

UNIVERSITÉ DU QUÉBEC À MONTRÉAL

GOLD MINERALIZATIONS AT THE SYENITE-HOSTED BEATTIE GOLD DEPOSIT  
AT DUPARQUET, NEOARCHEAN ABITIBI BELT, QUEBEC, CANADA

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BY  
LUDOVIC BIGOT

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UNIVERSITÉ DU QUÉBEC À MONTRÉAL

LES MINÉRALISATIONS AURIFÈRES DU GISEMENT ARCHÉEN DE BEATTIE À  
DUPARQUET, ABITIBI, QUÉBEC, CANADA

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MÉMOIRE  
PRÉSENTÉ  
COMME EXIGENCE PARTIELLE  
DE LA MAÎTRISE EN SCIENCES DE LA TERRE

PAR  
LUDOVIC BIGOT

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## AVANT-PROPOS

Ce mémoire est rédigé sous la forme d'un article scientifique qui sera soumis à la revue *Economic Geology* sous le titre *Gold Mineralizations at the Syenite-Hosted Beattie Gold Deposit at Duparquet, Neoarchean Abitibi Belt, Quebec, Canada*. Sa présentation est différente des mémoires habituellement présentées à l'UQAM; la rédaction est en anglais, et les figures et tableaux sont situés à la fin de l'article. Michel Jébrak, le directeur de maîtrise, est co-auteur de cet article. Ce format de mémoire a été choisi car il donne l'opportunité de transmettre de nouvelles connaissances à la communauté scientifique à l'échelle internationale.

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## RÉSUMÉ

Le gisement de Beattie est situé dans la ceinture de roches vertes de l’Abitibi, à proximité de la faille Porcupine-Destor, et est associé à une syénite porphyrique. Les corps minéralisés ont une ressource totale inférée de 2,77Moz contenu dans 56,2Mt de minerai à 1,53g/t d’or. Dans le gisement de Beattie, deux styles de minéralisation sont présents, à contrôle lithologique et à contrôle structural. La minéralisation à contrôle lithologique est limitée exclusivement à la syénite altérée en carbonates de fer. Cette minéralisation représente du minerai à basse teneur mais fort tonnage, dans lequel l’or est porté par la pyrite arsénifère et l’arsénopyrite. L’or est « invisible »; il se présente sous la forme de solution-solide dans le cœur poreux et riche en arsenic des pyrites. La minéralisation à contrôle structural est présente uniquement dans des zones de cisaillement adjacentes et incluses dans la syénite. Le minerai est à haute teneur dans des zones de brèches silicifiées à caractère hydraulique et tectonique, dans des veines de quartz cherteux, dans des veines polymétalliques intensément silicifiées, et dans des dykes de syénite trachytique très altérée. L’or y est visible sous forme d’électrum dans les microfractures des pyrites bréchifiées. Les deux styles de minéralisation et les altérations font partie d’une évolution paragénétique commune dans un environnement magmatique-hydrothermal. Des phases initiales oxydantes ont évolué en conditions réduites à la faveur de fluides magmatiques et externes riches en CO<sub>2</sub>. Ce changement d’état d’oxydation a entraîné la précipitation des sulfures et la cristallisation de l’or « invisible » dans la pyrite arsénifère et l’arsénopyrite. Une phase d’altération ultérieure, dominée par des fluides hydrothermaux riches en silice, a pour résultat la bréchification des sulfures et leur redistribution dans des corridors siliceux, ainsi que la remobilisation de l’or dans les microfractures des pyrites bréchifiées. Les compositions calculées de δ<sup>18</sup>O et δD sur les veines de quartz associées à cet événement tardif sont respectivement 7.67‰ et - 66‰, indiquant une dominance des fluides magmatiques avec une probable composante métamorphique. Cet événement syn-métamorphique (transition D2-D3) est associé à un fort enrichissement en Hg, Te, Mo, Ag, Sb, As, et Se. Le style de cristallisation de l’or dans le gisement de Beattie est une conséquence de la dominance des régimes en systèmes ouverts, tant au niveau de l’incorporation dans le magma de l’arsenic qui contrôle la précipitation de l’or « invisible », que de la circulation tardive de fluides hydrothermaux riches en silice qui sont à l’origine de la remobilisation de l’or. Le gisement de Beattie partage des affinités avec des gisements associés aux intrusions (*Intrusion Related Gold Deposit*), des gisements épithermaux à tendance neutre à intermédiaire, et des gisements de type mésothermaux.

MOTS CLÉS : Gisement d’or, porphyre Archéen, or invisible, remobilisation de l’or

## INTRODUCTION GÉNÉRALE

La fertilité aurifère des couloirs de déformation en Abitibi (e.g., Porcupine-Destor, Larder-Lake-Cadillac) est bien reconnue, et notamment avec le style de gisements d'or orogénique (Rabeau, 2010). Cependant, l'exploration de l'or en Abitibi dans ces corridors de déformation porte de plus en plus sur des gisements aurifères porphyriques (Legault et Lalonde, 2009; Lafrance, 2011). Cette tendance s'est concrétisée par la découverte du gisement de Canadian Malartic par Osisko, et sa mise en production au printemps 2011. Avec le développement du modèle de gisements associés aux intrusions (*Intrusion Related Gold Deposit* ; Lang et Baker, 2001), et une certaine remise en cause de la classification de nombreux gisements d'or orogéniques (Groves et al., 1998), les modèles génétiques ont évolué ou ont été précisés. Ainsi, une origine commune dans un large système magmatique-hydrothermal est proposé pour les gisements de Canadian Malartic, Beattie, ou encore Douay (Robert, 2001). Ces gisements présentent des particularités communes, telles que des minéralisations disséminées dans de grands volumes de roches alcalines porphyriques, ou encore des altérations à carbonate, séricite, et silice; mais ils ont aussi des caractères atypiques bien distincts, tels que le style de cristallisation de l'or, des phases différentes de minéralisations, ou encore des associations d'éléments qui mettent en avant l'hétérogénéité de gisements pourtant classés sous une même appellation.

Les récents travaux effectués sur la propriété de Beattie démontrent une dominance du caractère disséminé et à basse teneur de la minéralisation dans l'intrusion, mais aussi un contrôle structural bien marqué le long de failles cisaillantes avec des teneurs élevées en or (Dupéré, 2011). Lorsque la mine Beattie était en opération (1933-1956), l'exploitation se faisait uniquement sur les corps minéralisés à haute teneur le long des failles (Davidson et Banfield, 1944); 33 t d'or ont été produit de 9,26 Mt de mineraï à 3,86g/t Au. À la différence, le projet proposé par Corporation Minière Osisko et Clifton Star Resources Inc. portait sur une exploitation à faible teneur mais fort tonnage dans la syénite et ses bordures; les ressources inférées indiquent 83 t d'or pour 56.2 Mt de mineraï à 1.53g/t Au (Dupéré, 2011). De même que pour ces deux types d'opérations, les pensées sur le modèle génétique du gisement ont évolué. En effet, Davidson et Banfield (1944) considèrent le gisement de

Beattie comme mesothermal, ou possiblement épithermal, alors que les études plus récentes (Robert, 2001 ; Legault et al., 2005) regardent la minéralisation comme génétiquement liée à la syénite.

L'objectif de ce projet de recherche est d'approfondir la compréhension métallogénique du gisement de Beattie, en déterminant la nature et les contrôles de la minéralisation. Pour cela, différents volets de la géologie ont été étudiés. La caractérisation minéralogique de la minéralisation s'est faite par l'analyse de sections polies au microscope optique, au microscope à balayage électronique, au microscope électronique à transmission, ainsi qu'à la microsonde. Cette étude a permis d'établir une séquence paragénétique détaillée pour le gisement. Par ailleurs, des analyses géochimiques de la minéralisation ainsi que des analyses isotopiques ont permis de tirer des conclusions quant à la nature précise des minéralisations et des fluides. Un volet modélisation 3D, avec l'utilisation du logiciel Leapfrog 3D®, a été développé pour différents aspects du gisement, particulièrement les altérations. Cela a permis une visualisation de la géométrie des paramètres lithologiques et structuraux, ainsi qu'une quantification des volumes. La dernière étape est une étude structurale sur les failles et veines du gisement avec l'utilisation du logiciel T-Tecto®. Cela a abouti à une reconstitution du tenseur des contraintes pour les phases tardives de déformation. L'approche pluridisciplinaire de cette recherche a permis d'apporter des précisions d'ordre minéralogique, chimique, et temporel sur le modèle génétique proposé par Robert (2001).

Ce mémoire est présenté sous forme d'un article scientifique qui sera soumis dans une revue scientifique qui traite de la géologie économique. L'article a été rédigé par le premier auteur. Le second auteur est Michel Jébrak qui est le directeur de maîtrise. Son implication s'est faite par la génération de pistes de recherches, l'aide sur le terrain, le suivi de la recherche, la relecture de l'article, et le soutien financier.

Dans l'article, la problématique du projet de recherche est d'abord exposée. Ensuite, la géologie régionale est présentée. La géologie de la propriété est par la suite décrite et comprend la stratigraphie, la pétrologie de la syénite, la géologie structurale, et se termine par une typologie des veines. S'ensuit l'étude de la minéralisation : tout d'abord la typologie des styles de minéralisation, puis l'étude de la paragenèse métallique, l'étude des caractéristiques

de l'or et la géochimie de la minéralisation, vient ensuite l'analyse des altérations, suivit par la caractérisation des phases tardives cassantes et les résultats des analyses isotopiques. Une discussion aborde le volet précipitation et évolution de l'or, suivit par les nouvelles conclusions quant au modèle génétique proposé, et se termine par une comparaison avec d'autres gisements aurifères. La conclusion met en avant les points importants et les avancées réalisées au cours de ce projet de recherche.

Des données complémentaires sont présentent à la suite de l'article en appendice. Elles comprennent respectivement dans l'ordre, la méthodologie pour les analyses géochimiques et les résultats lithogéochimiques, l'étude de modélisation 3D, l'analyse des brèches, l'apport de la susceptibilité magnétique à l'exploration du gisement de Beattie, l'étude structurale des phases tardives cassantes, l'analyse quantitative des fluides d'altération, et l'analyse de l'or au microscope électronique à transmission.

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## CHAPITRE 1

### **GOLD MINERALIZATIONS AT THE SYENITE-HOSTED BEATTIE GOLD DEPOSIT AT DUPARQUET, NEOARCHEAN ABITIBI BELT, QUEBEC, CANADA**

#### **Abstract**

The Beattie deposit is located in the Abitibi greenstone belt, near the Porcupine-Destor Fault Zone, and is associated with a porphyric syenite. Several ore bodies possess a total inferred resource of 56.2 Mt at 1.53g/t gold, for a total of 2.77Moz. In the deposit, two distinct mineralization styles are present: lithologically-controlled, and structurally-controlled. Lithologically-controlled mineralization is limited to the iron-carbonatized syenite. Ore consists of bulk tonnage low-grade mineralization where gold-bearing arsenian pyrite and arsenopyrite are the dominant sulfides. Gold is “invisible”; it occurs as solid solution in the spongy As-rich core of pyrite. Structurally-controlled mineralization is present within shear zones which are within and adjacent to the syenite. This ore consists of relatively high-grade mineralization in silicified breccia with both hydraulic and tectonic features, cherty veins, polymetallic silica veins, and trachytic syenitic dykes. In this facies gold is visible as electrum filling microfractures of brecciated pyrite. The mineralization styles demonstrate a paragenetic evolution in a magmatic-hydrothermal environment. Initial oxidizing stages produced martite. Reducing conditions, marked by CO<sub>2</sub>-rich magmatic and extrenal fluids, led to sulfides precipitation and gold deposition as “invisible” in the As-rich pyrite and arsenopyrite. A later-stage alteration event, dominated by silica-rich hydrothermal fluids produced sulfides brecciation and their redistribution in silica corridors, and gold remobilization in microfractures of brecciated pyrite. Calculated composition for δ<sup>18</sup>O and δD on quartz veins associated with this late event are respectively 7.67‰ and - 66‰, indicating a dominance of magmatic fluids with a probable metamorphic component. This event is associated with a strong enrichment in Hg, Te, Mo, Ag, Sb, As, and Se. The speciation of gold in the Beattie deposit is a consequence of the dominance of open-system regimes, both

for the incorporation in the fluid of arsenic that controlled the early gold deposition, and for the circulation of silica-rich fluids that characterized the late fracture-controlled gold mineralization. The Beattie deposit share characteristics with intrusion-related gold deposit, low to intermediate sulfidation epithermal deposit, and mesothermal deposit.

**KEYWORDS:** Gold deposit, Archean porphyry, invisible gold, gold remobilization

## 1.1 Introduction

The Beattie gold deposit occurs on the north-side segment of the Porcupine-Destor Fault Zone (PDFZ) in the Archean Abitibi Sub-Province (Figure 1.1a), and is associated with a syenitic intrusion. The PDFZ is east-west trending for 200 km. It is one of two main breaks in southern Abitibi, along with the Cadillac – Larder Lake Fault Zone. With a total gold production about 2446 tons (Legault et al., 2005), the PDFZ hosts several gold mines including the Timmins mining camp (Figure 1.1a). Despite the large gold production along the PDFZ, several metallogenic models of deposits remain poorly understood or need to be clarified (Robert et al., 2005).

The Beattie deposit, named in the honor of its discoverer John Beattie in 1910, is located in Duparquet township, in northwestern Quebec, approximately 40 kilometers northwest of the city of Rouyn-Noranda. The mining production of the Beattie deposit started in 1933, followed in 1943 by the mining of the Donchester property, and then the Central Duparquet property, both located in the eastern part of the Beattie deposit (Figure 1.1b). The total production at the Beattie mine, until its closure in 1956, was 32 tons of gold extracted from 9 257 321 tons of ore material, at an average grade of 3.86g/t Au (Bevan, 2009). Arsenic and silver were recovered as secondary products. Recent exploration work, conducted by Osisko Mining Corporation and Clifton Star Resources Inc., was performed on the Beattie deposit. It included geological mapping with a focus on alteration and structural features (Figure 1.1b, Figure 1.2c), geophysical surveys (Figure 1.2a), and an intensive drilling program which established inferred resources of 2.77 Moz gold (56.2 Mt @ 1.53g/t Au).

In the Abitibi greenstone belt, the majority of gold deposits along main fault zones have been recognized as mesothermal (e.g., Val d'or) according to the orogenic gold model of Colvine (1989), and Groves et al. (1998). Hydrothermal mineralized fluids circulating along first and lower-order structures in accreted terranes would be the origin of gold-bearing quartz-carbonate veins and lodes, both typical of orogenic gold deposits. However, several examples of felsic intrusion-related gold deposits occur along major fault zones and share geological features with the orogenic gold model, including metal association, wall-rock alteration, ore fluids, and structural controls (Groves et al., 2003). In Abitibi, the spatial association between small porphyry felsic intrusions and gold deposits has been either interpreted as a structural relation in which the intrusion is acting as a competent trap for mineralized fluids (Colvine, 1989; Witt, 1992), or as a genetic link between gold and the porphyry intrusion (Cameron and Hattori, 1987; Robert, 2001). The former interpretation has been commonly used, whereas the latter interpretation, referring to complex magmatic processes which imply redox reactions, is less prevalent. On the basis of geological relations and geochronological constraints, Robert (2001) distinguished a new group of syenitic gold deposits in the Abitibi greenstone belt, including Beattie, Canadian-Malartic, Holt-McDermott, Young-Davidson, and Douay. These deposits share common characteristics regarding the age of crystallization, the spatial association, the depth of emplacement, the mineralization, and their origin: they crystallized between about 2680 and 2672 Ma near major faults; they occur in association with the Timiskaming sedimentary Group; they were emplaced at shallow depth, typically lower than 2km; the mineralization consists of disseminated sulfides and structural control is an important feature; they are mostly oxidized intrusions and a genetic link between these deposits is proposed. Despite this classification, two questions need to be clarified about the Beattie deposit: How was gold precipitated? What is the evolution of the deposit in terms of petrology, geochemistry, and deformation?

The goal of this paper is a better understanding of the metallogeny of the Beattie deposit by identifying the nature and the controls of the gold mineralization. This will also improve the genetic model proposed by Robert (2001) for syenite-hosted gold deposits in the Abitibi.

## 1.2 Regional geology

The Archean Abitibi sub-province is located within the south-eastern part of the Superior Province. It is the largest continuous volcano-plutonic belt of the Canadian Shield, extending 700 km long E-W and 300 km wide N-S (Figure 1.1a). A Proterozoic structure, the Grenville orogenic front, truncates this belt in its eastern part. The Abitibi greenstone belt shares several characteristics with Archean greenstone belts in the Yilgarn craton regarding the age and the type of deposits (Robert et al., 2005; Duuring et al., 2007).

The Porcupine-Destor Fault Zone (PDFZ) is interpreted as a tectonic suture of allochthonous terranes (Mueller et al., 1996; Daigneault et al., 2002). It represents a geological boundary between the North Volcanic Zone (NVZ) and the South Volcanic Zone (SVZ). Chown et al. (1992) interpret the NVZ as an intact arc segment with two volcanic cycles, whereas Dimroth et al. (1982) consider the SVZ as a homoclinal oceanic floor-to-arc succession. An alternative model proposes the PDFZ is a consequence of anticlines folds detachment in a large autochthonous terrane (Benn and Peschler, 2005; Bedart et al., 2012).

Boivin (1974) considers a prehnite-pumpellyite metamorphic facies in the Duparquet area. The regional metamorphism in the area of Rouyn-Noranda ranges the subgreenschist-facies to greenschist-facies and locally amphibolite-facies around plutons (Powell et al., 1995). The age of regional metamorphism can be bracketed between 2677 and 2643 Ma, with average pressures of approximately 200 MPa.

A polyphased tectonic event in the southern Abitibi sub-province is proposed by Bleeker and Parrish (1996), and Daigneault et al. (2002). A first contractional deformation (D1) is marked by tilting, folding and local thrusting. A second event (D2), involving a major north-south shortening, is responsible for the development of east-west grain of lithologies, upright folds and penetrative foliation. It evolved gradually into D3 dextral faults and shear zones mainly localised along major fault zones. These deformational events started at 2687 Ma and ended around 2662 Ma with a breaking period between about 2680 and 2672 Ma.

### 1.3 Local geology

The Beattie syenite extends E-W for a length over 3300m, and a width N-S about 500m (Figure 1.1b). Both syenite and mineralization dip 35 to 40° to the E. Several smaller syenitic stocks, spatially associated with the Beattie syenite, occur in the E-W trend in the area of Duparquet.

#### 1.3.1 Stratigraphy

The oldest unit in the area of Duparquet is the Kinojevis group where the main assemblage is the Deguisier formation (2718-2722 Ma; Zhang et al., 1993). Lithologies comprise abundant Fe-Mg tholeiitic massive and pillow basalts, with minor andesites and lapilli tuffs (Goutier and Lacroix, 1992). These volcanic rocks are interpreted as oceanic crust in a backarc rift (Kerrich et al., 2008). The Porcupine group (2692-2696 Ma) overlies previous units. It consists of greywacke, siltstone and mudstone indicating predominantly distal deposition by turbidity currents in a deep basin. U-Pb dating on zircon suggests that erosion of calc-alkaline to tholeiitic mafic volcanic rocks from the oldest assemblages of the south Abitibi greenstone belt would be the source for Porcupine group sediments (Ayer et al., 2002; Ayer et al., 2005). The youngest assemblage, the Duparquet formation, which belongs to the Timiskaming sedimentary group, consists of conglomerate to sandstone (Figure 1.1b) that record deposition in a stream dominated alluvial fan, fan-delta and/or braid delta adjacent to the sea or lakes (Mueller et al., 1996). These rocks are unconformably deposited on older assemblages. The Duparquet formation forms an easterly trending corridor along the Porcupine-Destor fault zone (Benn and Peschler, 2005; Peschler et al., 2006). Its maximal deposition age is  $2678.9 \pm 2.8$  Ma (David et al., 2006), and it lasts until 2672 Ma. This sedimentary period is known as the Timiskaming event. Based on U-Pb dating and petrological setting, David et al. (2006) suggest a proximal provenance for the volcanic and syenitic fragments.

A period of plutonism occurred during the Timiskaming event (Robert, 2001). The Beattie syenite (Figure 1.1b) has been dated at  $2681.6 \pm 1$  Ma (U/Pb on zircon; Mueller et al.,

1996) indicating a pre-Timiskaming age, however this date is comprised in the uncertainty bracket of the sedimentary event. The age constraint based on field geology remains obscure; work by Robert (2001) outlines evidences that the Duparquet formation sediments unconformably truncate the Beattie syenite (Figure 1.1b), however Graham (1954) reports numerous conglomerate xenolith in the syenite. This uncertainty of the relative age between the alkaline rocks and the sediments highlights their sub-contemporaneous formation.

### 1.3.2 Petrology of the Beattie syenite

The Beattie syenite is a composite stock; several types of syenite differ in texture, composition, and morphology. Bourdeau et al. (2011) recognized five distinct syenites composing the Beattie intrusion: (1) Unaltered syenite in the core of the Beattie intrusion. This unit is magnetite and amphibole-rich but gold free; (2) The Beattie syenite porphyry, the main syenitic unit of the intrusion in terms of volume. It is strongly altered in carbonate and sericite, and contains most of the gold mineralization; (3) The Central Duparquet syenite porphyry, located in southeastern part of Duparquet. Phenocrysts size increase about twice, ranging from 5 to 16 mm, compared to the Beattie syenite porphyry, and it contains some gold; (4) The megaporphyritic syenite, related in space to the Central Duparquet syenite, and characterized by significant feldspar megacrysts of 1 to 6 cm in size. This syenite is gold free; (5) Trachytic syenite dykes, composed of 85-95% feldspar, and consisting of feldspar laths in a red aphanitic matrix of orthoclase. This unit is usually strongly altered to carbonate and is gold-bearing. Furthermore, these dykes usually crosscut mineralized zones. Despite the variability of composition and texture of syenites and phenocrysts morphologies, all five syenite units share a common geochemical signature suggesting a same magmatic source.

The dominant intrusion, the Beattie syenite porphyry, is a porphyritic rock mainly characterized by light-grey K-felspar phenocrysts, microcline and orthoclase, 2 to 10 mm in size, set in a grey to reddish aphanitic groundmass of an aggregate of anhedral anorthoclase crystals (Figure 1.5a, b). Feldspar minerals comprise about 80% of all minerals in the syenite (Davidson and Banfield, 1944). The groundmass also contains some clinopyroxene relicts, but mostly hornblende which is largely altered to calcite, chlorite, and to a lesser amount

epidote; green muscovite (sericite) is also common in the matrix. Accessory minerals include euhedral titanite, epidote, titaniferous magnetite and apatite.

### 1.3.3 Structural geology

In the area of Duparquet, stratigraphic units are deformed (Figure 1.1b). Major folds with E-W axial planes occur in the Kinojevis group; Deguisier formation layers are upright with younging to the south (Goutier and Lacroix, 1992). Sedimentary rocks of the Duparquet formation have strongly accommodated the deformation. The main structural characteristics include: A N-S flattening with L/l ratio of about 4 to 5 as measured on clasts (Figure 1.3c), E-W to ENE-WSW tight synclinal folds plunging 40° to 60° to the west, and an ENE-trending cleavage which steeply dips to the south. In addition, stretching lineations plunge shallowly to the east. In the Beattie syenite, except for local flattening, deformation is not well recorded.

Faulting is a major structural characteristic in the area of Duparquet. The PDFZ trends N110 with a dip ranging from 50° to 80° to the south (Graham, 1954). The Beattie syenite is bound to the north and to the south by two faults which branch away from PDFZ (Figure 1.1b, Figure 1.2c). The former is the Beattie fault which is mainly a thrust fault with a dextral component; the latter is the Donchester fault which is regarded as a shear fault. Both faults dip steeply sub-vertically to the north. An E-W to ENE-WSW sub-vertical shear zone (named the “Fracture zone” from Davidson and Banfield, 1944; Figure 1.1b, Figure 1.2c, Figure 1.3b), of about 10 to 20 meters width, occurs within the intrusion. In the Beattie syenite, foliation only exists in the Fracture zone. Major conjugated NW-SE dextral and NNE-SSW sinistral strike-slip movements crosscut the deposit (Figure 1.1b, Figure 1.20). In addition to these main strike-slip faults, several minor faults are present in the deposit. Furthermore, jointing is common in the syenite; a widespread joint system steeply dips to the S and W (Figure 1.3a).

### 1.3.4 Veins

Veins are locally abundant, but are not a key feature of the deposit. Veins are mostly present within the syenite and the Timiskaming sediments; only few have been observed in the volcanic rocks. Studied veins have three main orientations: ENE-WSW, ESE-WNW and NNW-SSE (Table 1.5; Figure 1.20, Figure 1.21a).

ENE-WSW veins occur in the Fracture zone. They consist of mineralized cherty quartz veins steeply dipping to the south (Figure 1.6e, f). Quartz is subhedral and forms a mosaic of very fine equigranular crystals, usually 10 to 15  $\mu\text{m}$ .

ESE-WNW veins consist of flat extension barren quartz-carbonate veins, presenting open-space crystallisation, pseudo-colloform texture, and filled partially by chlorite (Figure 1.22a, b, c). Quartz is subhedral and forms a mosaic of medium grained crystals, ranging from 0.5 to 1.5 mm; the smallest quartz crystals form a cryptocrystalline groundmass. Interstitial sericite is locally present. Medium grained calcite is commonly present, and locally massive dissolution occurs (Figure 1.22c). These quartz-carbonate veins are irregular and their width range from 1 to 20 cm. They usually dip shallowly to the north. These veins locally display slight curves, suggesting crystallisation in ductile environment. Hydraulic breccia with sharp fragments of altered syenite is a common feature (Figure 1.22d).

NNW-SSE trending veins are mainly present in the Fracture zone within the syenite where sericitization, and locally silicification, is strong. Veins consist of tension veins filled with quartz and iron-carbonate (Figure 1.22e, f). These veins are not mineralized. Quartz is mostly subhedral and medium grained. Iron-carbonate consists mainly on euhedral to subhedral medium grained ankerite crystals that usually display a brown oxidized color on surface. Partial carbonate dissolution occurs in the strongest sericitization zone (Figure 1.22e, g). Most veins have a periodic spacing, with the same dimensions of about 10 to 20 cm length and 0.3 to 1 cm width; spacing usually ranges from 2 to 8 cm. Sinistral movements on NNW-SSE veins locally occur in the Fracture zone (Figure 1.22g). Cross-cutting evidences indicate NNW-SSE trending veins postdate ENE-WSW trending veins (Figure 1.22f).

## 1.4 Deposit geology

In the Beattie deposit, mineralization is hosted both in the Beattie syenite and in the Beattie and Donchester faults zones and the Fracture zone. Several mineralized zones within and adjacent to the Beattie syenite are recognizable (Figure 1.2b). The main ore body is located along the Beattie fault which marks the north contact between the Beattie syenite and the volcanic rocks of the Deguisier formation. The style of mineralization depends on the location in the deposit, either in the faults zones, or in the intrusion. All mineralization, regardless of their location, share a common paragenetic evolution in a magmatic-hydrothermal environment. Gold mineralization is generally correlated with the fineness of sulfides (pyrite and arsenopyrite) and their abundance.

### 1.4.1 Mineralization styles

Mineralization styles in the Beattie deposit are twofold: (i) lithologically-controlled, limited to the syenite, and (ii) structurally-controlled, associated with the Beattie and Donchester fault zones and the Fracture zone (Table 1.1; Figure 1.4). Disseminated ore is widespread in the deposit and occurs both in the lithologically-controlled facies and in the structurally-controlled facies. However, gold content, mineralogy, alteration, and volume of ore are different depending on the facies. Ore in the lithologically-controlled facies is almost the same everywhere in the Beattie syenite; ore in the structurally-controlled facies displays several types of mineralization including breccia, replacement, veins, and dyke.

#### 1.4.1.1 Lithologically-controlled facies

Lithologically-controlled facies occurs exclusively within the Beattie syenite porphyry, and out of the Beattie and Donchester faults zones and the Fracture zone. Rock consists of a grey massive syenite (Figure 1.5a), occasionally reddish (Figure 1.5b), resulting from medium to intense iron-carbonatization. Phenocrysts are usually smaller than 1 mm in size, and fine chlorite stockworks as well as small veins of calcite have developed (Figure 1.5a, b). Ore is always disseminated. This facies represents a bulk-tonnage of low-grade

material. Some mineralized zones are roughly continuous on 100m, and 83% of the overall ore in the deposit is below 2.5g/t Au (Dupéré et al., 2011). Mineralization generally grades from 0.3g/t Au to 2.5g/t Au; the sulfide concentration is about 5 to 8% (always <10%). The only important sulfides in the ore are gold-bearing arsenian pyrite and arsenopyrite; pyrite is about six times more abundant than arsenopyrite.

#### **1.4.1.2 Structurally-controlled facies**

Structurally-controlled facies comprises the majority of the historically mined ore in the Beattie deposit. The mineralization zones are present either along the Beattie and Donchester faults zones, or within the Fracture zone, or both. Most of the mineralization grades >2.5g/t Au, with regular anomalous zones above 10g/t Au; the sulfide content ranges from 10 to 20%, and high-silica value is typical. Dupéré et al. (2011) indicate that ore grading 5g/t Au and more account for only 4.5% of the total ore in the deposit. In the deposit, four types of mineralization belong to the structurally-controlled facies.

Breccia ore was the main mined ore in the Beattie deposit (Davidson and Banfield, 1944). Rock consists of a greyish blue to yellowish colored breccia of angular to sub-angular fragments cemented in an abundant silicified groundmass, and cut by a network of tiny quartz stringers (Figure 1.6a, b, d). The fragments are usually monogenic and consist of altered syenite or altered volcanic rocks. The particles size distribution is normal, ranging from a few mm to a few cm. The motion of the fragment is usually very low, indicating a low mobility; occasionally the dilatation increases and the fragments are matrix supported. The fabric is mostly inherited. The complexity of the fragments (i.e., irregularity of the margins of the clasts; Jébrak, 1997) is low, however some fragments can have a significant complexity (Appendice C). These characteristics provide a better understanding of the fracturing processes. Silicification is a key feature of the breccia ore (Figure 1.6b, c, d). The ore bodies are similar to dykes. They form sub-vertical narrow volumes extended E-W along the Beattie faults and the Fracture zone (Figure 1.4). Ore is both included in the syenite and in the volcanic rocks. The mineralization of breccia ore type consists of fine disseminated pyrite mainly, with small clusters of pyrite; grade is usually >5g/t Au.

Cherty quartz veins (Figure 1.6e, f) occur only within the Fracture zone among foliated syenite. Rock consists of irregular grey-blue cherty quartz veins of about 10 to 20 cm width. Pyrite grains are very fine and disseminated. Small amount of sericite is occasionally present in the veins. The mineralization of cherty quartz veins usually grades >10g/t Au.

Polymetallic veins occur along the Beattie fault and the Fracture zone, but are rare. They consist of Au-As-S-Cu-Hg-Te rich semi-massive veins and are related to intensive silicification (Figure 1.6g). The mineralogical assemblage comprises pyrite, enargite, tennantite-tetrahedrite, and coloradoite. These narrow veins comprise the highest metallic values in the deposit such as 119.5g/t Au and 341g/t Au.

Trachytic syenite dykes (Figure 1.6h) form small ore bodies, about 6 meters width and irregular in length, which truncate both the syenite and the Fracture zone. They occur exclusively near the Beattie and Donchester fault zones and the Fracture zone. Unlike breccia ore, cherty quartz veins, and polymetallic veins, high-silica content is not a significant characteristic of trachytic syenite dykes. Carbonatization and sericitization are the dominant alterations. Mineralization consists of fine disseminated sulfides, mainly pyrite with small amount of arsenopyrite. Trachytic syenite usually grades >5g/t Au. These mineralized and structurally-controlled syenitic dykes have been named *Bostonite* by O'Neill (1932). The alteration and ore of trachytic syenite dykes appear as the lithologically-controlled facies; however, their occurrence related to the Beattie and Donchester faults and the Fracture zone highlights its structural control.

#### **1.4.2 Magmatic-hydrothermal paragenesis**

The paragenetic sequence in the Beattie deposit is divided into four stages (Figure 1.7): (1) Fe-Ti assemblage, (2) martitization, (3) sulfidation, and (4) breccia facies. Ore mineral paragenesis presents an evolution from magmatic phase to hydrothermal event.

(1) Fe-Ti assemblage consists of subhedral porphyric K-feldspar, and anhedral K-feldspar groundmass, with subhedral titaniferous magnetite (0.1-0.25 mm; up to 2% Ti in magnetite) which accounts for more than 55% of all metallic minerals (Figure 1.8a). Euhedral titanite (0.15-0.35 mm) is usually twinned and widespread (Figure 1.8a), and comprises more

than 40% of all metallic minerals. Very fine inclusions of apatite (0.005 mm) with submicron barite crystals locally occur in titanite crystals (Figure 1.8b, c, d), and in the matrix. Anhedral ilmenite (<0.05 mm) is present as trace mineral, <2%. Leucoxene, 4-5%, results of partial alteration of titanite, and titaniferous magnetite to a lesser extent. This Fe-Ti-oxide rich phase is altered by subhedral chlorite and a low Ca-carbonatization which grows mainly within hornblende crystals.

(2) The martitization phase consists of the gradual oxidation of magnetite grains (Figure 1.9a). Fine hematite crystals, representing about 35% of metallic minerals, form as exsolutions in magnetite which accounts for 50% of the overall metallic minerals. Leucoxene is widespread with 15% of all metallic minerals. It appears under various forms associated with various alteration styles: as pseudomorphs of titanite (Figure 1.9b), as replacement of ilmenite, and as a coronitic texture around titaniferous magnetite.

(3) Sulfidation phase is characterized by the abundance of fine sulfides, mainly pyrite. Euhedral pyrite (0.02-0.125 mm; Figure 1.10a, b, c, d, e, f), locally subhedral, is disseminated within the stock. It is the most abundant sulfide comprising 75% of all metallic minerals. Three overgrowth generations of pyrite can be identified: (G1) gold-bearing porous arsenian pyrite (Figure 1.10a, b, c, d, f). (G2) and (G3) non-porous, arsenic-poor and gold free pyrite. Very fine subhedral arsenopyrite (<0.015 mm) accounts for 12% and appears mainly as inclusions in G1 pyrite (Figure 1.10a, b, c, d, f). Galena (<0.01 mm; 1%) and sphalerite (<0.01 mm; 1%) also occur as inclusion in G1 pyrite (Figure 1.10a, b, c, d, f). In the sulfidation stage, neither titanite pseudomorphs, nor iron-oxide crystals exist. They are broken-down and result in widely disseminated leucoxene (12%). Locally, hematite dust occurs in the matrix and gives the reddish-wine color of the syenite (Figure 1.5b). Carbonatization is the dominant alteration. Sulfidation phase is controlled by the intrusion as it happens exclusively within the syenite and out of fault zones. Sulfidation phase represents the lithologically-controlled facies.

(4) The breccia stage is dominated by the fracturing of rocks and the brecciation of sulfides within silica corridors of decimeter to meter in width. In these corridors, the sulfide concentration is about two times more abundant than in the sulfidation phase. The three generations of euhedral to subhedral pyrite (G1, G2, G3) are recognised and all have

undergone brecciation. Pyrite is still the dominant sulfide, composing 70% of the overall metallic minerals (Figure 1.11a, b). In addition, the margins of the pyrite usually display alteration zones with irregular shapes (Figure 1.11a, e). Very fine subhedral arsenopyrite (<0.015 mm) accounts for 10% of metallic minerals, and mostly occurs as inclusion in G1 pyrite. However, both pyrite and arsenopyrite have recrystallized in minor amount. In addition to galena (<0.01 mm; 1%) and sphalerite (<0.01 mm; 1%), new metallic minerals (0.002-0.01mm) are present in microfractures and in the porosity of the pyrite crystals, as well as in the matrix: chalcopyrite (Figure 1.11h), enargite (Figure 1.11g, j), tennantite-tetrahedrite (CuAsS-CuSbS; Figure 1.11a, i), coloradoite (HgTe; Figure 1.11a, j), hessite ( $\text{Ag}_2\text{Te}$ ; Figure 1.11a), and electrum (AuAg; Figure 1.11a, c, d, e, f). These minerals individually amount to less than 1%. Significant leucoxene concentrations (10%) happen locally. Barite crystals (0.02-0.03 mm; 2%) are present in the matrix. The breccia episode is controlled by the structure as it occurs only in the Beattie and Donchester faults zones, and the Fracture zone. Breccia stage represents the structurally-controlled facies.

A late-hydroxidation event led to goethite crystallisation, locally displaying coronitic texture around euhedral pyrite (Figure 1.11j), and ultimately pseudomorphic textures.

#### **1.4.3 Gold features and ore mineralogy**

Gold distribution in sulfides is directly related to the mineralization control, either lithological or structural. Gold texture evolves from “invisible” in pyrite and arsenopyrite to free-state, and gold concentration in host minerals ranges from insignificant to several hundreds of ppm. Furthermore, ore mineralogy changes significantly depending on gold features.

In the lithologically-controlled facies, gold appears in arsenian pyrite (G1) and arsenopyrite as “invisible” gold. This distribution is the dominant style in the Beattie deposit. Microprobe assays indicate that gold concentration usually increases from arsenian pyrite (G1) to arsenopyrite (Table 1.2; Appendix A); gold contents in arsenian pyrite (average 2% As, and up to 4.7% As) range from 80 to 270 ppm Au, whereas gold values range from 100 to 420 ppm Au in arsenopyrite. Arsenian pyrite is S deficient (50-51% wt%), whereas

arsenopyrite is characterized by lower As content (34.2 – 40.5 wt%) and higher content of S (21.5 – 27.6 wt%) and Fe (36.3 – 36.6 wt%). Significant amount of gold and arsenic coincide with the high porosity of G1 pyrite. Hence, a correlation between gold, arsenic, and the level of porosity is highlighted in G1 pyrite.

Transmission Electron Microscope data (TEM; Appendix G), in G1 arsenian pyrite (Figure 1.12a) of the lithologically-controlled facies, shows heterogeneity in the lattice fringes (Figure 1.12b) which highlights the porous characteristic of this pyrite. In addition, the range of grey to dark color in the pyrite (Figure 1.12b) indicates respectively lower and higher arsenic concentration. Despite a high magnification (x470000), no gold particles have been found in the studied gold-bearing thin sections.

In the structurally-controlled facies, gold has distinct features of occurrence and texture. Most pyrite and arsenopyrite host minor to negligible invisible gold content, regularly lower than 70 ppm Au (Table 1.3). Gold is mainly visible but small, ranging from 2 to 15  $\mu\text{m}$ . It appears as electrum ( $\approx$  85% Au, 15% Ag; Table 1.4) which crystallises by filling microfractures in the pyrite, and in rare occurrences appears as free-state in the silica corridors (Figure 1.11a, c, d, e, f). In this facies, pyrite is still carrying gold but arsenic is not a significant gold marker anymore (Figure 1.13a). New arsenopyrite is still As deficient (42 wt%), but less than in the lithologically-controlled facies.

Ore mineralogy and elemental association are substantially different in the lithologically-controlled facies than in the structurally-controlled facies. Cu, Pb, Zn, and Bi concentrations are approximately the same in both facies, with a tendency for enrichment for Bi and Zn in the lithologically-controlled facies, and a slightly increase for Cu in the structurally-controlled facies. Structurally-controlled facies are marked by a strong increase of Hg, Te, and Mo (Figure 1.14), and to a lesser degree of Ag, As, Au, Sb, and Se. Structurally-controlled facies are polymetallic facies. Mineral ratios change in ore bodies; when lithological control is predominant,  $\text{Au}/\text{Ag} \approx 1.2$ , whereas for structural control,  $\text{Au}/\text{Ag} \approx 0.8$ . On the basis of gold concentration, two populations may be recognized (Figure 1.13a, b): (1)  $\text{Au} < 4$  ppm: there is a linear correlation with As. In addition, when  $\text{Ag} < 2$  ppm gold is linearly correlated with Ag, whereas above this threshold Ag distribution is scattered. (2)  $\text{Au} > 4$  to 6 ppm: both As and Ag populations are uncorrelated and heterogeneously

distributed. Minerals crystallization is significantly different depending on the mineralization facies. Unlike the lithologically-controlled facies, where metallic minerals crystallize as micrometric to nanometric inclusions in G1 pyrite, crystallisation styles of ore minerals in the structurally-controlled facies are heterogeneous. They either occur as filling microfractures and porosity of pyrite for enargite, coloradoite, tennantite-tetrahedrite, hessite, electrum (Figure 1.11a, c, d, e, g, i), or as veins for enargite and coloradoite (Figure 1.11g, i), or as free-state usually in silica corridors for chalcopyrite, electrum, and late-barite (Figure 1.11a, f, h).

#### 1.4.4 Alteration

Alteration is widespread in the Beattie deposit (Figure 1.2c, Figure 1.15a, Figure 1.16a, Figure 1.17a, Figure 1.18a). The main alteration features are carbonatization, hematization, sericitization, and silicification. They always occur within the syenite, and along its margins. In addition, previous work identified a local sodic and potassic feldspathization along the Beattie fault. Furthermore, bleaching has been observed near ore bodies in the basalts (Davidson and Banfield, 1944). The four main alterations have been recognized in regard to their significant volume using 3D modeling with Leapfrog3D® (Appendice B), their enrichment or depletion of elements using isocon diagram (Grant, 2005), and their style of replacement.

Carbonatization is the dominant alteration in the deposit in terms of its volume. It comprises more than 151 million cubic metres of rock (Figure 1.16a), compare to 248 million cubic metres of syenite; 62% of this carbonatization happens in the syenite, and the remaining 38% is in the basalts and andesites of the Kinojevis group, and turbidites of the Porcupine group. Two carbonatization phases are recognized: (1) early, Fe-Ti assemblage-related. This carbonatization is a low degree of alteration consisting mainly of localised amphibole replacement. Altered rock is usually grey with a greenish tint. These altered rocks occur in small and distinct areas, near anomalous magnetic zones ( $>0.0293 \text{ nT/m}^2$ ; Figure 1.2a, c). (2) syn-sulfidation is the most abundant carbonatization phase of the deposit. This alteration consists of partial to strong replacement of silicates; it is a pervasive alteration.

Carbonatized syenite is usually grey with a locally reddish tint (Figure 1.16b). This second carbonatization event never occurs near magnetic halos; neither magnetite, nor hematite is found associated with this second carbonatization (Figure 1.7). Bourdeau et al. (2011) identified the composition of different types of carbonates; amphibole replacement consists of calcite, whereas the second carbonatization is dominated by iron-carbonates, mostly ankerite and siderite. Syn-sulfidation carbonatization led to CaO, C, and K enrichment of 200%, whereas Na<sub>2</sub>O is depleted by more than 80% (Figure 1.16c). Furthermore, ore minerals including As, Au, Mo, W, S, and Hg are enriched significantly in comparison to the least altered rock.

Hematization is the second largest alteration type in volume. It comprises more than 62 million cubic metres of rock (Figure 1.15a) with 94% of the hematization within the syenite. Hematization is recognized both as martite (martitization phase), and as a high concentration of hematite dust resulting from the destruction of martite and the remaining magnetite. Hematized rocks usually have a reddish-wine color (Figure 1.15b), typical of oxidized rocks. Even though hematite dust occurs in rare instances in some mineralized zones, hematization is not gold-related as most elements remain immobile (Figure 1.15c).

Sericitization is a smaller but widespread alteration comprising of 10 million cubic metres of rock (Figure 1.17a). Sericitized rock displays a green-color (Figure 1.17b), and sericitization occurs only in the syenite and the in Timiskaming sediments in the south (Figure 1.2c, Figure 1.17a). Bourdeau et al. (2011) identified sericite mineral as ferro-alumino-celadonite,  $KAl(Fe^{2+},Mg)Si_4O_{10}(OH_2)$ . Sericitization consists of replacements of potassic minerals, mainly K-feldspar, and is associated with a strong enrichment in Au, As, Te, S, Hg, Mo, and W (Figure 1.17c). K and Rb are both enriched by about 200%. Sericite alteration is controlled by permeability of microfracture and is thus diffuse in the Fracture zone in which green-colored syenite displays large channels of about 20 meters width. When strong sericitization happens, kaolinite is widespread but less abundant than sericite (Figure 1.19a).

Silicification is the smallest alteration in terms of volume. It comprises 4 million cubic metres of rock both in the syenite and in the volcanic and turbiditic rocks (Figure 1.18a); 80% of the apparent silicification is present in the syenite, and the remaining 20% is

in the host rocks. Silicified rock consists of a grey-blue pervasive silica alteration (Figure 1.18b) implying a partial to complete replacement of the host rock. High-grade mineralization is strongly associated with silicification in the Beattie deposit. Furthermore, silica alteration is associated with the strongest enrichment of elements among all alterations in comparison to the least altered rock (Figure 1.18c): SiO<sub>2</sub> increase of 130%; Au, Ag, As, Hg, Sb, Te, Mo, Se and S enrich from 250% to several thousand percent. Silica alteration is strongly controlled by structural features: breccia, veins, and faults. Unlike sericite alteration, silicification forms generally narrow volumes with sharp contacts. Field evidence of brecciated sericitised syenite fragments in silica corridors indicate sericitization pre-date silicification (Figure 1.19b).

Based on microscopic observations and field evidence, a relative chronology of alteration can be established : (1) Fe-Ti assemblage-related Ca-carbonatization, (2) hematization, (3) syn-sulfidation Fe-carbonatization, (4) sericitization, and (5) silicification.

#### **1.4.5 Late-faulting**

Major conjugated strike-slip faults with an oblique component crosscut the deposit: NW-SE dextral movement and NNE-SSW sinistral movements (Figure 1.1b, Figure 1.20). Within the syenite, several minor faults with two main fault dip orientations, S and W, are present (Figure 1.21b). However, fault orientation is not always homogenous. Dipping is usually steep, ranging from 60° to vertical. Oblique-slip is the dominant fault type. Most faults contain slickenlines generations with an average dip of about 50° (Figure 1.21b). Slickenlines are conserved locally in crystal fiber lineations of chlorite, tourmaline, or calcite. Fault breccia, with angular fragments cm-size and chlorite in matrix, occurs locally associated with important net slip of several meters (Figure 1.22h). Field evidence indicates ENE cherty veins and ESE flat extension veins have been affected by the late-faulting. No direct relation between ENE cherty and ESE flat extension veins has been observed; however, the different stress regimes required for their formation highlight that they could not have formed at the same time.

Stress tensor reconstruction applied on minor faults, using T-TECTO® (Appendice E), indicates a relative chronology of two main compressive regimes (Figure 1.21c): (1) NNW-SSE stress with  $\sigma_1$  plunging  $35^\circ$  to SSE. This stress episode could be the source of NW-SE and NNE-SSW trending strike-slip faults. Furthermore, this brittle compressive period matches with the NNW-SSE trending tension quartz veins considering their stress orientation. (2) ENE-WSW axis with  $\sigma_1$  plunging  $35^\circ$  to ENE. Sinistral movement observed on NNW-SSE trending tension quartz veins (Figure 1.22g) in the Fracture zone could be related to the ENE-WSW compressive event.

#### 1.4.6 Stable isotopes

The oxygen and hydrogen stable isotope data for fluids in quartz veins in the Beattie deposit are reported in Table 1.6 (Appendice A). ENE cherty veins  $\delta^{18}\text{O}$  values range from 12.2 to 16.1‰, with one anomalous value of 6.9‰ (n=6). Assuming a temperature of  $350^\circ\text{C}$  (cf. arsenopyrite geothermometer 1.5.1.2) for quartz precipitation, the oxygen isotope composition for hydrothermal fluids calculated for quartz analyses have a restricted range from 6.2 to 10.1‰ with an average value of 7.67‰ excluding the anomalous value of 0.9‰. The  $\delta\text{D}$  values vary from -53 to -83‰ (n=6). ESE veins  $\delta^{18}\text{O}$  values range from 11.7 to 14.9‰ (n=4). The same calculation for the oxygen isotope composition for hydrothermal fluids gives a range from 5.7 to 8.9‰, with an average value of 7.95‰. The  $\delta\text{D}$  are homogeneous, varying from -26 to -36‰ (n=4). NNW tension veins  $\delta^{18}\text{O}$  values range from 13.1 to 14.5‰ (n=4). The same calculation for the oxygen isotope composition for hydrothermal fluids returns a range from 7.1 to 9.5‰, with an average value of 8.25‰. The  $\delta\text{D}$  values range from -47 to -69‰ (n=4).

### 1.5 Discussion

Robert (2001) proposed a distinct classification of some syenitic gold deposits in the Abitibi belt, including Beattie, Canadian-Malartic, Holt-McDermott, Young-Davidson, and Douay. In this model, these deposits represent different components of a large magmatic-

hydrothermal system centered on composite syenitic stocks. New mineralogical studies coupled with geochemical, structural, and 3D modeling data provide a better understanding of the Beattie deposit, from the precipitation and the evolution of gold, to the genetic model.

### **1.5.1 Precipitation and evolution of gold in the Beattie deposit**

A bimodal distribution of gold has been recognized in the Beattie deposit: either invisible gold in arsenian-pyrite and arsenopyrite in the Beattie syenite, or visible gold in microfractures of brecciated pyrite in the Beattie and Donchester fault zones and the Fracture zone.

#### **1.5.1.1 Early invisible gold**

The paragenetic sequence (Figure 1.7) indicates that invisible gold is the only gold appearance in the initial stage of sulfidation. It is related to the As-rich event (Figure 1.13a). Mesothermal vein, epithermal vein, sediment-hosted (Carlin-type), and gold-bearing volcanic-hosted massive sulphide deposits commonly contain invisible gold in arsenopyrite and arsenian-pyrite (Cabri et al., 1989; Cathelineau et al., 1989; Cook and Chryssoulis, 1990; Fleet et al., 1993; Mumin et al., 1994; Palenik et al., 2004; Reich et al., 2005; Cepedal et al., 2008; Martin-Izard and Rodriguez-Terente, 2009; Sung et al., 2009). Arsenopyrite and arsenian-pyrite are the favoured host for invisible gold (Ashley et al., 2000). Fleet et al. (1993) reported concentration up to 1.4 wt% of gold in arsenian-pyrite, whereas Kovalev et al. (2011) described gold value up to 5.36 wt% in fine acicular-prismatic arsenopyrite. However, Cook et al. (2009) concluded that As is not essential for Au to enter in the pyrite structure, but textural criteria including areas of clustered tellurides inclusions, microsheared and brecciated pyrite, as well as pyrite recrystallization would be good factors for high gold values.

The role of arsenian-pyrite and arsenopyrite in scavenging gold has been long recognized, but the scavenging mechanism remains controversial. The most common mechanism is the substitution of elements. Cook and Chryssoulis (1990) proposed that As

substitution in pyrite leads to  $(\text{AsS})_3$  anion pairs. Charge could be balanced by substitution of  $\text{Fe}^{2+}$  by  $\text{Au}^{3+}$ ,  $\text{As}^{3+}$  or  $\text{Sb}^{3+}$ . Fleet et al. (1993) reported that gold is incorporated in arsenian-pyrite as metastable solid solution via As-rich growth surfaces. Cepedal et al. (2008) proposed a mechanism in which gold is removed from ore fluids by chemisorption at As-rich, Fe-deficient surface sites and incorporated into the sulfides as solid solution. Reich et al. (2005) and Martin-Izard and Rodriguez-Terente (2009) argued that Au values below the solubility limit represents structurally bound Au occurring as solid solution ( $\text{Au}^+$ ), whereas high Au values above the solubility limit correspond to nanoparticles of native gold ( $\text{Au}^0$ ). Nanoparticles are usually about 2nm in size (Genkin et al., 1998). Kovalev et al. (2011) suggested the Au-enrichment in arsenopyrite would result of the high defectiveness of its structure, determined by Fe, As, S proportions. Regardless of arsenic, Sung et al. (2009) and Bi et al. (2011) concluded that a positive correlation between Au and Te indicates that gold occurs as submicroscopic Au-bearing telluride inclusions in the host mineral. In the Beattie deposit, there does not appear to be any evidence that associate gold with telluride when gold is invisible. In addition, using TEM data in the gold-bearing arsenian pyrites (Figure 1.12a, b), no gold nanoparticles have been found. However, microprobe assays indicate significant gold values in these pyrites. Hence, the gold speciation for invisible gold in the Beattie deposit is believed to be in solid solution ( $\text{Au}^+$ ) resulting from substitution processes.

In the Beattie deposit, gold-bearing G1 pyrite is characterized by textural characteristics such as high porosity and the local presence of a significant number of micro-inclusions (Figure 1.10a, b, c, d, f). These features indicate that the pyrite has been affected by a new hydrothermal event. The original G1 pyrite would have been transformed into an Au-bearing arsenian-pyrite with spongy areas and metallic inclusions of arsenopyrite, galena, and sphalerite. This stage would mark the enrichment in Au and As. The fluid circulation would thus have been enriched in Au and As compared to the fluid which led to original G1 pyrite crystallization. These features suggest a coupled dissolution-reprecipitation process (Putnis, 2002) in which areas of original G1 pyrite and/or original metallic inclusions are dissolved, followed by reprecipitation. High porosity appears to be an efficient trap for the accumulation of metallic particles. Similar crystal textures are recognized in other deposits (Mumin et al., 1994; Cepedal et al., 2008; Cook et al., 2009; Sung et al., 2009; Linnen, 2012).

Sung et al. (2009) argued this dissolution-reprecipitation process may have been important in upgrading the ore grade by scavenging gold from the hydrothermal fluid.

Several factors including T,  $f_{O_2}$ ,  $f_{S_2}$ , pH, Eh, and fluid composition control the ability of fluids to transport and deposit Au and As (Cathelineau et al., 1989). In the Beattie deposit Au-rich arsenopyrite and arsenian-pyrite crystallize together (Figure 1.7). Atomic proportions indicate arsenopyrite in G1 pyrite is about 27% As (33 atomic % is the regular stoichiometric value for arsenopyrite), which is common for invisible gold-bearing arsenopyrite (Genkin et al., 1998; Vaughan and Kyin, 2004). Using an arsenopyrite geothermometer (Barton, 1969; Kretschmar and Scott, 1976; Sharp et al., 1985), temperature and  $f_{S_2}$  are unquantifiable due to probable disequilibrium crystallization conditions (Figure 1.24). However, Kretschmar and Scott (1976) and Sharp et al. (1985) considered that gold-bearing pyrite and arsenopyrite crystallization is favoured at temperatures inferior to 410°C. The pyrite-arsenopyrite association gives an indication of the conditions of invisible gold deposition with regard to the oxygen and sulfur composition, and the pH (Figure 1.23a, b). The environment would be reduced (Log  $a\sum S$  ranging from -2 to 0, and Log  $f(O_2)$  ranging from -28.8 to -31) with a slightly lower pH compared to initial stages probably caused by the arrival of reduced fluids in the system. Many investigators have shown that Au is abundantly carried as bisulfide complexes (e.g.,  $Au(HS)_2^-$ ) in hydrothermal solutions (Cathelineau et al., 1989; Brown et al., 2003; Cepedal et al., 2008). Desulfidation of the hydrothermal phase caused by fluid-rock interaction, as well as redox variations coupled with a decrease of the oxygen fugacity and pH would result in the destabilization of the Au-bisulfide complexes and the Au deposition. Alternatively, since gold and arsenic could be transported as thioarsenide complexes ( $AuAsS_2$ ), changes in the oxidation-state, pH increase, and/or decrease of  $f_{S_2}$  could cause the simultaneous precipitation of gold and arsenic (Romberger, 1986). Whereas Cameron and Hattori (1987) suggested magmatic oxidized fluid would be the best carrier of gold, Mikucki (1998) considered that reducing conditions are expected to decrease Au solubility and favor the coprecipitation of Au with the newly crystallized pyrite and arsenopyrite. A new model proposes that chalcophile melts (i.e., Bi-rich melts) would act as a gold scavenger from hydrothermal fluid (Ciobanu et al., 2006; Tooth et al., 2011). This could explain the geochemical affinity between gold and low-melting point chalcophile elements (e.g., Bi at 271°C) recognized in several gold deposits. However, the Bi content in the Beattie deposit is

low (usually <2ppm). Consequently, the hypothesis of gold precipitation due to a scavenging from Bi-rich melts does not seem realistic in the Beattie deposit. In fact, the intimate relationship between Au and As in G1 pyrite and arsenopyrite suggests that a thioarsenide complexes ( $\text{AuAsS}_2$ ) would be the most likely gold carrier, as oppose to a bisulphide complexes ( $\text{Au}(\text{HS})_2^-$ ).

### 1.5.1.2 Late fracture control of gold mineralization

After the crystallization of the invisible gold in arsenian-pyrite and arsenopyrite, an intense silica-rich magmatic-hydrothermal phase led to sulfide redistribution in the Beattie and Donchester fault zones and the Fracture zone. This redistribution is indicative of the progressive increase of gold concentration in the main faults zones. Minor recrystallizations of pyrite and arsenopyrite have been described in association with this hydrothermal stage (Figure 1.7), however they are gold-free. The process of gold remobilization is presented in Figure 1.25. Gold in solid solution migrated into microfractures of the brecciated pyrite, and ultimately into the groundmass, where it reprecipitated with silver to form electrum (Figure 1.11a, f). Only minor to insignificant values of invisible gold remain in the arsenian-pyrite and arsenopyrite (Table 1.3). Contrary to the invisible gold, visible gold is not correlated to high As values, but Au content appears to coincide with Ag (Figure 1.13a, b). The expulsion mechanism of invisible gold from the lattice bound of arsenian-pyrite and arsenopyrite and its redeposition as visible gold has been recognized by previous investigators (Bonnemaison and Marcoux, 1990; Mumin et al., 1994; Morey et al., 2008; Cook et al., 2009). The most common phenomena for the mechanism of expulsion could have been the combined action of deformation and hydrothermal circulation, the increase grade of metamorphism, the decrease of refractory properties of host minerals, the decrease in grain size, the increase of fluid temperature and  $f_{\text{S}2}$ , and the increase recrystallization of host minerals and their fracturing/brecciation. In the Beattie deposit, two end-models are highlighted: either open system changes or closed system changes. The open system changes are seen as circulation of external fluids which bring new elements, whereas closed system changes are recognized as mineral transformations only. In the Beattie deposit, post depositional processes associated

with strong hydrothermal alteration and crystallization of new minerals indicate that predominant open system changes affected significantly the speciation of gold.

The hydrothermal conditions of T,  $f_{O_2}$ ,  $f_{S_2}$ , pH, Eh and fluid type can be determined using phase diagrams. The new arsenopyrite is still As deficient with atomic proportions that indicate 28 to 30.5% As. Hence, using an arsenopyrite geothermometer (Barton, 1969; Kretschmar and Scott, 1976; Sharp et al., 1985), temperature of recrystallization is thought to be  $\leq 350^{\circ}\text{C}$  (Figure 1.24). In addition, the silica alteration, as well as low  $f_{S_2}$  ( $<-8.5$ ), sulfate crystallization (e.g., barite) and late-hydroxidation (e.g., goethite) indicate the oxidizing of hydrothermal conditions at low pH (Figure 1.23a, b).

The silica-rich magmatic-hydrothermal phase has carried and deposited new elements along the Beattie and Donchester fault zones, and the Fracture zone (Figure 1.25). These elements include Hg, Te, Mo, Ag, Sb, As. Pyrite is brecciated and clustered with local alteration zones at its margins. Similar to the dissolution-reprecipitation process which occurred in G1 pyrite for the invisible gold, microfractures and remaining porosity acted as traps for new metallic minerals such as enargite, coloradoite, tennantite-tetrahedrite, hessite, and electrum (Figure 1.11a, c, d, e, g).

### 1.5.2 Proposed genetic model

Based on field evidence, mineralogical and geochemical studies, new conclusions can be drawn on the evolution of the Beattie deposit with regards to the timing of gold deposition with deformation and alteration.

(1) Folding and tilting of the Kinojevis volcanic layers ( $D_1$ ) was followed by the Timiskaming event, a period of basin formation and sediment deposition which was contemporaneous to the emplacement of alkaline intrusions. The Beattie syenite was emplaced during the last stage of this first deformation period (Mueller et al., 1996; Legault et al., 2005). Mineralogical data suggest that initial syenitic stages were uniformly oxidized (e.g., magnetite, sulfates). Liquid immiscibility remains a possible explanation for entrapment of apatite as inclusion in silicates. Then, an excess of Ba and S in the immiscible  $\text{PO}_4$  liquid (Panina and Motorina, 2008) would have formed barite inclusions in the apatite. The

evolution of the system consists of an in-situ oxidation, the hematization that was not metasomatism-related as the element concentration is similar to the initial phase. At this stage, no gold mineralization was present.

(2) The subsequent step was dominated by CO<sub>2</sub>-rich reduced fluids compared to the initial stages (Figure 1.23b). Destabilization of previous Fe and Ca-rich minerals (e.g., magnetite, hematite, and titanite) would have been in part the source of elements for iron-carbonate crystallization. A more reduced and S-rich fluid would have produced pyrite, arsenopyrite, galena, and sphalerite. Invisible gold-bearing arsenian pyrite and arsenopyrite are the key minerals of this stage. Consistent with field evidence which indicates that the sediments unconformably truncate mineralized stockworks in the Beattie syenite (Robert, 2001), and which does not display any invisible gold mineralization in the Duparquet formation sediments, the invisible gold must have been deposited before the Timiskaming sediment deposition (pre-D2). This mineralization step was slightly Bi-rich in comparison to the remainder of the paragenetic sequence (Figure 1.14). This highlights the dominance of magmatic fluids (Baker et al., 2005). The source of arsenic in the Beattie deposit is still controversial, however, regional studies (Dinel et al., 2005; *J.Goutier, pers. comm.*) indicate that the sediments of the Porcupine group would be As-rich. This could be the source for As in the As-rich gold deposits near this sedimentary unit such as the Beattie deposit. Therefore, contrary to well-known gold deposits in alkaline rocks where metals originate from magmatic fluids only (Mutschler, 1985), the concentration of gold and arsenic in the Beattie deposit would have resulted from mixing between external fluids and magmatic fluids. It was an open system regime in which this mixed Au-As-rich fluid would have resulted in the sulfidation stage, at which point precipitation occurred. Hence, the early gold-bearing fluids, that were responsible for the crystallization of the invisible gold in the lithologically-controlled mineralization, are interpreted as mixed magmatic and external fluids having circulated during the pre-major shortening event (D2).

(3) The remaining stages in the genetic model happened during the transition D2-D3 at a period that reached the metamorphic peak. Both the silica-rich magmatic-hydrothermal fluid circulation and the movement along the Beattie and Donchester fault and the Fracture zone were very intense. The sulfides were redistributed within the faults zones and gold was

remobilized in microfractures of the brecciated pyrite. The breccia in the Beattie deposit can be classified by the propagation and the mobility processes (Appendice C; Jébrak, 1997): A primary breccia mechanism comprised a hydraulic propagation which was associated with an insignificant mobility process considering that the motion is in-situ. A subsequent breccia formed through tectonic propagation coupled with grinding/milling mobility. This second breccia process is commonly observed in the deposit along the faults zones. It led to a decrease of the fragment size, and an increase in the motion of the fragments. These hydrothermal and tectonic processes favoured the expulsion of the invisible gold from lattice bounds of arsenian sulfides, and its precipitation as visible gold in the microfractures of brecciated pyrite. The NNW-SSE tension veins, which post-date mineralized breccia and ENE cherty veins, coincided with the late north-south shortening in Abitibi (Late-D3). Hence, the silica and metallic-rich magmatic-hydrothermal event, which caused the sulfide redistribution within the faults zones and the gold remobilization, happened between D2-D3 transition and possibly until late D3 deformation.

Based on 3D modeling data and chemical characteristics of sericitized rock at 300°C, 313.4 million cubic metres of fluid is required to alter 9.9 million cubic metres of rock (Appendice F). The required volume proportion of fluid is roughly the same as the Butte deposit, where 100 billion cubic metres of magmatic fluid are required to alter 4.5 billion cubic metres of rock (Cathles and Shannon, 2007). Therefore, the sericite alteration in the Beattie deposit is similar to a closed system dominated by magmatic fluids in a porphyry deposit. Also, it is believed that a portion of the potassium came from the alteration of pre-existing feldspars. The timing of sericite crystallization is thought to be early during the transition D2-D3. Unlike the sericitization, the silicification is clearly an open system as the strong increase in silica (enrichment of 17.74% in silica) could not come from the magmatic system. Based on 3D modeling data and chemical characteristics of silicified rock at 300°C, 388 million cubic metres of fluid are required to silicify 4 million cubic metres of rock. Silicification is most likely occurred during peak metamorphism, when the hydrothermal circulation in the plumbing was most intense.

The veins clearly formed before (e.g., ENE cherty veins, ESE flat extension veins) or during (e.g., NNW tension veins) the late-D3 deformation (Table 1.5). Regardless of the type

of vein stable isotope signature of  $\delta^{18}\text{O}$  ( $>5.7$  and  $<10.1\text{\textperthousand}$ ) is roughly similar, whereas the  $\delta\text{D}$  value is more discriminating (Table 1.6; ENE cherty veins average -66, ESE flat extension veins  $<-36$ , NNW tension veins average -60). McCuaig and Kerrich (1998) considered that  $\delta^{18}\text{O}$  values greater than  $+8\text{\textperthousand}$  cannot be magmatic fluids alone. Hence, data for ENE cherty veins suggests that magmatic fluids could be the dominant source for the fluids (Figure 1.26); however, a metamorphic component is likely to be present too. NNW tension veins share both magmatic and metamorphic characteristics, whereas ESE flat extension veins only have metamorphic fluids signature (Figure 1.26). Age constraints on the field and isotope data highlight the metamorphic feature of the ESE flat extension veins. They appear similar to the gold-bearing veins of the Sigma deposit (Robert, 1994) as they emplaced during metamorphic peak. However, the flat extension veins are not mineralized at the Beattie deposit. McCuaig and Kerrich (1998) argued that the range in  $\delta\text{D}$  values may be due to the presence of heterogeneous fluids or the nature of the fluids inclusion (primary vs. secondary). The calculated oxygen and hydrogen isotope for fluids in Archean lode gold deposit was restricted to 6 to  $11\text{\textperthousand}$   $\delta^{18}\text{O}$ , and -30 to  $-80\text{\textperthousand}$   $\delta\text{D}$  (McCuaig and Kerrich, 1998; Neumayr, 2007). In addition, Legault et al. (2005) identified  $\delta^{18}\text{O}$  between 13.2 and  $14.7\text{\textperthousand}$  for deposits similar to the Beattie deposit along PDFZ. These values are consistent with those from the Archean quartz veins in the Beattie deposit.

The obliquity of the two stress tensor ( $\sigma_1$  plunging  $35^\circ$ ; Appendix E; Figure 1.21c) suggests ESE tilting of the mineralized syenite after the late-faulting event. Tilting hypothesis is supported by 3D modeling in which the  $35^\circ$  plunging of the units is visible (Appendix B; Figure 1.27); the plunging was observed in the Beattie underground mine as well (Davidson and Banfield, 1944). The gold mineralization is still present at the base of the model (Figure 1.27), that supposes the mineralization continue in depth. Hence, the tilting could have had a significant part in the preservation of the deposit.

### 1.5.3 Comparison

The classification of the Beattie deposit is not unequivocal. Davidson and Banfield (1944) argued that the Beattie deposit should be classified as mesothermal or epithermal,

whereas Robert (2001) and Legault et al. (2005) classified the Beattie deposit as an intrusion-related gold deposit.

According to this study, the Beattie deposit shares characteristics with several types of deposit. Regarding the felsic host rock of the mineralization, the dominance of lithologically-controlled facies in which mineralization is disseminated and in stockworks, the relatively low-grade bulk-tonnage mineralization related to sulfides, the As-rich assemblage, and the sericite and carbonate alteration, the Beattie deposit looks like an intrusion-related gold deposit in accordance with Lang and Baker (2001), and Hart (2005, 2007). However, some features are atypical of intrusion-related gold deposits, particularly the oxidizing environment, the pervasive alteration, and the characteristics of the structurally-controlled facies. The mineralization in the structurally-controlled facies is significantly different. The very low Au/Ag ratio  $\approx 0.8$ , the strong enrichment in Hg, Mo, Te, Ag, As, and Sb, the metallic mineralogy (e.g., enargite, tennantite-tetrahedrite, coloradoite, hessite, electrum), the breccia ore and mineralized veins, the strong silicification, sericitization, and local kaolinization, the trachytic and pseudo-colloform textures, and the clear control of faults suggest that the Beattie deposit finds analogies with low to intermediate sulfidation epithermal deposits, according to Sillitoe and Hedenquist (2003) and Corbett (2009). However, some of these features are also typical of the apical part of mesothermal deposit (Groves et al., 1998). The coexistence of intrusion-related gold deposit signatures, epithermal deposit features, and mesothermal deposit characteristics in a same environment would be the result of an evolution from a magmatic dominant system to a shallow hydrothermal dominant system. Two cases are possible: (1) The mineralized intrusion (lithologically-controlled facies) would have been overprinted by mesothermal shear zones (structurally-controlled facies) as a result of the D2-D3 metamorphic event. (2) In a large magmatic environment, the porphyry system (lithologically-controlled facies) would have been overprinted by an epithermal system (structurally-controlled facies) within shear and faults zones during the D2-D3 contractional event. This transition from porphyry to epithermal has already been recognized (Barron, 1998; Richards, 1992; Müller et al., 2002; Williams-Jones and Heinrich, 2005; Sillitoe, 2010; Cooke et al., 2011), and can provide important metal remobilization and reprecipitation events. This latter case appears the most probable for the Beattie deposit.

Based on the geometry of the deposit and the mineral association and occurrences, some Archean Canadian gold deposits share analogous characteristics with the Beattie deposit. Two examples in the Abitibi belt include the Douay deposit and the Young-Davidson deposit, along the major Casa-Berardi-Douay-Cameron fault zone in the Joutel gold district and along the Larder-Lake-Cadillac fault zone in the Matachewan gold district, respectively. The Douay deposit appears to be the most similar deposit to the Beattie deposit. It is an oxidized porphyritic syenite-hosted gold deposit in which gold-mineralization is pyrite related and associated with silicification, carbonatization, albitization, and sericitization. Ore bodies consist of disseminated low-grade material set both in the syenite and along the sheared lithological contacts (Dupéré and Duplessis, 2007). The mineral paragenesis consists of early magnetite and hematite which evolved in gold-bearing pyrite, followed by a cataclastic event which led to liberation of gold (Gauthier and Harnois, 1994). The Young-Davidson deposit is an oxidized syenite-hosted gold deposit which intruded the Timiskaming sediments. Mineralization is structurally-controlled as veins. Several generations of pyrite are recognized and some are gold-bearing. The first generation displays a spongy core with a myriad of metallic inclusions. Gold occurs as visible both as inclusion and in the fractures of the pyrite (Linnen, 2012). The Archean Beattie deposit also shares analogous features with more recent deposits, more particularly the Mesozoic and Cenozoic gold deposits associated with the alkaline magmatism in the American cordillera (Mutschler, 1985), or even with the Oligocene Cripple Creek deposit in Colorado (Thompson et al., 1985).

## 1.6 Conclusion

The ore assemblage from the Beattie deposit is dominated by disseminated pyrite, both As-rich and As-free, and arsenopyrite. Gold appears as both invisible and visible. Invisible gold occurs in solid-solution in the spongy As-rich pyrite (G1) and arsenopyrite within the strongly iron-carbonatized Beattie syenite. This mineralization facies is lithologically-controlled. Visible gold is associated with a later-stage alteration event, dominated by silicification, which produced sulfide redistribution within the faults zones

(e.g., the Beattie fault, the Donchester fault, the Fracture zone) and the veins (e.g., ENE-WSW cherty), and gold remobilization in microfractures of brecciated pyrite. Gold crystallized as electrum, and arsenic is not positively correlated to gold in this case. This mineralization facies is structurally-controlled, and related to significant enrichment in Hg, Te, Mo, Ag, Sb, As, and Se.

The deposit evolved from oxidizing (e.g., magnetite, sulfates, hematite) to reducing (e.g., pyrite, arsenopyrite) conditions. In addition, the hydrothermal system progressed from a CO<sub>2</sub>-rich magmatic fluid widely present in the deposit, to Si-rich magmatic-hydrothermal fluids limited to the fault zones. These differences in distribution of alteration highlight a progressive reorganization of the hydrothermal system. Maturity of an evolving fluid, marked by changes of redox condition, pH,  $f_{\text{S}2}$ , and a fluid mixing would have caused the decrease of the gold solubility in the fluid and then its precipitation as invisible state in the As-rich sulfides within the Beattie syenite. One unique magmatic gold event occurred in the deposit, and is associated with the first crystallization of pyrite and arsenopyrite. Subsequent physico-chemical modifications (e.g., increase in temperature and  $f_{\text{S}2}$ , brecciated textures), in an active tectonic and hydrothermal environment during metamorphic peak, would have caused the expulsion of about 15% of the overall gold from lattice bounds, and their redeposition as visible gold within the E-W fault zones. The speciation of gold in the Beattie deposit is a consequence of the dominance of open system regimes, both for the incorporation of arsenic that control early gold deposition, and for the circulation of hydrothermal silica-rich fluids which dominated the late fracture-control gold mineralization.

This study has provided a greater understanding of the metallogenesis of the Beattie deposit. However, several aspects need to be clarified, particularly the precise structural setting of the deposit, the exact age of the gold event (Re-Os dating on pyrite) as well as nature of the gold-bearing pyrite ( $\delta^{34}\text{S}$ ), or even the possible link with the lamprophyres.

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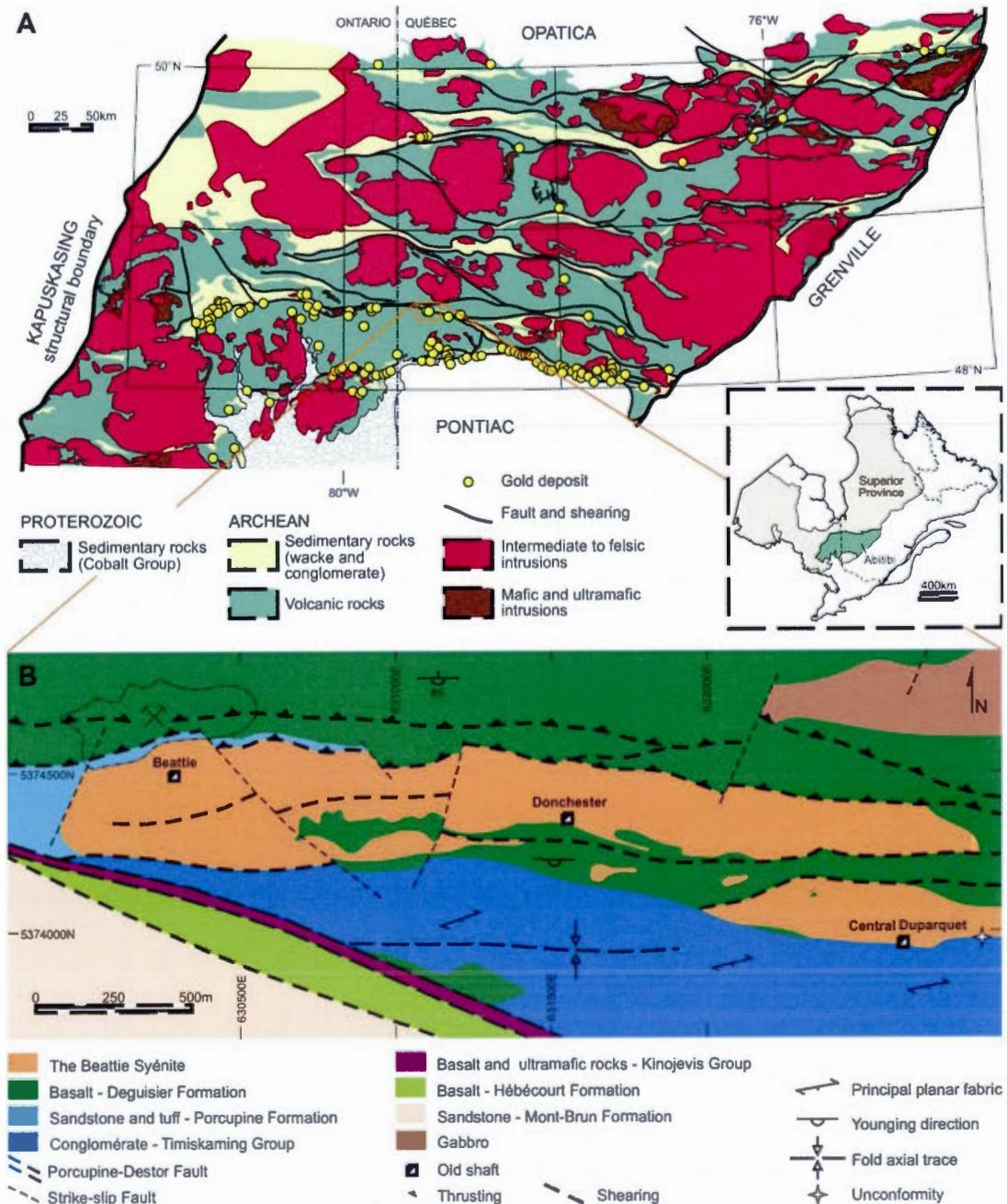
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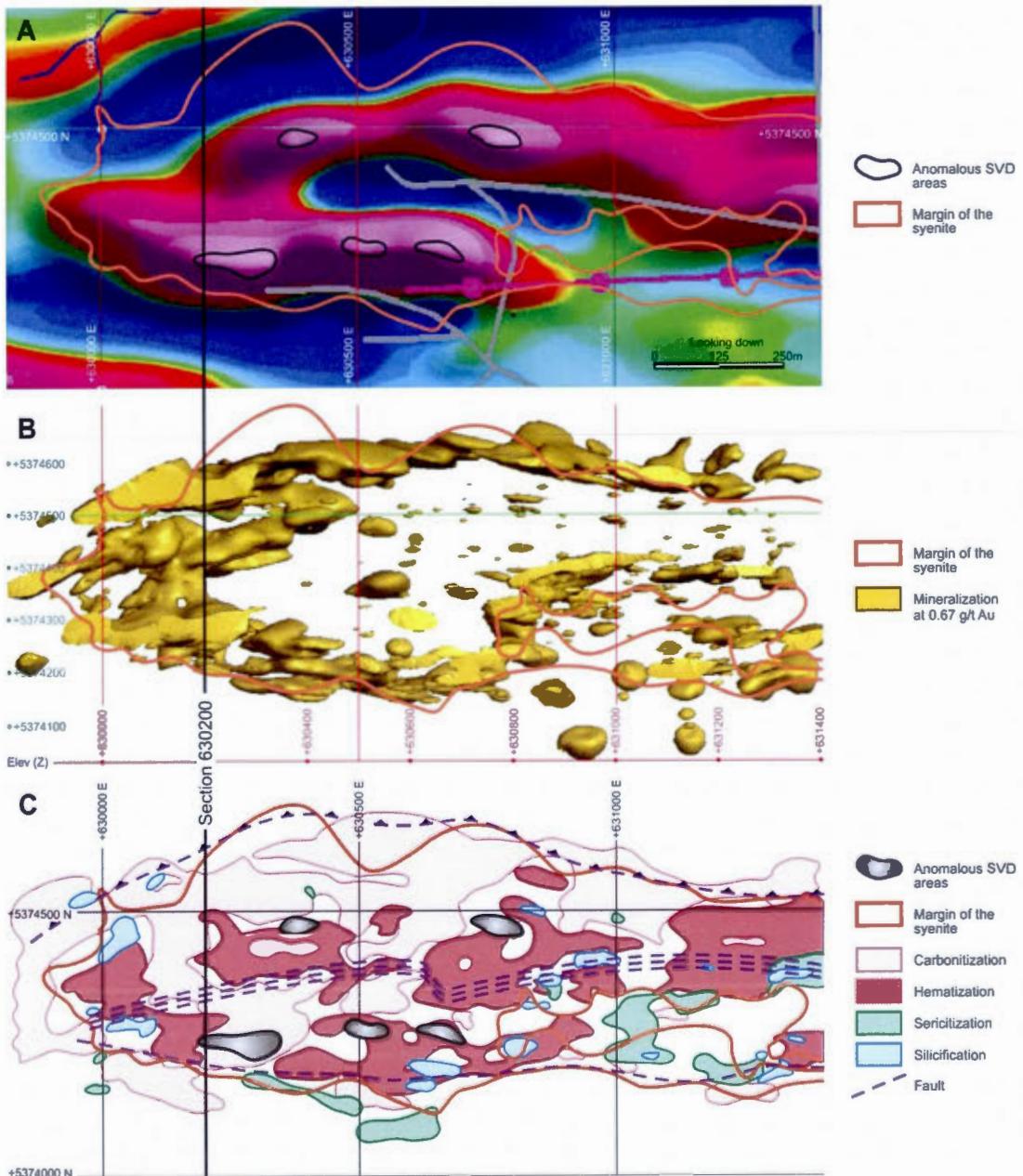
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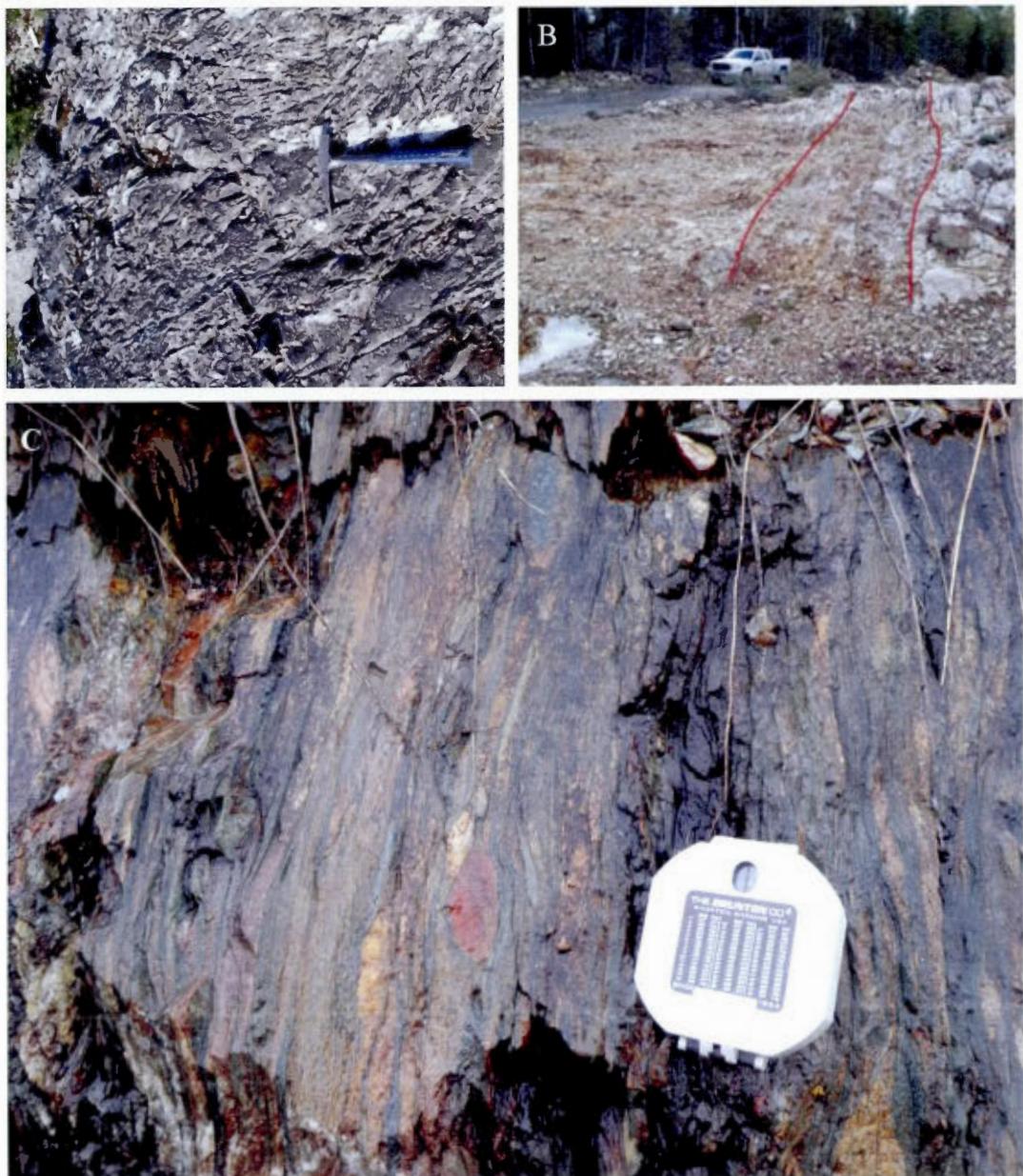
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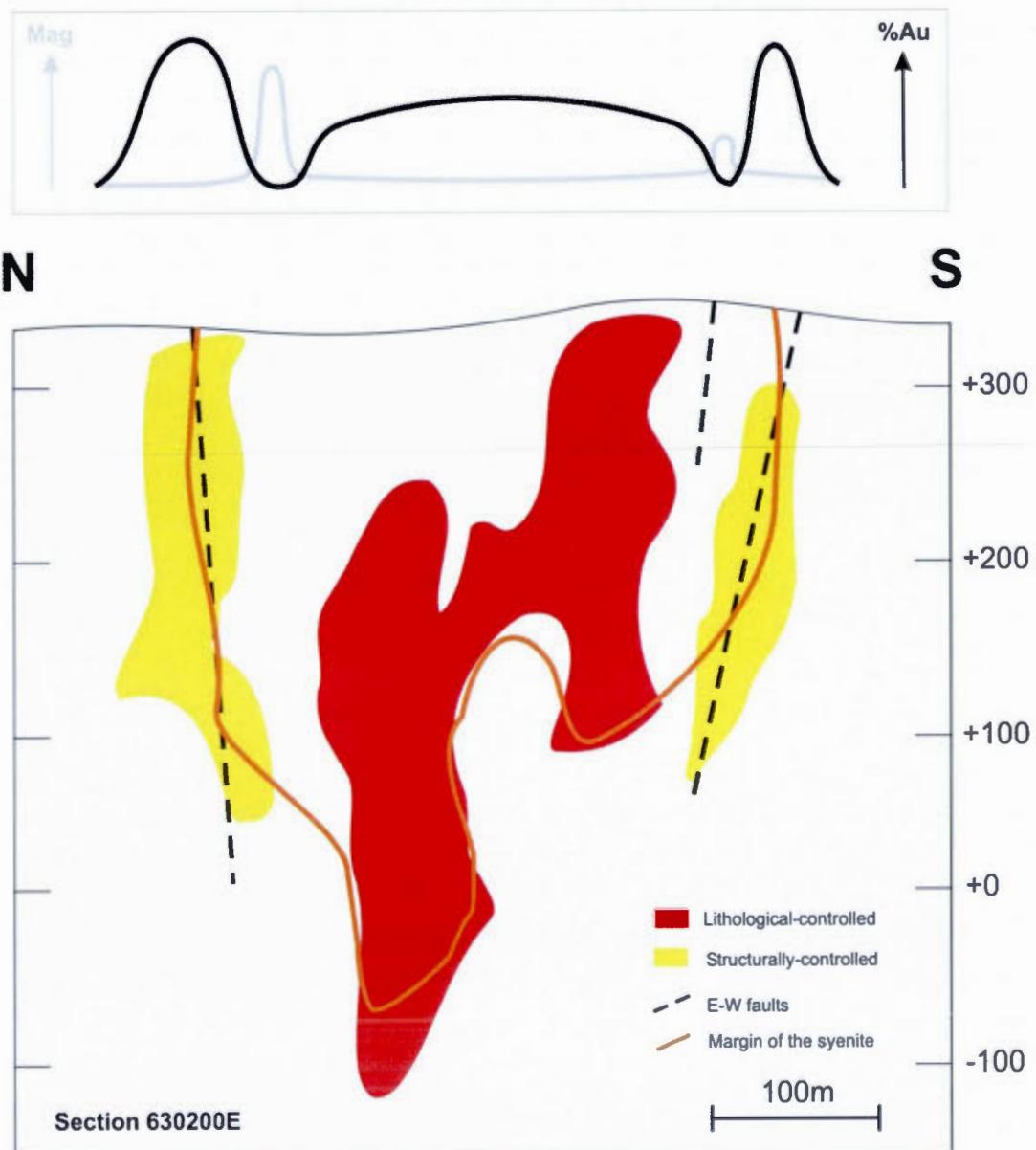
**Figure 1.1.** (a) Simplified geology of the Abitibi greenstone belt (modified from Doucet et al., 2000). (b) Geology of Duparquet property (modified from Mueller et al., 1996, and Bourdeau et al., 2011).

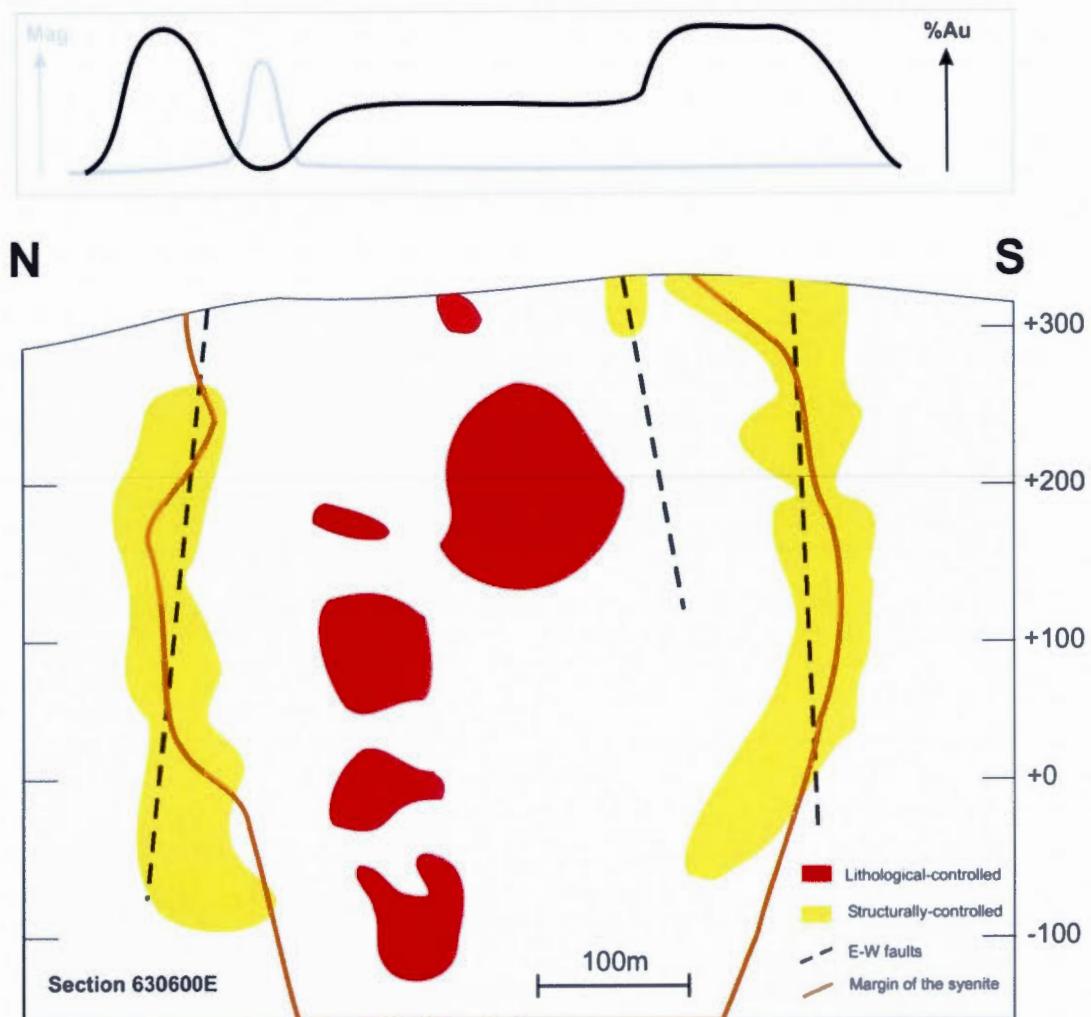


**Figure 1.2.** Comparison at the same scale: (a) Airborne Geophysics (SVD). (b) Mineralized zones in the Beattie deposit at 0.67g/t Au. (c) Alteration halos on the surface, with the Beattie fault (north), the Donchester fault (south), and the Fracture zone (center of the syenite), and with anomalous SVD areas.

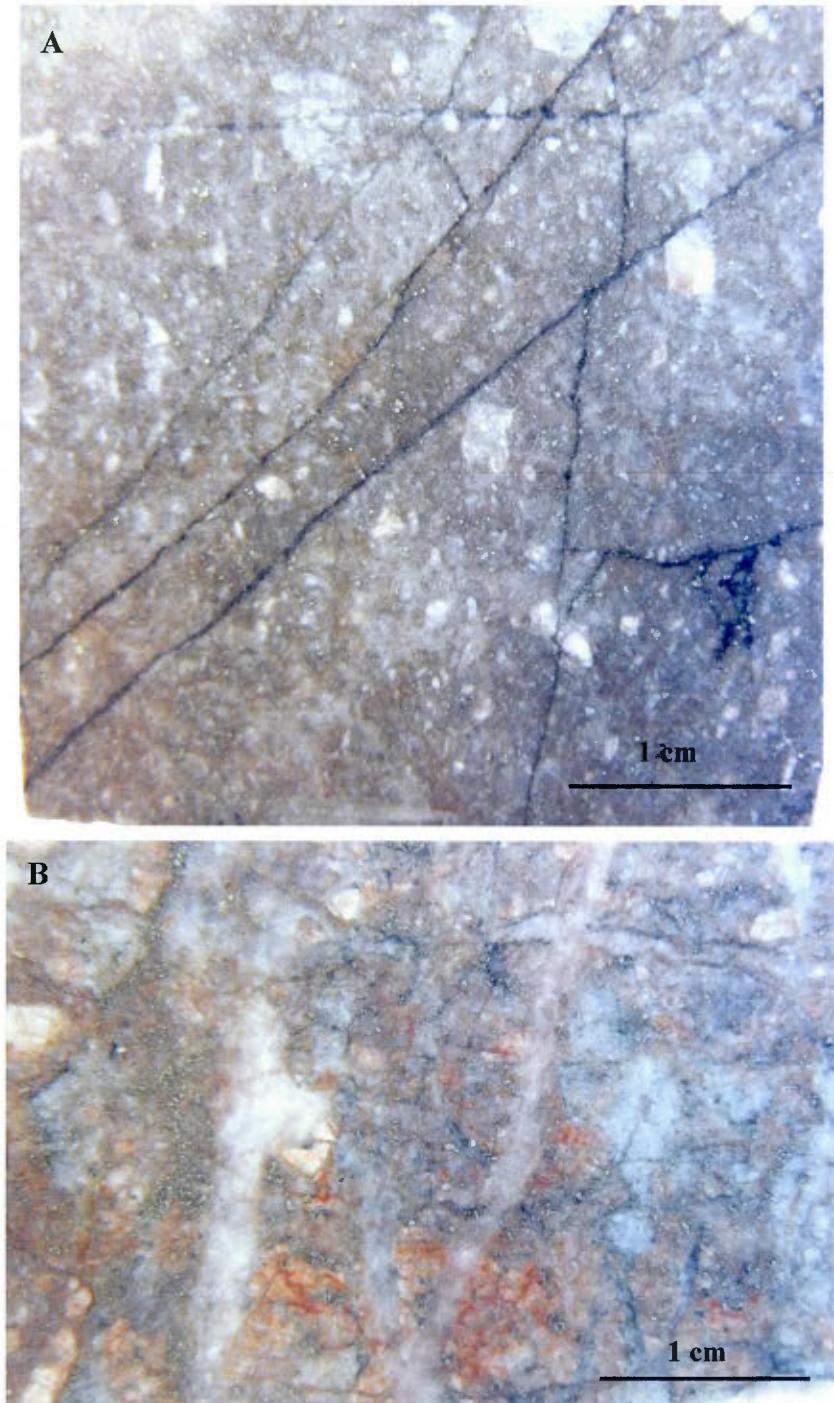


**Figure 1.3.** Photographs illustrating structural particularities of the Beattie deposit. (a) Joint network well developed. (b) The Fracture zone ENE trending. (c) E-W flattening in the Duparquet sediment formation.

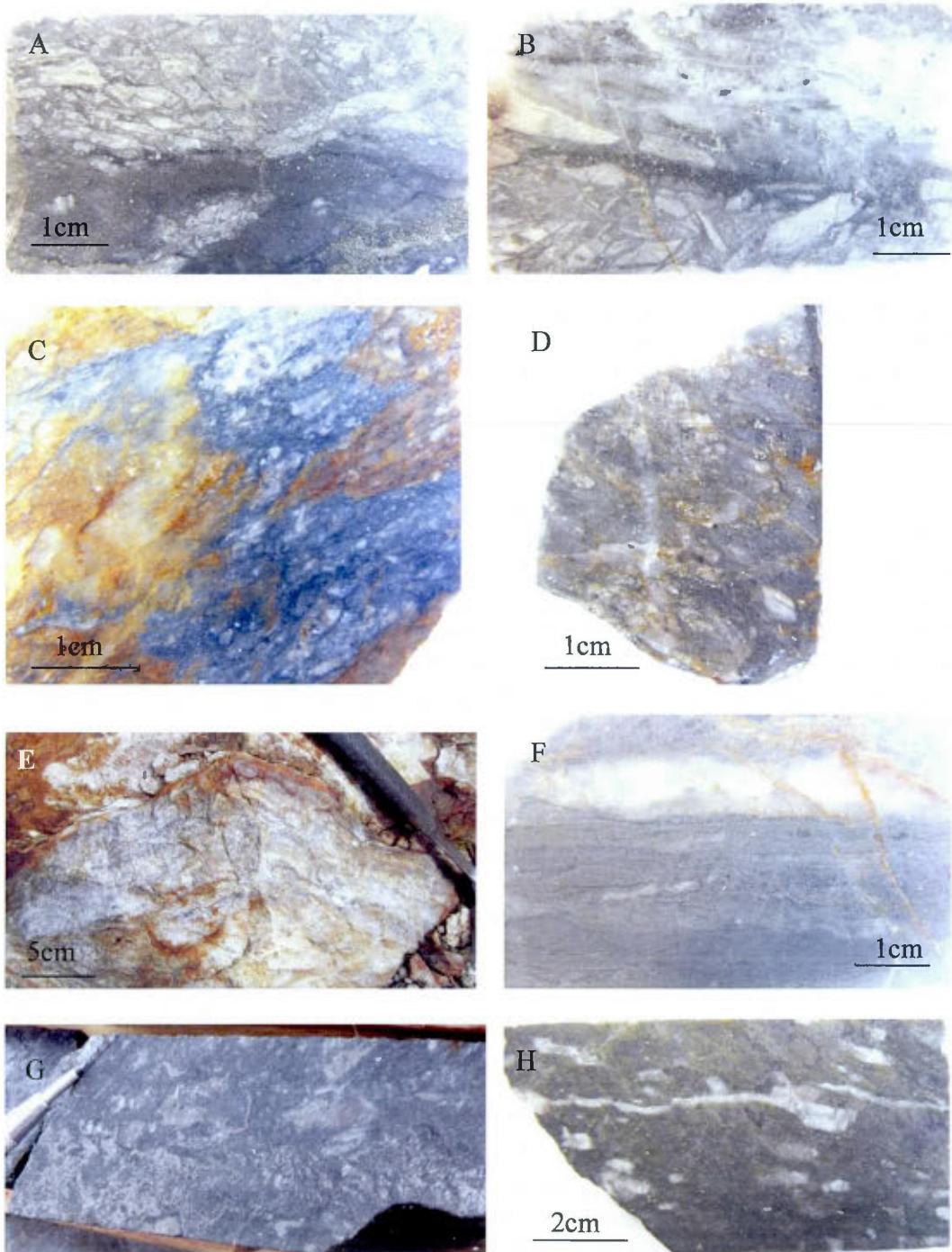




**Figure 1.4.** Two cross sections in the Beattie deposit (630200E, and 630600E) showing the control of the mineralization, lithological and structural. Cartoon based on the 2008, 2009, and 2010 drilling program.

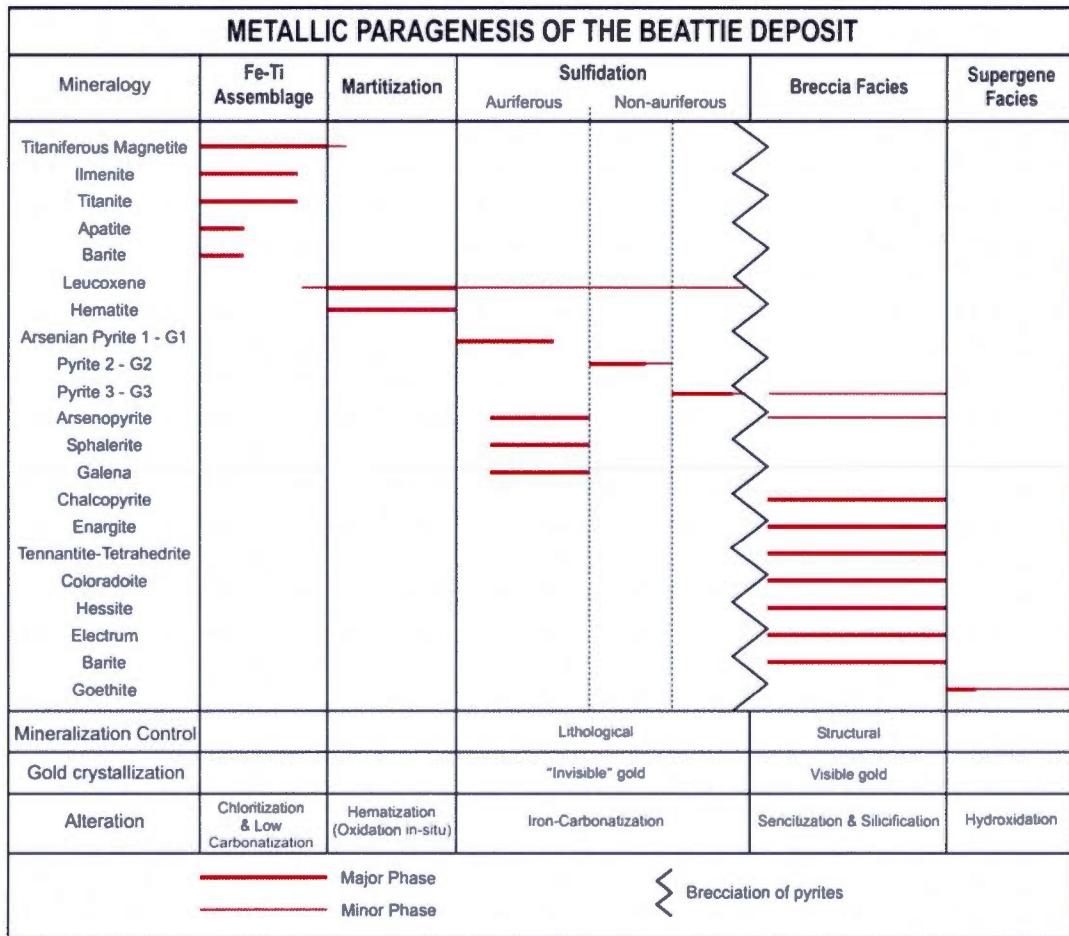


**Figure 1.5.** Photographs of the syenite illustrating the lithologically-controlled facies. (a) Grey massive syenite, carbonatized, with disseminated sulfides (1.5g/t Au). (b) Reddish syenite with disseminated sulfides, and cut by carbonate veinlets (2.4g/t Au).

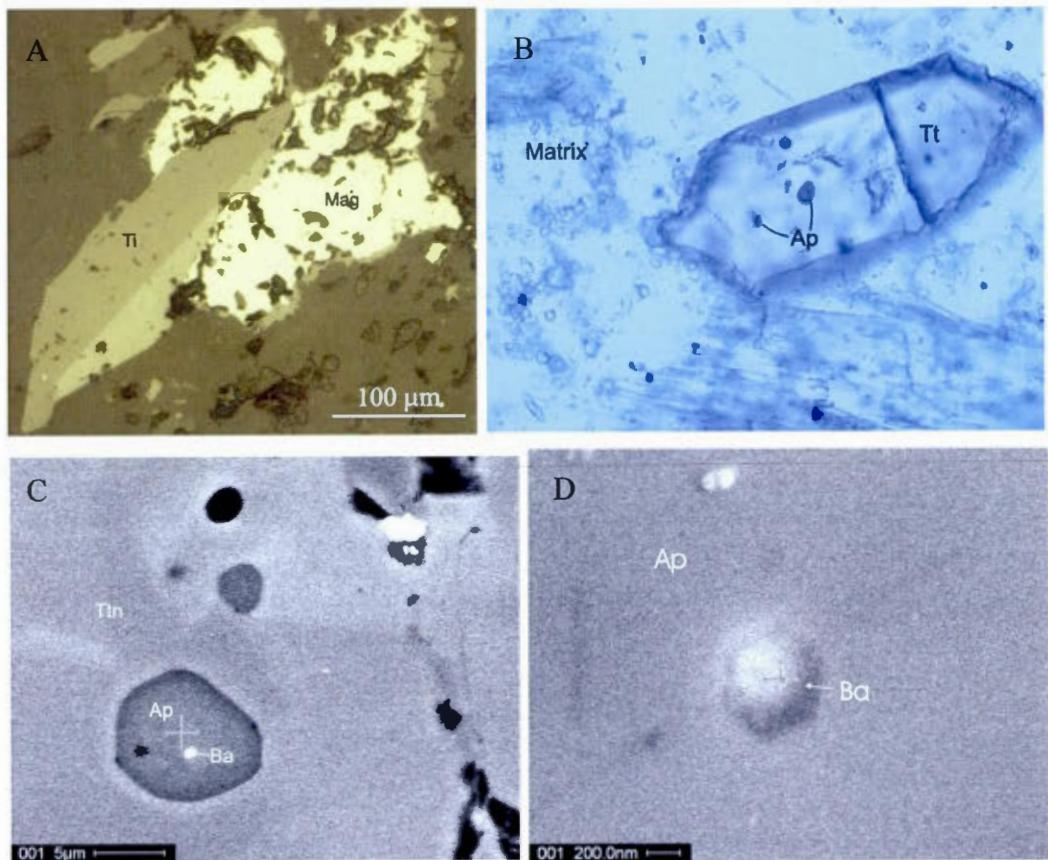


**Figure 1.6.** Photographs of rock samples illustrating the structurally-controlled facies. (a), (b), (c), and (d) Breccia ore with disseminated and small cluster of sulfides, very strong silicification (A: 21.9g/t Au; B: 17.45g/t Au; C: 9.7g/t Au; D: 18.6g/t Au). (e) and (f) Cherty quartz veins with fine sulfides (10.6g/t Au). (g) Semi-massive polymetallic high-grade vein

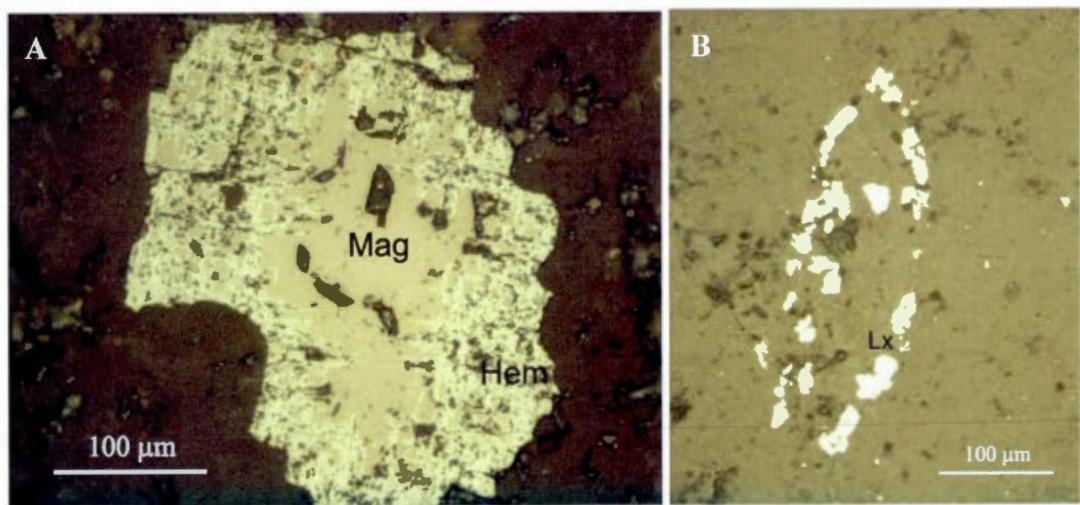
(120g/t Au), related to important silicification. (h) Trachytic syenite dyke with preserved phenocrysts in a carbonatized matrix (similar as the grey massive syenite – Fig.5A), and with disseminated sulfides (5.6g/t Au).



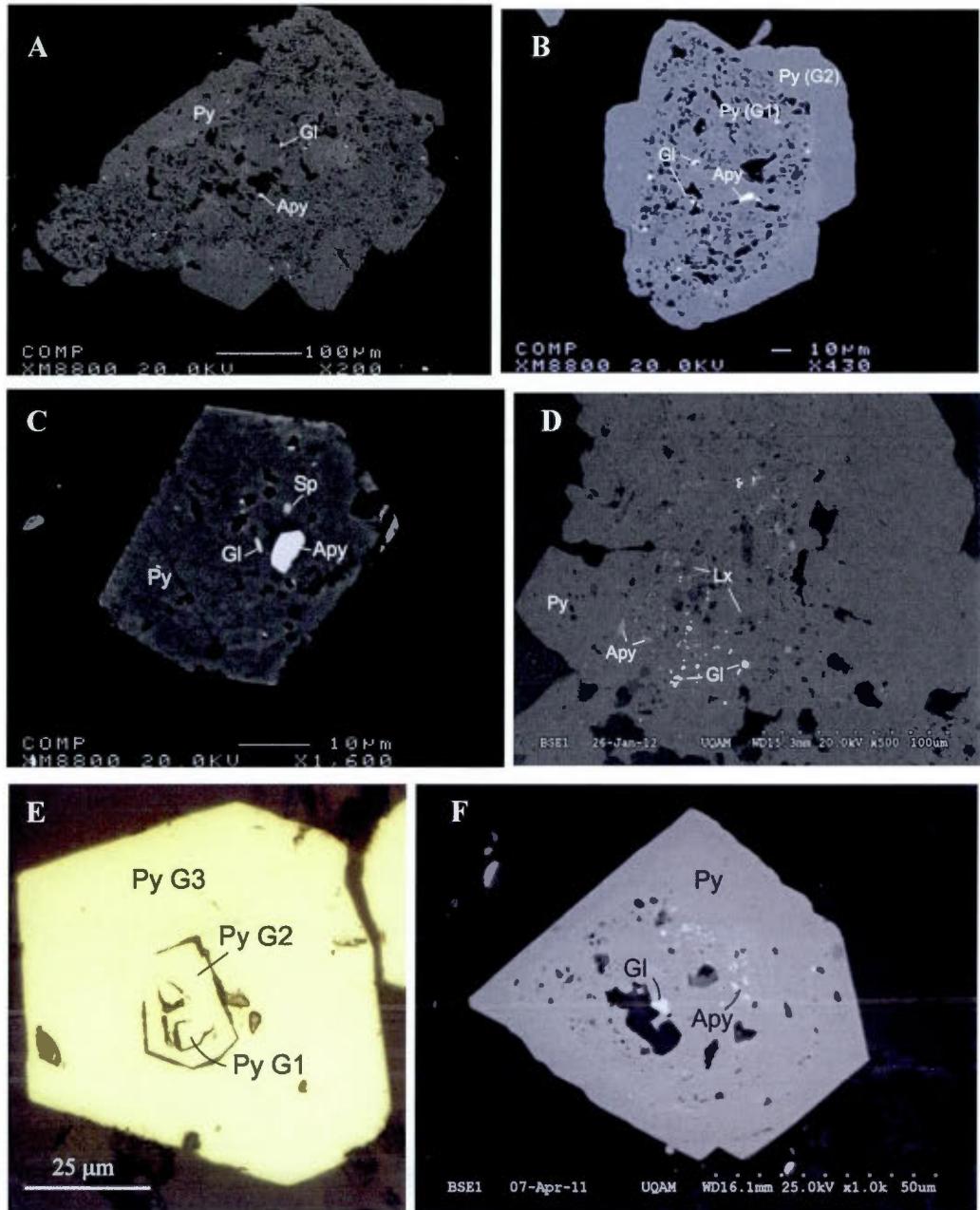
**Figure 1.7.** Metallic paragenesis of the Beattie deposit.



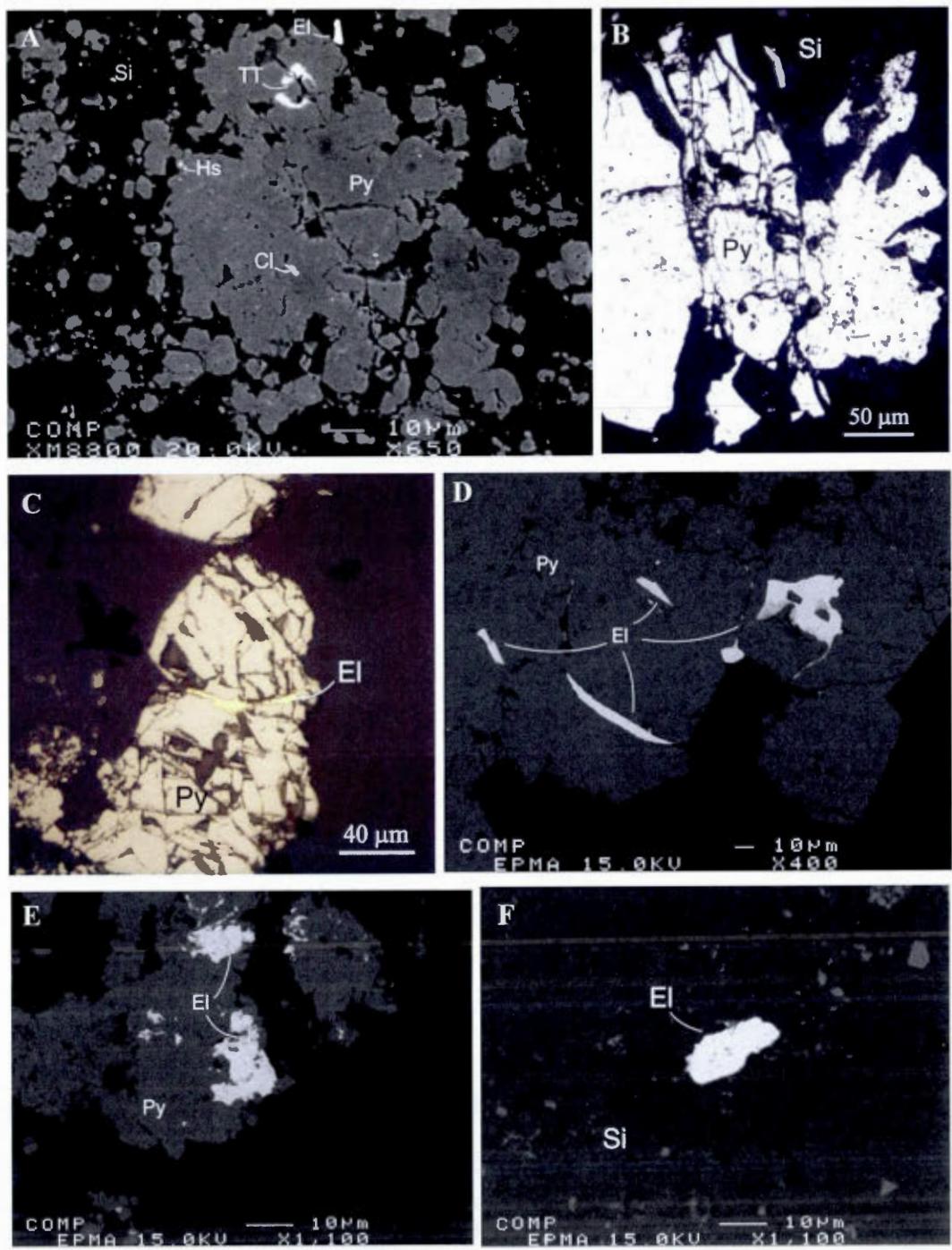
**Figure 1.8.** Microphotographs of the Fe-Ti assemblage. (a) Magnetite and titanite. (b) titanite crystal with mineral inclusions of apatite and barite. (c) and (d) Barite in inclusion within an apatite crystal included in titanite.

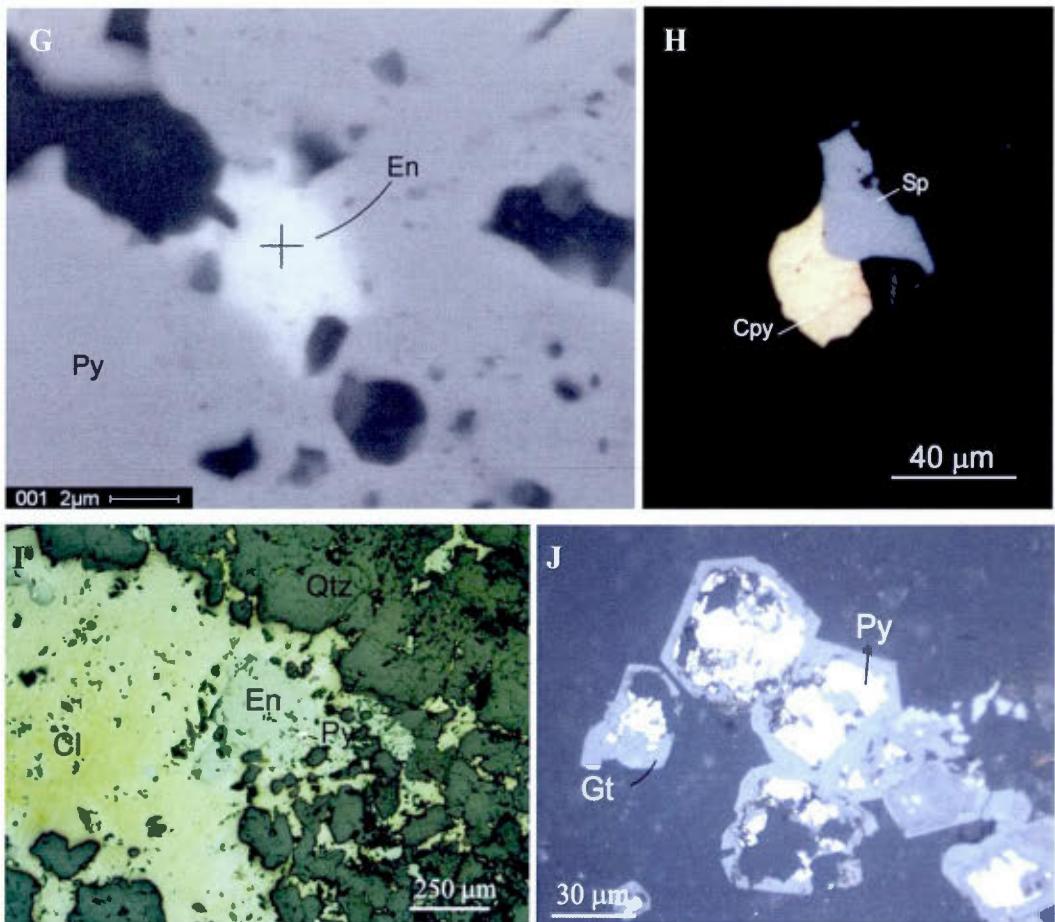


**Figure 1.9.** Microphotographs of the Martitization phase. (a) Exsolution of hematite in a magnetite crystal. (b) Leucoxene as pseudomorph of titanite crystal.

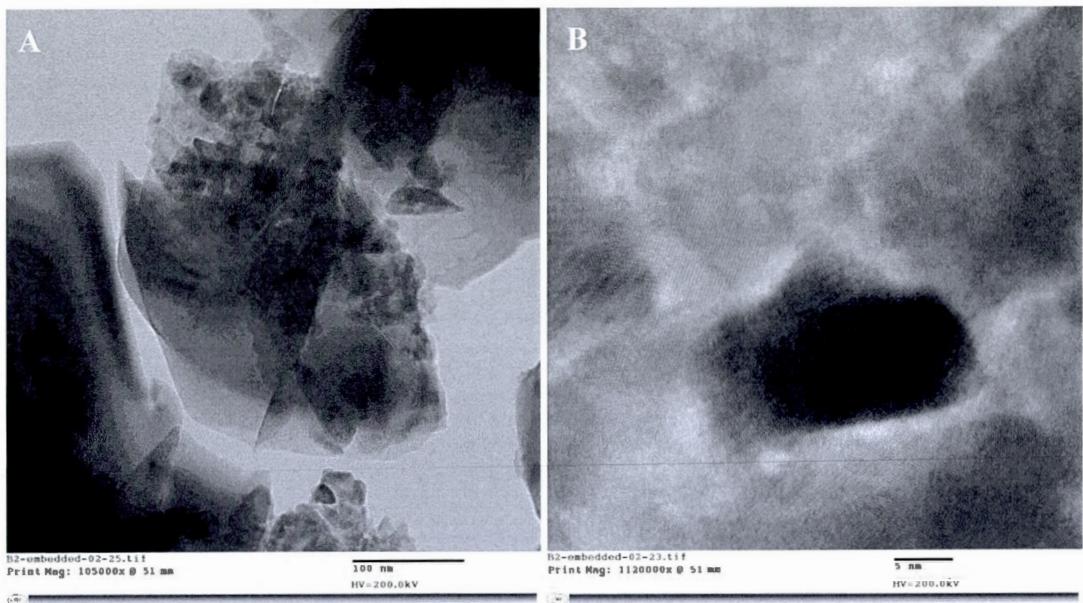


**Figure 1.10.** Microphotographs of the Sulfidation phase (lithologically-controlled facies). (a) and (b) Typical gold-bearing spongy pyrite (picture A: 200 ppm Au in mineral; picture B: 180 ppm Au in mineral) with metallic inclusions of arsenopyrite and galena (G1 Pyrite); G2 overgrowing pyrite gold-free and without porosity. (c) G1 pyrite with inclusions of arsenopyrite (420 ppm Au in mineral), galena, and sphalerite. (d) Pyrite's core with a myriad of inclusions of leucoxene, galena, and arsenopyrite, in addition to porosity. (e) Three generations of pyrite. (f) Porous pyrite core (G1) with arsenopyrite and galena inclusion, and G2 clean pyrite.

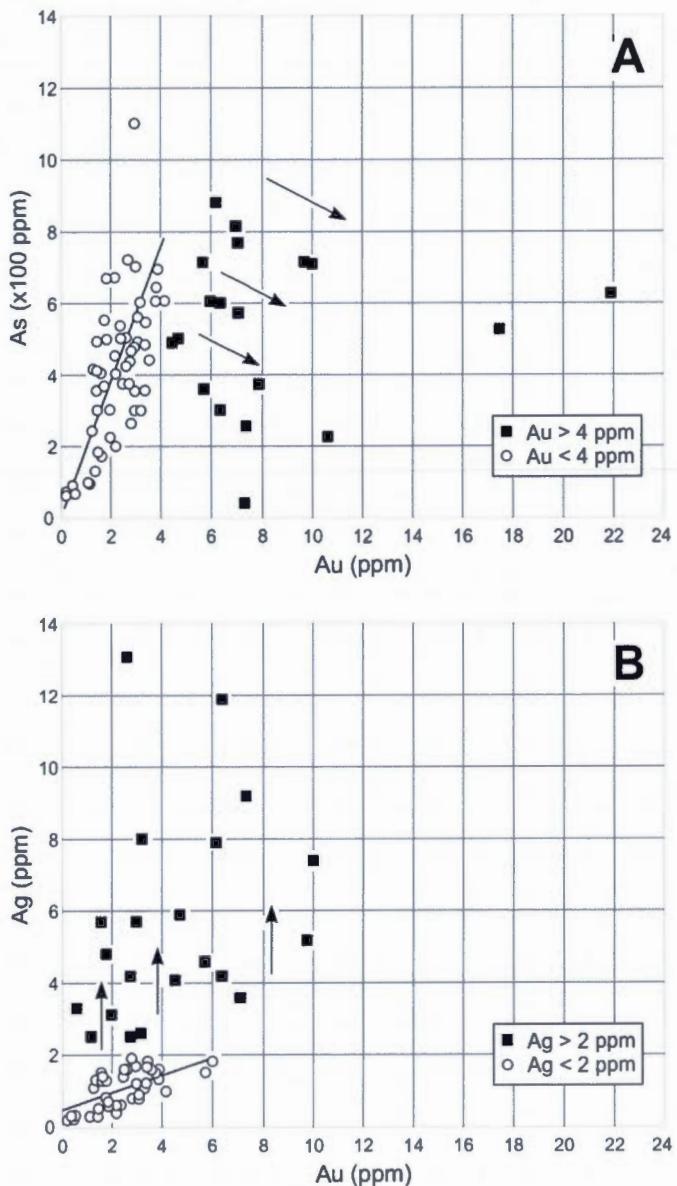


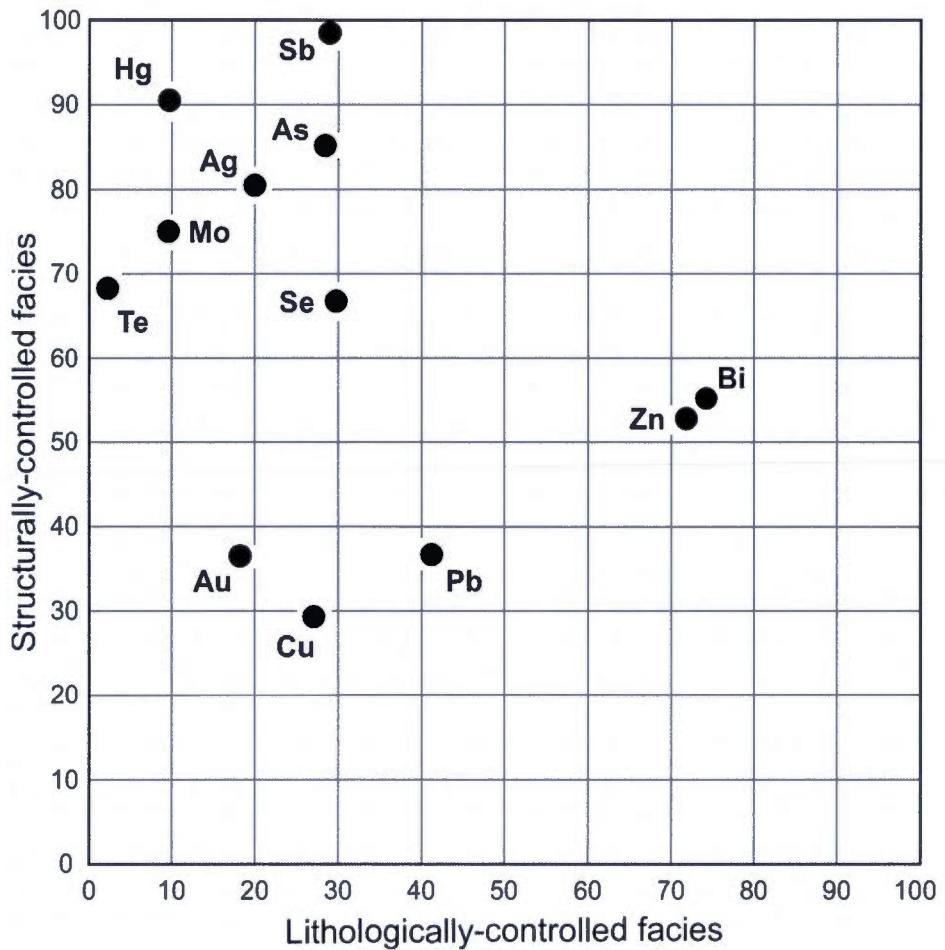


**Figure 1.11.** Microphotographs of the Breccia phase (structurally-controlled facies). (a) Brecciated pyrite with crystals of tennantite-tetrahedrite, coloradoite, and hessite in microfractures. Electrum is free in the matrix. (b) Hydraulic brecciation of pyrite in silica corridors. (c), (d), and (e) Electrum crystal in microfractures and porosity of brecciated pyrite in silica corridors. (f) Free electrum in the groundmass within silica corridors. (g) Enargite crystal trapped in porosity of pyrite. (h) Chalcopyrite and sphalerite in the matrix. (i) Enargite, coloradoite, and pyrite as semi-massive veins with high-silica content. (j) Coronitic texture of goethite around pyrite.

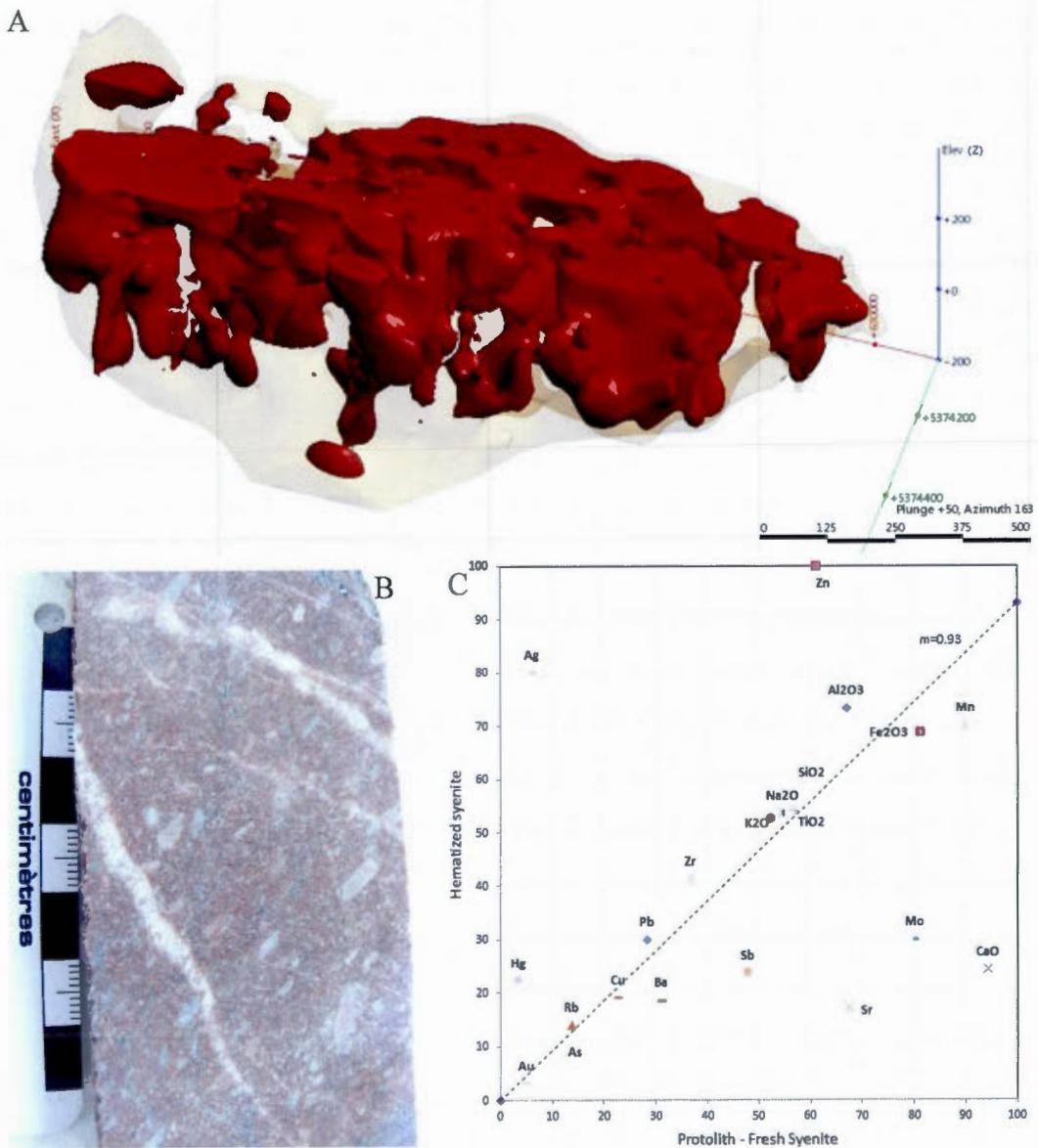


**Figure 1.12.** Microphotographs of arsenic-rich pyrite in the lithologically-controlled facies with TEM. (a) As-rich pyrites crystals. (b) Close up of (a): lattice fringes are very heterogeneous; the dark zone is an arsenic-rich portion of the pyrite.

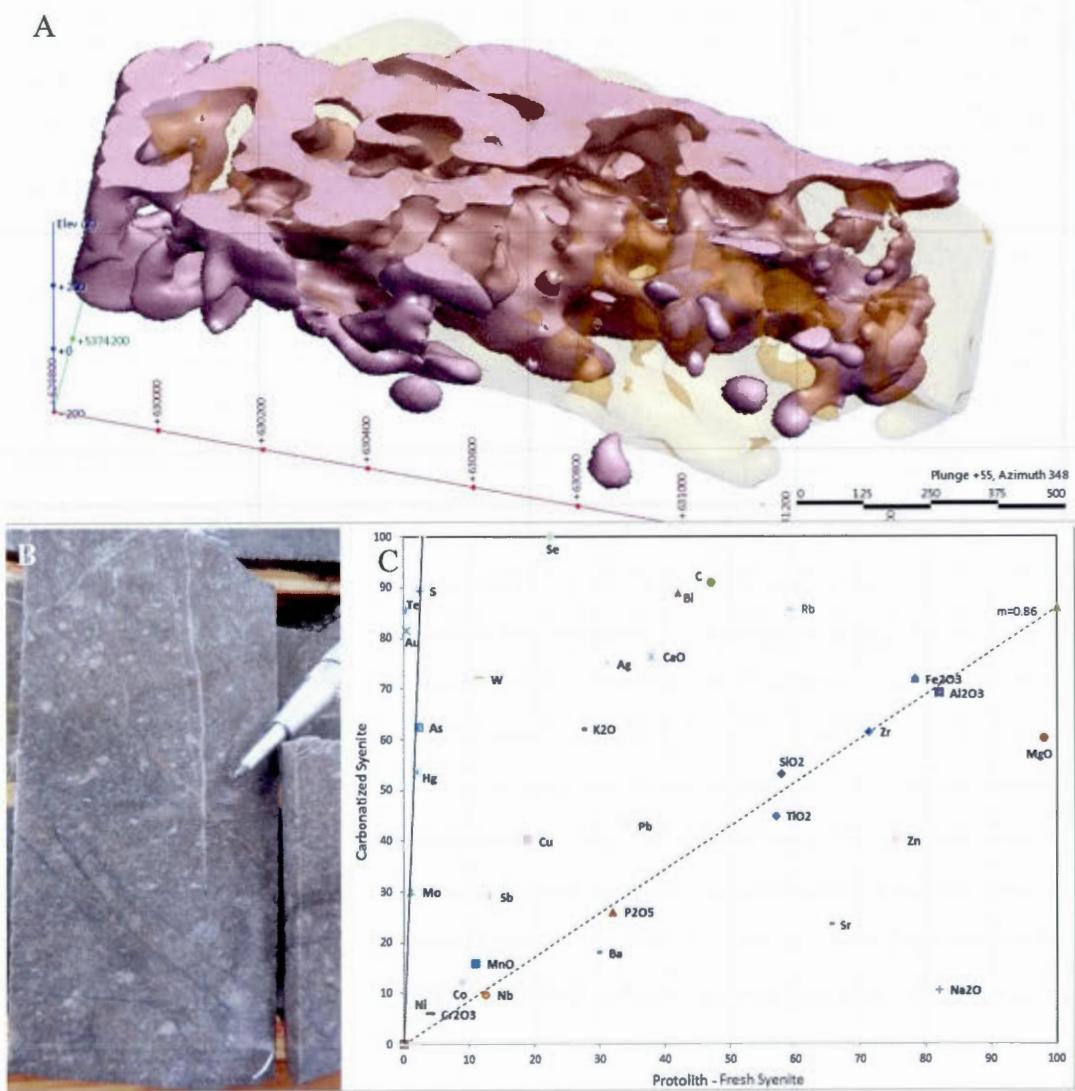




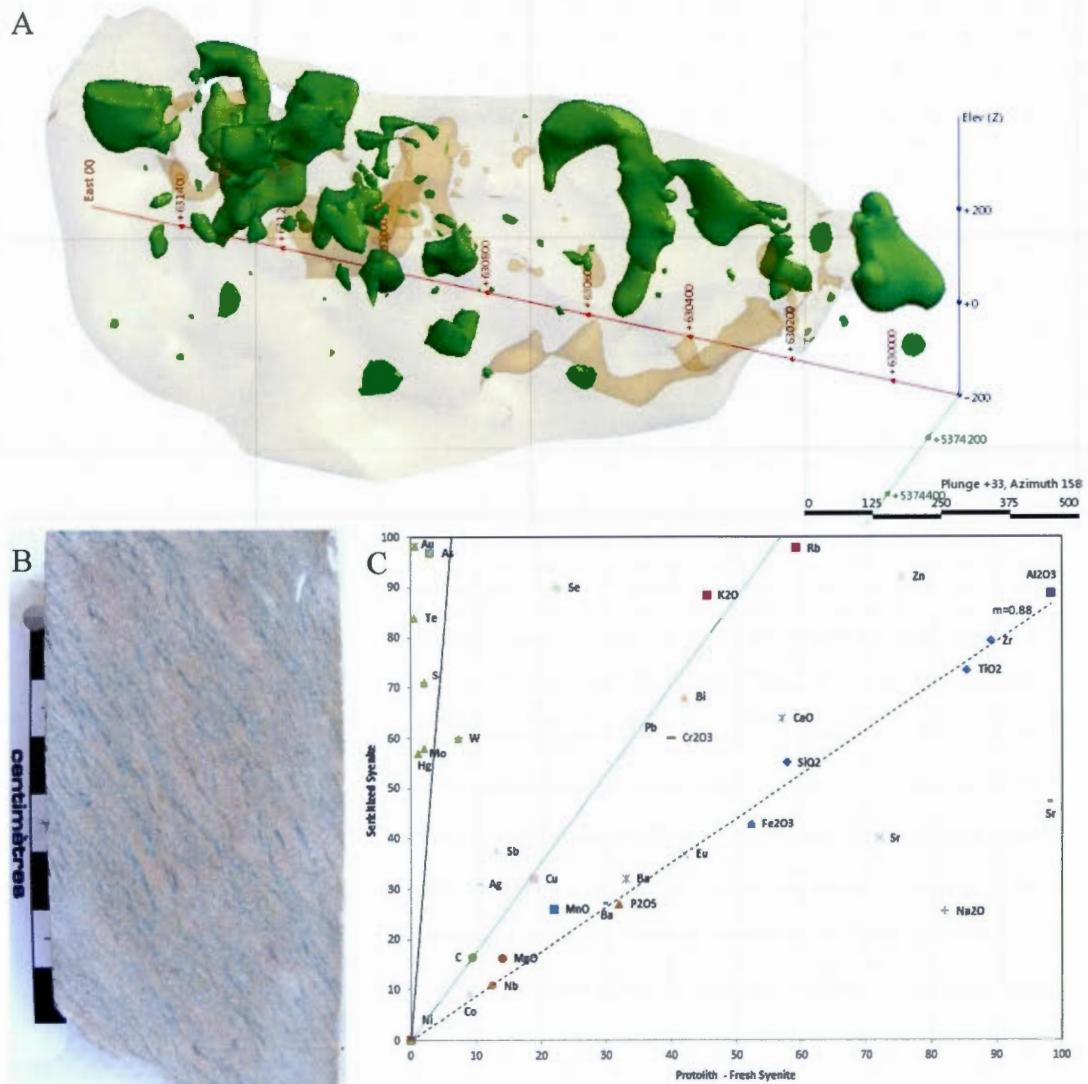
**Figure 1.14.** Comparison of the two mineralization styles in the Beattie deposit in terms of element concentration. It is an Isocon diagram (Grant, 2005). Same elements have been compared in the two mineralization styles, their ratio is plotted in the diagram. It is a quantitative estimation of changes in concentration depending on the mineralization style.



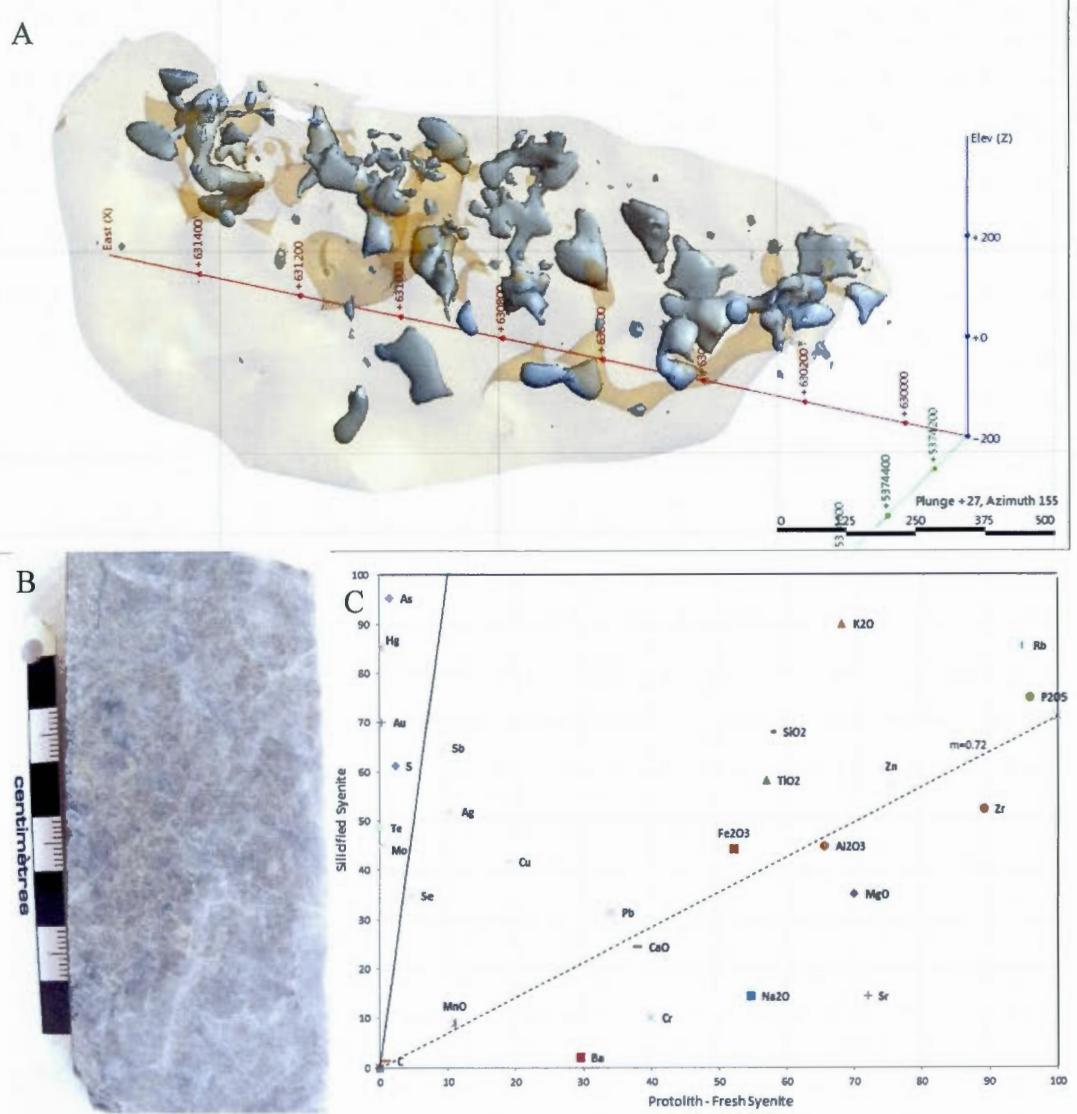
**Figure 1.15.** Hematization. (a) 3D model of hematized rocks – view plunges  $50^{\circ}$  to N163. (b) Reddish hematized syenite. (c) Isocon diagram of hematization.



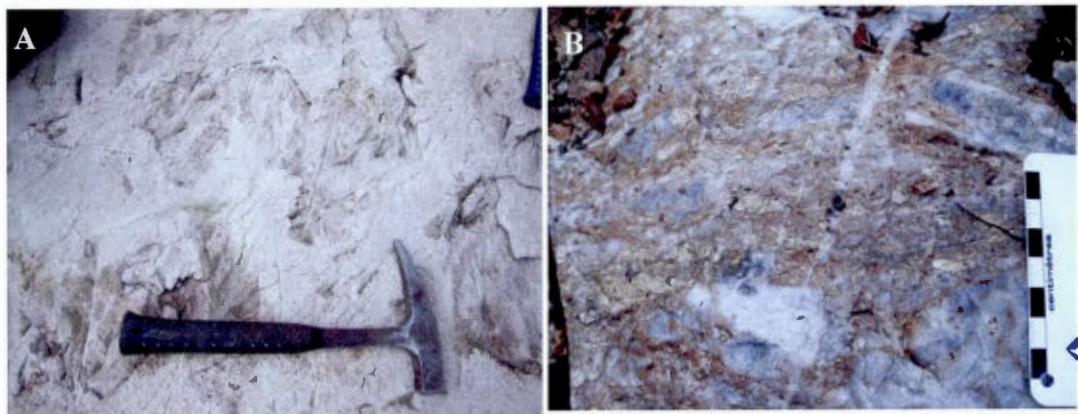
**Figure 1.16.** Carbonatization. (a) 3D model of carbonated rocks – view plunges 55° to N348. (b) Grey carbonatized syenite. (c) Isocon diagram of carbonatization.



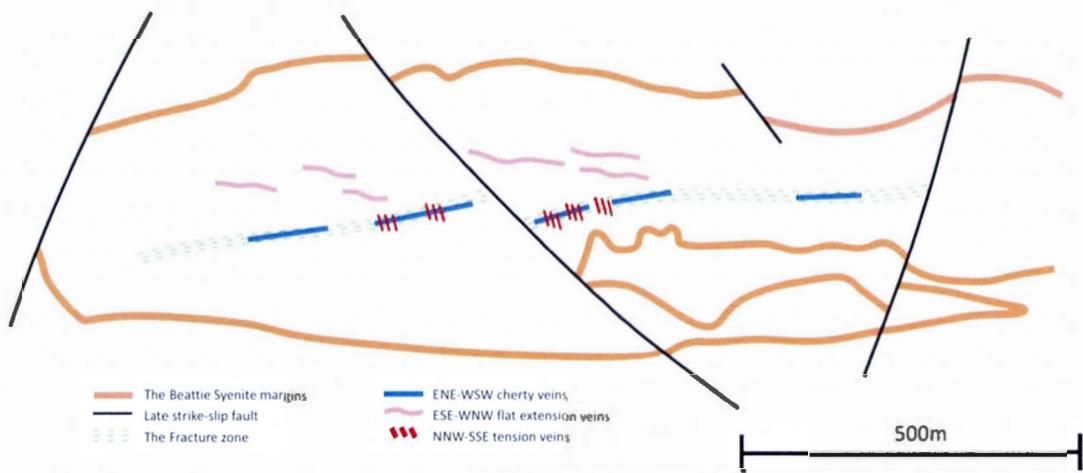
**Figure 1.17.** Sericitization. (a) 3D model of sericitized rocks – view plunges 33° to N158. (b) Green sericitized syenite. (c) Isocon diagram of sericitization.



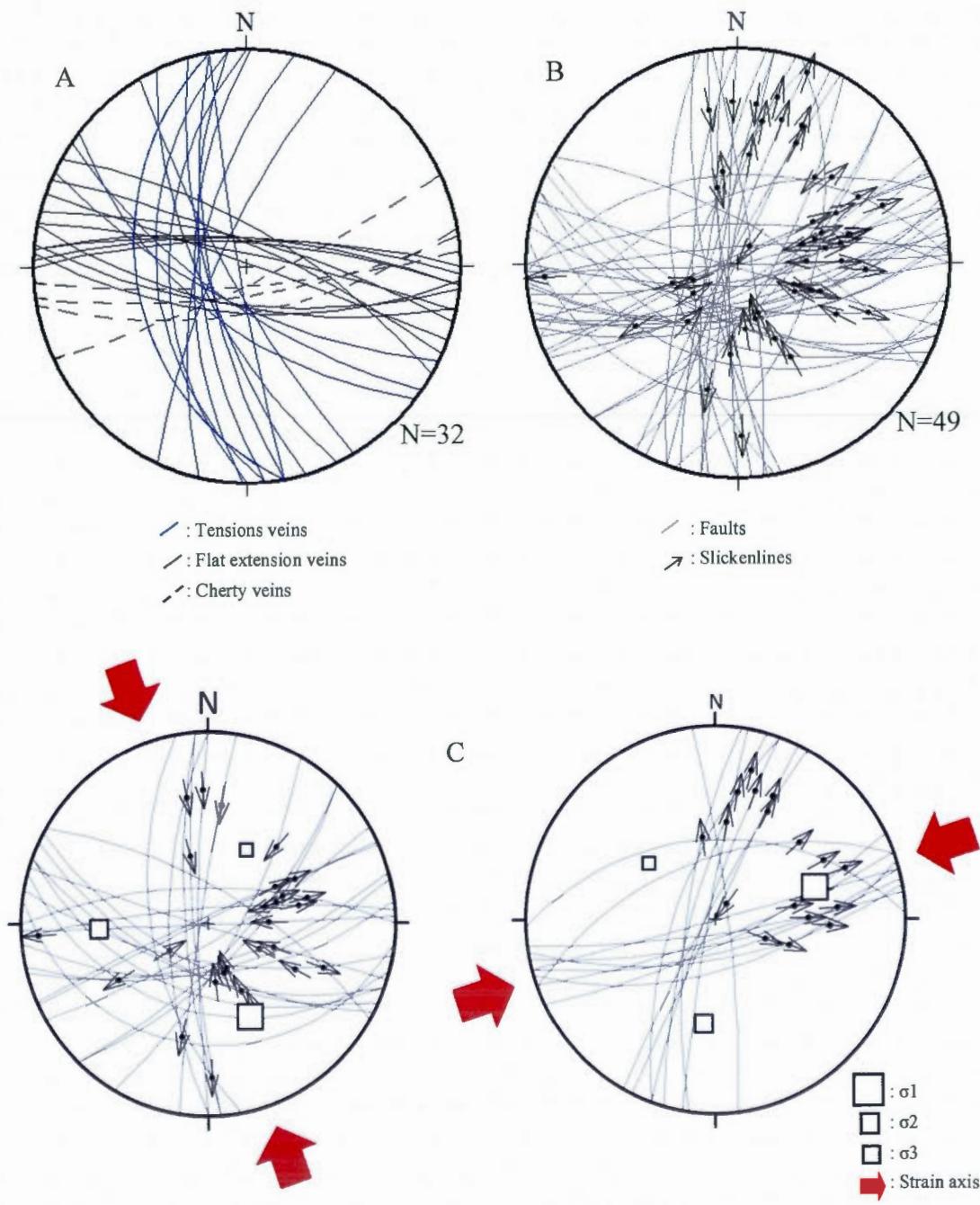
**Figure 1.18.** Silicification. (a) 3D model of silicified rocks – view plunges 27° to N155. (b) Grey silicified syenite. (c) Isocon diagram of silicification.



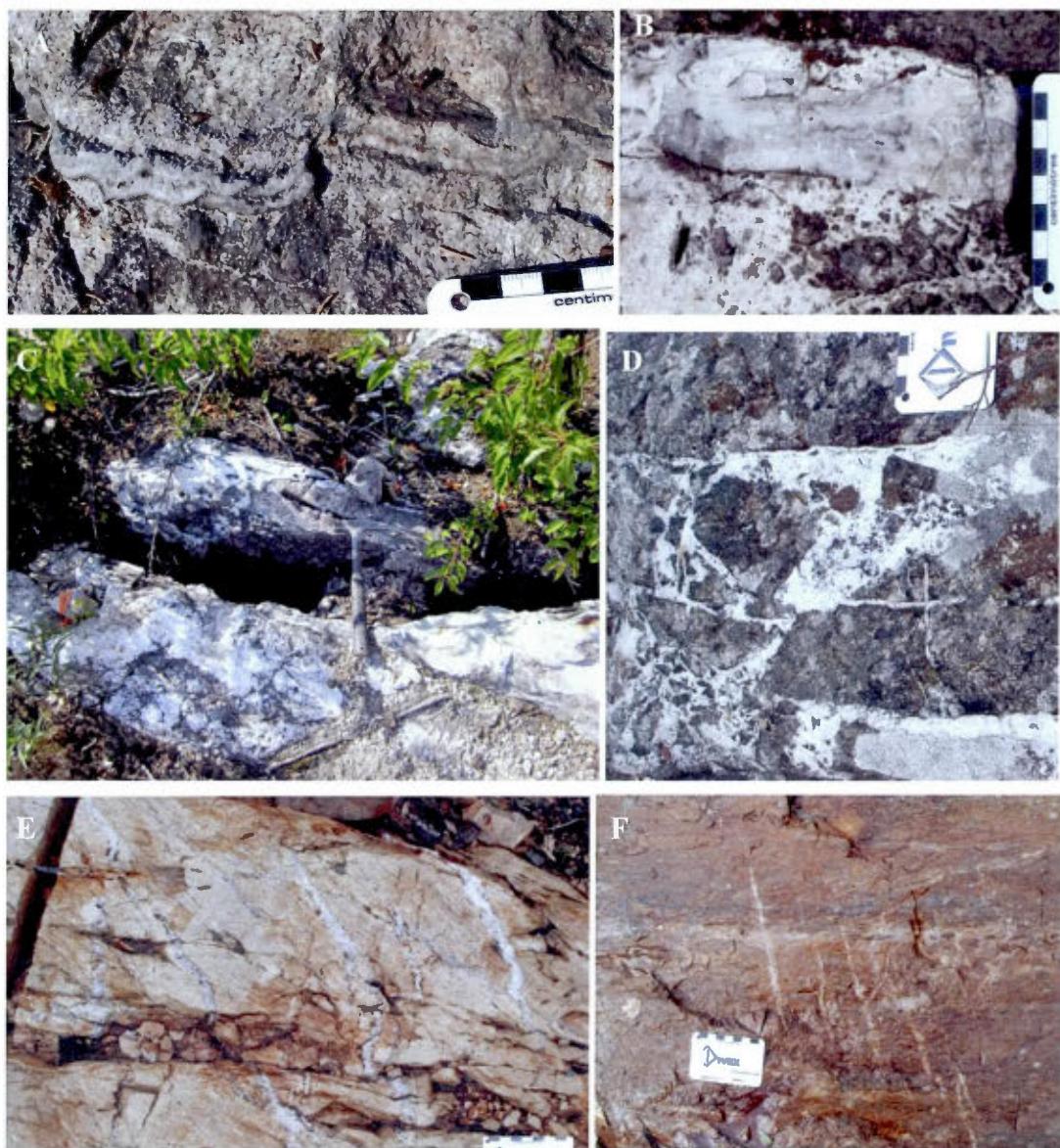
**Figure 1.19.** (a) Kaolinization, related to strong sericitization. (b) Fragments of sericitized syenite in silica corridors.

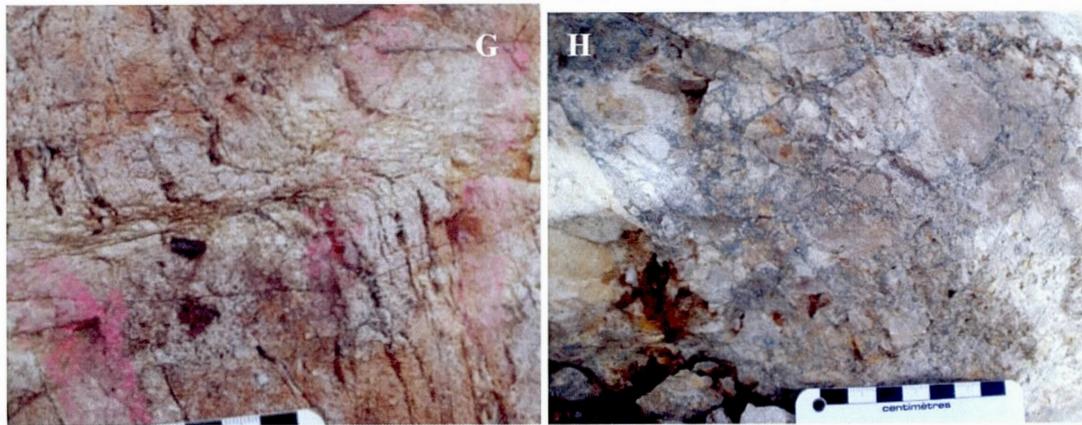


**Figure 1.20.** Cartoon of the different styles of veins and their location in the Beattie deposit.

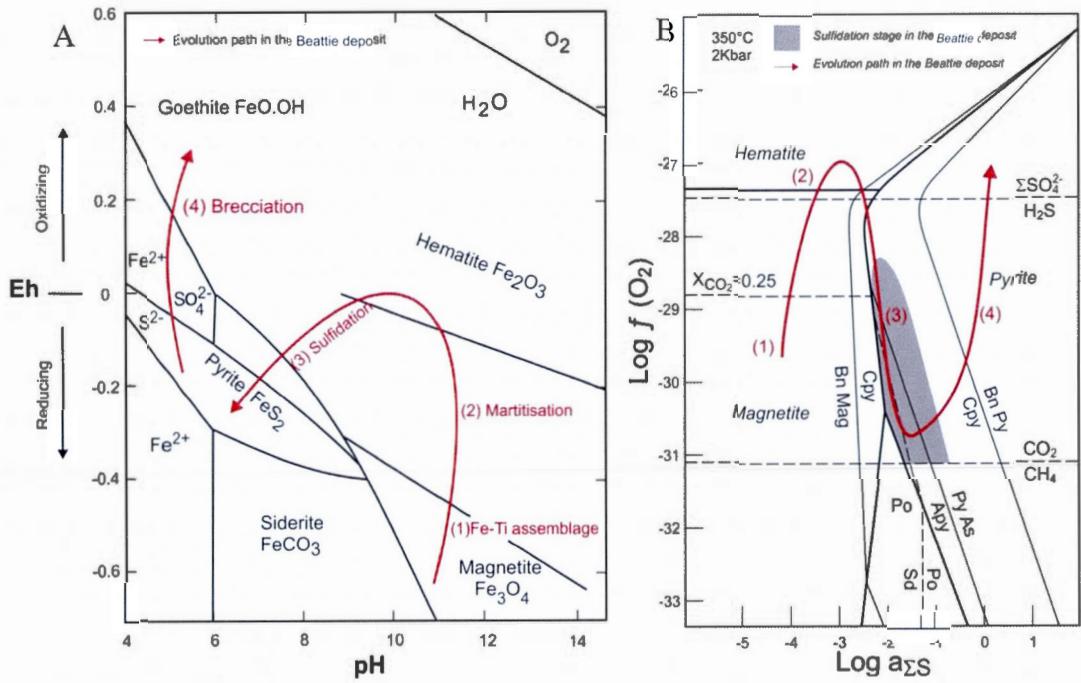


**Figure 1.21.** Structural analysis in the Beattie deposit. (a) Stereonet of veins. (b) Stereonet of minor late faults with slickenlines orientation. (c) Paleostress reconstruction using T-Tecto®: phase 1, strain axis NNW-SSE with  $\sigma_1$  plunging  $35^\circ$  to SSE; phase 2, strain axis ENE-WSW with  $\sigma_1$  plunging  $35^\circ$  to ENE.

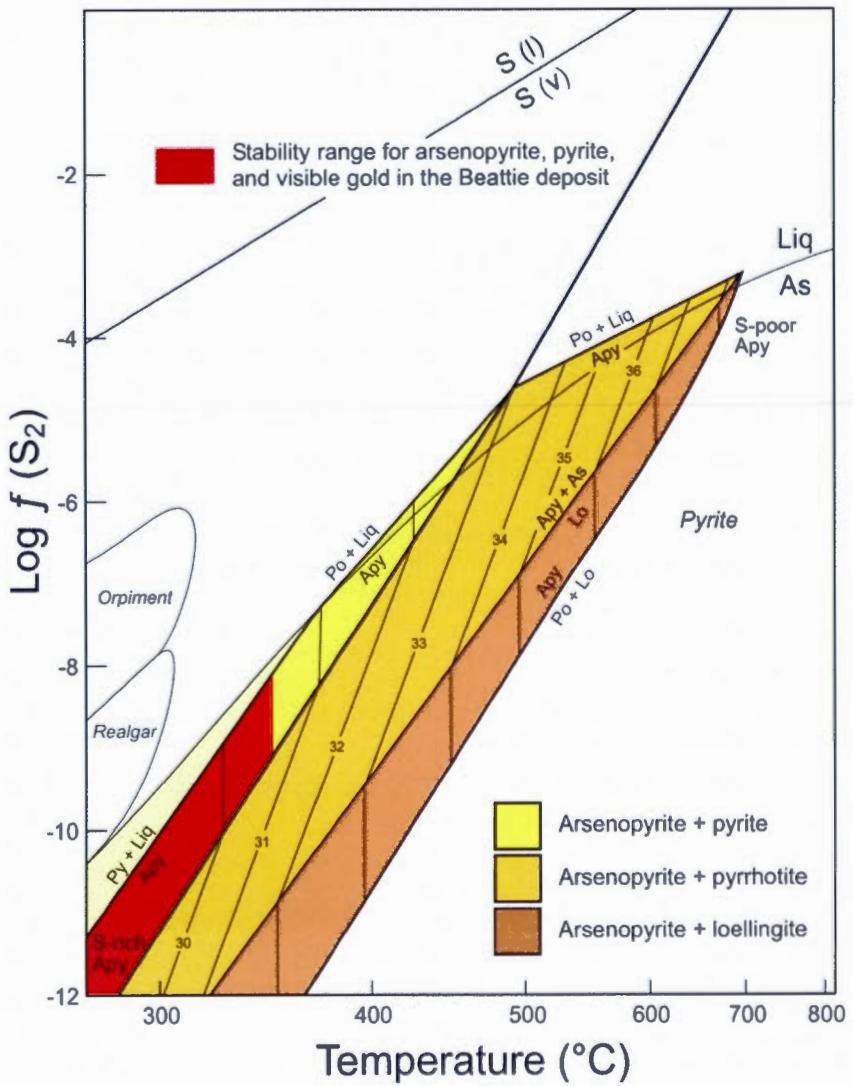




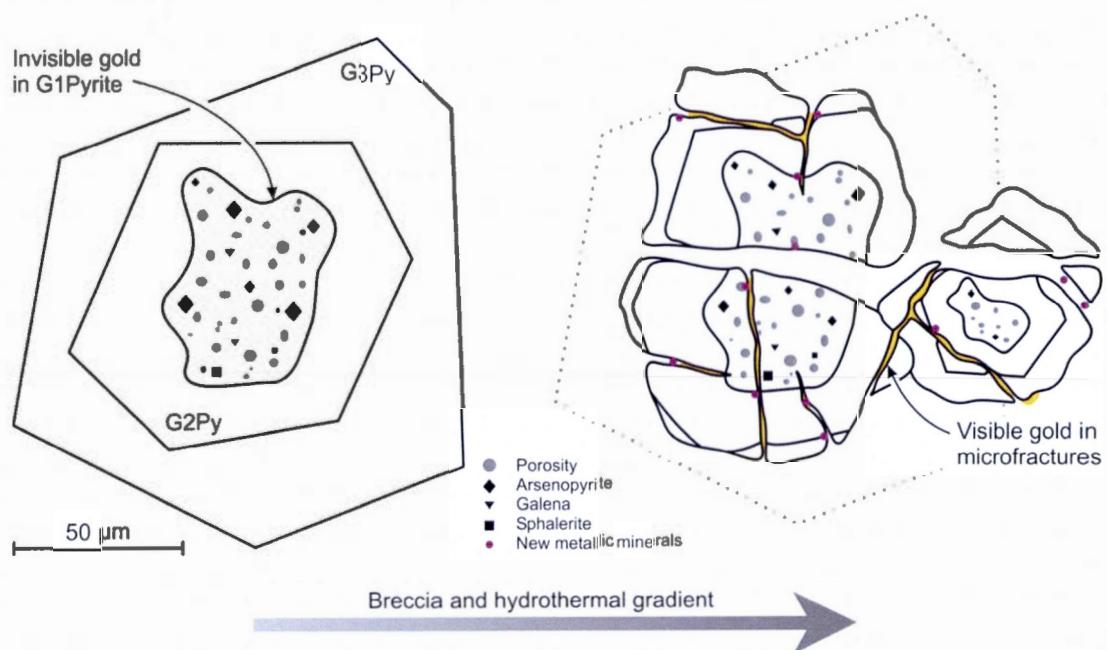
**Figure 1.22.** Photographs illustrating some of the vein styles. (a) ESE-WNW veins with open space crystallization. (b) ESE-WNW veins with pseudo-colloform texture. (c) ESE-WNW veins with a strong carbonate dissolution. (d) ESE-WNW vein with hydraulic breccia. (e) NNW-SSE tension veins near the Fracture zone. (f) NNE-SSE tension veins which cut ENE-WSW cherty veins. (g) NNW-SSE tension veins associated with carbonate dissolution, and cut by ENE-WSW shear. (h) Breccia fault related to an important net-slip on the NNE-SSW sinistral fault.



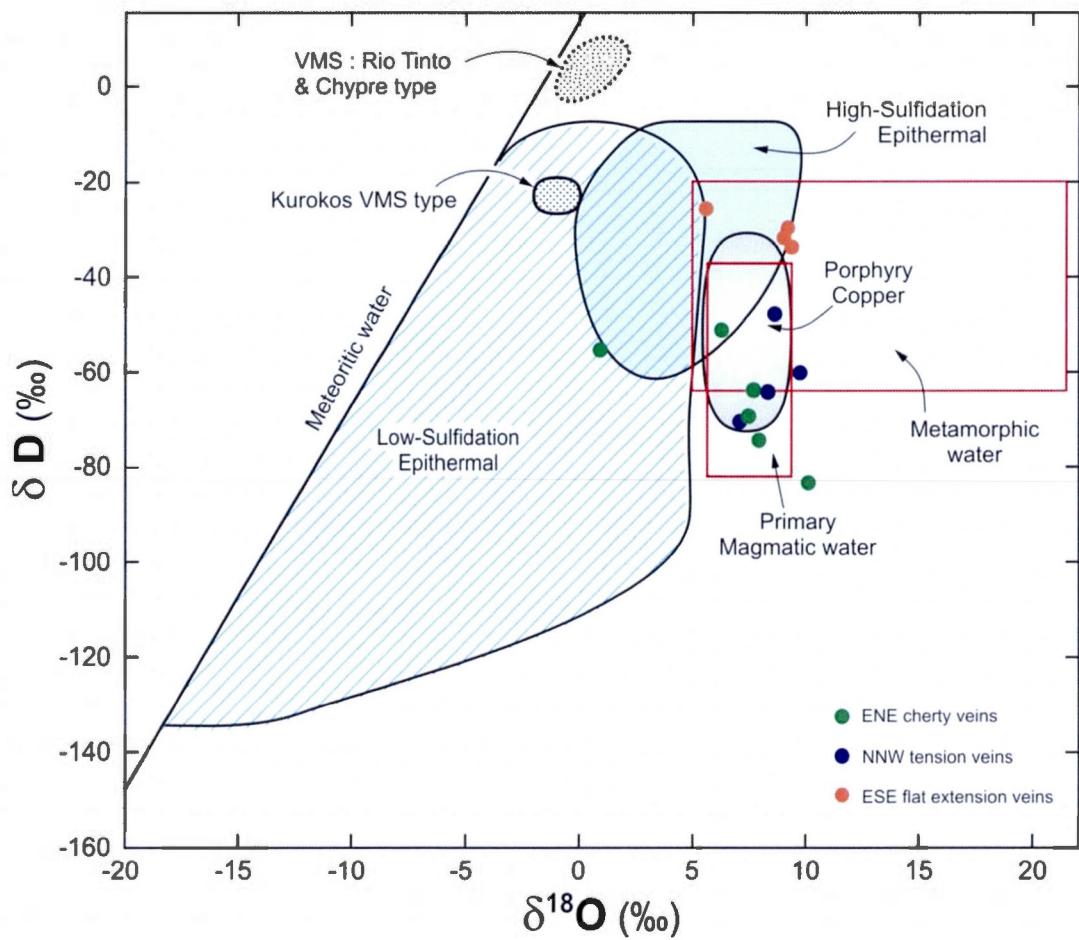
**Figure 1.23.** Evolution of the chemical conditions in the Beattie deposit. (a) Eh vs pH stability diagram, modified from Périssonier (2001). (b)  $f(O_2)$  vs  $\log a_{\Sigma S}$  stability diagram, modified from Mikucki and Roberts (2004).



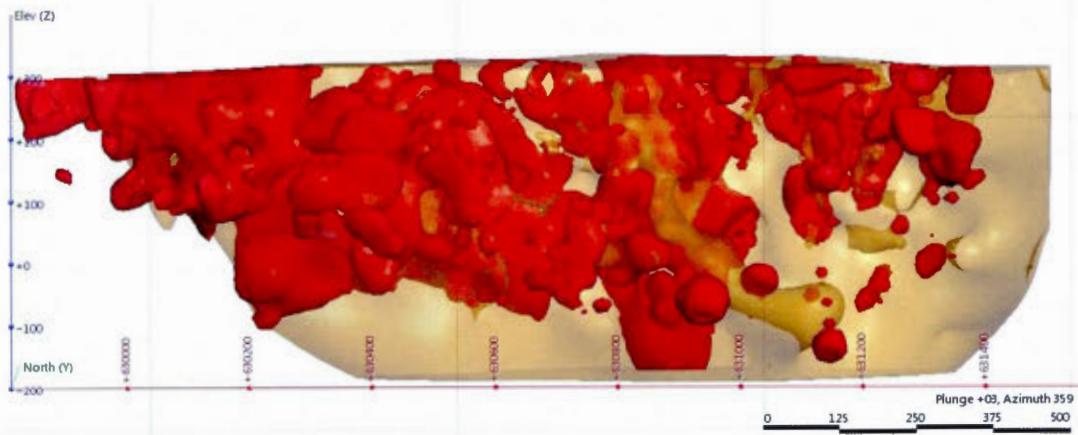
**Figure 1.24.** The average composition of magmatic-hydrothermal arsenopyrite of the breccia stage is plotted. Phase diagram redrawn from Barton (1969), Kretschmar and Scott (1976), and Sharp et al. (1985). Apy: arsenopyrite, Py: pyrite, Lo: loellingite. Atomic proportions of Arsenopyrite are indicated.



**Figure 1.25.** Cartoon of gold evolution in the Beattie deposit. Left: invisible gold in arsenian pyrite (G1) and arsenopyrite. Gold is in solid solution (yellow grid) in lattice bounds. Right: brecciated pyrite with several microfractures and new-porosity. Gold (yellow stripes) is remobilized and redeposited as visible in microfractures. New metallic minerals (Tennantite-tetrahedrite, enargite, hessite, coloradoite, chalcopyrite, barite) crystallize in the microfractures. There are two brecciated pyrites instead of only one in the left cartoon, it illustrates the increase of concentration of pyrite (two times more abundant).



**Figure 1.26.** Calculated isotopic composition for fluids in the Beattie deposit (colored spots) in comparison to other types of deposit. Modified from Jébrak and Marcoux (2008).



**Figure 1.27.** View to the north of ore bodies at 0.67 g/t Au (red) and the syenite (light brown). Evidence of plunging about  $35^\circ$  to ESE.

**Table 1.1.** Main characteristics of the two mineralization styles in the Beattie deposit.

	Lithologically-controlled facies	Structurally-controlled facies
<b>Nature of the Mineralization</b>	disseminated sulfides	disseminated sulfides, small clusters of sulfides
<b>Structure</b>	massive, stockwork	replacement, breccia, veins, dyke, silica corridors
<b>Settings of the Mineralization</b>	limited to the Beattie syenite	in second order breaks and shear zones adjacent and within the Beattie syenite
<b>Alteration</b>	iron-carbonatization	silicification, sericitization
<b>Metallic Mineralogy</b>	PY, APY, GL, SP	PY, APY, GL, SP, CP, EN, TT, CL, HS, EL
<b>Elemental Association (+Au)</b>	As, Zn, Pb, Ag	As, Zn, Pb, Ag, Te, Hg, Mo, Cu, Sb, Se
<b>Gold Occurrence</b>	invisible gold included in lattice bounds of arsenian pyrite and arsenopyrite	electrum in microfractures and porosity of brecciated pyrite
<b>Au Grades</b>	<2.5g/t Au	>2.5g/t Au ; regularly higher than 5g/t Au
<b>Sulfide Contents</b>	5 to 8%	10 to 20%
<b>Au/Ag</b>	1.2	0.8
<b>Process</b>	Magmatic stage : syn-sulfidation	Magmatic-Hydrothermal stage : breccia facies

**Abbreviations** PY:Pyrite; APY:Arsenopyrite; GL:Galena; SP:Sphalerite; CP:Chalcopyrite; EN:Enargite; TT:Tennantite-Tetrahedrite; CL:Coloradoite; HS:Hessite; EL:Electrum

**Table 1.2.** Microprobe assays on pyrite in the lithologically-controlled facies (EMP wt %)

Sample ID	Ag %	Se %	Fe %	S %	As %	Hg %	Te %	Au %	Cu %	Total %
1A-2	0.016	0.018	45.5	50.8	3.80	0.020	0.000	0.027	0.049	100.26
1A-3	0.000	0.012	46.3	53.3	1.02	0.000	0.000	0.009	0.001	100.65
A1-1	0.004	0.009	45.4	52.1	1.69	0.035	0.000	0.012	0.280	99.45
A1-2	0.000	0.015	45.7	50.4	4.40	0.022	0.005	0.013	0.028	100.61
A1-3	0.000	0.017	46.7	53.4	0.381	0.040	0.015	B.D.L	0.036	100.60
B2-1	0.009	0.015	47.1	53.6	0.014	0.003	0.008	B.D.L	0.000	100.74
B2-2	0.008	0.023	46.1	51.4	2.57	0.002	0.000	0.020	0.006	100.10
B2-3	0.022	0.015	46.2	52.2	1.35	0.008	0.000	0.015	0.011	99.83
B2-4	0.000	0.004	46.7	52.6	0.439	0.000	0.015	B.D.L	0.000	99.69
A4-2	0.002	0.013	45.6	50.0	4.71	0.019	0.006	0.018	0.173	100.46
A4-3	0.000	0.008	47.0	53.3	0.058	0.011	0.012	B.D.L	0.007	100.38
11-APY	0.000	0.210	35.2	20.6	37.8	0.008	0.009	0.042	0.008	93.96
6-1	0.011	0.007	46.1	51.2	2.87	0.029	0.000	0.008	0.023	100.21
6-2	0.000	0.017	46.2	52.0	1.95	0.004	0.002	0.008	0.015	100.15
6-3	0.000	0.004	46.8	53.3	0.020	0.000	0.000	B.D.L	0.000	100.12
11-1	0.001	0.011	46.1	53.4	0.162	0.022	0.000	B.D.L	0.000	99.74
11-2	0.004	0.009	45.7	51.9	1.92	0.000	0.000	0.011	0.004	99.58

**Table 1.3.** Microprobe assays on pyrite in the structurally-controlled facies (EMP wt %)

<b>Sample ID</b>	Ag %	Se %	Fe %	S %	As %	Hg %	Te %	Au %	Cu %	Total %
1-1	0.000	0.131	35.9	21.4	42.5	0.061	0.019	B.D.L	0.000	100.08
1-2	0.008	0.138	34.9	21.5	42.4	0.061	0.032	0.007	0.000	99.02
62-1	0.037	0.014	46.0	52.8	0.551	0.000	0.000	B.D.L	0.362	99.80
62-2	0.022	0.007	45.4	51.5	2.45	0.005	0.000	B.D.L	0.909	100.24
62-3	0.000	0.008	46.5	53.0	0.498	0.027	0.000	B.D.L	0.001	100.01
86-2	0.025	0.088	46.9	52.9	0.485	0.017	0.000	B.D.L	0.004	100.36
62-1	0.002	0.011	46.0	52.2	1.75	0.008	0.000	B.D.L	0.050	100.03
62-2	0.008	0.010	41.5	47.6	1.64	0.000	0.011	B.D.L	0.024	91.07

**Table 1.4.** Microprobe assays on electrum in the structurally-controlled facies (EMP wt %)

Sample ID	Au %	Ag %	Total %
El-1	85.71	15.23	100.940
El-2	84.75	15.24	99.990
El-3	87.30	12.68	99.980
El-4	90.44	9.000	99.440
El-5	85.22	15.10	100.320
El-6	85.19	14.96	100.150
El-7	83.35	15.80	99.150
El-8	83.87	14.52	98.390
El-9	83.39	16.06	99.450
El-10	85.19	15.25	100.440
El-11	85.05	15.53	100.580
El-12	83.50	14.50	98.710

**Table 1.5.** Main characteristics of the three veins style in the Beattie deposit

	<b>ENE-WSW</b>	<b>ESE-WNW</b>	<b>NNW-SSE</b>
<b>Location</b>	In the Fracture zone	Out of the Fracture zone	In the Fracture zone
<b>Dip</b>	steep to the south	shallow to the north	medium to steep to the west
<b>Structure</b>	massive	flat extension, open-space, pseudo-colloform	tension veins, periodic
<b>Morphology</b>	straight, narrow, irregular	sinuous, occasionnaly large	thin, occasionnally folded and sheared
<b>Composition</b>	cherty quartz, few sericite	quartz, carbonate (calcite to ankerite), few sericite	quartz, carbonate (brown ankerite)
<b>Carbonate Dissolution</b>	none	yes, locally total	yes, partial
<b>Gold Occurrence</b>	fine disseminated in sulfides, >10 g/t Au	none	none
<b>Breccia-related</b>	breccia ore spatially related	hydraulic breccia	none
<b>Relative Chronology</b>	early	early (different from ENE veins)	late

**Table 1.6.** Isotopic results for quartz veins in the Beattie deposit.

Sample	Vein type	$\delta\text{O18}$ (‰ VSMOW)	$\delta\text{O18}$ fluid at 350°C (‰ VSMOW)	$\delta\text{D}$ (‰ VSMOW)
6-A	ENE-WSW cherty	13.35	7.30	-67
4-B	ENE-WSW cherty	13.71	7.67	-74
Iso2	ENE-WSW cherty	13.2	7.2	-63
Iso4	ENE-WSW cherty	12.2	6.2	-53
Iso8	ENE-WSW cherty	16.1	10.1	-83
Iso10	ENE-WSW cherty	6.9	0.9	-57
1-C	ESE-WNW flat	14.56	8.52	-32
Iso1	ESE-WNW flat	14.9	8.9	-36
Iso6	ESE-WNW flat	14.8	8.8	-30
Iso9	ESE-WNW flat	11.7	5.7	-26
4-D	NNW-SSE tension	14.25	8.21	-63
Iso3	NNW-SSE tension	15.5	9.5	-60
Iso5	NNW-SSE tension	13.1	7.1	-69
Iso7	NNW-SSE tension	14.3	8.3	-47

## CONCLUSION GÉNÉRALE

Les études minéralogiques couplées à la géochimie ont apporté des avancées significatives dans la compréhension de l'évolution du gisement aurifère de Beattie. Des phases initiales oxydantes ont évolué en conditions plus réduites à la faveur de mélanges de fluides magmatiques (i.e., Bi) et externes (i.e., As dans les sédiments) riches en CO<sub>2</sub> qui ont entraîné la précipitation de l'or dans la structure cristalline de la pyrite arsénifère et l'arsénopyrite sous forme de solution solide. Deux phases subséquentes de pyritisation sont dépourvues d'or et d'arsenic. Cela permet de proposer qu'une phase unique d'apport en or est à l'origine de l'anomalie aurifère à Duparquet. Cet or invisible dans les sulfures arsénifères pourrait représenter jusqu'à 85% du mineraï. Il est étroitement associé à la syénite de Beattie très carbonatisée à carbonates de fer. Une phase d'altération ultérieure, dominée par des fluides magmatique-hydrothermaux (i.e., isotope O et H) riches en silice, entraîne une bréchification des sulfures et leur redistribution dans des corridors siliceux à l'intérieur des zones de failles E-W, ainsi qu'une remobilisation de l'or dans les microfractures des pyrites bréchifiées. L'or expulsé de la structure cristalline de la pyrite et de l'arsénopyrite précipite dans les pièges texturaux de la pyrite avec l'argent sous forme d'éléctrum. L'or est désormais visible au microscope. La géothermométrie appliquée à l'arsénopyrite indique des températures de cristallisation inférieures à 350°C pour l'évènement magmatique-hydrothermal associé à l'or visible. La dominance de régimes en systèmes ouverts a directement affecté le style de cristallisation de l'or, tant au niveau de l'incorporation dans le fluide de l'arsenic qui contrôle la précipitation de l'or invisible, que la circulation tardive de fluides riches en silice qui sont à l'origine de la remobilisation de l'or.

Le modèle génétique, initialement proposé par Robert (2001) a pu être précisé. La mise en place de l'intrusion suivit d'un mélange de fluides magmatiques et externes, qui aboutit à la maturité du magma évolué et à la précipitation de l'or dans les sulfures arsénifères, est contraint entre D1 et D2. L'évènement de redistribution des sulfures et de remobilisation de l'or est syn-métamorphique durant la transition D2-D3. Il est clair désormais que la silicification pervasive des roches spatialement associées aux failles E-W est le dernier évènement d'altération (D3), alors que la séricitisation est plus précoce.

Plusieurs évidences indiquent que le gisement de Beattie a des affinités avec les gisements associés aux intrusions (*Intrusion Related Gold Deposit*; Lang et Baker, 2001; *Syenite Associated Disseminated Gold Deposit*; Robert, 2001), mais aussi avec des gisements de types épithermaux (Sillitoe et Hedenquist, 2003), et possiblement mésothermaux (Groves, 1998). Globalement, le gisement de Douay apparaît comme le plus similaire à Beattie, tant sur le plan pétrologique, que sur le plan métallogénique et structural (Gauthier et Arnois, 1994; Dupéré, 2007).

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## APPENDICE A

MÉTHODE ANALYTIQUE POUR LA GÉOCHIMIE, ET RÉSULTATS D'ANALYSES  
LITHOGEOCHIMIQUES

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### **Méthodologie pour les analyses lithogéochimiques**

Cinquante-deux échantillons ont été sélectionnés à partir de zones d'affleurement et de forage dans le gisement de Beattie. Des analyses sur roche totale et éléments traces ont été réalisé au laboratoire ACME de Vancouver, et ALS de Val d'or. Les concentrations des éléments majeurs et traces ont été déterminées par Inductively Coupled Plasma Mass Spectrometry (ICP – MS), et par Inductively Coupled Plasma Emission Spectrometry (ICP – ES) suivant la fusion lithium metaborate/tetraborate et la digestion avec un acide nitrique.

### **Méthodologie pour les analyses à la microsonde**

Dix lames minces polies préalablement étudiées au microscope optique ont été sélectionnées pour l'analyse microsonde. Ces échantillons sont représentatifs des quatre phases de la paragénèse minérale du gisement de Beattie. Les analyses ont été effectuées au laboratoire de microsonde de l'université McGill. L'appareil utilisé est un JXA JEOL-8900L dans lequel sont intégrés les spectromètres WDS et EDS. Les analyses ont été réalisées sous 20kV, avec un courant de faisceau de 50nA, et un diamètre de faisceau de 3µm. Les éléments Au, As, Ag, Fe, S, Cu, Zn, Hg, Se, et W ont été mesurés dans des pyrites et des arsénopyrites. Les seuils de détection étaient respectivement pour les éléments mentionnés, 0,0065% après 540 secondes, 0,0491% après 60 secondes, 0,035% après 60 secondes, 0,034% après 10 secondes, 0,0121% après 10 secondes, 0,0229% après 60 secondes, 0,0382% après 60 secondes, 0,0337% après 340 secondes, 0,0156% après 60 secondes, et 0,0248% après 100 secondes. Pour les analyses sur électrum, le comptage était de 20 secondes pour l'or et pour l'argent.

### **Méthodologie pour les analyses d'isotopes stables d'oxygène et d'hydrogène**

Quatorze échantillons de quartz pur provenant des trois types de veines dans le gisement de Beattie ont été analysés au laboratoire des isotopes stables de l'université Queen's, Ontario. Les isotopes d'hydrogène sont mesurés avec un spectromètre DELTA <sup>plus</sup> XP. L'analyse est faite par Thermochemical Elemental Analyser (TC/EA) à haute

température. Pour les isotopes d'oxygène, le matériel est préparé séparément sur une ligne d'extraction silicatée, puis mesuré avec un spectromètre de masse Finnigan Mat 252.

#### Résultats lithogéochimiques :

SAMPLE	D006556	D009304	144103	D007580	144103	D007601	04-A	D047015	01-A	D000398	D023252
SiO <sub>2</sub> %	55.1	57.9	61.1	56.1	61.1	58.4	54.66	53	82.67	72.9	70.9
Al <sub>2</sub> O <sub>3</sub>	14.8	11.5	11.8	15.55	11.8	13.8	17.32	13.75	5.46	9.42	8.56
Fe <sub>2</sub> O <sub>3</sub>	4.29	6.64	4.14	4.74	4.14	4.3	4.99	4.17	4.77	2.63	3.3
CaO	4.26	4.3	6.54	4.23	6.54	4.39	3.83	4.88	0.3	2.25	3.88
MgO	1.63	1.37	1.28	1.74	1.28	1.7	1.47	2.19	0.09	0.48	1.12
Na <sub>2</sub> O	1.7	0.21	0.98	0.22	0.98	0.86	3.75	2.31	0.38	0.15	0.36
K <sub>2</sub> O	8.84	6.43	4.07	5.24	4.07	4.71	5.43	7.45	3.79	7.02	5.62
TiO <sub>2</sub>	0.49	1.24	0.43	0.56	0.43	0.47	0.54	0.47	0.16	0.31	0.28
MnO	0.13	0.2	0.11	0.13	0.11	0.12	0.11	0.1	0.03	0.06	0.14
P <sub>2</sub> O <sub>5</sub>	0.27	0.39	0.29	0.32	0.29	0.26	0.31	0.27	0.27	0.16	0.16
C	1.66	1.6	2.03	1.64	2.03	1.65		1.91		0.62	1.27
S	1.42	1.92	0.43	0.09	0.43	0.06		0.91		1.56	1.33
LOI	6.18	6.96	7.78	7.66	7.78	7.59	7.1	7	1.9	2.78	5.3
Total	98.1	97.2	98.8	96.8	98.8	96.8	99.5	96	99.78	98.3	99.8
Ba ppm	2720	436	1360	2270	1360	1460	2028	2540	790	959	1185
Ce	220	66.8	168	245	168	197	250	218	93.3	130	133
Eu	3.69	2.21	3.17	4.35	3.17	3.87	4.07	3.65	1.65	2.43	2.58
Cr	10	10	10	10	10	10		10		20	10
Rb	195.5	137	105.5	163.5	105.5	131	132.3	165.5	68.5	144.5	107.5
Sr	947	413	680	474	680	410	926	1010	144.6	227	422
Sn	1	2	1	1	1	1	1	1	0.6	1	1
Sr	947	413	680	474	680	410	926	1010	144.6	227	422
Th	22.8	2.6	17.2	23.5	17.2	19.25	29.4	22.7	9	14.9	13.35
U	6.78	9.33	8.29	6.38	8.29	6.95	6.6	6.33	3.9	4.48	4.12
V	144	191	190	148	190	81	105	192	130	179	191
W	12	47	11	10	11	4	2.7	13	6.8	8	8
Y	29.9	46.7	24.6	33.4	24.6	27.3	35.3	29.5	17.8	18.9	20.7
Zr	317	255	246	344	246	307	376	301	124.3	234	186
As	300	500	93.3	3.5	93.3	6.7	8.2	200	713.8	>250	300
Au	1.965	4.71	1.13	0.028	1.13	0.038	0.0051	2.18	9.708	7.86	3.17
Bi	0.68	0.24	0.87	0.43	0.87	0.65	0.3	0.6	0.9	0.37	0.36
Hg	1.45	1.69	1.595	0.043	1.595	10.35	0.05	1.105	2.91	>25.0	22.3
Sb	3.74	12.5	10	2.8	10	1.6	1	2.35	4.1	26.6	10.3
Se	1.8	2.9	1.8	0.7	1.8	0.5	0.8	1.5	2.4	4	4.7
Te	1.68	4.36	1.66	0.01	1.66	0.52		0.56		>250	16.85
Ag	3.1	5.9	2.5	0.9	2.5	0.7	0.06	2	5.2	149	8
Cu	32	88	35	22	35	64	26.3	25	21.7	168	25
Mo	57	103	67	0.6	67	1	0.3	18	497.8	14	67
Ni	4	3	4	6	4	6	2.2	2	5.4	18	3
Pb	62	25	89	18	89	20	34.9	38	57.2	22	30
Zn	92	60	70	96	70	73	56	64	46	34	96

D000396	06-A	D009165	D013071	D036693	D039219	D023142	D035021	D048208	D005117	D000486
65.5	92.2	67.2	49.2	41.5	53.4	63.7	58.5	59.1	51.3	56.3
3.08	2	13.3	13.2	10.9	13.6	11.65	10.25	12.4	12.9	13.15
2.03	1.78	4.1	3.92	3.19	5.33	6.64	6.44	10.9	12.65	7.38
1.16	0.14	0.86	12.15	19.95	7.76	0.97	4.35	3	7.13	4.81
0.19	0.09	0.27	1.4	0.62	2.29	0.29	1.16	0.85	2.6	1.96
0.05	0.02	0.4	2.23	3.81	2.69	0.2	0.56	0.06	1.27	0.7
2.16	0.89	9.53	3.84	1.6	5.11	8.91	6.94	5.49	2.41	5.56
0.09	0.08	0.42	0.48	0.35	0.47	1.24	1.18	1.4	1.4	0.71
0.03	0.01	0.03	0.15	0.11	0.2	0.08	0.15	0.05	0.31	0.16
0.07	0.06	0.31	0.24	0.23	0.28	0.43	0.37	1.11	0.3	0.11
0.31	0.05	0.17	3.07	4.28	1.99	0.47	1.37	0.59	2.13	1.61
1.59	0.56	2.34	1.44	2.05	0.56	2.8	3.37	2.38	0.7	2.05
17	1.09	2.96	10.8	14.25	8.17	3.86	5.94	4.49	9.37	7.07
91.4	98.4	99.7	97.8	96.7	99.6	98	95.9	98.9	101.5	98
236	124	2080	705	438	1790	300	194.5	445	324	248
43.1	49.4	179	191	159.5	210	59.7	47.8	66.7	34.2	19.7
0.91	1.23	2.91	3.31	2.62	3.66	2.01	1.92	2.71	1.83	0.78
10	30	20	10	10	10	10	6	6	10	50
42.7	23.7	172.5	98.4	49.5	110	154.5	111.5	118.5	70.1	127.5
82	36.5	384	663	1045	1145	129	110.5	318	122	78.2
3	1	1	1	1	2	1	3	1	1	1
82	36.5	384	663	1045	1145	129	110.5	318	122	78.2
4.28	1.81	20.4	20.1	16.8	20.6	2.46	1.82	2.59	1.2	0.86
1.26	3.64	9.08	6.45	4.91	5.61	3.19	0.67	1.75	0.43	0.26
51	260	147	206	334	178	142	188	497	136	220
2	3	10	9	11	11	47	17	48	9	8
6.7	6.7	26	29.1	22.4	31.6	41.7	37.1	55.3	34.3	15.2
110	30	287	284	237	307	253	226	258	169	108
>250	223	600	700	300	223	1900	14600	5500	>250	>250
120	10.6	3.14	2.97	6.35	1.995	7.04	5.41	6.64	1.29	1.895
1.29	0.3	0.88	0.56	0.46	0.62	0.16	0.05	0.06	0.04	0.06
>25.0	170	2.28	1.385	50	0.741	7.96	0.372	0.594	>25.0	>25.0
>250	41.1	3.76	6.3	7.1	2.57	7.62	15.35	14.25	3.41	24
107	2.7	3.2	2.6	3.7	1.4	5.1	5.8	4.3	1.2	3.5
>250	117.5	1.99	5.97	24.1	0.66	6.47	0.99	0.75	12.65	11.85
3580	44.6	2.6	5.7	4.2	2.1	2.2	1.1	2.7	4.4	2.5
7920	125	31	22	25	22	124	67	89	38	70
29	175	103	11	42	4	13	8	81	12	1
3	6	5	3	3	2	6	1	8	7	48
22	35	49	23	17	29	69	17	50	3	6
1040	38	53	54	17	119	201	24	87	101	71

D0003 74	D0235 42	D0392 78	D0393 28	D0003 51	56626 0	56629 2	D0057 42	D0058 85	D0094 78	D0057 31
57.6	53.3	58.2	57.2	53.2	56.8	53.4	57.8	56.3	56.6	57.2
16.2	15.15	14.3	14.95	13.8	14.7	14.9	15.8	15.5	15.7	15.65
4.78	4.17	4.46	5.59	4.79	4.54	4.77	5.55	4.44	5.28	5.36
3.97	7.82	3.24	3.88	7.63	3.42	5.38	5.04	5.88	3.8	4.32
0.59	0.57	0.96	1.04	0.86	1.22	1.55	1.5	1.13	1.56	1.92
0.51	0.51	1.6	1.72	0.7	1.45	0.59	4.96	4.21	5.22	5.5
11.65	10.65	9.09	9.56	10.35	9.87	11.95	4.97	4.65	4.07	3.92
0.54	0.5	0.44	0.5	0.45	0.47	0.48	0.61	0.53	0.56	0.59
0.14	0.13	0.15	0.23	0.16	0.13	0.18	0.13	0.13	0.12	0.15
0.28	0.27	0.27	0.3	0.26	0.26	0.29	0.37	0.28	0.31	0.38
1.13	1.67	1.15	1.45	1.82	1.25	1.7	0.92	1.66	0.65	1.46
1.04	1.68	0.69	0.89	1.49	0.49	1.57	0.23	0.14	0.04	0.04
4.28	5.56	4.34	4.9	2.69	3.6	4.38	4.38	6.71	3.2	5.81
101	99	97.5	100.5	95.2	96.8	98.2	101.5	100	97	101.5
1415	1880	2890	2360	1810	2380	1770	2910	2130	3240	2870
240	203	204	221	193	202	212	242	231	246	243
4.06	4.09	3.57	3.88	3.8	3.87	3.95	4.29	4.01	4.31	4.32
10	10	10	10	10	10	10	10	10	10	10
223	189	176.5	179.5	171	182.5	180	102	122	97	101
461	1005	772	1135	706	628	511	2150	1075	1730	1420
1	3	1	1	1	1	1	2	2	2	2
461	1005	772	1135	706	628	511	2150	1075	1730	1420
25.1	21.4	21.7	23.3	19.85	21.1	22.7	23.3	23.1	23.6	23.9
6.46	5.73	6.05	7.66	4.33	5.64	6.5	6.47	6.24	6.33	6.39
163	97	124	219	67	105	105	170	145	154	160
15	10	10	17	9	8	9	4	7	2	4
32.2	28.9	29.2	32.3	27.6	27.3	28.2	34.7	33.6	35.1	33.9
345	332	315	316	307	334	342	335	331	345	330
400	400	172	300	>250	180	>250	67.5	25.4	10.5	15.4
1.565	2.16	1.61	2.93	2.72	1.51	2.5	0.555	0.014	0.003	0.003
0.73	1.12	0.66	0.52	0.89	0.24	0.93	0.74	0.53	0.43	0.65
1.66	1.27	0.722	0.862	1.34	5.55	5.04	0.886	0.114	0.059	0.128
3.77	2.41	2.23	4.3	2.9	1.91	4.15	2.75	0.69	1.34	0.95
1.4	1.7	1.2	1.6	2	1.1	2.1	1.2	0.7	0.9	0.7
0.8	0.68	0.48	0.95	2.14	0.68	0.93	0.62	0.04	0.006	0.01
1.3	1.9	1.4	2.3	2.5	1.5	1.6	3.3	1.4	2.2	2.1
31	20	32	23	40	30	29	27	28	16	30
4	6	11	2	30	5	25	1	0.6	0.6	0.6
5	5	3	5	8	6	7	5	2	4	5
32	35	39	38	43	35	33	85	30	40	72
60	34	59	72	40	59	94	102	54	116	73

D01283 9	D04737 9	14419 2	D02333 8	D04712 4	D01072 0	D04582 6	D03809 4	01-B -	D02334 8	03-A -
57.4	60.7	54.6	56.1	55.1	57.4	61	62.3	60.2	56.9	58.6 1
15.45	13.5	14.85	15.4	16.8	14.15	17.75	17.3	18.4	16.05	16.7 4
4.97	4.12	5.12	4.8	5.16	4.55	4.9	4.57	5.55	5.03	5.41
4.09	6.22	4.21	3.84	4.8	4.18	2.34	0.89	0.49	4.45	3.14
1.55	1.19	1.92	1.4	1.88	0.77	0.77	0.48	0.55	1.14	1.66
5.44	4.63	5.38	5.07	5.05	3.47	4.38	4.5	4.44	5.46	5.47
3.99	3.51	4.5	4.59	4.47	4.97	5.15	7.18	7.38	3.87	5.22
0.55	0.45	0.55	0.51	0.51	0.48	0.61	0.5	0.58	0.56	0.57
0.12	0.12	0.13	0.15	0.08	0.12	0.03	0.05	0.08	0.13	0.09
0.29	0.22	0.35	0.3	0.29	0.26	0.37	0.17	0.34	0.3	0.34
0.83	1.43	1.53	1.5	1.2	1.38	0.45	0.54		0.94	
0.03	0.19	0.03	0.03	0.14	0.25	0.12	0.07		0.04	
3.65	5.89	4.76	5.15	5.53	5.19	3.09	2.5	1.5	3.97	2
98	101	96.8	97.7	100	95.8	100.5	100.5	99.5	98.4	99.2 2
3230	4710	2520	1875	2300	2060	2000	2100	244 3	2820	3109
237	196.5	234	231	237	228	272	243	259	246	239. 1
4.17	3.55	4.22	4.51	3.99	3.91	4.46	3.81	4.35	4.27	4.13
10	10	10	10	10	10	10	10		10	
95	84	104	112	93.5	140	138.5	161.5	168	99.3	137. 6
1595	1450	1430	1085	1240	728	551	478	581	1660	2258
2	1	2	2	2	1	2	2	2	2	2
1595	1450	1430	1085	1240	728	551	478	581	1660	2258
23	20.6	23.5	23.7	24.3	23.1	27.2	21.9	28.8	24.4	25.9
6.16	5.84	6.14	6.21	6.83	6.12	7.81	5.85	5.4	6.82	6.7
147	113	114	96	149	136	143	122	143	151	115
2	6	2	3	6	7	5	7	13	2	0.9
34.7	31.5	32.4	33.3	32.9	30.4	36.4	30.5	33.2	35.6	33.5
338	284	311	380	342	320	370	388	377	345	368. 5
11.1	8.8	15.4	7.6	15	47.5	16.8	21.1	92.2	5.6	12.6
0.003	0.01	0.003	0.003	0.006	1.735	0.012	0.017	0.48	0.005	0.02 2
0.39	0.51	0.66	0.19	0.23	1.43	0.59	0.69	0.8	0.44	0.4
0.032	0.094	0.161	4.76	0.116	50	0.128	0.217	1.14	0.053	0.05
0.74	0.76	2.8	0.83	0.7	1.9	2.86	1.31	1.6	1	1.6
0.6	0.6	0.7	0.6	0.6	1.5	0.7	0.7	0.3	0.6	0.3
0.01	0.03	0.01	0.23	0.03	22.3	0.006	0.02		0.01	
1.7	1.9	1.8	1.4	1.6	4.8	0.8	0.8	0.3	2	0.06
21	18	35	27	17	29	30	23	27.8	15	22.8
0.6	0.6	0.6	1	0.6	111	0.6	0.6	1.5	0.6	1.6
2	3	4	6	4	6	6	4	5.4	4	6.4
41	29	30	27	19	28	26	43	38.2	40	28.3
99	81	65	84	105	73	60	89	41	90	61

D034717	D010576	D048108	D048158	D035064	D023192	05-A	04-C
59.4	59.7	55.5	60.4	56.2	57	75.57	59.6
18.3	12.15	12.85	15.75	16.1	17.65	11.75	20.5
4.59	8.49	8.81	7.63	11.3	7.26	1.93	4.31
0.81	2.36	5.21	1.52	1.74	2.88	0.95	0.48
0.37	0.92	1.05	4.23	2.34	2.47	0.33	0.56
5.37	0.51	3.69	0.87	0.69	4.68	3.31	4.39
5.27	3.95	2.07	2.78	3.45	2.91	3.01	6.85
0.54	1.29	1.27	0.62	0.7	0.83	1.03	0.63
0.07	0.14	0.16	0.04	0.2	0.09	0.03	0.02
0.16	0.41	0.41	0.2	0.23	0.13	0.39	0.33
0.32	1.21	1.8	0.4	0.59	0.62		
0.03	1	0.27	0.17	0.78	0.39		
1.1	5.13	5.93	3.96	4.4	3.67	1.6	2
96.3	95.1	97	98.1	97.5	99.7	99.92	99.6
1855	221	150	569	672	604	166	1928
232	65.6	71.3	61.5	142.5	22.3	56.4	251
4.06	2.16	2.37	1.15	2.9	0.89	1.75	4.01
10	6	6	450	390	70		
140	138.5	74.3	94.4	86.9	80.4	75	193
567	121	308	185	110	373	150.5	474
2	2	3	1	1	1	3	2
567	121	308	185	110	373	150.5	474
20.3	2.51	2.59	6.01	13.3	1.01	2.5	31.7
4.87	0.93	0.64	1.84	3.55	0.49	1.9	5.4
85	228	101	179	140	138	49	134
2	17	12	3	5	5	3.7	7.3
27.9	49	51.3	13.9	29	15.9	36.3	28.2
412	268	270	118	262	112	244.2	411
8.1	300	39.4	68.4	300	22.2	15.1	10.6
0.003	3.21	0.008	0.008	0.344	0.008	0.0061	0
0.33	0.06	0.07	0.29	1.62	0.18	0.1	0.5
0.318	0.927	0.083	0.08	0.296	7.92	0.12	0.03
0.79	5.12	1.08	2.13	5.67	0.81	7	0.5
0.5	1.4	0.6	0.4	1.1	0.4	0.3	0.3
0.006	0.38	0.006	0.04	0.35	0.48		
0.8	0.9	0.3	0.3	0.3	0.6	0.3	0.06
19	82	88	51	74	9	78.7	6
0.6	6	1	1	2	5	1	0.8
5	4	3	222	202	37	1.5	2.3
30	7	2	16	15	4	7.7	21.1
105	120	119	91	86	92	26	49

## APPENDICE B

### MODÉLISATION 3D DANS LE GISEMENT DE BEATTIE

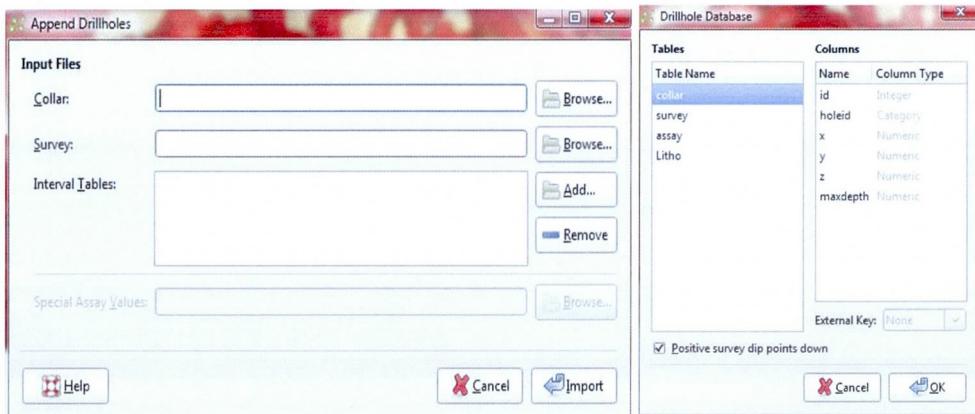
## Introduction

La modélisation 3D apparaît comme un outil incontournable pour la visualisation de différents paramètres, notamment la minéralisation, l'altération, la structure, et la lithologie. Le logiciel Leapfrog 3D ® est un logiciel axé sur l'exploration minière. Il est développé par la société ARANZ Geo qui est une division de Applied Research Associated New Zealand Limited (ARANZ). L'intérêt de Leapfrog 3D, en comparaison avec ses compétiteurs directs tel que gOcad notamment, est la facilité et la rapidité des traitements de modélisation 3D.

Dans le gisement de Beattie, de nombreux paramètres semblent influencer la minéralisation, qu'ils soient structuraux, lithologiques, ou géochimiques. Notre intérêt est de visualiser les relations, mais aussi d'utiliser le logiciel à son plein potentiel pour mettre en évidence des composantes non visibles initialement. En appliquant par exemple un poids à une faille, elle va influencer selon la valeur du poids la quantité de déformation dans le gisement. Ces outils permettent l'affinement du modèle. D'autres paramètres pertinents sont les calculs des volumes, ou encore la possibilité de réaliser des films à partir du modèle 3D.

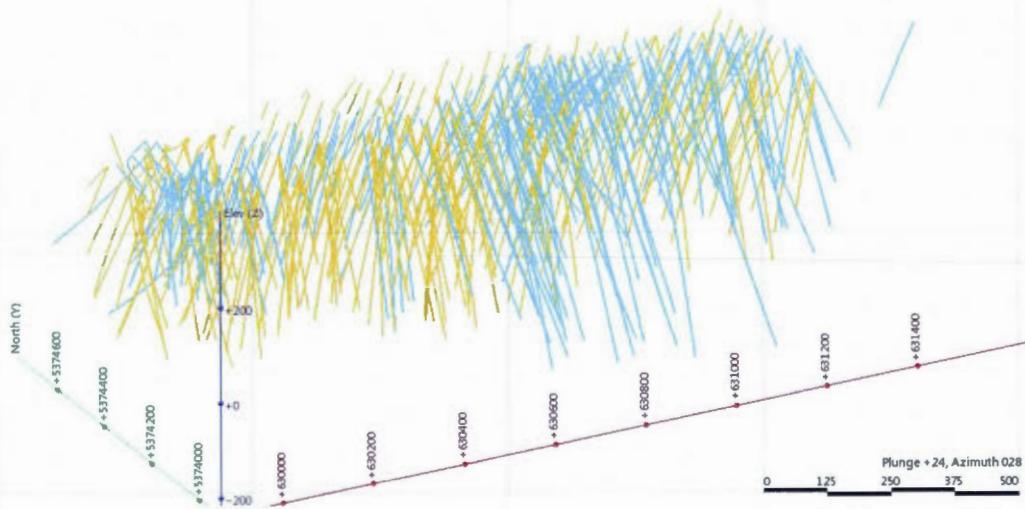
## Méthode

L'importation de données se fait à partir de tableur excel convertit en format CSV. Trois tables essentielles sont à intégrer pour la réalisation d'un modèle de base, *collar* (holeID, x, y, z, max depth), *survey* (holeID, depth, dip, azimuth), et *assay* (holeID, from, to, assay) (Figure 1).



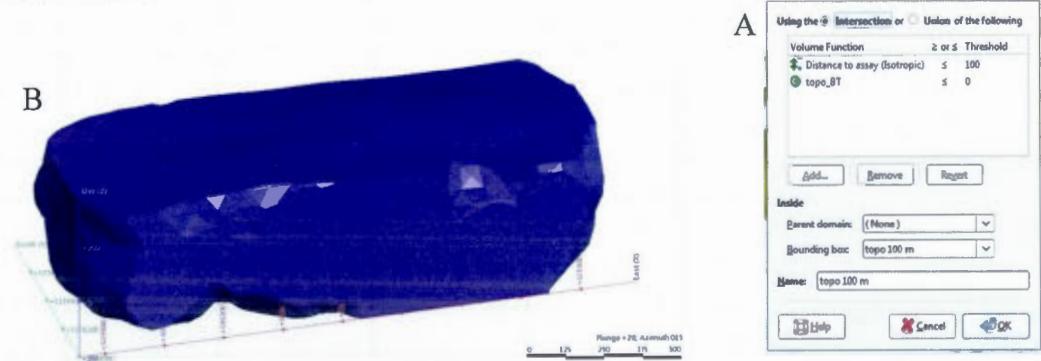
**Figure 1.** Importation de la base de données

Une fois la base de données importée, les forages sont immédiatement visualisable (Figure 2).



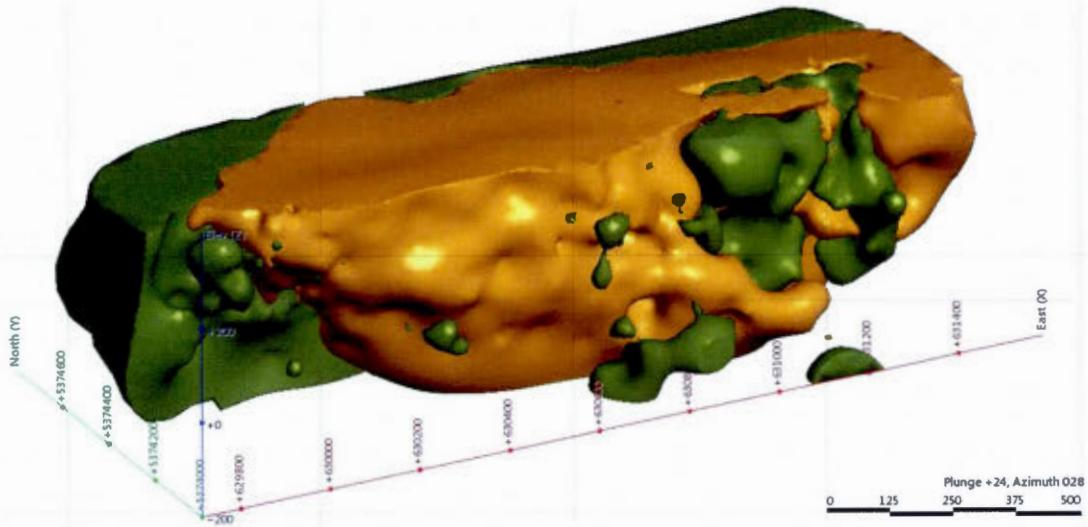
**Figure 2.** Visualisation des forages

Des domaines qui définissent la zone d'influence des forages doivent être déterminés par l'utilisateur. De cette valeur dépend la forme et le volume des corps qui vont être créées. La figure 3 présente que le domaine est contraint sous la surface et la zone d'influence des forages s'étend à 100m au maximum. Il s'agit du domaine que nous avons choisi pour réaliser l'étude.

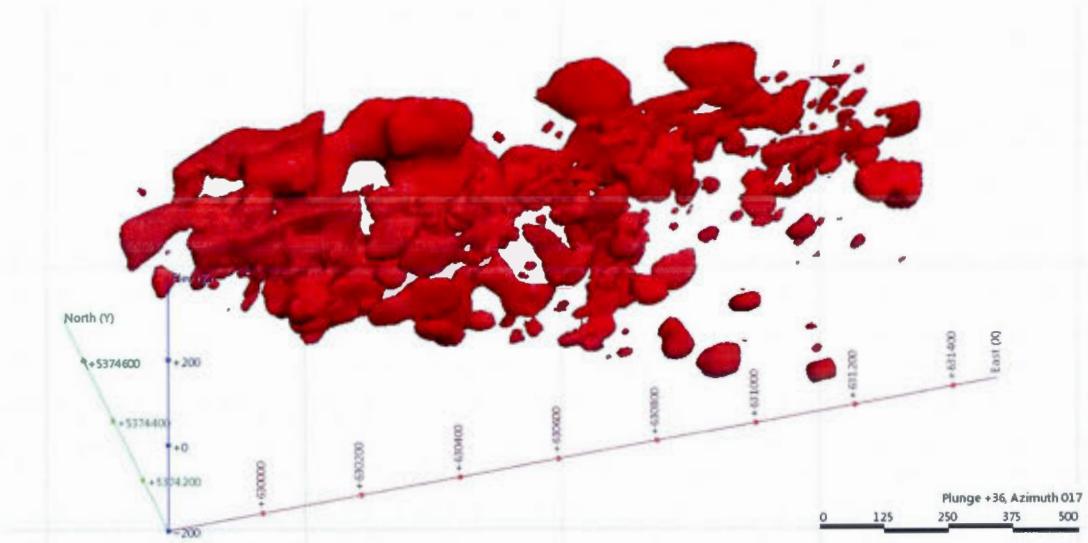


**Figure 3.** (a) Définition du domaine d'étude et du champ d'influence des forages. (b) visualisation du champ d'influence.

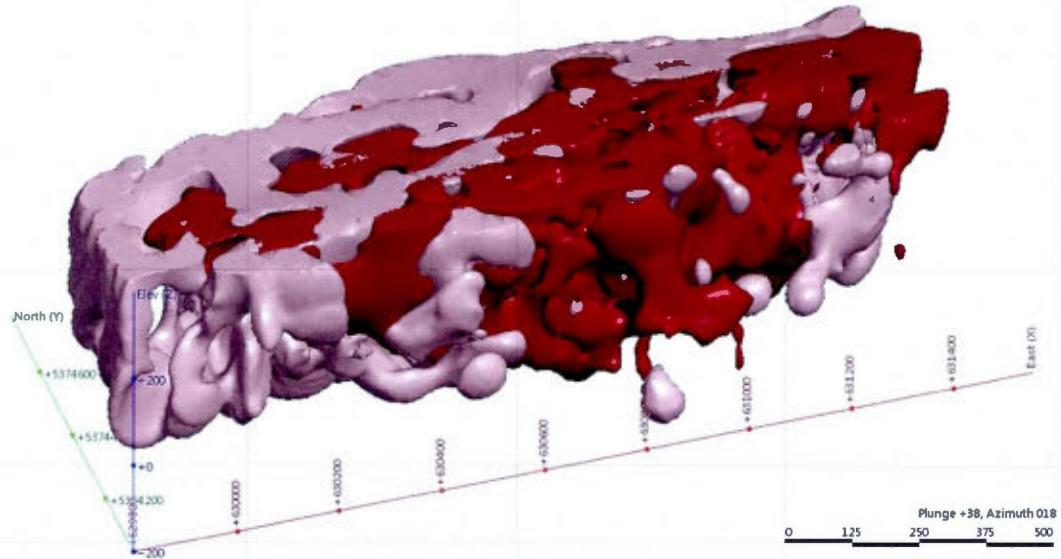
Après la définition du domaine d'étude, les volumes sont intégrés et rendu visibles : Les altérations, les zones minéralisées en fonction de la teneur donnée, et les lithologies (Figure 4,5,6).



**Figure 1.** Lithologies : volcanites du Kinogévis en vert, et la syénite de Beattie en orange.

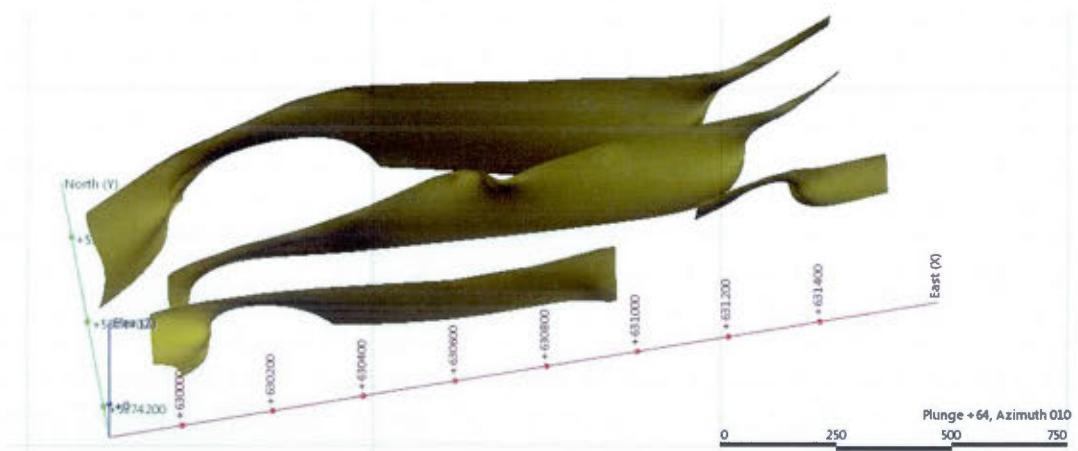


**Figure 2.** Zones minéralisées pour une teneur de 0.67g/t Au.



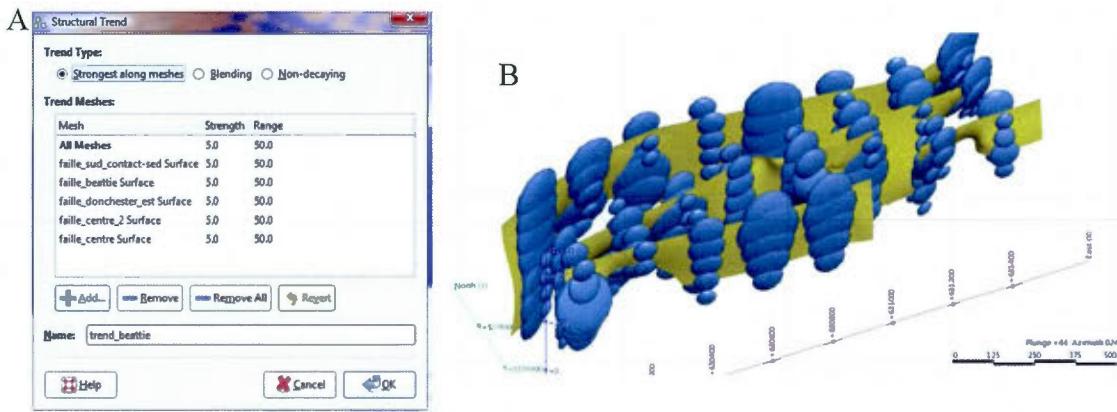
**Figure 3.** Zones d'altération : carbonatation en rose, hématisation en rouge vin.

Un module de dessin est intégré dans le logiciel. De nombreux objets structuraux peuvent ainsi être dessinés, notamment les failles (Figure 7). Il s'agit d'une interprétation en profondeur de la géologie en surface. Un moyen plus précis pour intégrer la structure est d'avoir une base de données de la structure à partir des forages. Il s'agit d'une base de données similaire à *assay* avec l'information sur la structure à chaque intervalle.



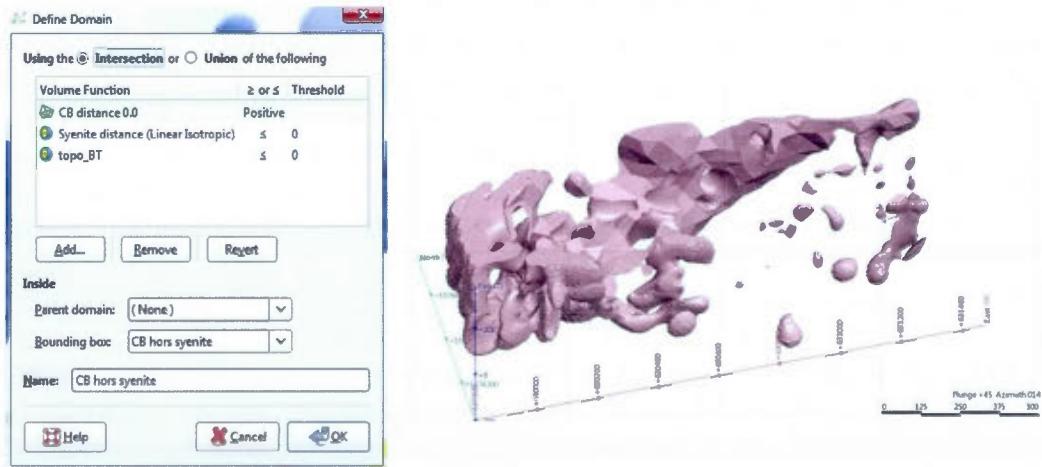
**Figure 4.** Système de failles dessiné à partir de la géologie de surface.

Lorsque la structure est dessinée, il est possible d'affiner le modèle 3D en le soumettant à l'influence d'une faille. Un poids et une zone d'influence va être donné à chaque faille (Figure 8a). La déformation sera d'autant plus intense qu'un corps est proche de la zone de faille (Figure 8b).



**Figure 8.** Structural trend, zone d'influence de chaque faille. (a) Définition du control structural – réglage des paramètres. (b) Visualisation de l'intensité de la déformation de long des failles.

Il est possible de définir des sous-domaines pour faire du traitement de données plus précis. Pour prendre un exemple, on veut connaitre le volume de l'altération à carbonate à l'extérieur de la syénite (Figure 9).



**Figure 9.** Sous domaine de carbonatation à l'extérieur de la syénite.

La phase ultime du traitement du modèle 3D est l'exportation d'image et de vidéo. Cette application est très intéressante dans le cadre de présentation du projet. Une vidéo du modèle 3D du gisement de Beattie est disponible. Pour l'obtenir, veuillez contacter l'auteur à l'adresse suivante : ludovic.bigot@gmail.com

## APPENDICE C

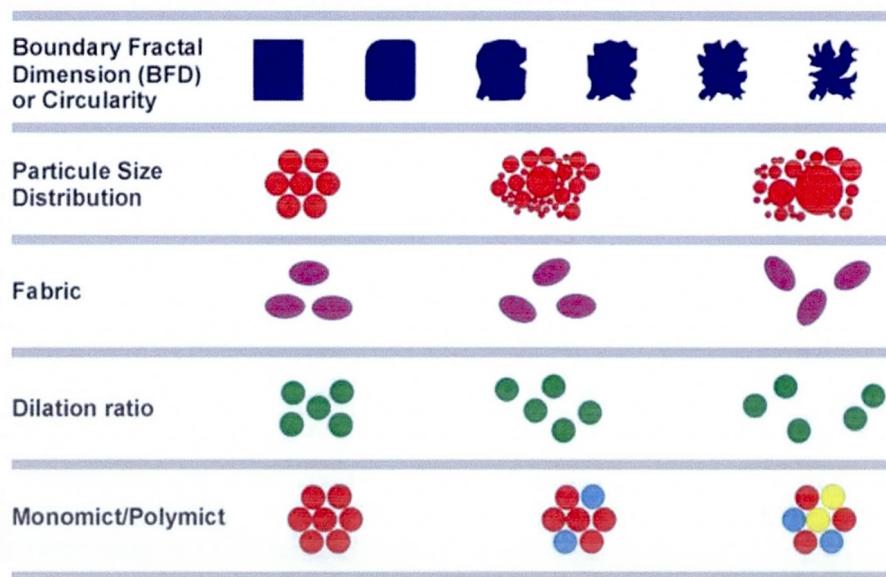
### ANALYSE DES BRÈCHES DU GISEMENT DE BEATTIE

## Introduction

Les brèches sont des objets géologiques qui sont communément présent dans les gisements. Elles sont présentent aussi bien dans les gisements magmatiques (Porphyres, épithermaux, IOCG), profonds et superficiels, que dans les gisements hydrothermaux (veines et shear zones), ou encore dans les gisements sédimentaires (MVT). La propagation des fluides à l'origine de la bréchification sont très variées (Jébrak, 1997): hydraulique, tectonique, explosion, et décompression. En association à ces processus de fracturation, la mobilité des fragments résulte de différents mécanismes: le broyage, l'effondrement, la fluidisation, et la dissolution.

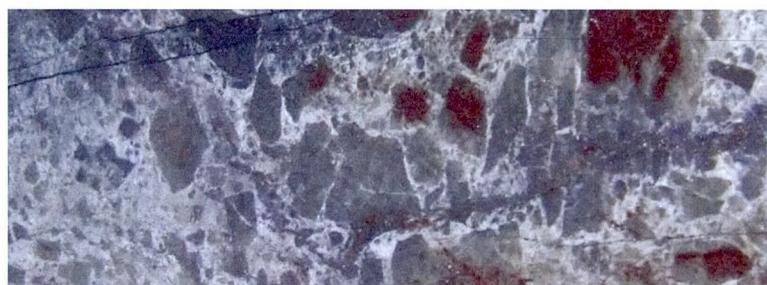
## Méthode

Jébrak (1997) propose une classification des brèches selon deux paramètres principaux, la description des clastes et du ciment, et la description générale de l'organisation de la brèche. Des paramètres non-dimensionnels sont nécessaires pour décrire la brèche (Figure 1), tels que la nature des clastes, l'espacement entre les clastes, la présence d'une fabrique, la distribution des particules, et la circularité ou complexité du claste.

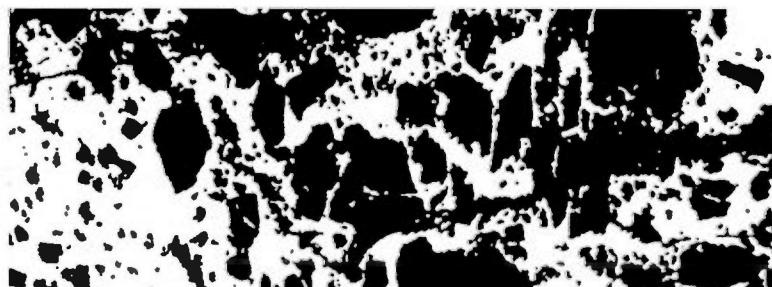


**Figure 1.** Paramètres non-dimensionnelles pour la classification des brèches.

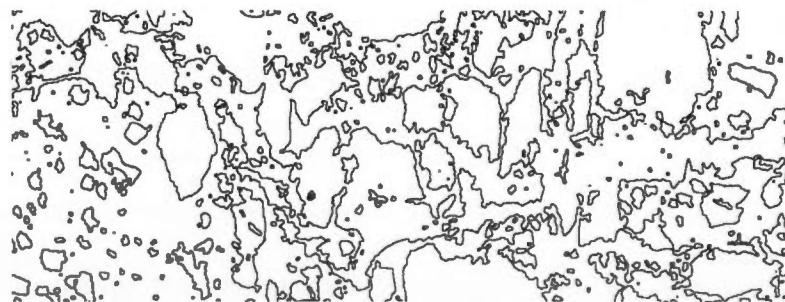
En connaissant cette méthode d'analyse, il faut un échantillon pour l'analyse. Cet échantillon est une brèche minéralisée d'une teneur de 6g/t Au (Figure 2; B08-78 ; 95.6m). Le traitement se fait par analyse d'image en utilisant le logiciel imageJ. Le traitement consiste en une séparation des fragments et de la matrice (Figure 3). Pour ce, le contraste et les couleurs sont retouchés. Puis le logiciel fait l'analyse des particules (résultats reportés dans le tableau.2). L'analyse se fait sur les fragments, et consiste-en des mesures de longueur, largeur, périmètre, et de circularité entre autres. Une fois l'analyse complète, l'image de brèche est observable sous forme de contour de fragments.



**Figure 2.** Brèche minéralisée du gisement de Beattie – exemple pour l'analyse.

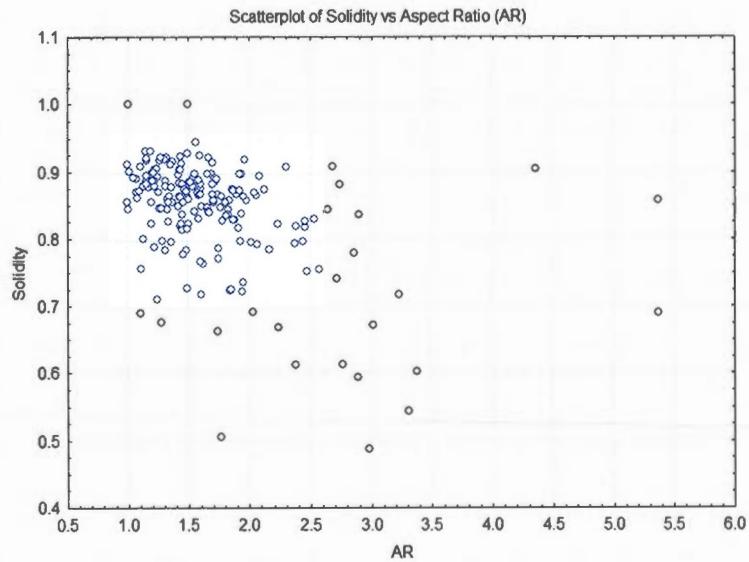


**Figure 3.** Image (Fig.2) retouchée pour l'analyse.

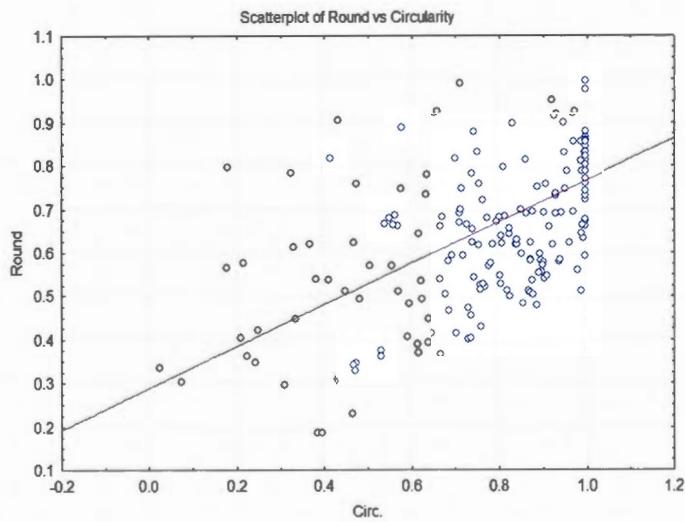


**Figure 4.** Contour des fragments isolés.

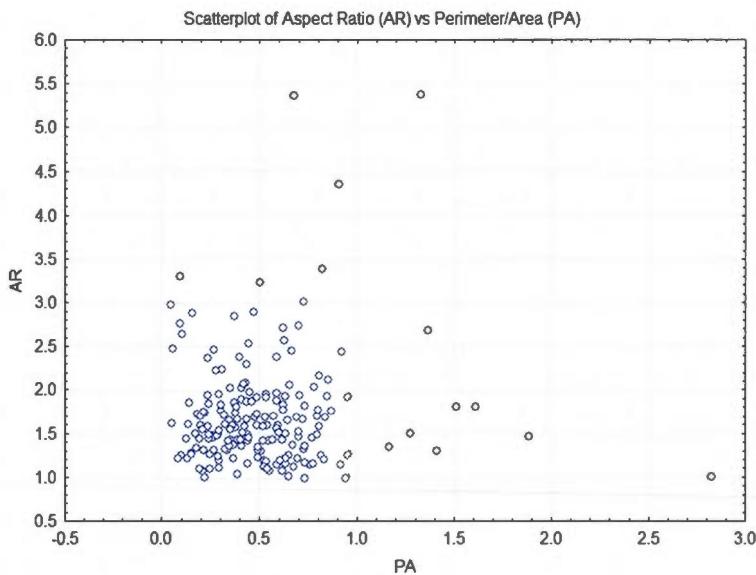
### Traitement des résultats de l'exemple de la brèche de Beattie



**Figure 5.** Solidité vs l'aspect ratio



**Figure 6.** Arrondissement vs la circularité (complexité des fragments)



**Figure 5.** Aspect Ratio vs Périmètre/Surface

**Table 1.** Résultat de l'analyse de l'exemple de brèche à Beattie

Parameter	Beattie breccia
Mono/Poly	Mono -syenite or volcanite
Dilatation Ratio	low to medium
Fabric Intensity	inherited to medium
Particles Size Distribution	normal - mm to cm
Aspect Ratio	medium - 1.66
Solidity	0.85
Perimeter/Area	0.59
Circularity	low

### Conclusions

Des paramètres de propagation hydraulique et tectonique peuvent être interprétés comme à l'origine de la fracturation de la roche. Pour la propagation hydraulique, la mobilité est très faible, considérée in-situ. Pour la propagation tectonique, la mobilité des clastes semble résulter du broyage étant donné la variation de la taille des fragments. La propagation hydraulique est nécessairement précoce, suivie par des processus tectonique. Les fluides auraient donc d'abord circulé dans les zones de faiblesse des failles Beattie, Donchester, et de

la Fracture zone. Par la suite, les mouvements tectoniques sur ces mêmes failles auraient entraîné une seconde bréchification.

#### Résultats de l'analyse des fragments de brèche par ImageJ :

	Area	Mean	Min	Max	X	Y	Perim.	Major	Minor	Angle	Circ.	AR	Round	Solidity	PA
1	200690	255	255	255	649.396	168.18	10176.56	871.94	293.05	175.459	0.02	2.975	0.336	0.487	0.051
2	425	255	255	255	226.832	9.164	90.184	24.176	22.382	41.323	0.66	1.08	0.926	0.892	0.212
3	6	255	255	255	377	0.833	9.071	3.718	2.055	0	0.92	1.809	0.553	0.857	1.512
4	6456	255	255	255	752.638	40.277	673.879	101.53	80.964	21.58	0.18	1.254	0.797	0.714	0.104
5	6	255	255	255	725	0.833	9.071	3.718	2.055	0	0.92	1.809	0.553	0.857	1.512
6	25	255	255	255	734.98	1.58	21.314	8.227	3.869	173.87	0.69	2.127	0.47	0.877	0.853
7	324	255	255	255	833.29	7.722	86.426	24.791	16.64	144.717	0.55	1.49	0.671	0.787	0.267
8	21	255	255	255	859.643	2.167	15.899	5.538	4.828	140.392	1	1.147	0.872	0.933	0.757
9	951	255	255	255	1123.48	20.885	138.711	47.534	25.473	94.861	0.62	1.866	0.536	0.877	0.146
10	12	255	255	255	255	6	11.314	3.909	3.909	0	1	1	1	0.857	0.943
11	14	255	255	255	197.714	9.143	13.314	4.746	3.756	24.358	0.99	1.264	0.791	0.848	0.951
12	51	255	255	255	822.48	15.343	27.556	10.202	6.365	95.775	0.84	1.603	0.624	0.879	0.540
13	24	255	255	255	180.375	18.167	16.728	6.111	5.001	109.605	1	1.222	0.818	0.906	0.697
14	62	255	255	255	210.597	23.161	29.799	11.569	6.824	95.653	0.88	1.695	0.59	0.899	0.481
15	53	255	255	255	669.538	31.104	32.87	13.532	4.987	51.784	0.62	2.713	0.369	0.741	0.620
16	36	255	255	255	694.75	34.778	22.728	9.426	4.863	83.003	0.88	1.939	0.516	0.9	0.631
17	49	255	255	255	841.153	36.582	26.142	10.304	6.055	93.75	0.9	1.702	0.588	0.916	0.534
18	12	255	255	255	1086	35	11.314	3.909	3.909	0	1	1	1	0.857	0.943
19	366	255	255	255	653.396	51.309	91.154	26.104	17.852	115.612	0.55	1.462	0.684	0.78	0.249
20	746	255	255	255	672.763	69.582	229.907	41.004	23.164	68.165	0.18	1.77	0.565	0.506	0.308
21	246	255	255	255	1103.92	49.695	72.912	18.715	16.736	13.375	0.58	1.118	0.894	0.759	0.296
22	20	255	255	255	695.2	45	16.971	7.014	3.631	129.806	0.87	1.932	0.518	0.8	0.849
23	21	255	255	255	111.405	47.024	16.142	6.258	4.272	118.684	1	1.465	0.683	0.857	0.769
24	17	255	255	255	687.794	49.324	14.728	6.19	3.497	159.699	0.99	1.77	0.565	0.85	0.866
25	28	255	255	255	606.286	54	18.142	6.024	5.918	90	1	1.018	0.982	0.903	0.648
26	19	255	255	255	75.342	55.658	14.728	5.301	4.564	135	1	1.161	0.861	0.884	0.775
27	50	255	255	255	576.52	58.42	36.284	13.838	4.6	33.324	0.48	3.008	0.332	0.671	0.726
28	186	255	255	255	105.129	66.871	55.598	16.845	14.059	140.337	0.76	1.198	0.835	0.825	0.299
29	32	255	255	255	59.438	63.656	20.142	6.883	5.919	179.302	0.99	1.163	0.86	0.889	0.629
30	160	255	255	255	1185.98	71.888	56.77	20.318	10.026	89.619	0.62	2.026	0.493	0.798	0.355
31	21	255	255	255	82.595	65.119	16.728	6.763	3.954	119.62	0.94	1.711	0.585	0.857	0.797
32	110	255	255	255	632.955	70.218	42.042	15.612	8.971	82.686	0.78	1.74	0.575	0.87	0.382
33	26	255	255	255	689.577	66.538	18.142	7.134	4.64	69.548	0.99	1.537	0.65	0.881	0.698

34	111	255	255	255	595.05	72.086	49.355	14.549	9.714	85.571	0.57	1.498	0.668	0.73	0.445
35	122	255	255	255	578.221	74.025	41.456	14.334	10.836	83.864	0.89	1.323	0.756	0.924	0.340
36	39	255	255	255	711.423	73.192	25.799	11.055	4.492	125.168	0.74	2.461	0.406	0.821	0.662
37	33	255	255	255	1153.53	70.379	21.556	9.321	4.508	167.522	0.89	2.068	0.484	0.868	0.653
38	52	255	255	255	1105.23	77.615	32.627	13.05	5.073	99.676	0.61	2.572	0.389	0.759	0.627
39	31	255	255	255	1200.89	74.919	20.142	7.055	5.594	178.535	0.96	1.261	0.793	0.873	0.650
40	44	255	255	255	533.864	78.682	24.385	7.802	7.181	58.717	0.93	1.087	0.92	0.863	0.554
41	72	255	255	255	55.736	81.694	48.527	22.174	4.134	30.3	0.38	5.364	0.186	0.689	0.674
42	163	255	255	255	612.715	83.739	53.355	16.618	12.488	22.209	0.72	1.331	0.751	0.847	0.327
43	1956	255	255	255	1159.74	102.51	214.208	81.128	30.698	158.056	0.54	2.643	0.378	0.843	0.110
44	46	255	255	255	295.957	86.63	23.556	8.761	6.686	155.705	1	1.31	0.763	0.92	0.512
45	179	255	255	255	709.366	91.494	52.527	16.875	13.506	47.751	0.82	1.249	0.8	0.895	0.293
46	47	255	255	255	797.734	88.011	27.799	10.617	5.636	68.77	0.76	1.884	0.531	0.832	0.591
47	12	255	255	255	94	90	11.314	3.909	3.909	0	1	1	1	0.857	0.943
48	75	255	255	255	535.58	94.687	33.213	11.647	8.199	107.244	0.85	1.421	0.704	0.898	0.443
49	22	255	255	255	677.773	93	16.142	5.741	4.879	90	1	1.176	0.85	0.88	0.734
50	631	255	255	255	596.614	114.99	156.167	32.016	25.094	95.856	0.33	1.276	0.784	0.676	0.247
51	12	255	255	255	175	95	11.314	3.909	3.909	0	1	1	1	0.857	0.943
52	12	255	255	255	278	97	11.314	3.909	3.909	0	1	1	1	0.857	0.943
53	12297	255	255	255	698.546	155.66	683.394	159.87	97.936	163.107	0.33	1.632	0.613	0.766	0.056
54	203	255	255	255	772.224	104.75	59.012	19.642	13.159	22.844	0.73	1.493	0.67	0.864	0.291
55	111	255	255	255	660.662	104.43	39.213	14.239	9.926	55.82	0.91	1.435	0.697	0.925	0.353
56	28	255	255	255	638.714	102	18.142	6.024	5.918	90	1	1.018	0.982	0.903	0.648
57	552	255	255	255	569.205	120.94	134.953	36.1	19.469	110.21	0.38	1.854	0.539	0.728	0.244
58	292	255	255	255	183.51	115.03	70.184	24.32	15.287	93.698	0.75	1.591	0.629	0.868	0.240
59	122	255	255	255	322.385	107.34	42.284	16.047	9.68	141.919	0.86	1.658	0.603	0.9	0.347
60	115	255	255	255	348.561	112.34	57.941	21.729	6.738	92.538	0.43	3.225	0.31	0.717	0.504
61	25	255	255	255	580.42	106.18	18.142	6.511	4.889	177.444	0.95	1.332	0.751	0.862	0.726
62	78	255	255	255	292.577	114.74	34.87	12.035	8.252	173.8	0.81	1.458	0.686	0.839	0.447
63	663	255	255	255	106.934	129.04	120.368	33.64	25.094	133.481	0.58	1.341	0.746	0.829	0.182
64	35	255	255	255	1152.53	120.73	21.556	7.991	5.577	76.427	0.95	1.433	0.698	0.886	0.616
65	96	255	255	255	542.729	125.28	39.456	15.226	8.028	121.127	0.78	1.897	0.527	0.869	0.411
66	13	255	255	255	65.885	124.89	11.899	4.38	3.779	135	1	1.159	0.863	0.897	0.915
67	42	255	255	255	1105.86	127.07	24.971	9.26	5.775	79.012	0.85	1.603	0.624	0.884	0.595
68	229	255	255	255	1095.3	142.29	60.77	20.627	14.135	72.591	0.78	1.459	0.685	0.884	0.265
69	191	255	255	255	516.081	142.66	54.77	18.988	12.808	126.363	0.8	1.483	0.675	0.876	0.287
70	34	255	255	255	549.5	138.24	20.142	7.074	6.12	#####	1	1.156	0.865	0.919	0.592
71	185	255	255	255	6.468	149.16	69.598	25.921	9.087	96.562	0.48	2.853	0.351	0.779	0.376
72	477	255	255	255	192.798	153.75	118.125	25.881	23.466	176.638	0.43	1.103	0.907	0.689	0.248

73	29	255	255	255	213.466	140.67	20.385	8.466	4.362	131.823	0.88	1.941	0.515	0.841	0.703
74	6	255	255	255	69	140.5	7.657	3.385	2.257	90	1	1.5	0.667	1	1.276
75	16	255	255	255	90.5	141	13.314	4.963	4.105	0	1	1.209	0.827	0.889	0.832
76	36	255	255	255	359.5	147	21.314	8.468	5.413	90	1	1.565	0.639	0.947	0.592
77	20	255	255	255	529.5	146	18.385	7.893	3.226	137.043	0.74	2.446	0.409	0.8	0.919
78	16	255	255	255	1074	146.5	13.314	4.963	4.105	90	1	1.209	0.827	0.889	0.832
79	901	255	255	255	60.085	176.9	213.522	52.218	21.969	139.623	0.25	2.377	0.421	0.613	0.237
80	12	255	255	255	507	154	11.314	3.909	3.909	0	1	1	1	0.857	0.943
81	594	255	255	255	1137.8	166.18	105.296	33.718	22.431	11.349	0.67	1.503	0.665	0.825	0.177
82	12	255	255	255	522	158	11.314	3.909	3.909	0	1	1	1	0.857	0.943
83	45	255	255	255	147.3	165.3	23.799	8.285	6.916	134.999	1	1.198	0.835	0.9	0.529
84	37	255	255	255	539.5	169.26	22.971	9.324	5.052	145.619	0.88	1.846	0.542	0.86	0.621
85	42	255	255	255	1102.24	171.45	24.142	9.578	5.583	101.584	0.91	1.716	0.583	0.894	0.575
86	151	255	255	255	523.076	177.81	49.941	17.6	10.924	101.976	0.76	1.611	0.621	0.851	0.331
87	827	255	255	255	196.525	186.63	127.924	36.761	28.644	18.282	0.64	1.283	0.779	0.848	0.155
88	28	255	255	255	151.786	177.61	18.728	7.002	5.092	97.047	1	1.375	0.727	0.918	0.669
89	20	255	255	255	1151	179.5	16.142	6.363	4.002	26.921	0.97	1.59	0.629	0.833	0.807
90	32	255	255	255	588.156	182.16	20.142	7.07	5.763	96.969	0.99	1.227	0.815	0.889	0.629
91	6	255	255	255	598.5	182	7.657	3.385	2.257	0	1	1.5	0.667	1	1.276
92	21	255	255	255	1143.98	184.41	16.142	6.258	4.272	28.684	1	1.465	0.683	0.857	0.769
93	74	255	255	255	20.838	188.89	33.456	13.661	6.897	148.08	0.83	1.981	0.505	0.86	0.452
94	34	255	255	255	1112.88	189.24	21.556	8.074	5.362	78.677	0.92	1.506	0.664	0.861	0.634
95	77	255	255	255	5.708	192.05	41.213	13.841	7.083	33.443	0.57	1.954	0.512	0.74	0.535
96	651	255	255	255	121.2	227.07	132.51	36.449	22.741	118.437	0.47	1.603	0.624	0.769	0.204
97	107	255	255	255	58.855	211.56	39.456	12.904	10.557	68.079	0.86	1.222	0.818	0.881	0.369
98	56	255	255	255	391	213.54	27.799	9.676	7.369	90	0.91	1.313	0.762	0.889	0.496
99	12	255	255	255	819	210	11.314	3.909	3.909	0	1	1	1	0.857	0.943
100	1712	255	255	255	169.762	242.33	315.304	61.517	35.434	131.422	0.22	1.736	0.576	0.663	0.184
101	26	255	255	255	208.538	215.42	18.142	7.134	4.64	159.548	0.99	1.537	0.65	0.881	0.698
102	6	255	255	255	25	216.5	7.657	3.385	2.257	90	1	1.5	0.667	1	1.276
103	12	255	255	255	77	219	11.314	3.909	3.909	0	1	1	1	0.857	0.943
104	199	255	255	255	76.575	235.73	61.012	21.58	11.741	90.721	0.67	1.838	0.544	0.847	0.307
105	65	255	255	255	397.238	227.38	28.971	9.801	8.444	138.233	0.97	1.161	0.861	0.922	0.446
106	12	255	255	255	28	227	11.314	3.909	3.909	0	1	1	1	0.857	0.943
107	151	255	255	255	382.944	232.29	49.113	18.188	10.571	162.361	0.79	1.721	0.581	0.888	0.325
108	25	255	255	255	60.18	233.82	17.556	6.852	4.645	135	1	1.475	0.678	0.877	0.702
109	21	255	255	255	409.5	237.5	15.314	5.171	5.171	0	1	1	1	0.913	0.729
110	12	255	255	255	337	240	11.314	3.909	3.909	0	1	1	1	0.857	0.943
111	446	255	255	255	38.993	259.04	95.397	29.7	19.12	89.36	0.62	1.553	0.644	0.844	0.214

112	116	255	255	255	794.052	248.97	45.113	14.575	10.134	148.086	0.72	1.438	0.695	0.823	0.389
113	43	255	255	255	813.221	247.76	23.556	8.842	6.192	31.36	0.97	1.428	0.7	0.905	0.548
114	6	255	255	255	0.833	249	9.657	3.718	2.055	90	0.81	1.809	0.553	0.857	1.610
115	503	255	255	255	335.335	263.72	111.882	33.485	19.126	119.177	0.51	1.751	0.571	0.789	0.222
116	49	255	255	255	347.541	253.54	24.385	8.796	7.093	96.8	1	1.24	0.806	0.907	0.498
117	357	255	255	255	120.847	262.74	79.497	21.431	21.21	146.066	0.71	1.01	0.99	0.847	0.223
118	3798	255	255	255	1107.85	291.69	337.605	76.846	62.928	155.156	0.42	1.221	0.819	0.791	0.089
119	278	255	255	255	141.759	274.33	69.841	22.931	15.436	99.978	0.72	1.486	0.673	0.874	0.251
120	103	255	255	255	783.432	272.49	39.113	14.38	9.12	44.947	0.85	1.577	0.634	0.873	0.380
121	17	255	255	255	962.088	268.38	13.899	5.229	4.14	74.828	1	1.263	0.792	0.919	0.818
122	230	255	255	255	999.509	273.62	61.698	19.608	14.935	4.734	0.76	1.313	0.762	0.881	0.268
123	19	255	255	255	805.342	270.66	14.728	5.301	4.564	135	1	1.161	0.861	0.884	0.775
124	456	255	255	255	979.649	283.95	91.054	31.514	18.424	92.78	0.69	1.71	0.585	0.86	0.200
125	5708	255	255	255	369.35	342.69	566.441	141.72	51.28	92.446	0.22	2.764	0.362	0.613	0.099
126	10	255	255	255	0.9	273	13.657	5.848	2.177	90	0.67	2.686	0.372	0.909	1.366
127	24	255	255	255	1025.67	276.21	18.728	7.909	3.863	172.154	0.86	2.047	0.488	0.873	0.780
128	205	255	255	255	49.978	289.15	61.355	22.617	11.541	92.223	0.68	1.96	0.51	0.867	0.299
129	501	255	255	255	1046.49	289.17	118.711	35.295	18.073	7.877	0.45	1.953	0.512	0.726	0.237
130	19	255	255	255	313.342	282.34	14.728	5.301	4.564	45.001	1	1.161	0.861	0.884	0.775
131	12	255	255	255	662	282	11.314	3.909	3.909	0	1	1	1	0.857	0.943
132	79	255	255	255	1018.97	287.93	32.627	11.695	8.601	90.555	0.93	1.36	0.735	0.913	0.413
133	81	255	255	255	296.377	292.24	35.456	13.911	7.414	51.47	0.81	1.876	0.533	0.876	0.438
134	121	255	255	255	32.822	302.22	48.284	19.161	8.041	113.529	0.65	2.383	0.42	0.823	0.399
135	170	255	255	255	10.618	306.63	50.77	18.282	11.84	115.561	0.83	1.544	0.648	0.895	0.299
136	483	255	255	255	123.084	317.74	104.711	32.823	18.736	106.033	0.55	1.752	0.571	0.774	0.217
137	87	255	255	255	174.121	307.56	36.284	12.953	8.552	16.631	0.83	1.515	0.66	0.888	0.417
138	84	255	255	255	439.619	309.01	36.042	13.067	8.185	77.911	0.81	1.596	0.626	0.87	0.429
139	12	255	255	255	687	305	11.314	3.909	3.909	0	1	1	1	0.857	0.943
140	26	255	255	255	93.577	310.46	18.142	7.134	4.64	110.452	0.99	1.537	0.65	0.881	0.698
141	307	255	255	255	304.663	320.19	75.497	23.852	16.388	69.967	0.68	1.455	0.687	0.816	0.246
142	17200	255	255	255	1091.31	399.51	1710.209	269.01	81.41	5.201	0.07	3.304	0.303	0.543	0.099
143	293	255	255	255	847.336	326.47	79.012	30.315	12.306	97.769	0.59	2.463	0.406	0.83	0.270
144	22	255	255	255	1169	314.5	16.142	6.173	4.537	148.799	1	1.361	0.735	0.88	0.734
145	119	255	255	255	1044.69	319.9	43.456	15.595	9.716	97.356	0.79	1.605	0.623	0.869	0.365
146	56	255	255	255	25.25	319.27	27.556	10.678	6.678	107.917	0.93	1.599	0.625	0.926	0.492
147	36	255	255	255	788.583	319.11	22.385	9.028	5.077	49.846	0.9	1.778	0.562	0.867	0.622
148	19	255	255	255	91.763	319.97	15.899	6.41	3.774	101.564	0.94	1.699	0.589	0.884	0.837
149	41	255	255	255	411.5	324.5	22.142	7.892	6.615	134.999	1	1.193	0.838	0.932	0.540
150	34	255	255	255	30.647	329.06	21.799	8.475	5.108	55.87	0.9	1.659	0.603	0.85	0.641

151	58	255	255	255	779.397	331.69	28.385	11.319	6.524	9.464	0.91	1.735	0.576	0.892	0.489
152	55	255	255	255	144.118	335.85	27.213	9.725	7.201	176.121	0.93	1.351	0.74	0.866	0.495
153	17666	255	255	255	727.791	403.27	1033.661	235.95	95.33	1.8	0.21	2.475	0.404	0.756	0.059
154	52	255	255	255	1126.02	338.62	27.213	9.724	6.809	36.286	0.88	1.428	0.7	0.867	0.523
155	391	255	255	255	20.63	350.88	77.497	26.874	18.525	126.085	0.82	1.451	0.689	0.914	0.198
156	41	255	255	255	531.061	340.45	24.142	10.13	5.153	171.763	0.88	1.966	0.509	0.921	0.589
157	120	255	255	255	179.267	350.26	44.284	14.503	10.535	94.545	0.77	1.377	0.726	0.857	0.369
158	203	255	255	255	113.204	353.26	63.255	24.053	10.746	138.975	0.64	2.238	0.447	0.825	0.312
159	97	255	255	255	77.809	349.21	43.698	17.701	6.977	178.69	0.64	2.537	0.394	0.833	0.450
160	117	255	255	255	3.218	359.73	55.799	20.785	7.167	100.869	0.47	2.9	0.345	0.836	0.477
161	82	255	255	255	198.878	351.65	33.213	12.252	8.522	126.806	0.93	1.438	0.696	0.906	0.405
162	19	255	255	255	1064.66	348.66	14.728	5.301	4.564	45.001	1	1.161	0.861	0.884	0.775
163	370	255	255	255	305.305	358.57	90.811	26.521	17.763	22.443	0.56	1.493	0.67	0.818	0.245
164	215	255	255	255	224.063	363.09	81.154	22.572	12.128	80.933	0.41	1.861	0.537	0.729	0.377
165	388	255	255	255	523.969	360.51	76.426	23.407	21.105	122.851	0.84	1.109	0.902	0.91	0.197
166	2004	255	255	255	285.322	394.54	322.919	85.805	29.737	125.948	0.24	2.885	0.347	0.593	0.161
167	13	255	255	255	1090.12	357.89	11.899	4.38	3.779	45	1	1.159	0.863	0.897	0.915
168	57	255	255	255	327.605	363.62	30.627	11.798	6.152	161.307	0.76	1.918	0.521	0.82	0.537
169	993	255	255	255	66.358	379.33	133.054	39.256	32.207	167.973	0.71	1.219	0.82	0.901	0.134
170	153	255	255	255	877.958	370.69	63.255	19.889	9.794	178.005	0.48	2.031	0.492	0.691	0.413
171	113	255	255	255	157.881	371.13	44.284	15.514	9.274	144.102	0.72	1.673	0.598	0.843	0.392
172	1	255	255	255	135.5	367.5	2.828	1.128	1.128	0	1	1	1	1	2.828
173	26	255	255	255	95.538	371.15	18.971	7.785	4.252	35.792	0.91	1.831	0.546	0.839	0.730
174	6	255	255	255	1067.67	370.33	8.485	3.14	2.433	45	1	1.291	0.775	0.8	1.414
175	24	255	255	255	247	373.5	17.314	7.142	4.279	90	1	1.669	0.599	0.923	0.721
176	14	255	255	255	449	377.5	13.314	5.862	3.041	53.23	0.99	1.928	0.519	0.875	0.951
177	30	255	255	255	348.4	380.07	20.728	8.065	4.736	84.323	0.88	1.703	0.587	0.87	0.691
178	26	255	255	255	1211.73	378.92	20.971	8.476	3.905	142.824	0.74	2.17	0.461	0.788	0.807
179	22	255	255	255	1042.18	380.23	17.556	7.071	3.961	27.905	0.9	1.785	0.56	0.83	0.798
180	89	255	255	255	978.455	387.5	34.87	10.915	10.382	144.694	0.92	1.051	0.951	0.894	0.392
181	90	255	255	255	1052.73	389.83	39.213	15.481	7.402	75.941	0.74	2.091	0.478	0.896	0.436
182	114	255	255	255	439.018	392.05	43.698	16.257	8.928	140.753	0.75	1.821	0.549	0.838	0.383
183	1749	255	255	255	462.561	419.13	244.936	59.96	37.139	55.186	0.37	1.614	0.619	0.722	0.140
184	61	255	255	255	174.057	397.86	32.042	9.385	8.276	137.335	0.75	1.134	0.882	0.803	0.525
185	63	255	255	255	875.468	396.14	30.971	12.435	6.451	170.894	0.83	1.928	0.519	0.9	0.492
186	10	255	255	255	1214.1	399	13.657	5.848	2.177	90	0.67	2.686	0.372	0.909	1.366
187	1289	255	255	255	111.665	419.27	168.995	48.757	33.661	135.946	0.57	1.448	0.69	0.826	0.131
188	162	255	255	255	145.846	405.07	56.77	16.758	12.308	127.811	0.63	1.362	0.734	0.798	0.350
189	12	255	255	255	78	403	11.314	3.909	3.909	0	1	1	1	0.857	0.943

190	12	255	255	255	837	403	11.314	3.909	3.909	0	1	1	1	0.857	0.943
191	119	255	255	255	197.256	407.42	50.184	17.731	8.545	156.679	0.59	2.075	0.482	0.796	0.422
192	34	255	255	255	236.647	406.15	20.971	6.837	6.332	77.196	0.97	1.08	0.926	0.872	0.617
193	700	255	255	255	13.464	419.71	108.77	33.691	26.454	144.357	0.74	1.274	0.785	0.923	0.155
194	76	255	255	255	887.539	413.63	35.213	13.468	7.185	117.423	0.77	1.874	0.534	0.831	0.463
195	125	255	255	255	324.028	416.95	44.042	15.734	10.116	91.487	0.81	1.555	0.643	0.899	0.352
196	36	255	255	255	512.056	413.67	21.799	7.408	6.187	171.674	0.95	1.197	0.835	0.857	0.606
197	563	255	255	255	411.699	430.51	122.468	30.747	23.314	120.402	0.47	1.319	0.758	0.787	0.218
198	26	255	255	255	165.846	417.15	17.556	6.12	5.409	45	1	1.131	0.884	0.881	0.675
199	484	255	255	255	60.579	431.86	134.953	37.102	16.609	159.441	0.33	2.234	0.448	0.669	0.279
200	12	255	255	255	521	422	11.314	3.909	3.909	0	1	1	1	0.857	0.943
201	149	255	255	255	511.802	430.5	51.113	16.418	11.555	108.78	0.72	1.421	0.704	0.851	0.343
202	45	255	255	255	903.656	425.83	25.556	9.079	6.311	99.211	0.87	1.439	0.695	0.865	0.568
203	16	255	255	255	523	429.5	13.314	4.963	4.105	90	1	1.209	0.827	0.889	0.832
204	9	255	255	255	359.389	431.83	10.485	3.919	2.924	110.301	1	1.34	0.746	0.857	1.165
205	45	255	255	255	227.211	436.99	24.385	7.956	7.202	46.719	0.95	1.105	0.905	0.874	0.542
206	19	255	255	255	140.658	436.34	14.728	5.301	4.564	135	1	1.161	0.861	0.884	0.775
207	93	255	255	255	854.758	439.35	40.87	14.079	8.41	177.853	0.7	1.674	0.597	0.842	0.439
208	51	255	255	255	606.245	439.66	30.042	12.435	5.222	136.309	0.71	2.381	0.42	0.797	0.589
209	24	255	255	255	96.5	438	17.314	7.142	4.279	0	1	1.669	0.599	0.923	0.721
210	3	255	255	255	285.833	437.17	5.657	2.365	1.615	135	1	1.464	0.683	0.857	1.886
211	53	255	255	255	10.689	442.16	27.556	10.361	6.513	173.673	0.88	1.591	0.629	0.891	0.520
212	60	255	255	255	320.25	442.43	49.355	16.06	4.757	172.186	0.31	3.376	0.296	0.603	0.823
213	86	255	255	255	1146.49	441.88	37.556	15.87	6.9	179.829	0.77	2.3	0.435	0.91	0.437
214	47	255	255	255	493.989	442.39	28.142	9.534	6.277	11.686	0.75	1.519	0.658	0.887	0.599
215	48	255	255	255	375.604	443.13	33.556	12.944	4.721	0.356	0.54	2.742	0.365	0.881	0.699
216	20	255	255	255	86.8	443.25	15.899	6.168	4.128	170.604	0.99	1.494	0.669	0.93	0.795
217	33	255	255	255	1103.32	443.71	29.899	13.524	3.107	1.861	0.46	4.353	0.23	0.904	0.906
218	6	255	255	255	307	444.17	9.071	3.718	2.055	0	0.92	1.809	0.553	0.857	1.512
219	6	255	255	255	514	444.17	9.071	3.718	2.055	0	0.92	1.809	0.553	0.857	1.512
220	18	255	255	255	1088.33	444.11	23.899	11.094	2.066	179.345	0.4	5.37	0.186	0.857	1.328

## APPENDICE D

### APPORT DE LA SUSCEPTIBILITÉ MAGNÉTIQUE À L'EXPLORATION DU GISEMENT DE BEATTIE

## Introduction

Les travaux minéralogiques ont montré que les concentrations en or sur la propriété de Beattie semblent associées à la destruction de magnétite, et dans une certaine mesure l'absence d'hématite.

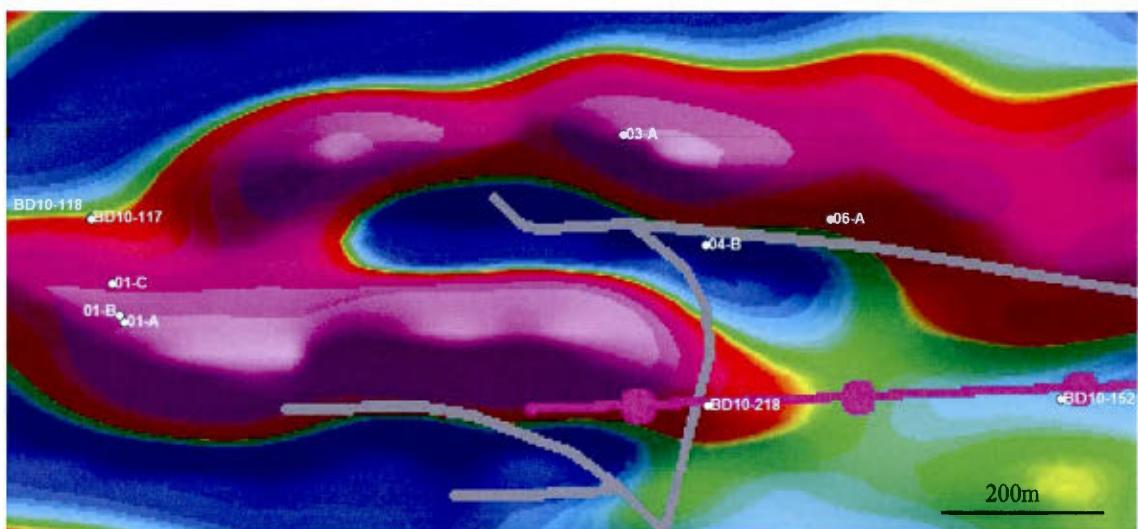
Un levé géophysique aéroporté à haute résolution, commandé par Corporation Minière Osisko, a été réalisé sur la propriété de Beattie. L'objectif de cette étude était de mettre en évidence les zones minéralisées, à l'échelle du gisement (Wares, 2010). L'utilisation d'un susceptibilimètre magnétique est efficace sur une plus faible échelle; c'est le cas pour les carottes de forage qui ont un volume connu. L'intérêt de la présente étude est de caractériser par les mesures de susceptibilité magnétique les différents faciès d'altération de la syénite de Beattie, et de les comparer avec les teneurs en or. La connaissance des valeurs types de chaque faciès d'altération serait un moyen complémentaire au loggage pour distinguer les roches.

## Méthode

Les mesures de susceptibilité magnétique ont été réalisées sur douze demi-carottes de forages et six échantillons de surface provenant de la propriété Beattie à Duparquet (Figure 1). Les carottes de forages sont de type NQ; la largeur est 4,6cm pour environ 10cm de longueur; les échantillons de surface ont des volumes comparables aux demi-carottes de forage. Six faciès lithologiques ont été distingués : le faciès de syénite/volcanite silicifiée-bréchifiée-remplacée (SISY-BRSY-RESY), le faciès de syénite fraîche très peu altéré à chlorite-carbonate (INSY), le faciès de syénite séricitisée (SRSY), le faciès de syénite carbonatée (CBSY), le faciès de syénite hématisée (HMSY) un peu carbonaté (transition entre martite et carbonatisation), et le faciès de syénite martitisée (HMSY).

Les résultats de susceptibilité magnétiques ont été associés aux teneurs en or dans chaque faciès; des observations pétrographiques ont permis une bonne distinction de deux faciès d'alteration : le faciès de syénite martitisée non aurifère, et le faciès de syénite hématisée (dusty hematite) qui peut être associé à des valeurs en or.

Le susceptibilimètre magnétique utilisé est une sonde MPP-EM2S+ - sonde multi-paramètres - de la compagnie Instrumentation GDD inc. La sonde détecte la susceptibilité magnétique avec une précision de 10-6 SI, ainsi que les valeurs de conductivité relative et absolue (MHOS/M). Les mesures ont été faites dans le laboratoire de recherche en géologie économique de l'UQAM. Le « bruit magnétique », ou bruit de fond, durant l'analyse était de 0,0001 SI.



**Figure 6.** Localisation des échantillons de surface et des trous de forage (BD...) une carte géophysique SVD de Duparquet.

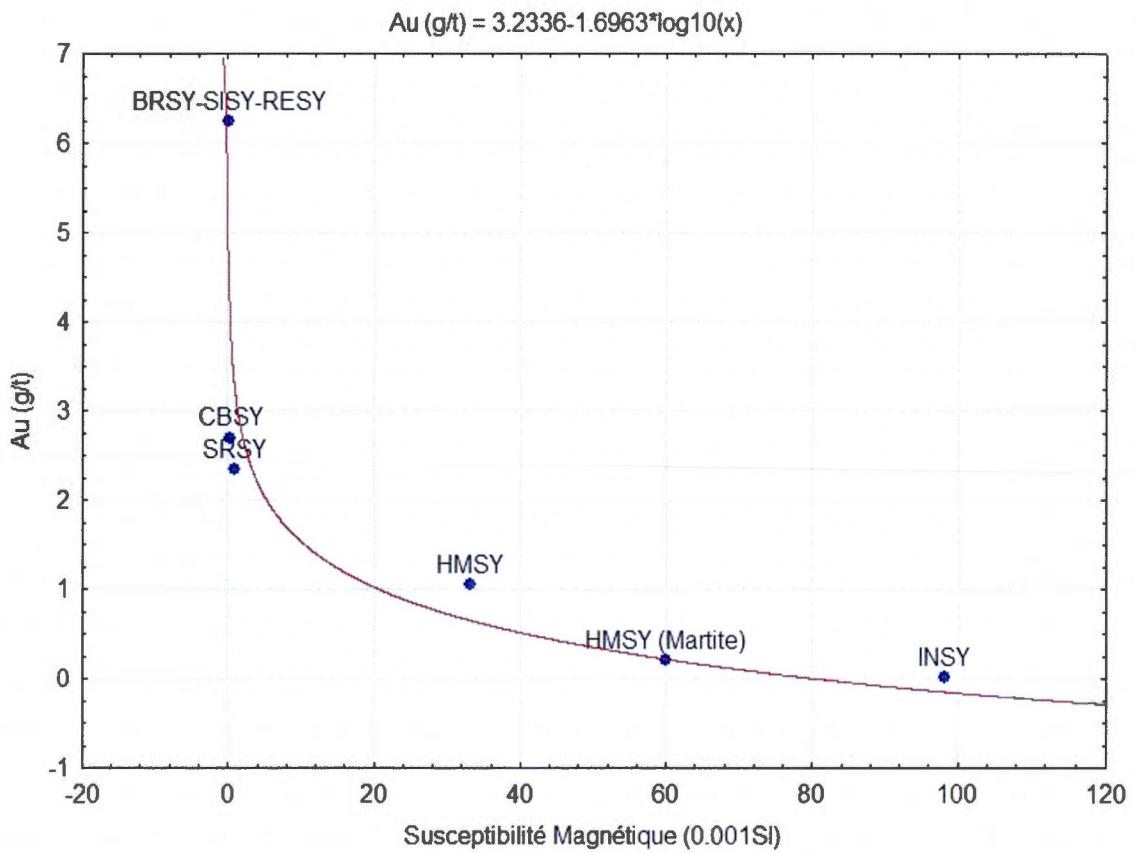
## Résultats

La susceptibilité magnétique varie grandement dans les faciès lithologiques de Beattie (Table 1); il y a un ordre de grandeur de quasiment 1000 entre le faciès de syénite silicifiée-bréchifiée-remplacée, et le faciès de syénite fraîche.

Les observations pétrographiques ont permis de constater que le faciès martitisé est dépourvu de pyrite, alors que dans le faciès hématisé il y a une faible quantité de pyrite (inférieur à 2%).

**Table 1.** Comparaison des différents faciès d'altération à Beattie

Faciès d'alteration	moyenne Au (g/t)	Susceptibilité Magnétique (0,001 SI)	Texture de Minéralisation	Type de Minéralisation
Syénite/volcanites silicifiée-bréchifiée-rempacée (SISY-BRSY-RESY - dans shear zones)	6,24	0,12	pyrite (15%) fine cataclasée dans des corridors siliceux	très aurifère; inclus dans pyrite, parfois à l'état libre
Syénite carbonatée (CBSY - dans intrusif)	2,7	0,43	pyrite (8%) disséminée dans la roche	aurifère; inclus dans pyrite et arsénopyrite
Syénite sérichtisée (SRSY - dans shear zones)	2,35	1,05	pyrite (2%) souvent concentrée dans des corridors à séricite	aurifère; inclus dans pyrite et arsénopyrite
Syénite hématisée (HMSY - dans intrusif)	1,06	33,25	hématite disséminée, faible pyrite (inf 2%)	Aurifère parfois; inclus dans pyrite
Syénite martitisée (HMSY - dans intrusif)	0,2	60	martite (5%) disséminée dans la roche	très faible aurifère
Syénite fraîche peu altérée à chlorite (INSY - dans intrusif)	0,0224	98,2	magnétite (5%) disséminée dans la roche	non aurifère



**Figure 2.** Teneur en or versus la susceptibilité magnétique mesurée dans les différents faciès d'altération

### Discussion

La distribution des sulfures, disséminée dans l'intrusion ou contrôlée dans des failles cisaillantes, n'est pas déterminante pour la valeur de la susceptibilité magnétique. C'est la concentration en fer dans la roche (naturelle ou par altération) qui contrôle les valeurs de susceptibilité magnétique. Plus elle est forte, plus la susceptibilité magnétique sera élevée. Une étude minéralogique a mis en évidence les fortes concentrations en fer, marquées par la magnétite (supérieur à 55% des minéraux métalliques), dans les faciès non altérés de la syénite de Beattie. Ainsi, à Beattie, plus l'altération est prononcée (marquée par la disparition des oxydes de fer magnétite et hématite), plus la valeur de susceptibilité magnétique est faible.

À partir des mesures de susceptibilité magnétique, il serait possible d'établir une évaluation qualitative de l'altération des roches (Figure 3). La méthode serait basée sur une numérotation de 1 à 5 selon la valeur de susceptibilité magnétique : (1) susceptibilité magnétique d'environ  $100 \cdot 10^{-3}$  SI. La roche est fraîche, pas ou très peu altérée; (2) susceptibilité magnétique d'environ  $60 \cdot 10^{-3}$  SI. La martitisation témoigne du départ du fer; (3) susceptibilité magnétique d'environ  $30 \cdot 10^{-3}$  SI correspondant à l'hématisation de la roche sous forme de dusty hématite résultant de la destruction de l'hématite avec un peu de carbonatation; (4) susceptibilité magnétique entre 1 et  $0,4 \cdot 10^{-3}$  SI correspondant à la carbonatation et la séricitisation sans dusty hematite; (5) susceptibilité magnétique inférieure à  $0,4 \cdot 10^{-3}$  SI correspondant au remplacement total de la roche par la silice, il n'y a plus de fer sauf dans les sulfures.

Susceptibilité magnétique ( $10^{-3}$ SI)	100	60	30	$<1$ et $>0,4$	$<0,4$
Degré d'altération	1	2	3	4	5
Évolution de l'altération	frais (INSY)	martitisation (HMSY)	hématisation (HMSY)	carbonatation & sericitisation (CB-SRSY)	remplacement par silice (SI-RESY)
<i>intensité d'altération</i>					

Figure 3. Méthode qualitative pour caractériser l'altération à Beattie

Cette méthode de caractérisation de l'altération, basée sur des paramètres géophysiques, peut être un outil efficace pour les roches de la syénite de Beattie. L'utilisation du logiciel NORMAT (Piché and Jébrak, 2004), dont les fondements sont géochimiques, peut être utilisé de façon complémentaire à la méthode géophysique. La rareté de roches à quartz normatif dans le secteur de Beattie a notamment permis à Legault et al. (2005), avec le logiciel NORMAT, de déceler les zones fortement altérées par silicification. Ce sont les mêmes zones identifiées dans ce rapport comme ayant un degré d'altération très avancé, numéro cinq.

L'étude de la paragénèse métallique du gîte de Beattie montre que l'or, et la pyrite, ne sont pas associés au faciès de martitisation mais sont tardifs par rapport à la martite. Ici, la présence d'or associée au faciès d'hématisation (=carbonatation), et la faible quantité de pyrite, témoignent que l'altération à hématite est tardive par rapport à la martitisation.

### Conclusions

Dans le gisement de Beattie, la minéralisation aurifère n'a aucune affinité avec la magnétite; en revanche, l'or se trouve parfois en faibles concentrations (1g/t) dans les roches hématisées associé à un degré de carbonatation. La magnétite est un métallotecte négatif pour l'or à Beattie. La susceptibilité magnétique dans les roches du gisement de Beattie est inversement corrélée (décroissance logarithmique) avec la teneur en or et l'intensité d'altération des roches. Une susceptibilité magnétique d'environ  $100 \cdot 10^{-3}$  SI correspond à une roche non altérée avec une teneur en or d'environ 0,02g/t Au. Une susceptibilité magnétique d'environ  $30 \cdot 10^{-3}$  SI correspond à une altération « partielle » (dusty hématite) avec une teneur en or d'environ 1g/t Au. Une susceptibilité magnétique inférieure à  $0,4 \cdot 10^{-3}$  SI correspond à une altération très intense (remplacement) avec une teneur en or supérieure à 6,2g/t Au. Ces valeurs de susceptibilité magnétiques peuvent servir d'indicateurs pour le caractère aurifère des roches à Beattie.

Une nouvelle méthode de caractérisation de l'altération des roches par les mesures de susceptibilité magnétique peut être développée pour le gisement de Beattie. Il s'agit d'attribuer un numéro à la roche selon les valeurs de susceptibilité magnétique. Plus les valeurs sont élevées (proche de  $100 \cdot 10^{-3}$  SI), plus les numéros tendent vers 1 qui caractérise l'état de fraîcheur des roches; plus les valeurs sont faibles (inférieur à  $0,4 \cdot 10^{-3}$  SI), plus les numéros tendent vers 5 qui caractérise l'état d'altération très intense des roches, marqué par la silicification pervasive et le remplacement total.

## APPENDICE E

### ANALYSE STRUCTURALE DES PHASES TARDIVES CASSANTES

## **Introduction**

Lorsque la mine Beattie était en activité de 1936 à 1954, l'exploitation était exclusivement concentrée au niveau de la faille Beattie marquant le contact entre la syénite au sud et les roches volcaniques mafiques au nord. Des forages effectués en 2009 et 2010 par les compagnies minières Osisko et Clifton Star ont mis en évidence que les minéralisations aurifères à fortes teneurs se distribuent le long de failles, soulignant un contrôle structural important. L'étroite relation entre la minéralisation aurifère et les failles nécessite une étude structurale approfondie afin d'établir une meilleure compréhension de l'évolution structurale à Beattie.

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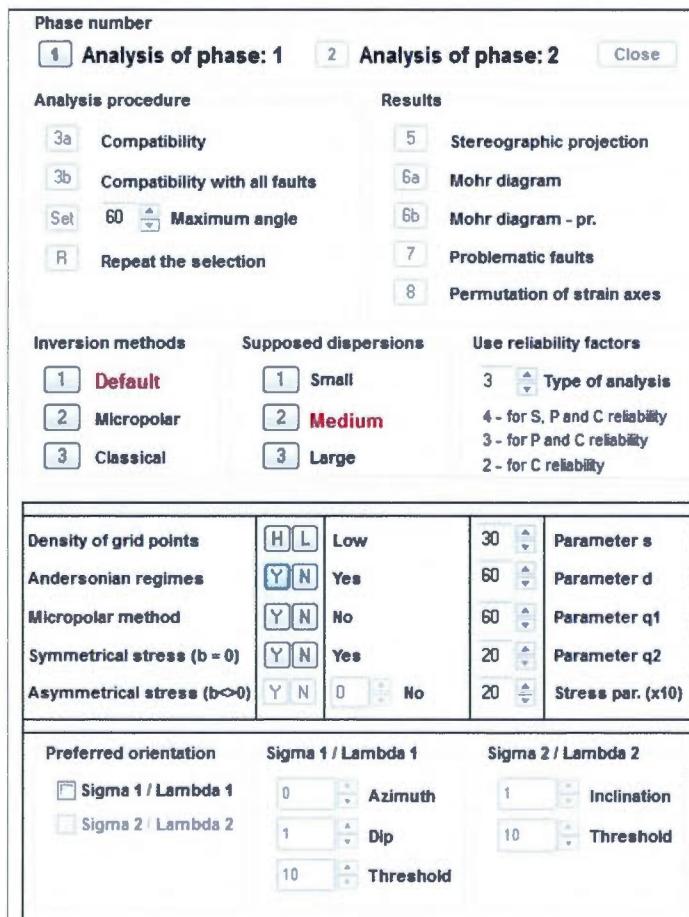
## **Méthode**

### *Échantillonnage*

Des observations structurales basées sur les relations de recouplement et la géométrie des objets ont été faites sur des affleurements à proximité de la Fracture zone dans la syénite de Beattie. Des attitudes de plans de failles et de stries ont été collectées sur ces mêmes affleurements. La prise de mesure s'est faite rigoureusement; (1) déterminer l'indice de certitude de la mesure – C-certain, P-probable, S-supposé, (2) déterminer le type de mouvement sur la faille – N-normal, I-inverse, D-dextre, S-senestre, (3) déterminer l'azimuth de la faille, (4) déterminer le pendage de la faille, (5) déterminer le plongement de la strie le long du plan de faille, (6) déterminer l'orientation de la strie. Des mesures de veines et de joints ont aussi été faites.

### *Intégration des données*

L'analyse des données s'est faites avec le logiciel d'analyse de failles T-TECTO 3.0® basé sur l'inversion de données et permettant la reconstitution du tenseur des contraintes. Lorsque les données sont intégrées au logiciel (Table 1, Table 2), il faut fixer des paramètres (Figure 1) propres à chaque cas. Les paramètres qui influencent le plus le modèle sont le type d'inversion et le paramètre de dispersion. Les paramètres secondaires sont le facteur de certitude des mesures, l'angle de friction, le type de régime, et le paramètre de contrainte entre autres.



**Figure 1.** Paramètres à fixer pour effectuer une reconstitution du tenseur des contraintes - logiciel T-TECTO

#### Paramètres fixés

Les paramètres ont été fixés selon les observations de terrain. Le type d'inversion utilisé est la méthode conventionnelle basée sur la symétrie des contraintes. Le paramètre de dispersion est moyen car il y a des orientations dominantes dans les plans de failles et non une distribution totalement hétérogène qui aurait indiqué une dispersion large. Le facteur de certitude des mesures est fixé à 3, ce qui signifie que seuls « certain » et « probable » sont conservés dans l'analyse. Le paramètre de dispersion est fixé à 30, valeur arbitraire pour les dispersions moyennes. L'angle de friction est ici fixé par défaut à une valeur de 60 degré. Étant donné qu'il y a plus d'une strie sur un même plan de faille, il faut considérer la

réactivation de ces failles, ce qui dans ce cas exclut le régime Andersonian qui implique uniquement les failles néoformées.

**Table 1.** Mesures des failles et stries associés

C I 134 66 55 N	1.00E+0000	1.00E+0000	1
C I 178 89 54 E	1.00E+0000	1.00E+0000	2
C I 197 88 49 E	1.00E+0000	1.00E+0000	3
C D 335 88 35 E	1.00E+0000	1.00E+0000	4
S D 120 72 35 N	1.00E+0000	1.00E+0000	5
S I 134 70 60 N	1.00E+0000	1.00E+0000	6
C D 336 86 46 E	1.00E+0000	1.00E+0000	7
C D 258 73 46 E	1.00E+0000	1.00E+0000	8
C D 356 81 38 E	1.00E+0000	1.00E+0000	9
S I 266 81 51 W	1.00E+0000	1.00E+0000	10
P D 360 69 35 E	1.00E+0000	1.00E+0000	11
C D 190 67 38 E	1.00E+0000	1.00E+0000	12
C D 187 25 12 W	1.00E+0000	1.00E+0000	13
C D 30 69 25 E	1.00E+0000	1.00E+0000	14
C D 195 40 54 W	1.00E+0000	1.00E+0000	15
C D 200 49 52 E	1.00E+0000	1.00E+0000	16
C D 205 60 57 E	1.00E+0000	1.00E+0000	17
C D 200 59 75 E	1.00E+0000	1.00E+0000	18
C D 167 62 85 E	1.00E+0000	1.00E+0000	19
C D 174 60 84 E	1.00E+0000	1.00E+0000	20
P D 165 66 65 E	1.00E+0000	1.00E+0000	21
S S 335 88 65 W	1.00E+0000	1.00E+0000	22
S S 150 79 45 E	1.00E+0000	1.00E+0000	23
S S 260 56 15 S	1.00E+0000	1.00E+0000	24
P S 208 65 77 W	1.00E+0000	1.00E+0000	25
S S 270 66 30 S	1.00E+0000	1.00E+0000	26
S S 175 60 63 W	1.00E+0000	1.00E+0000	27
C S 180 78 56 W	1.00E+0000	1.00E+0000	28
C S 270 85 16 N	1.00E+0000	1.00E+0000	29
C S 280 80 17 N	1.00E+0000	1.00E+0000	30
C S 262 82 18 N	1.00E+0000	1.00E+0000	31
P I 83 82 82 N	1.00E+0000	1.00E+0000	32

P S 108 51 4 N 1.00E+0000 1.00E+0000 33  
 C D 296 89 19 N 1.00E+0000 1.00E+0000 34  
 C D 278 73 46 N 1.00E+0000 1.00E+0000 35  
 P S 172 81 38 E 1.00E+0000 1.00E+0000 36  
 C D 280 86 36 N 1.00E+0000 1.00E+0000 37  
 C D 290 66 14 N 1.00E+0000 1.00E+0000 38  
 C S 165 70 65 E 1.00E+0000 1.00E+0000 39  
 P D 299 85 26 N 1.00E+0000 1.00E+0000 40  
 P S 165 89 33 E 1.00E+0000 1.00E+0000 41  
 P D 290 84 20 N 1.00E+0000 1.00E+0000 42  
 C D 286 76 23 N 1.00E+0000 1.00E+0000 43  
 P S 160 79 45 E 1.00E+0000 1.00E+0000 44  
 P S 163 60 55 E 1.00E+0000 1.00E+0000 45  
 P S 170 66 30 E 1.00E+0000 1.00E+0000 46  
 P S 154 75 18 E 1.00E+0000 1.00E+0000 47  
 P D 358 45 35 E 1.00E+0000 1.00E+0000 48  
 P D 340 55 35 E 1.00E+0000 1.00E+0000 49

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**Table 2.** Mesures de veines et joints

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T 218 60  
 T 271 62  
 T 254 69  
 T 215 70  
 T 240 80  
 T 206 70  
 T 265 65  
 T 265 44  
 T 260 86  
 T 245 65  
 T 260 35  
 T 270 50  
 T 283 64  
 T 260 70  
 T 295 75  
 J 191 69  
 J 176 80  
 J 024 78

J 015 76

J 030 70

J 186 70

J 002 75

J 005 75

J 155 81

J 360 76

J 020 88

J 006 77

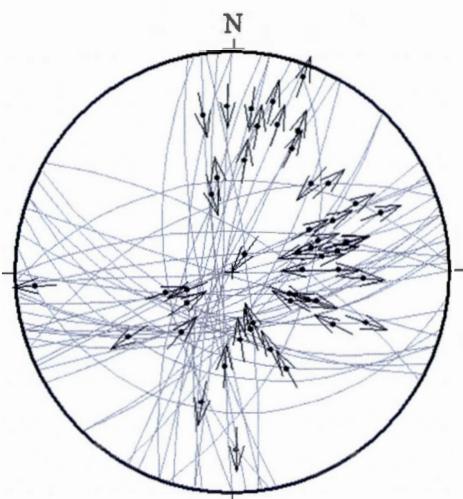
J 170 66

J 184 66

J 172 75

J 012 83

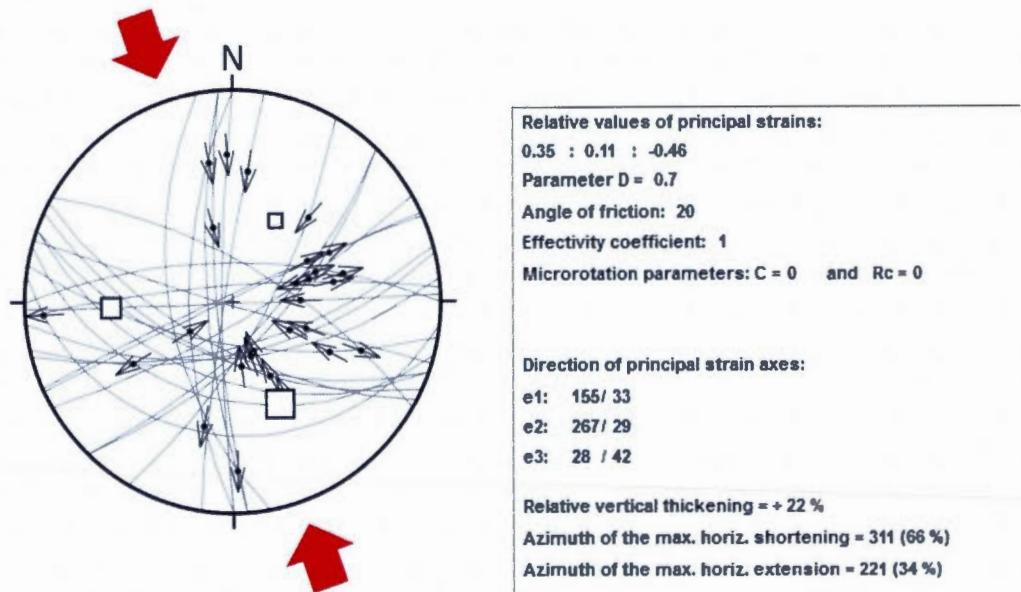
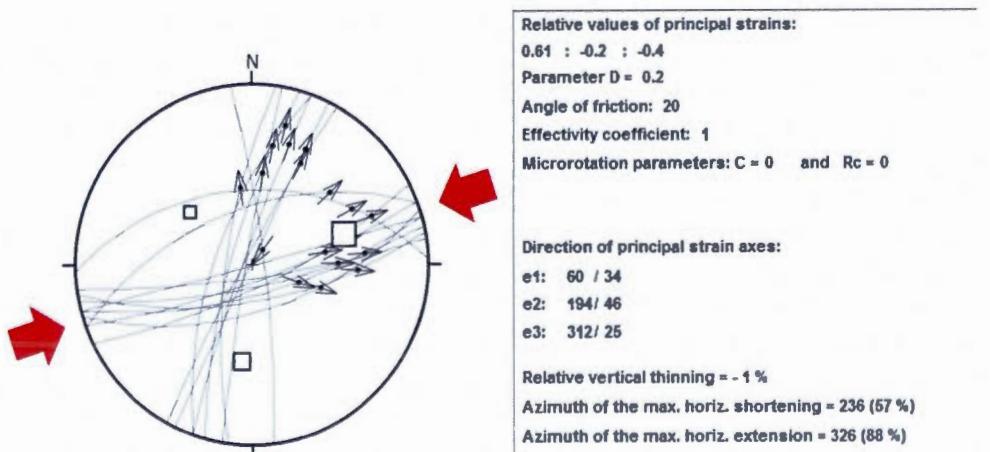
#### Traitement des données



**Figure 2.** Stéréonet de toutes les failles mineures tardives avec les stries

#### *Reconstitution du tenseur des contraintes*

Le logiciel T-TECTO distingue deux phases structurales : (1) un axe de raccourcissement maximal NNO-SSE avec la contrainte maximale  $\sigma_1$  plongeant de 33 degrés vers le N155 (Figure 3a,b), (2) un axe de raccourcissement maximal ENE-OSO avec la contrainte maximale  $\sigma_1$  plongeant de 34 degré vers le N060 (Figure 4a,b).

**Figure 3.** Phase 1**Figure 4.** Phase 2

## APPENDICE F

### CALCULS DES VOLUMES DE FLUIDE POUR LES ALTÉRATIONS

Résultats des volumes de fluides des altérations avec les paramètres :

Objet	Volume (m <sup>3</sup> )	densité (rho)	Masse de Roche altérée (t)	Gain en éléments (%)	Solubilité des éléments (ppm)	volume minimal de fluide (m <sup>3</sup> )	Ratio Eau/Roche
Syénite	247 980 000	2.7	669 546 000				
Hématisation	62 399 000	2.6	162 237 400	Oxydation in-situ			
Carbonatation	151 470 000	2.66	402 910 200	3.83	10	154 314 606 600.0	1 018.78
Silification	4 068 600	2.69	10 944 534	17.74	500	388 312 066.3	95.44
Sericitisation	9 924 300	2.79	27 688 797	4.29	379	313 416 725.9	31.58

Solubilité des minéraux à 300°C : Quartz et Calcite d'après Dolejs et Manning (2010); Séricite d'après Mexias et al. (2005). Le volume de carbonatation est global, il comprend les 2 phases d'altération : (1) à calcite, (2) à carbonate de fer.

## APPENDICE G

### ANALYSE DE L'OR INVISIBLE AU MICROSCOPE ÉLÉCTRONIQUE À TRANSMISSION

## **Introduction**

La cristallisation de l'or essentiellement sous forme invisible est un problème de majeure pour une étude minéralogique. En effet, l'observation se fait généralement au microscope optique et microscope à balayage électronique (MEB), mais très rarement avec d'autres outils. Comptenant qu'il y a de l'or dans les pyrites arsénifères et arsénopyrite, mais que sa dimension est nécessairement inférieur au micron (limite du MEB), il a fallu trouver une méthode d'analyse alternative. Seul le microscope électronique à transmission (TEM) permet une étude très fine grâce à des grossissements jusqu'à 470000 fois, ce qui donne une échelle d'environ 2 nanomètres.

## **Méthode**

Les analyses ont été réalisées au département de physique de l'université de McGill, au sein du FEMR (Facility for Electron Microscopy Research). Les manipulations, incluant préparation d'échantillons et utilisation du TEM, ont été faites par Dirk Schuman, étudiant en post-doc avec le professeur Hojatolla Vali.

### *Préparation*

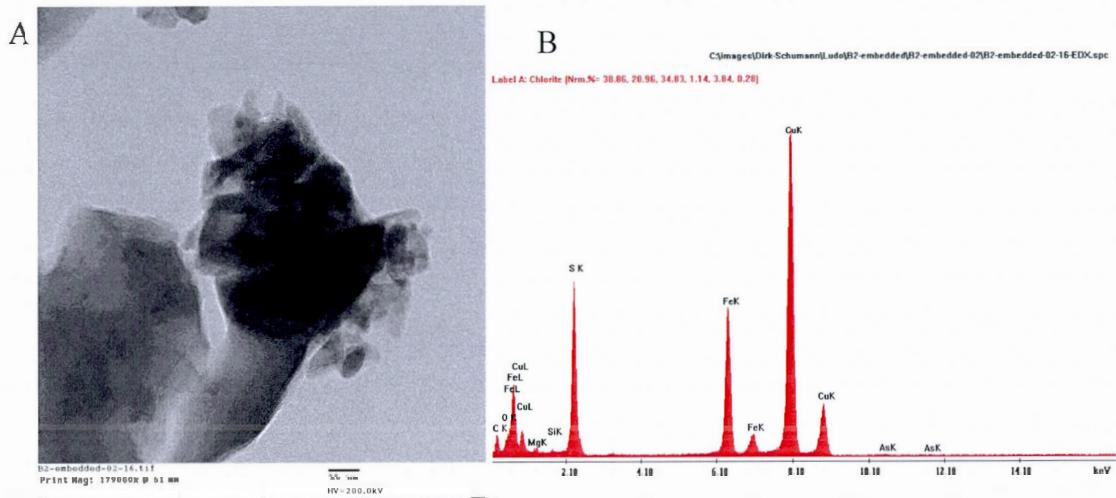
Le support initial est une lame mince polie régulière (échantillon B2 – faciès à contrôle lithologique à 2,4g/t Au). Cette lame est trempée dans un acide durant 24 heures afin de retirer l'époxy. Par la suite, nous avons découpé avec un scalpel des surfaces d'environ 3mm de diamètre contenant les pyrites à analyser. Une fois les échantillons sélectionnés, il faut réduire l'épaisseur de la section polie à 80 nm. Pour cela, nous avons utilisé le procédé « Ion-milling » qui consiste à envoyer un faisceau sur une petite partie de l'échantillon afin d'en réduire son épaisseur. Malheureusement, compte tenu de la forte concentration en carbonate dans la roche, cette méthode fut un échec car le faisceau détruisait bien plus rapidement les carbonates (la matrice), alors que les pyrites étaient moins affectées. Il en résultait un déséquilibre entre l'épaisseur des pyrites, supérieur au 80 nm requis, et les carbonates quasiment totalement détruit. Il a fallu trouver une autre méthode de préparation. Cette méthode consiste à placer les échantillons découpés et sélectionnés dans une résine. Puis les techniciens du FEMR ont fait de très fines sections d'épaisseur de 80 nm requis pour

placer l'échantillon dans l'appareil. Cette méthode s'est révélée bien meilleur dans la préservation à la fois de la matrice carbonatée et des pyrites. Une fois les sections découpées, elles sont placées sur une grille (« grid ») en cuivre d'environ 1mm de diamètre. Ce support en cuivre avec l'échantillon est ensuite inséré dans le TEM.

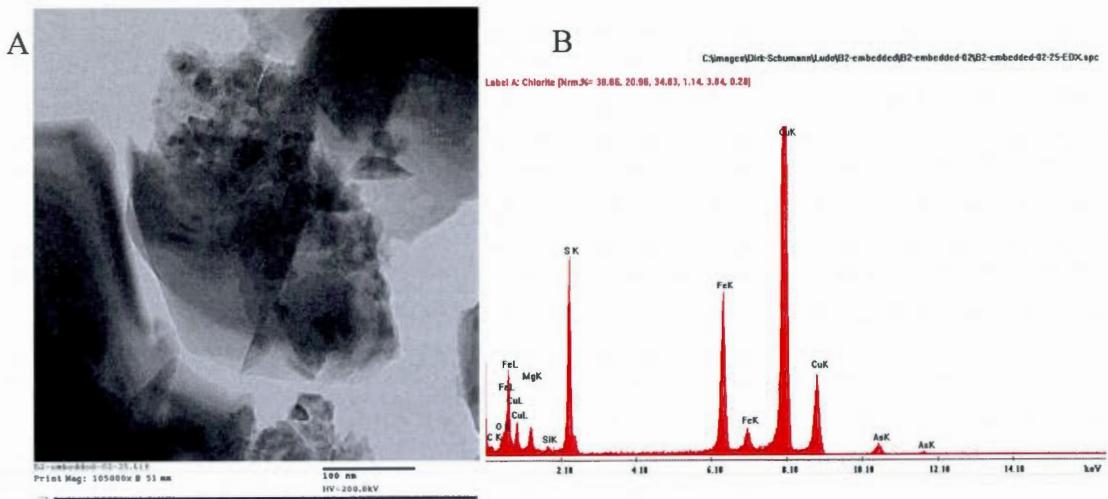
#### *Procédé analytique*

Le TEM utilisé est un Philips CM200. Le voltage appliqué est de 200 kV. Les autres propriétés de l'appareil sont les suivantes : AMT XR-80C 2k x 2k CCD, avec Camera System et EDAX Genesis EDS. Les analyses se faisait généralement à des grossissements entre 300000 fois et jusqu'à 470000 fois.

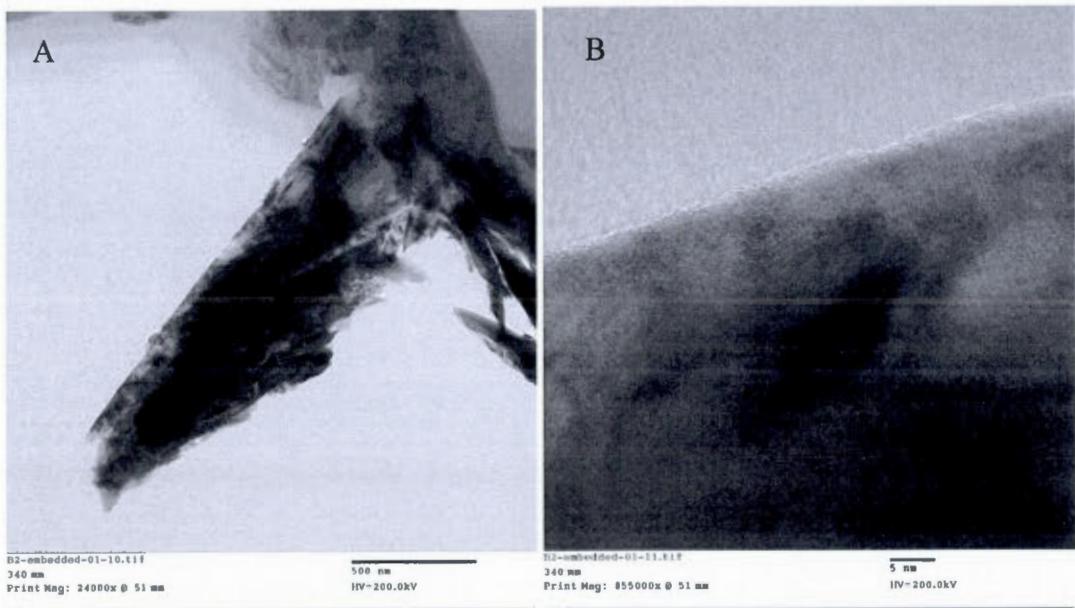
#### Résultats

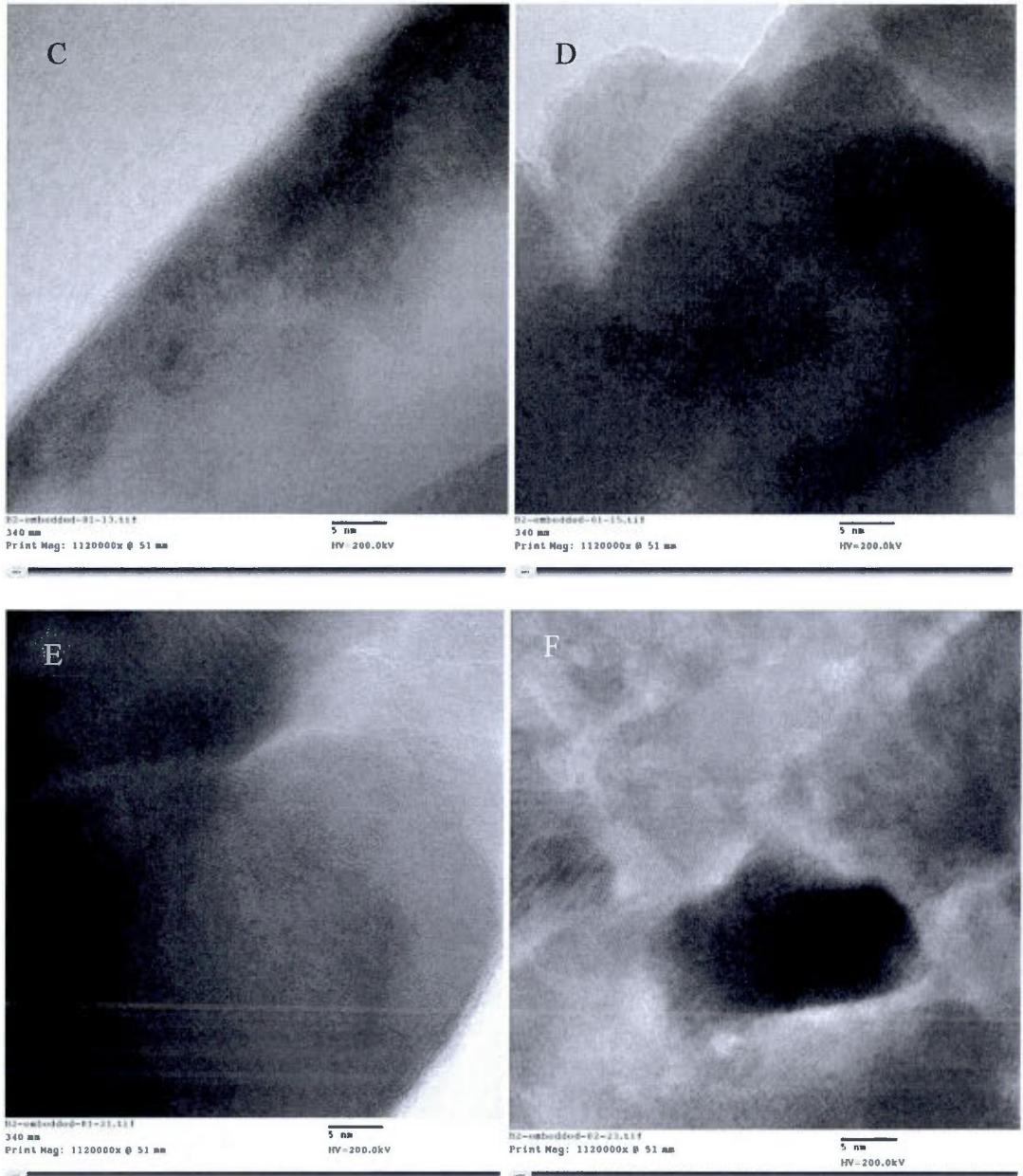


**Figure 1.** (a) Image d'une pyrite non arsénifère. (b) Analyse élémentaire avec EDAX; à noter que les anomalies en Cu sont dus à la grille de cuivre qui perturbe en partie le signal.



**Figure 2.** (a) Pyrite arsénifère, aurifère, et poreuse. (b) Analyse élémentaire de la pyrite. Noter les pics en arsenic, et l'absence des pics en or. Cela s'explique par le degré de détection de l'EDAX de l'ordre du %. En revanche les analyses à la microsonde, avec un seuil de détection de 60ppm Au, avaient confirmées la présence d'or dans cette pyrite.





**Figure 3.** (a) pyrite. (b), (c), (d), et (e) fort grossissement sur (a); visualisation des structures cristallines de la pyrite; les différentes teintes de gris indiquent des changements de composition. (f) Structure cristallines d'une autre pyrite; hétérogénéité des structures bien visibles.

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