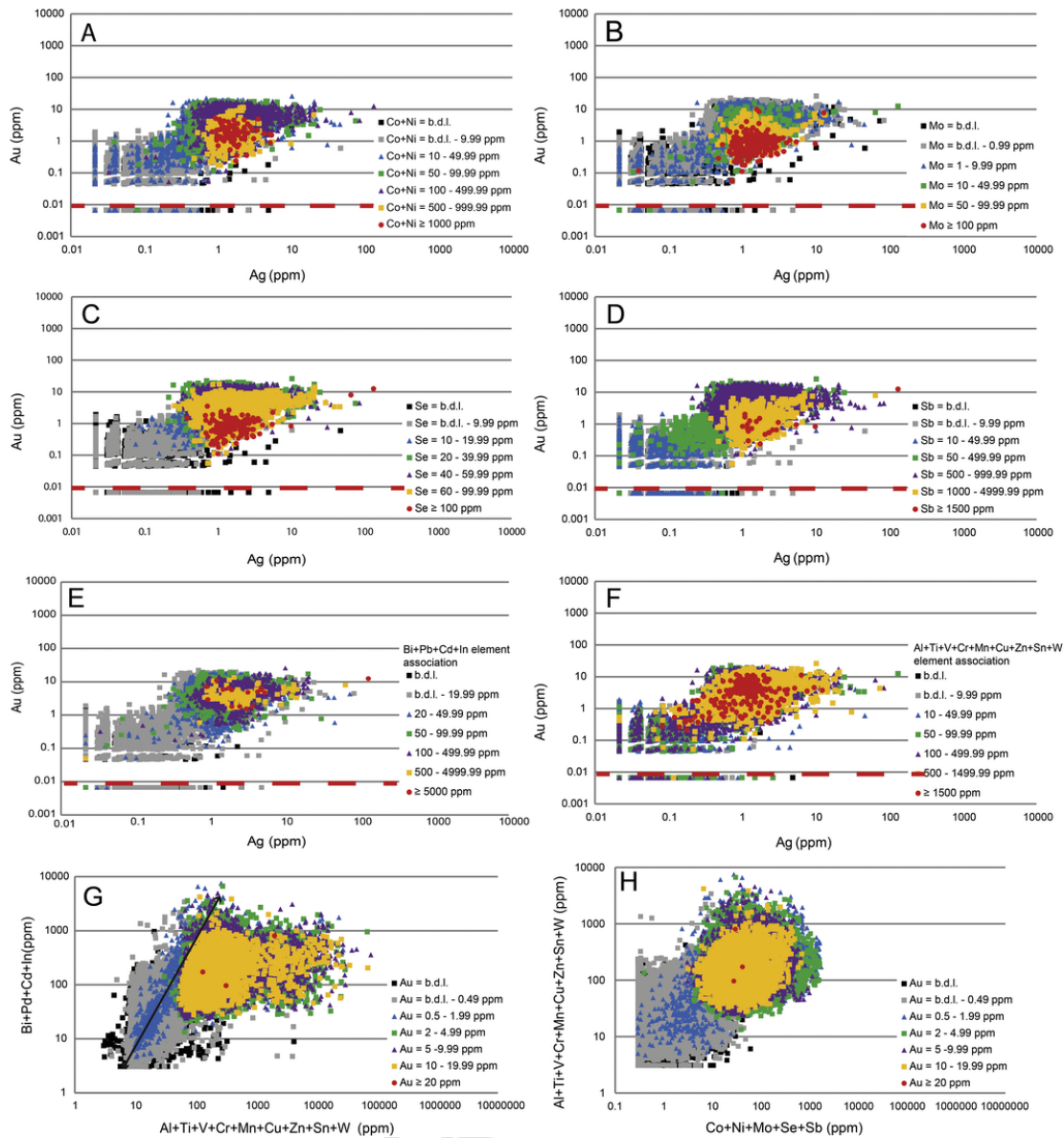


Fig. 16. Binary element plots (in ppm) based on the TSD for the mapped arsenopyrite grain from The Ovens gold deposit (Nova Scotia). Note that 42,810 data points are summarized in each of these plots and the points are color coded (i.e., cold to hot) in order to reflect the enrichment in either an element or elemental association. A) to F) Binary plots (in ppm) of Au versus Ag. G) F1 (Bi-Pb-Cd-In) versus F2 element associations (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) concentrations. H) F2 element association (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) versus Co-Ni-Mo-Se-Sb concentrations. Note that the black line (14G) highlights a strong correlation between the identified element associations. The dashed red horizontal line represents the detection limit for Au. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

clusions) and potentially relates to the upgrading of Ag tenors through CDP processes as well (mostly 1000 ppm and up to 10,000 ppm in Dufferin deposit) as fracture fillings. As noted previously, the enrichment of Se in some cases may suggest precipitation of silver selenide (i.e., naumannite), but this remains to be confirmed with future work involving imaging at the nanoscale. Importantly, this event appears not to show Au input and may have had a downgrading effect on the primary gold mineralization. Note that this event shows highly variable Au:Ag ratios (i.e., 1:10, 1:100, 1:1000, 1:10000).

5.3. Implications of Au:Ag ratios

As discussed previously, the Au versus Ag plots from the studied deposits indicate that, where best defined by the abundance of

data, the two gold events (i.e., primary versus secondary) show similar Au:Ag ratios of 10:1. This observation is consistent with the fact that both gold events likely relate to similar metamorphic processes and similar fluid compositions (i.e., an aqueous carbonic composition, $X_{\text{CO}_2} = 0.10\text{--}0.2$ and 5–10 wt% equiv. NaCl), as noted discussed previously (e.g., see review in Kontak and Horne (2010)). Thus, the consistent Au:Ag ratio suggests a similar source, transporting fluid, and chemical conditions for primary gold and similarly for the Au upgrading event which preserves the initial Au:Ag ratio during Au remobilization and its re-precipitation. Moreover, as demonstrated by Cole and Drummond (1986), this ratio can be used as a thermodynamic proxy. Thus, a low temperature ($< 250^\circ\text{C}$), low Cl^- content, moderate to high pH (≥ 5) and high $\Sigma\text{H}_2\text{S}/\Sigma\text{SO}_4$ ratio ($< 10^5$) are suggested for the Au-bearing fluids and in addition suggests that

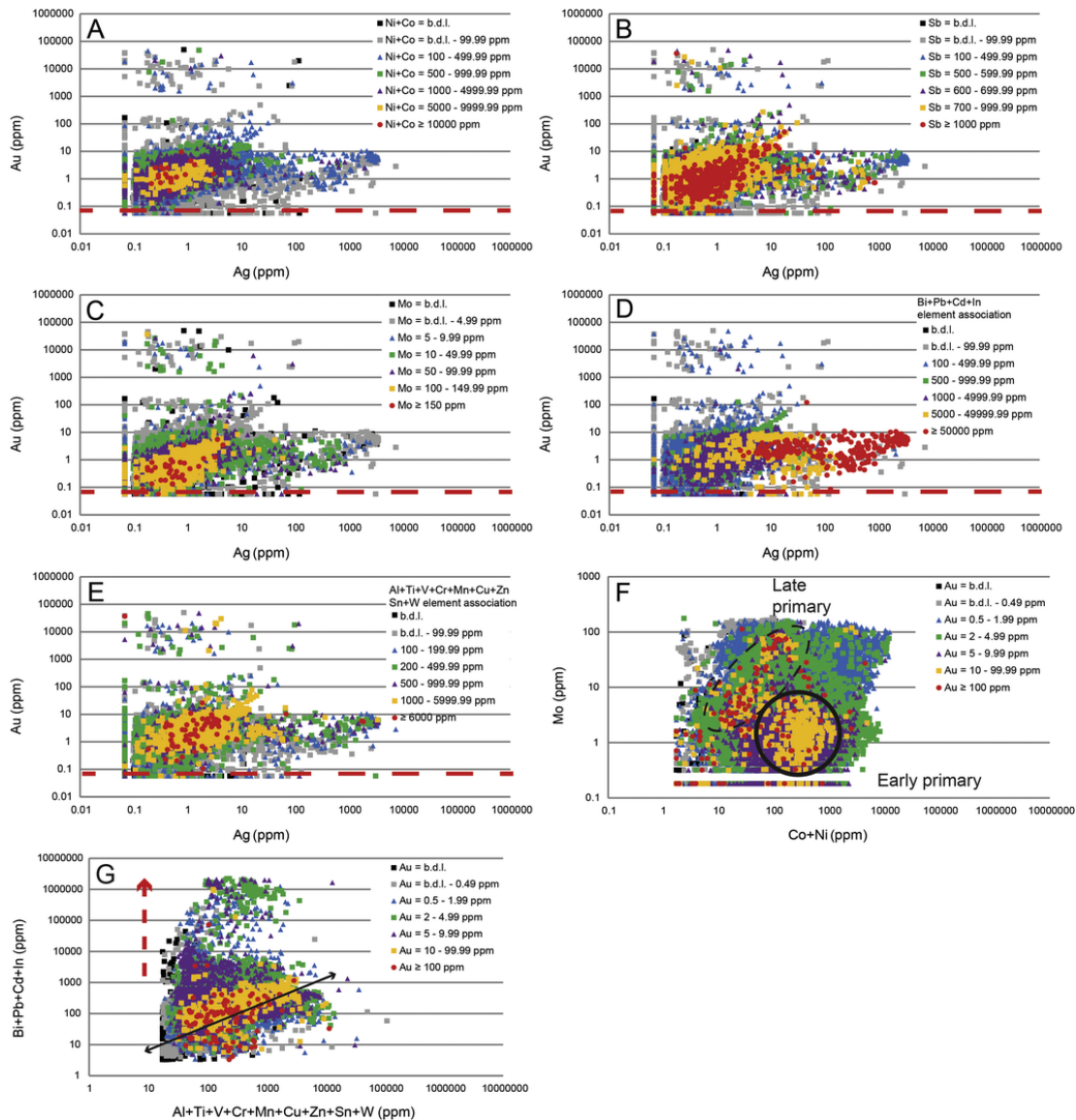


Fig. 17. Binary element plots (in ppm) based on the TSD for the mapped arsenopyrite grain from the Dufferin gold deposit (Nova Scotia). Note that 35,325 data points are summarized in each of these plots and the points are color coded (i.e., cold to hot) in order to reflect the enrichment in either an element or elemental association. A) to E) Binary plots (in ppm) of Au versus Ag. F) Mo versus Co—Ni concentrations. G) F1 (Bi-Pb-Cd-In) versus F2 element associations (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) concentrations. Note that dashed red line (15G) is inferred to suggest a subsequent remobilization and upgrading of Au tenor in the F1 element association, whereas the black line (15G) highlights a strong correlation between element associations. The dashed red horizontal line represents the detection limit for Au. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

gold was likely transported as an $\text{Au}(\text{HS})_2^-$ complex. Whereas the latter temperature is generally slightly lower than inferred from most of the fluid inclusion studies of the Meguma deposits (i.e., 300–350 °C; see above), the geochemical and isotopic features of the deposits are commensurate with the other parameters.

Independently, the fluid responsible for the Bi-Pb-Cd-In-Ag element assemblage shows variable Au:Ag ratios that range from 1:10 to 1:10000. This Ag event is not associated with new introduction of Au in the system, but only with remobilization of Au of the first gold event.

Importantly, it should be noted that the Au:Ag ratios observed could potentially also be influenced by systematic resampling of previously ablated material in the form of ejecta or condensate on the sample surface. However, each dataset includes a minimum of 28,000 data points (i.e., Cochrane Hill dataset) and each deposit shows an in-

ternally consistent trend and ratio values. Thus, the demonstrated consistency of the Au:Ag ratios of the analyzed materials is considered to validate the interpretations presented herein.

5.4. Influence of host rock stratigraphy and other reservoirs

The influence of the immediate wall rock to mineralized veins and, at a broader scale, the host stratigraphy on the geochemical signature of the mineralizing fluids has previously been discussed for the Meguma gold deposits based on extensive analysis of $\delta^{34}\text{S}$ for sulfides, $\delta^{13}\text{C}$ and $^{87}\text{Sr}/^{86}\text{Sr}$ for carbonates, $\delta^{18}\text{O}$ for quartz and in-situ LA-ICP-MS trace- element chemistry of carbonates (Kontak and Smith, 1989; Sangster, 1992; Kontak and Kerrich, 1995, 1997; Kontak and Jackson, 1999; Kontak et al., 2011). The $\delta^{34}\text{S}$ values of sulfides from veins, wall rock and regional stratigraphy show a simi-

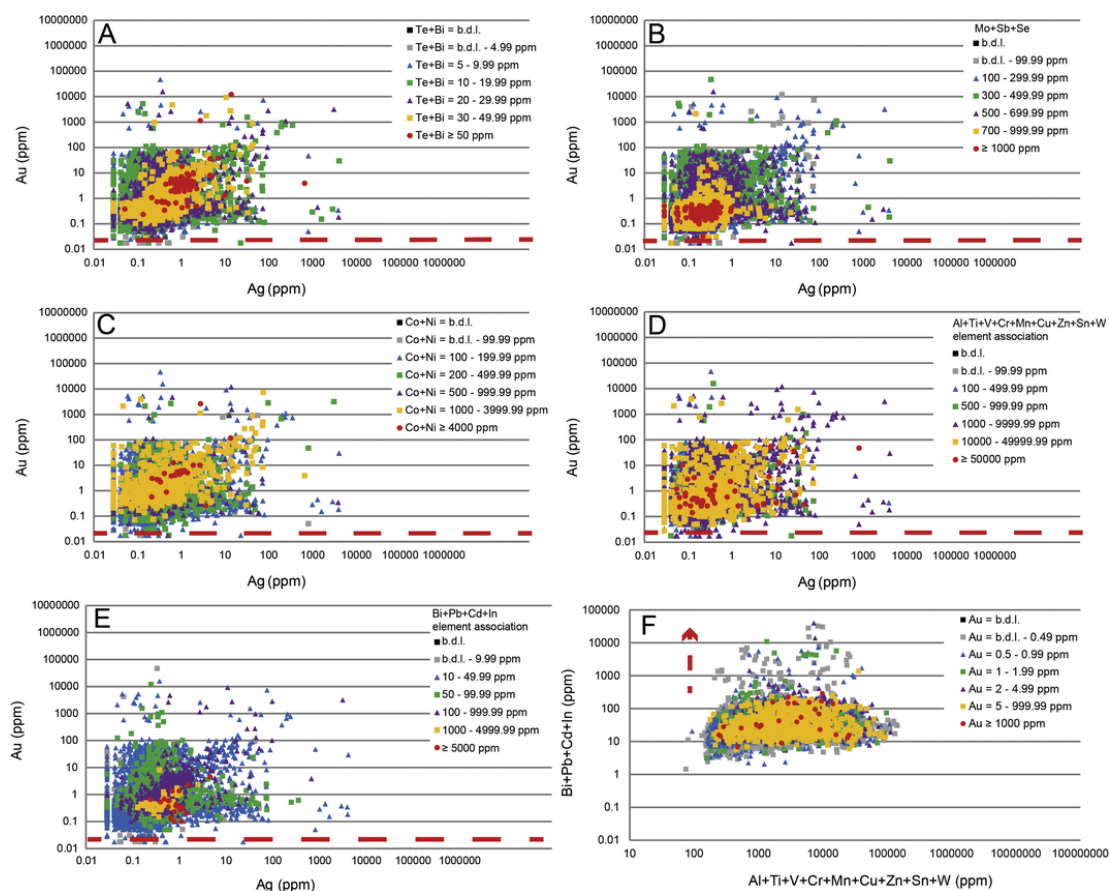


Fig. 18. Binary element plots (in ppm) based on the TSD for the mapped arsenopyrite grain from the Cochrane Hill gold deposit (Nova Scotia). Note that 28,660 data points are summarized in each of these plots and the points are color coded (i.e., cold to hot) in order to reflect the enrichment in either an element or elemental association. A) to E) Binary plots (in ppm) of Au versus Ag. F) F1 (Bi-Pb-Cd-In) versus F2 element associations (Al-Ti-V-Cr-Mn-Cu-Zn-Sn-W) concentrations. Note that dashed red line (16F) is inferred to suggest a subsequent remobilization of Au tenor along F1 and associated with these elements. Note that the horizontal dashed line represents the detection limit for Au. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

lar systematic increase from +10 to +30‰ and suggest sulfur is locally sourced (i.e., Meguma stratigraphy). In addition, very low $\delta^{13}\text{C}$ values (to ca. -25‰) are suggestive of a biogenic carbon source for vein carbonate and, given that this overlaps the signatures for graphite and diagenetic carbonate in the host rocks, this signature also provides a strong case favoring extensive influence of the host stratigraphy. Further evidence of a wall-rock influence is provided from the highly radiogenic Sr signal recorded in some vein carbonates, in addition to an overall upwards increase in $\delta^{18}\text{O}$ for vein quartz which was suggested to reflect an extensive fluid:rock interaction based on modelling (Kontak et al., 2011). Lastly, the trace-element signature of vein carbonate has been interpreted to reflect a variety of influential parameters (e.g., changing T, redox state, mineral assemblages) in addition to different reservoirs including non-Meguma stratigraphy (see discussion in Kontak and Jackson (1999)). All of these observations are consistent with the macro-scale observation of abundant arsenopyrite in both vein and wall rock settings in these deposits demands a source of As, which has been argued to be locally sourced in the carbon- and redox-sensitive element-rich black sedimentary wall rocks to the both the Meguma deposits (e.g., Graves and Zentilli, 1982; Smith and Kontak, 1986; Kontak and Smith, 1989; Sangster, 1992) and similar settings elsewhere (Bierlein and Crowe, 2000; Thomas et al., 2011).

In order to further assess the influence of the immediate host stratigraphy on the arsenopyrite geochemistry, a compilation

of whole-rock geochemical data from the western part of the Goldenville and Halifax groups (White, 2010) has been used in the Au versus Ag plots already referred to. Referring first to the Au versus Ag plot, we use the Beaver Dam TSD dataset as broadly representative of the Meguma deposits (Fig. 22). As can be seen in the latter diagram, the whole-rock analyses overlap the Ag-rich trend (up to 100 ppm) with minor Au as defined by the TSD data. In addition, we have already shown above that this latter trend defines a Bi-Pb-Cd-In-Ag element association using the TSD data for arsenopyrite from several Meguma gold deposits. Consequently, this observation suggests that the later stage fluid responsible for this trend and its elemental association may have been strongly influenced by either the Meguma metasedimentary host rocks or a reservoir similar to them geochemically.

5.5. Implications for gold metallogeny in the Meguma terrane

It has been noted in earlier discussions of the Meguma gold deposits (e.g., Kontak et al., 1990a, 2011; Kontak and Horne, 2010) that the veins record similar mineralizing fluids on the basis of their low salinities, volatile chemistry ($X_{\text{CO}_2}=0.1-0.2$) and $\delta^{18}\text{O}_{\text{H}_2\text{O}}$ values, in addition to inferred temperature (ca. $\leq 300-400^\circ\text{C}$) of formation. These features are noted even though at least two vein forming events are recognized, although the only demonstrated example of an earlier deposits based on absolute dating is The Ovens (Morelli et al., 2005

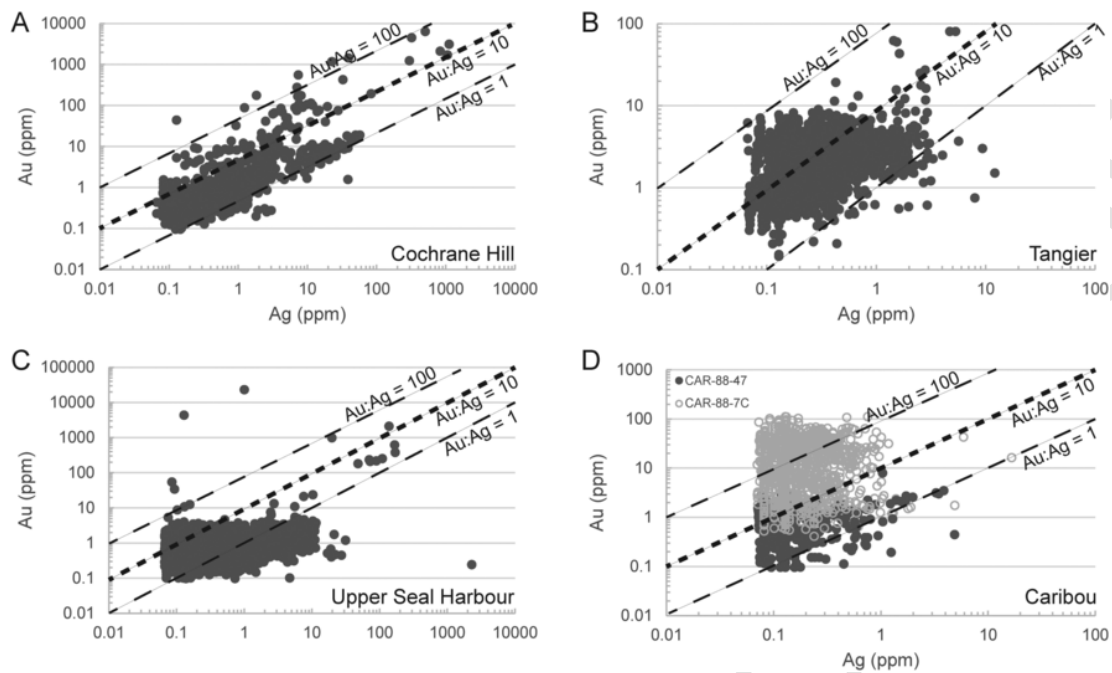


Fig. 19. Binary element plots of Au versus Ag plots (in ppm) for TSD from traverses performed on arsenopyrite grains. A) Sample CH-K092 from the Cochrane Hill gold deposit; B) Sample TG-01 from the Tangier gold deposit; C) Sample DM-91-12 (including 2 traverses) from the Upper Seal Harbour gold deposit; and D) Samples CAR-88-47 (including 3 traverses) and CAR-88-7C (including 2 traverses) from the Caribou gold deposit. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

). These two gold mineralizing events correspond to regional metamorphic and later widespread magmatic processes, respectively. That a similar elemental paragenesis is recorded in the different deposits, which is clearly shown in the summary of elemental paragenesis (Fig. 12), is strong evidence that similar fluids infiltrated these areas even if generated at different times. The arsenopyrite elemental data provide therefore important insight into aspects of the gold metallogeny of these deposits not previously considered. In the first case, the geochemical similarity indicates that if two mineralizing events are indeed present, then a broadly similar fluid in terms of its minor and trace element chemistry can be generated during different tectono-thermal processes. The data also indicate that given that the analyzed samples cover a large area (i.e., 300 km strike length), the ore-forming process is remarkably similar and thus must reflect processes that are repeatable in time and space.

6. Conclusions

Quantitative element distribution maps and traverse analysis of arsenopyrite grains from eight bedding-concordant quartz vein systems in the Lower Paleozoic Meguma gold deposits localized through the Meguma terrane (Nova Scotia, Canada) have been used to investigate a variety of aspects of these gold deposits. The integration of the element maps, petrography and SEM-EDS imaging and analysis with time slice domain (TSD) provide a new and powerful means to assess elemental paragenesis in these and other gold systems globally. Meanwhile, the TSD data provide a means to examine the geochemical signature of the sulfide analyzed and also the chemical evolution of the mineralized system. Importantly, the methodology used in this study is equally applicable to other (i.e., non-gold) hydrothermal ore deposit settings.

In this study elemental paragenesis was first determined for each deposit setting. Results indicate that, despite covering a large geographic area (i.e., 300 km long) and several deposits, the paragenesis

is similar and hence it follows that like fluids and processes operated across this terrane. Since the samples used reflect two distinct thermo-tectonic events (i.e., regional metamorphism and widespread felsic(-mafic) magmatism), the results imply that fluids with similar metal contents can be generated at more than one time, which has important implications for gold metallogeny.

The use of the TSD provides the means to generate meaningful multi-element diagrams. The Au versus Ag plot is used to track Au mineralization and in this case two or more gold events are recognized. The early gold event has up to 100 ppm Au present in the sulfides, likely as invisible Au or nanoparticles. The subsequent gold events are recorded by an upgrading due to zone refining of the early Au with it either having a similar Au:Ag ratio or a much higher one. This plot also revealed a distinct Ag rich event with associated Bi-Pb-Cd-In association.

This study represents the first application of LA-ICP-MS mapping to gold metallogeny by studying representative samples across a large part of a gold mineralized terrane. Whereas some of the interpretations may be controversial, the results clearly demonstrate that the method has the potential to provide insight into processes not previously possible. It is hoped that this project provides the impetus for other such studies to be undertaken both in gold deposits and other hydrothermal settings.

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.gr.2019.11.011>.

Uncited references

Acknowledgments

This work is a continuation of Kontak's ongoing studies of the Meguma gold deposits, which was supported from 1986 to 2006 by the Nova Scotia Department of Natural Resources and in recent years by NSERC Discovery Grants to Kontak. The present study is an out-

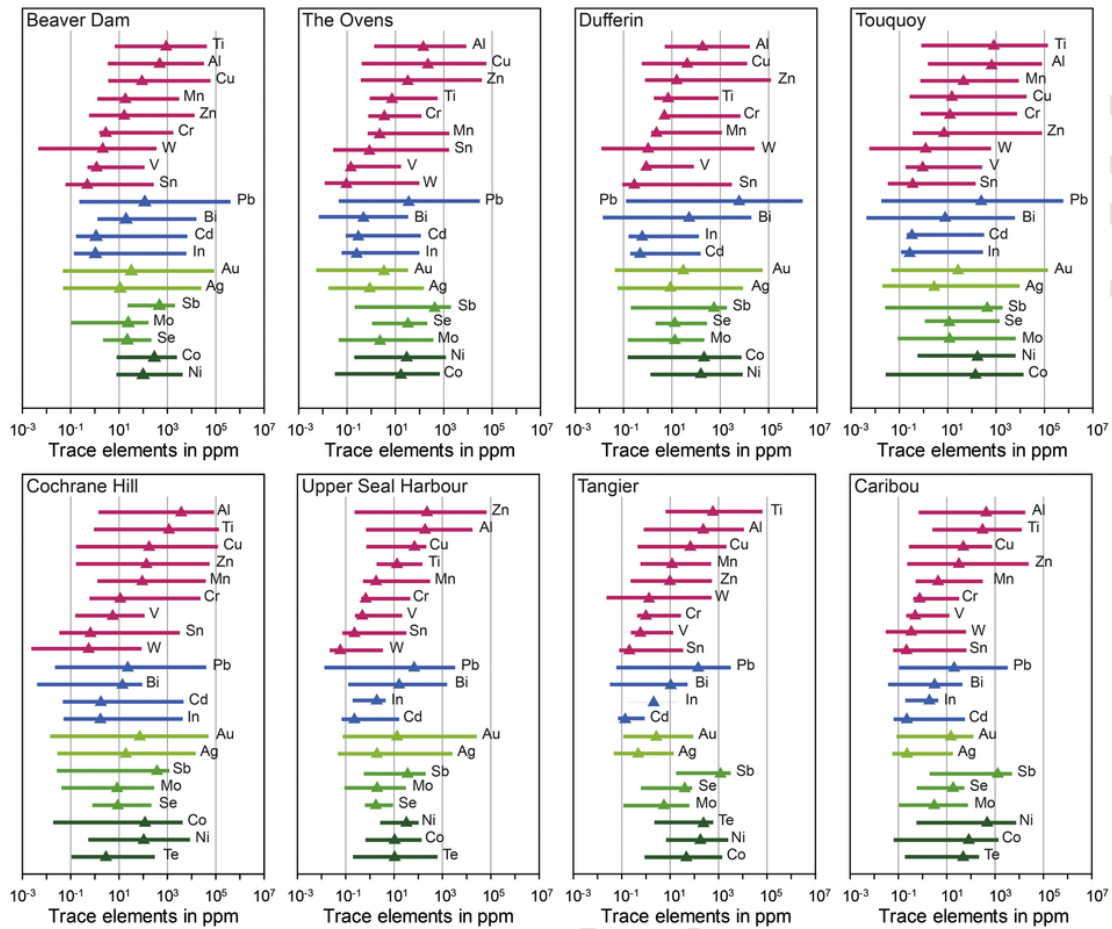


Fig. 20. A summary of the range and mean trace element concentration data for the TSD data for arsenopyrite from the studied deposits. Elements are classified and color coded by their element association and paragenesis. The length of the line shows the range of the element values whereas the triangle represents the mean. Note that there are distinct differences in both the mean and range for Au and Ag values among the different deposits.

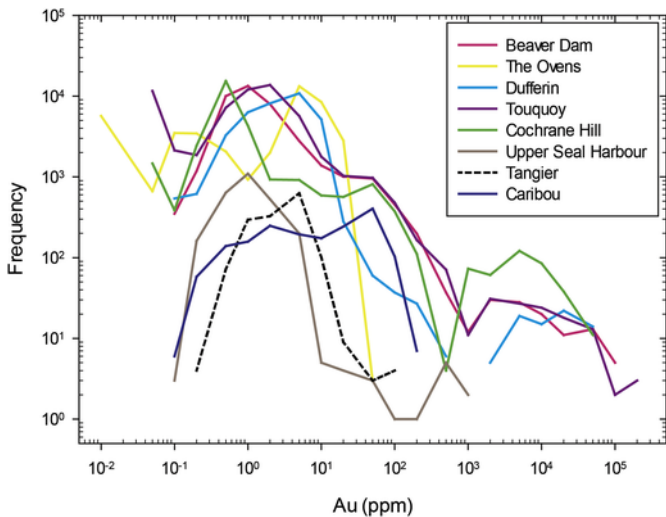


Fig. 21. Frequency of Au concentration (in ppm) in arsenopyrite for the eight studied deposits. Note the very similar trend of gold concentration for all eight samples analyzed at low values (i.e., <100 ppm) and that only some of the arsenopyrite grains record high gold values (i.e., >1000 ppm). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

growth of the Geological Survey of Canada's TGI-4 (Targeted Geoscience Initiative) which supported development of the methodology used in this paper (i.e., LA ICP-MS mapping and TSD approach) which were in part presented in earlier contributions by the same authors. We acknowledge that the Touquoy and Beaver Dam samples analyzed were used as part of the MSc thesis project by L. Chen at the University of Alberta.

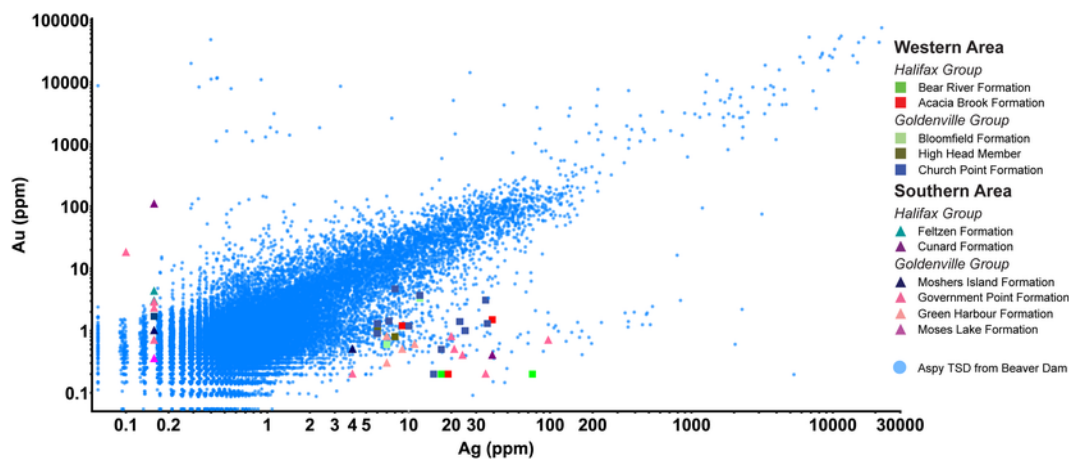


Fig. 22. Binary plot of Au versus Ag (in ppm) for TSD for arsenopyrite from the Beaver Dam deposit ($n=39,797$; blue points) compared to whole-rock data for the Goldenville and Halifax groups from the western part of the Meguma terrane shown in various colors (see legend; data from White, 2010). Note how the whole-rock data overlap with some of the arsenopyrite data which defines the Ag-rich trend seen in many of the binary Au—Ag plots presented. See the text for further discussion of the significance of this plot. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

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