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NEW OROGENIC TYPE GOLD OCCURRENCES IN THE UYANGA ORE KNOT (CENTRAL MONGOLIA)

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ARTICLE INFO ABSTRACT Article history: The Uyanga ore knot district of the Khangay metallogenic zone are hosted by the lower-middle Devonian volcanogenic-sedimentary Erdenetsogt formation. About 40

lower-middle Devonian volcanogenic-sedimentary Erdenetsogt formation. About 40 samples were collected from the host rocks, veins and quartz veins in the Uyanga ore knot district in 2016. The new Burgetei, Ult and Senjit gold occurrences were studied. The quartz-sulfide, gold-arsenic and gold-antimony-mercury mineralization are determined in the berecitization, silicification, limonitization and glauconization altered metasomatic zones within the Uyanga ore knot districts. The rocks of the Erdenetsogt formation have an irregular gold content: 0.96 g/t Au is determined in quartz vein taken from trench of the Burgetei occurrence (BG-7/16), Au content is highest up to 3.5 g/t in the quartzite-jasper (Ult-7/16 and Ult-9/16) cut by quartz veins in the Ult occurrence. The Senjit occurrence represents Au-Hg-Sb epizonal level of orogenic gold deposits structure with highest Hg content up to 851 ppm. This year no sulfide minerals were found in the siltstone of this occurrence. The Au content of arsenical pyrite of the Burgetei and Ult is below the detection limit by electron microprobe analysis. The Au content of arsenopyrite of the Ult occurrence is highest (up to 238 ppm). Therefore, the ore-mineral assemblages in the gold occurrences reflect the differences between the three explored sites, formed in the course of fluid evolution during the fluid-rock interaction. Variable concentrations of indicative elements (As, Te, Sb, Hg) and their ratios confirm this fact. The geodynamic position, the type of the hydrothermal alteration of both igneous and sedimentary rocks, textures and mineral assemblages, the mineralization sequences are consistent with orogenic classification for the Burgetei, Ult and Senjit gold occurrences.

Keywords: pyrite, arsenopyrite, oxidation, fluid-rock interaction

INTRODUCTION

The classical theory implies that orogenic Au deposits form during the late stage of orogeny and are typically hosted by rocks of greenschist to amphibolite metamorphic grade that may be spatially related to granitoid intrusions (Groves et al., 1998; Goldfarb et al., 2001). Consensus on even the most fundamental features of an orogenic gold system model, however, remains elusive (Wyman et al., 2016). According to

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(Goldfarb et al., 2001) orogenic gold-forming events continued to be extremely widespread during Paleozoic time along the extensive margins of both Gondwana and the Paleo-Tethys Ocean basin. Goldfarb, Groves (2015) have rewired the current models for orogenic gold formation: "Using the principle of Occam's razor that the hypothesis with the least number of assumptions is the best, the simplest, most elegant, universal model is that all orogenic gold deposits formed throughout time from devolatilization of oceanic crust and associated carbonaceous, sulfidic sediments with goldenriched pyrite late in the history of subduction when local plate motion was perturbed". Orogenic gold deposits can form over a variety of depths, from as shallow as 3 km to as deep as 20 km, typically during late orogenic shifts from compressional transpressional to or transtensional regimes. Arsenopyrite and pyrite represent the most abundant sulphide minerals that occur in association with the Au mineralization, which formed from low salinity, CO₂-bearing, reduced and near-neutral pH fluids that show a wide range in fluid temperatures from about 200 to 500°C (Goldfarb et al., 2005; Keith et al., 2018). A comparison of the above data of leading scientists of the world and obtained by us in recent years on new ore Mongolian gold occurrences made it possible to make an assumption about the genetic belonging of the latter.

The vast territory of Mongolia occupies a large part of the Central Asian Orogenic Belt (CAOB). The Khangay-Khentey belt is located in central Mongolia and records a progressive accretionary orogeny of the CAOB from the Middle Cambrian to the Early Mesozoic (Erdenesaikhan et al., 2013). The geology and petrology namely of the Khangay region were reported during 1:50 000 scale geological mapping (Lhundev et al., 1994). The primary ore occurrences of the Uyanga gold ore knot district of the Khangay gold zone are hosted in volcanogenic-sedimentary rocks of the lowermiddle Devonian Erdenetsogt formation (Fig. 1). The volcanogenic rocks are metamorphosed condition greenschist facies under The geochemical characteristics of Middle Paleozoic greenstones are of particular interest for understanding the geodynamic evolution of the Khangay belt. Orolmaa and other (2008) investigated the Permian-Triassic granitoid magmatism and metallogeny of the Khangay upland (Fedorova,1975). They have indicated that gold mineralization in the area that is confined within the Khangay gold belt appears to be independent of the granitoid intrusions.

Among other gold ore occurrences of unknown of genesis. а group gold-quartz vein mineralization such as Burgetei, Ult, Senjit, Huren Khoshuu were identified in the eastern tributary of the Ongi Gol river, along the Ovor Ult, Booroljuut rivers valley within the Ult-Booroljuut ore placer deposits of the Uyanga ore knot district in the Khangay gold metallogenic zone (Dejidmaa, 1996, 2012; Lhundev et al., 1994). The Uyanga group of primary gold occurrences are associated with the gold quartzsulfide mineralization in the hydrothermal metasomatisms of shear zones (Altanzul and Baasandorj, 2014, Khishgee and Akasaka, 2015).

In this paper, new information concerning the Burgetei, Ult, and Senjit gold occurrences of the Uyanga gold ore knot is presented. In terms of the geological conditions, mineral assemblages, chemical composition of ore minerals, ratios of indicator elements and isotopic investigation an assumption is made to clarify the genetic processes of the ore occurrences formation, because they are perhaps the typical orogenic gold occurrences. It is appropriate to refer to the opinion of Mongolian scientist: "All gold deposits of Mongolia belong to granite-related deposits, i.e. epithermal and orogenic gold deposits. Mineralization age of gold deposits depends on geodynamic condition of each specified area, however most significant genetic type among them is orogenic gold deposits (Delgertsogt, 2017).

GEOLOGY OF THE STUDY AREA

The study area is located in the southeast of the Tsetserleg turbidite accretion terrain in the Khangay-Khentey orogenic structure of the Mongol-Okhotsk fold belt (Fig. 1) (Tomurtogoo, 2002; 2008). Tsetserleg accretion terrain is composed of the lower-middle Devonian chert-basalt-sandstone Erdenetsogt, the middle-upper

Devonian sandstone-siltstone Tsetserleg and the carboniferous polymictic sandstonelower siltstone Dzargalant formations (Amantov et al., 1970; Tomorchodor et al., 1990; Lhundev et al., 1994; Tomurtogoo 2002; Badarch et al., 2002). These Paleozoic rocks are separated from each other mainly by thrust faults (Fig. 1c) of the Ult-Booroljuut gold ore-placer mineralization of the Uyanga ore knot district (Dejidmaa, 2012) within the Tsetserleg terrain in the Khangay-Khentey orogenic system (Borisenko et al., 2006; Orolmaa et al., 2008).

Within this ore knot district, except the number of gold-placer deposits occurred along Ult-Booroljuut river valley, the primary occurrences of the Burgetei, Ult, Senjit gold-quartz veins are the lower-middle Devonian hosted in Erdenetsogt formation with various alteration zones (Togtokh et al., 1986; Lhundev et al., 1994). The chert-basalt layer of the lowermiddle Devonian Erdenetsogt formation hosting the above-mentioned gold primary occurrences conformably beds inside the terrigenic formation. The primary gold occurrence was subsequently metasomatized with alteration zones and with variable thickness in different gold occurrence and usually anticline folded with terrigenic layer, cut together bv gabbrodiorite dykes (Lhundev et al., 1994).

A number of hydrothermal alterations (quartz



Figure 1. Geology and natural resource map of Ult area of the Uyanga district (c) (modified from Lhundev et al., 1994) Location of study area: Central Asia orogenic belt (a) Tsetserleg accretion terrain of Khangai-Khentei Orogenic Belt (b) (Tomurtogoo, 2012)

veins, quartzite, mineralization belt of faults hvdrothermal altered rocks) zone. were identified along the fault in the area of gold primary occurrences. The ore mineralization is accumulated mainly in quartz veins and zone of tectonic breccia. In the hydrothermal berezite altered rocks (quartz, carbonate, sericite, pyrite) those zones, arsenopyrite, sometimes of chalcopyrite, the inset and vein of galenite and sulfates have been identified in the goldmercury hosting fine grain sugar like quartzite in the jasper-volcanogenic-terrigenic layer of the Erdenetsogt formation (Lhundev et al., 1994). Ore bodies in gold occurrences are consists of gold-sulfide-quartz gold-quartz, ore Therefore. associations. gold primary occurrences of ore knot district have a complex structure and hosted in the lower-middle Devonian volcanogenic-terrigenous Erdenetsogt formation.

SAMPLING AND ANALYTICAL METHODOLOGY

This work used the field data sampled from the exploration trenches in the Burgetei. Ult and Senjit gold occurrences of the Uyanga ore knot district in 2016. About 40 rock samples were collected from the host rocks, veins and quartz veins. About 20 samples were analyzed for petrography, mineralogy and geochemical analysis. Minerals were separated into heavy and light fractions for mineralogical analysis. Microprobe analysis was carried out by electron scan microscope with Energy Dispersive X-ray Spectroscopy method (SEM-EDX) at the laboratory of substance analysis of the School of Geology and Mining Engineering, Mongolian University of Science and Technology. Binocular was used for the separation of pyrite and arsenopyrite monominerals from the other sulfide minerals. In addition, samples were analyzed by electron scan microscope LEO 01430VP, by electron probe microanalyzer Camebax-Micro, JEOL JXA-8100 and atomabsorption methods in the V.S. Sobolev Institute of Geology and Mineralogy, SB RAS. Gold was determined according to the custom prepared method by electron-probe microanalyzer JEOL JXA-8100 at 20kV and 200 nA with the extended diameter of up to 10-15 mm electron beam. Gold analysis were carried out by atomic absorption spectrometer (AAS) with 55 AA Agilent instrument in the SGS-IMME Co Ltd., Mongolia, rare element analysis was carried out by ICP OES method with Optima 7300DV Perkin Elmer-pair inductive plasma spectrometer. Sulfur isotope analyses were carried out at the Multi-elemental and Isotope V.S. Sobolev analysis CCA. Institute of Mineralogy, Geology and SB RAS (Novosibirsk). The procedures overview is given in (Ariunbileg et al., 2016; Pal'yanova et al., 2016).

RESULTS

Occurrences type and ore mineralization

(46° Gold occurrence Burgetei 34'30.00"N /102°15'5.00"E) is located in the valley of Burgetei and Ovor Burgetei slope junction, at the top of 2300 m watershed hill on the center of the NE-NW arched anticlinal fold of the volcanogenic-terrigenic Erdenetsogt formation. Siliceous rock layers of this formation with varying thick host basalt and its alteration zone is cross cut by NW stretched 0.5-0.8 m thick microdiorite dykes, quartz veins and Sericitization. berecitization. veinlets. limonitization alteration of the Burgetei occurrence are well observed at 1, 2, 3 trenches which cross approximately in 200 m wide. Host sediments, basalt, diorite dyke and veins near the gold-bearing quartz veins of the Burgetei occurrence are all changed by the hydrothermalmetasomatic alteration (Lhundev et al., 1994). Surface weathering, erosion and oxidation intensity clearly shows metasomatic alteration zone.

The siltstone of volcanogenic-terrigenic Erdenetsogt formation cut veins and structures of metasomatic alteration; ore controlling transition brittle-ductile zone with the deformation is clearly seen in the second trench of the Burgetei occurrence (Fig. 2a). Deposits and occurrences of the orogenic gold are formed in such a type of structures, i.e. orogenic gold deposits are emplaced during collisional events throughout much of the middle to upper crust. Oxidized quartz veins (1-10 cm) and veinlets of the Burgetei well observed in the exploration trenches (sample BG-7/16 contain 0.86 ppm of gold). Except (BG-1/16, BG-7/16), quartz, carbonate and iron oxides, ore minerals such as pyrite, antimonite and hematite are observed in the quartz veins (1-10 cm).

Carbonate altered (magnesite) metasomatic ring is observed within the basalt in outside of alteration zone that cut by quartz veins (Fig. 2b). Ore-bearing grey, cryptocrystallic crystal, altered rock (BG-2/16) is composed of sericite, limonite altered sericite, quartz, iron oxide and pyrite. The main body is composed of very small sericite aggregates that form dendritic like structures (Fig. 3), inside of which the cluster of iron oxide and completely oxidized pyrite relicts are observed. These small aggregate quartz veins with ore mineralization are distributed in a spherulite shapes with varying sizes.



Figure 2. Transform fault of the brittle-ductile deformation of the quartz veins in the Burgetei exploration trench (a). Carbonate veinlets in the volcanites of the Burgetei occurrence trench (b)



Figure 3. The growth of beam shaped sericite aggregate in the Burgetei, Sample BG- 2/16 (a) and sample BG-4/16 (b). P – Pyrite crystals

Mineralogical analysis of this sample identified the intense limonitization of chromite and serpentine. Therefore, it can be accepted by that the primary rock with limonitization and sericitization in the mineralization zone was basalt. Except the berecitization zone of the

exploration trench (BG-3/16), the sericite, quartz, zircon and ore minerals composed from oxidized pyrite and arsenopyrite are observed inside the sheared and altered grey siltstone with the abundant iron oxide and cryptocrystalline in the brecciated, metasomatic zone.

The sericitizated quartz metasomatite with pyrite (BG-4/16, Au content 0.02 ppm) are spread like veins and lens. Pyrite is almost not involved in the oxidation, it is in a good crystalline form, sometimes grouped into larger (1.5-2.3 mm) frame, and rutile granules are observed. Pyrite cube crystal aggregate of larger size are spread along the veins and fractures. Such ore minerals take less than 25-30% of total. These crystals are chosen for the microprobe study.

Dyke of granodiorite porphyry, microdiorite (BG-6/16, BG-8/16) is intensely altered, and composed of plagioclase (60-65%), crystalline amphibole (20-25%) and quartz (10-12%). The main background part has а micro hypidiomorphic structure. They have partially and completely substituted by sericite, chlorite, carbonate, epidote aggregates. This indicates the berezite alteration of the granodiorite body in the Burgetei occurrence. Mineral association of the Burgetei occurrence are classified into high, low and without sulfide types.

Mineralogical analysis revealed the alteration minerals such as sericite, chlorite, muscovite, and Fe-Mn carbonates, except the serpentine in the Burgetei volcanogenic rock samples. The SEM-EDS analysis identified ore minerals such as pyrite, antimonite and gold. The lowermiddle Devonian volcanic rocks of the Burgetei occurrence are intensely berecitizated and even barite and nickel arsenide were formed in some samples except the ilmenite, chromite, siderite, chlorite and sericite. The observed high content of the platinum group element iridium, except the monazite and xenotime in the sample, suggests its plume origin enriched by ocean tholeiitic basalt (Erdenesaikhan et al., 2013). By the influence of the metasomatic alteration, the microdiorite, granodiorite-porphyry dykes host the pyrite, carbonates, chlorite, monazite and xenotime.

Gold occurrence Ult (46°34′55" /102°17′20") was observed in the east of hydrothermalmetasomatic alteration zone that stretches arching in NE-NW where the irregular distributed berecitization, pyritization and limonitization hosted in siliceous siltstone with the lower layer andesite-basalt, basalt layers of the lower-middle Devonian Erdentsogt formation (Lhundev et al., 1994).

Gold mineralization of the Ult occurrence is represented by the argillic altered dark siltstone were observed in fractured iron hat (gossan) formed from the sulfide oxidation on the exploration trench at the 2300 m altitude. It might be interpreted as pipe to the deep. This alteration associated with the mineralization is observed also in the jasper-quartzite-basalt formation. Small size quartz-sulfide veins are observed here. Microdiorite (Ult-1/16) is composed of plagioclase 50-55%, quartz 15-18% and secondary minerals (sericite-carbonatechlorite 25-30%). Sometimes initial color minerals were completely substituted by carbonates, chlorite and iron oxides (Fig. 4). The light grey (enriched by iron oxide), fine



Figure 4. Metasomatized microdiorite sample Ult-1/16, magnification 40x (a). SEM-EDX- electron images, where Mn - and Mnz – monazite (b).

grained sericite and argillic siltstone rock cut by quartz veins (Ult-2/16, Ult-5/16) contains quartz -sericite narrow veinlets. In addition, electron microscope analysis revealed quartz, barite and carbonate veins in it. It is composed of dark micro fauna (radiolaria), aleurolitic mass in argillic rock with pyrite, ore minerals - pyrite and arsenopyrite, hornblende and carbonates. Non-oxidized pyrites of the Ult occurrence are mainly observed in the siltstone and jasper.

In the Ult occurrence, the siltstone with intense metasomatic alteration, with argillic layers, quartzite, predominantly sericite, chlorite, carbonates and pyrite altered by berecitization in jasper, and rarely occur monazite and ilmenite in accessories minerals were formed as revealed by the scan electron microscope (SEM-EDX). Limonitization involved in the surface oxidation and predominantly Cu and Zn oxidation minerals.

Gold occurrence Senjit (46°33′50″/102°18′20″) is represented by the clayey schist, brecciation, and argillic alteration that are intensely developed along the NE directed transform fault inside the fine grain, green grey sandstonesiliceous siltstone branch of the Erdenetsogt formation. Moreover, volcanogenic rocks are altered to the green grey color glauconites (Ca hosting water alumosilicates). All these conditions suggest their formation in a shallow ocean environment. SEM-EDX analysis identified zircon and ilmenite. Scandium in the zircon mineral proves that alkali volcanogenic rock was involved to glauconite under the shallow ocean environment. For this gold ore occurrence, there is less information than the other two. However, it cannot be missed, since it represents a certain Au-Hg-Sb epizonal level (Goldfarb and Groves, 2015).

Geochemical study

Geochemical assay data processing was carried out on the Uyanga gold occurrences (Table 1). It is determined, that at the Burgetei, Ult, Senjit primary occurrences, the berecitization, pyritization, silicification, limonitization and diorite, granodiorite dykes in the siliceousvolcanogenic-terrigenic layers of the Erdenetsogt formation have an irregular gold content. For instance, 0.96 g/t Au is determined in quartz vein taken from trench of the Burgetei occurrence (BG-7/16). Au content is highest up to 3.42 g/t in the quartzite-jasper (Ult-7/16) cut by quartz veins in the Ult occurrence, and 0.05-0.12 g/t Au content is defined at the Senjit occurrence. Another sample Ult 9/16 similar to the Ult-7/16 was analyzed only for gold (not listed in the Table 1). It turned out that it contained 3.5 g/t gold. Thus, the volcanogenicterrigenic layers of the Uyanga group occurrences altered by metamorphism and metasomatic processes contain visible content of Au.

Relatively high content of As, Sb, Hg, S, Sc, Cr, Ni, Co, V, Mn, Mg, Fe that are accumulated in association with the gold mineralization in comparison with the clarke content of the metamorphic and magmatic rocks were defined in the volcanogenic-terrigenic layers of the Erdenetsogt formation (Table 1). The content of the above ore element is different.

For instance, the geochemical analysis revealed the high content of As in the Burgetei BG-2/16) occurrence (532)ppm in simultaneously with Co, Ni and Fe (50 ppm, 152 pm and 10% respectively). Although Bi and Te concentrations are below the detection limit of ICP MS, high content of W was detected in three of seven samples. Such an association of elements are consistent with the deeper-earlier formation of the Burgetei gold occurrence. Because the increased content of antimony was determined in the quartz vein (BG-7/16) and in microdiorite-porphyrite (BG-8/16), the the Burgetei could be attributed to the Au-As±Sb one. The microdiorite of the Burgetei and Ult occurrences are similar in the contents of Mn, Cr, Fe, Ni. At the same time, the highest contents of Mn, As and Au are characteristic of the Ult tectonic structures with intensive metasomatic transformations, namely siltstone berecitization sericitization, with and limonitization (Ult-6/16) and quartzite-jasper (Ult-7/16). Because the increased content of mercury (up to 86 ppb) was determined in these two samples, the type of this ore occurrence could be attributed to the Au-As±Hg one. A distinctive feature of the Senjit siltstones is high Sb and Hg contents, namely 33-39 g/t and 8.5-

Occurrence				Burgetei					U	lt			Ser	ıjit	
Sample	BG-1/16	BG-2/16	BG-3/16	BG-4/16	BG-6/16	BG-7/16	BG-8/16	Ult- 1/16	Ult-2/16	Ult- 3/16	Ult- 5/16	Ult- 6/16	Ult- 7/16	Sn-5/16	Sn-6/16
Host rocks	Quartz viens with limoni- tization	Aleurolite With sericitiza- tion	Gabbro-diorite	Sericitiza-tion With pyrite (sericito-lite)	-ətro-ibonsrĐ ətiryhqroq ariens	гиэіч хтилу	Microdio-rite- Microdio-rite	Micro-diorite	Aleuroli-te -iticitias atriciti zation	Quartz viens	ətilor-uəlA ətiryq thiw	Aleuroli-te with sericitiza- tion-beresiti-	Jasper Jaspe-roid/ Jaspe-roid/	Aleuro-lite with argillite- roitez	Aleurol-ite with argillite- roitsz
Rock forming	elements (w	t. %)													
Al	3.83	6.43	6.2	5.59	5.6	0.48	5.66	6.38	4.72	0.35	4.94	1.05	0.9	1.44	1.03
Ca	7.35	2.7	3.96	2.33	1.52	0.05	2.99	2.77	0.16	0.02	0.49	0.12	0.09	0.1	0.02
Fe	6,67	10,02	5,69	6,29	3,02	1,22	4,47	4,49	2,58	0,72	3.36	>15	>15	1,97	1,52
Mg	1,73	1.37	1,98	0,35	1,13	0,05	3,08	3,32	0,75	0,03	0,98	0,08	0,05	0,07	0,06
Ti	0.05	0.1	0.13	0.14	0.41	0.01	0.06	0.04	0.23	<0.01	0.26	0.03	0.04	0.05	0.05
Κ	1.24	2.09	2.46	2.53	2.23	0.12	1.83	1.99	2.24	0.09	3.03	0.18	0.18	0.35	0.29
Na	0.17	0.31	1.65	2.93	3.67	0.02	2.31	3.4	2.69	0.03	2.79	0.02	<0.01	0.01	0.02
Trace element	s (ppm)														
Mn	2991	2597	1583	1124	565	408	837	785	259	78	1034	4175	3121	69	79
Ba	291	543	807	634	1068	65	1017	982	656	48	1471	219	173	610	440
Be	0.7	1.5	2.1	1.5	2.2	<0.5	1.1	1.4	2.1	<0.5	2.6	1.2	1.1	0.8	0.7
Cd	б	7	∇	7	$\vec{\nabla}$	1	∇	∇	∇	∇	∇	24	27	∇	\sim
Co	41	50	25	24	11	7	21	19	7	1	7	2	∇	$\overline{\nabla}$	1
Cr	342	339	110	152	91	421	426	336	45	204	39	60	70	115	241
Cu	12	61	35	54	10	12	32	4	13	5	34	75	58	66	43
Ni	107	152	84	89	11	15	78	117	11	8	8	28	24	10	14
Pb	6	7	3	9	25	58	11	17	17	9	41	15	19	19	13
La	7	13	30	14	12	2	19	27	28	$\overline{\nabla}$	26	10	15	18	11
Li	28	48	44	11	38	4	54	99	26	18	33	15	9	11	28
Ρ	0.12	0.17	0.23	0.25	0.06	<0.01	0.06	0.07	0.06	<0.01	0.07	0.05	0.07	0.07	0.03
Sc	13.1	14.4	14.2	8.6	5.4	0.8	8.7	7.8	5	<0.5	4.7	5.8	8.1	6.7	3.6
Sr	377	94	394	197	334	11	574	565	195	7	192	58	41	132	54
Λ	97	177	147	106	58	9	142	101	72	4	69	81	95	71	44
M	21	<10	16	<10	<10	<10	13	11	<10	<10	≤ 10	<10	<10	<10	<10
Υ	11	10	16	6	8	2	6	8	11	∇	10	6	14	10	9
Zn	113	137	108	114	91	38	84	86	65	8	116	67	73	8	8
Zr	14	28	35	20	125	Ŷ	85	101	108	3	137	13	17	18	13
Ore-forming e	lements (ppr	(u													
Au	<0.01	<0.01	<0.01	0.02	<0.01	0.96	<0.01	<0.01	<0.01	<0.01	<0.01	0.67	3.42	0.05	0.12
\mathbf{As}	233	532	88	139	24	179	38	73	21	46	24	1865	2062	25	32
\mathbf{Sb}	6	Ŷ	8	10	14	20	19	14	9	8	8	Ŷ	Ŷ	33	39
Hg	0.1	0.1	0.09	0.09	0.32	0.1	0.36	0.51	0.12	0.09	0.08	0.62	0.86	8.51	3.13
S%	< 0.01	<0.01	<0.01	0.13	<0.01	0.01	0.01	0.01	<0.01	0.01	0.04	0.07	0.06	0.02	0.01
Sn is ev	erywhere l	ess than 10) ppm; Bi	is everywł	tere less th	an 10 ppr	n; Ag and	Mo are e	everywhei	e less the	in 2 ppm				
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Table 1. Trace elements content of the Uyanga gold mineralization at the Burgetei, Ult, and

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Figure 5. (a) Concentrations of Sb, Hg vs. As and (b) Mg, Mn vs. Fe in rocks of the Uyanga orogenic gold occurrences

3.1 g/t respectively. The Au-Hg-Sb association in the Senjit occurrences is taking place and present the epizonal part of the pluton (the shallowest zone). In the Senjit ore occurrence, a band of argillized rocks with gold-bearing quartz veins containing jasper are characteristic, but sulfide minerals were not found during in situ sampling. These results suggest, that ore element contents vary for each gold primary occurrences of the Uyanga gold mineralization knot depending on the type of metasomatites, of deformation amplitude tectonic and weathering grade.

As orogenic gold deposits are emplaced during collisional events throughout much of the middle to upper crust, the correlation of important elements is interesting. Fig. 5 shows the correlation of As and Sb (circles), Hg (squares) in the left part (A) and correlation of Fe and Mg (circles) or Mn (squares) in different rocks of the Uyanga ore knot.

There is no clear correlation between arsenic and mercury and antimony, rather one can see their almost constant contents with the increase for arsenic. In this case, the Ult occurrence is clearly distinguished by the highest arsenic in the rocks (and the highest gold content). Iron and Mg, Mn correlate positively at low iron concentrations, at the highest Fe, the correlation is weak. The Senjit occurrence with highest Sb and Hg versus low As contents just like low Mg and Mn versus low Fe content belongs to the top of the structure zone (black circles and squares, Fig. 5).

Pyrite and arsenopyrite analysis

Pyrite and arsenopyrite dominate in sulfide

minerals composition of the Uyanga gold primary occurrence. Pyrite is stable under various physicochemical conditions and its refractory behavior post-depositional to metamorphism and its near-ubiquity makes it suitable for micro-analytical studies to reconstruct ore-forming processes through space and time (Keith et al., 2018). Assemblages containing arsenopyrite and arsenical pyrite are common in orogenic gold deposits formed at greenschist facies conditions. Gold is observed in both minerals but is more abundant in arsenopyrite. The fact that measured values in arsenopyrite and pyrite differ by two orders of magnitude is concordant with empirical evidence for incorporation of greater amounts of Au into arsenopyrite relative to pyrite (Cook et al., 2013). Chemical composition of pyrites of the Burgetei and Ult ore occurrences are shown in Table 2.

As shown in Table 2, the S content in pyrites varies from 51.4 to 53.0 wt.%, with a mean of 52.67 wt.% (Burgetei) and from 51.19 to 52.85 wt.%, with a mean of 52.40 wt % (Ult). The S contents of most of the samples are lower than the theoretical values (53.45 wt%). Hence, As-S substitution apparently occurs in pyrite. From all the analyzed trace-elements the highest As impurity (6.5 wt.%) was observed in the tabular pyrites of the Ult occurrence (as in its wall rocks). The high As content there is in the Burgetei occurrence (4.4 wt.%) simultaneously with highest Ni (445 ppm) and Sb (81 ppm) contents. The concentration of gold in the studied pyrites is below the detection limit, specified in Table 3 (78 ppm). Arsenical pyrite is of great environmental interest because its

Element/		Fe	S	Ni	Cu	As nom	Se	Te	Bi	Sb	Au
Occurrence		wt.%	wt.%	ppm	ppm	As ppm	ppm	ppm	ppm	ppm	ppm
	Min	43.76	51.4	n.d.	n.d.	n.d.	n.d.	45	n.d.	n.d.	n.a.
Burgetei n=60	Max	46.91	53.01	5735	297	12900	348	589	1171	357	n.a.
	Mean	46.39	52.67	445.18	52.85	4411.5	81.62	325.1	166.35	81.61	n.a.
	Min	45.61	51.19	n.d.	n.d.	1513	n.d.	59	n.d.	n.d.	n.a.
Ult n=30	Max	46.69	52.85	1207	292	23400	409	597	1212	130	n.a.
	Mean	46.43	52.40	114.7	96.2	6553.13	78.43	343	312.03	6.27	n.a.

Table 2. Pyrite chemical composition of the Burgetei and Ult

Table 3. Arsenopyrite chemical composition of the Ult occurrence

Element / Occurrence		Fe, wt.%	S wt,%	As, wt.%	Ni, ppm	Cu, ppm	Se, ppm	Te, ppm	Bi, ppm	Sb, ppm	Au, ppm	Ag ppm
	Min	33.65	18.98	45.41	n.d.	n.d.	n.d.	264	268	n.d.	n.d.	n.d.
Ult (n=13)	Max	34.13	19.86	47.1	3043	307	513	1124	3146	293	238	10
	Mean	33.85	19.54	45.97	293.9	41.54	216.1	661.5	1911	59.54	107.9	1,5

(n= number of analyses; n.d.= not detected; n.a.= not analysed)

oxidative dissolution can release significant amounts of As and trace metals into the environment.

Orogenic Au pyrite has intermediate Te and Se characteristically contents but shows ล significant Te variation over a small range of Se (Keith et al., 2018, Deditius et al., 2014). It should be noted that in the ore occurrences of the Burgetei and Ult, the contents of Te and Se and, accordingly, their ratios are very similar 0.24 ± 0.1 (Fig. 4). The average content of Te (325 - 343 ppm) corresponds to the tellurium of reference world wild orogenic gold deposits. The question of the tellurium incorporation in pyrite requires further study. It can be controlled by the crystal-chemical properties of pyrite (solid solutions) or due to the telluride inclusions.

Trace-element content (Te, Se, Bi, Sb, Cu, Au, Ag) in the arsenopyrite is quite different. Since gold was clearly detected in arsenopyrite, a detailed electron probe microanalysis (13 points) of arsenopyrite of the Ult-samples from quartz vein was performed. This analysis shown limited variation with respect to Fe:As:S ratio. At the same time, the average gold content in recrystallized, coarse-prismatic arsenopyrite is

108 ppm with a maximum of 238 ppm, and minimum 0 ppm. Tellurium content reaches of 0.11wt.% in the arsenopyrite. The high concentration of Bi (up to 0.3 wt.%) and Se (up to 0.05 wt.%) is characteristic of arsenopyrite, but silver is detected only in two points (103 and 97 ppm).

The concentrations and ratios of As and Se, Te could be used to infer ore-forming processes. However, the mechanisms controlling the distribution of Te and Se are not understood well enough (Cook et al., 2013; Keith et al., 2018). Belousov and co-authors data (Belousov et al., 2018) on pyrite indicate two end-member associations in orogenic gold deposit ores: Au-As ores (pyrite As/Te > 200) and Au-Te ores (pyrite As/Te <10). Undoubtedly, the discussed gold ore occurrences are of the first type.

Diagrams in Fig. 6 show the concentrations of Te and Se vs. As in some interesting pyrites and arsenopyrite of the studied occurrences. Significant amount of the thin needle-like pyrites contains Te in the range of 0.004 - 0.06 wt.%; much more tellurium is found in arsenopyrite, the range is 0.03-1.1 %. Not all the samples studied contain selenium, but all of the above with respect to tellurium also applies to



Figure 6. Concentrations of (a) Te and (b) Se vs. As in pyrite (left axis) and arsenopyrite (squares, right axis) from orogenic Au occurrences

the selenium-arsenic relationship. More high concentration of As is observed in the majority of prismatic pyrite crystals from the BG-5/16 and (quartz vein) Ult-8/16 (volcanic metasomatite with jasper and silicification) in comparison with the BG-4/14 (sericite with pyrite). There is no correlation between the content of arsenic and Te. In the arsenopyrite of Ult-3/16 sample (quartz vein) an inverse observed, which is correlation is very characteristic of orogenic gold deposits.

The area in the form of a rectangle is limited by the contents of Te of 200-600 ppm and As of 100 – 10000 ppm. Keith et al. (2018) proposed that the association of Te and As is controlled by the crystal-chemical properties of pyrite due to their preferential incorporation on the $(1\ 1\ 0)$ surfaces in the cubic crystal system. It is noteworthy that some orogenic Au pyrites show elevated Te concentrations and therefore those pyrites host telluride inclusions. The black dashed line in Fig. 3a of the paper above defines the solubility limit for Te solid solution in pyrite as a function of As, which is based on the equation presented by Reich et al. (2005) for the Au-As system. Tellurium concentrations above the solubility line indicate telluride inclusions in pyrite, e.g. at 10,000 ppm of As, Te should be low than 200 ppm. Selenium most likely occurs in solid solution in the corresponding pyrites due to isovalent S substitution. No systematic correlation exists between As and Se in pyrite and arsenopyrite.

DISCUSSION

Sulfide minerals are important indicators of hydrothermal mineralization, and the stable isotopes of sulfur offer a useful tool to constrain the origins of sulfide (Kresse et al., 2018). Although the total sulfur isotopic composition is influenced by temperature, pH value, oxygen fugacity and ion activity, the study of sulfur isotope abundances gives an idea regarding the modes of formation and subsequent histories of sulfur bearing materials. The isotopic content analysis of sulfur in pyrites from Ult-5/16 sample (siltstone with pyrite) and Ult-9/16 sample (quartzite-jasper, Au-content is 3.5 g/t) shows the mantle origin of Uvanga mineralization: δ^{34} S is -6.94 ‰ and -0.91 ‰ respectively. The ranges of δ^{34} S values found in nature for a number of different forms of sulfur are referenced in (Thode, 1970; Brueckner et al. 2014). Different forms of sulfur-bearing materials such as basic sills, primary igneous rocks and volcanic gases show distinctly narrower ranges of ³⁴S values, which tend to be disposed symmetrically about zero. This distinction provides a basic diagnostic tool in sulfur isotope geochemistry of the Uyanga ore mineralization. Recent data (Ward et al., 2017) have revealed that the sulfur isotope composition has a δ^{34} S range of -7.7 to +9.8% during formation of the Wallaby orogenic gold deposit. Given that changes in pH or temperature are minimal, all changes in sulfur isotope and trace element concentrations can be attributed to changes in oxygen fugacity. The

increasing in δ^{34} S is most likely due to a reduction in oxygen fugacity. It was suggested by (Ward et al., 2017) from observations, that next generation of pyrite has a higher δ^{34} S, and rims of pyrite grains have a lower δ^{34} S then a core. Moreover, they have discovered similarities between the rims of pyrite generation 1 crystals and the cores of pyrite generation 2 crystals.

Host sediments of the Uyanga ore knot, basalt, (grano) diorite dykes and other rocks are all changed by the hydrothermal-metasomatic alteration near the gold-bearing quartz veins. Surface weathering, erosion and oxidation intensity clearly shows metasomatic alteration zone. Sometimes initial rock-forming color minerals were completely substituted by carbonates, chlorite and various iron oxides. Limonitization is involved in the surface oxidation of the Cu and Zn minerals. Nevertheless, non-oxidized pyrites sometimes were observed both in the siltstone jasper and in the zones of sericitization. Pyrite cube crystal aggregates spread along the veins and fractures. Thus, we were able to distinguish two generations of pyrite. Further data for sulphides from orogenic gold occurrences would be required to fully evaluate the importance of Devonian metamorphic fluid contributions to the sulfur isotope budget of the Uyanga ore knot.

The ore mineralization is accumulated mainly in quartz veins and zone of tectonic breccia. In the Uyanga gold ore knot, pyrite and arsenopyrite are the most abundant sulfides during the ore forming stage in comparison with chalcopyrite, galena and sphalerite. It indicates that the mineral association of the three new occurrences is relatively simple. Chalcopyrite was identified in the berezite altered rocks, the inset and vein of galena and sulfates have been found in the gold-mercury fine grain sugar like jasperquartzite. Within the ore-controlling zone with brittle-ductile deformation, ore minerals such as antimonite and hematite were observed in the quartz veins and even barite and nickel arsenide were found in some samples. It is widely known that Au concentrations are generally correlate As in arsenical pyrite, but with gold concentration in the studied pyrites was observed below the detection limit by electron microprobe, therefore not given in the paper. Consistently detectable gold in arsenical pyrite should be determined by LA-ICP-MS (Cook et al., 2013).

To better understand the relationship between bulk metasomatite rocks vs. sulfide minerals chemistry, a triangle diagram showing Co-As-Sb-contents of host rocks was constructed (Fig. 7). Siltstones from the Senjit occurrence is generally enriched in Sb (Hg) (blue squares), As tends to show higher and lower contents in the



Figure 7. Compositions of the metasomatic alteration rocks at the three gold occurrences of the Uyanga ore district as a triangle diagram showing indicative elements

Burgetei occurrence (green squares), some rocks of the Ult do not contain Co and Sb (below the detection limit) and high content of gold (red squares and diamond in the angle As).

The most notable changes from the Burgetei to the Ult pyrites chemistry are increase in As concentrations (0.44 and 0.65 wt.%), gradual increase in Cu, Bi concentrations, and decrease in Ni, Sb contents. The first increasing (As) mirror the fact that under more reduced conditions pyrite enrichment in As is connected with the replacement of the S atoms by As. There is no significant correlation between the content of arsenic and Te in pyrite. In the arsenopyrite of Ult-3/16 sample (quartz vein) an inverse correlation is observed, which is very characteristic of the orogenic gold deposits.

Recent data on the geodynamic conditions of the Khangay-Khentey orogenic structure, wall rock alteration and metasomatite composition of the Uyanga gold ore knot, mineral assemblages, chemical compositions of sulfide minerals including characteristic set of impurities (Se, Te, Sb, Hg) in conjunction with the detailed review and analysis of the references below, allowed us to attribute three primary occurrences to the orogenic Au. The varying content of ore elements such as As, Sb, Hg, S indicates the different erosion level of ore formation: Au-As±Sb Burgetei, Au-As±Hg Ult and Au-Hg±Sb occurrences. This is evidenced by variations in the content of Co, Ni, Fe, Mn, W that are

characteristic for the deeper hypozonal horizon. Fluid sources and gold deposition mechanisms could be discussed only tentatively. It was suggested that ore mineralization from the three new gold occurrences has been formed via a gradual change in redox conditions within an evolving fluid. Probably the desulfidation took place too at the final stage (the sulfide minerals are very rare at the Senjit occurrence). The intensive fluid-rock interaction in the Khangay-Khentey mobile orogenic area established the conditions required to igneous and metamorphic Fe-minerals oxidation, (hydr) oxides, chlorite and carbonates precipitation. The relative abundance of Fe (hydr) oxide minerals in the Uyanga host rocks confirm this idea. Negative δ^{34} S isotope values are common in oxidizing environments. This oxidizing environment destabilize Au-bisulfide complexes and trigger gold deposition.

CONCLUSIONS

The primary ore occurrences of the Uyanga gold ore knot district of the Khangay gold zone are hosted in volcanogenic-sedimentary rocks of the lower-middle Devonian Erdenetsogt formation. New information on the Burgetei, Ult, and Senjit gold occurrences of the Uyanga gold ore knot is presented.

The ore mineralization of these occurrences is accumulated mainly in zones of tectonic breccia and quartz veins, pyrite and arsenopyrite are the most abundant sulfides in comparison with chalcopyrite, antimonite, galena and sphalerite. Intensively oxidized and non-oxidized pyrites of different stages were distinguished confirming the variation in oxidation conditions.

The content of metalloids (As, Sb, Te), transition metals (Hg and iron group elements Mn, Fe, Co, Ni) increases in ore bodies of Uyanga gold primary occurrences. The contents of Cu, Pb, Zn and Sn are lower than average content of earth crust. This is the distinct geochemical characteristic of the orogenic gold type deposit (Eilu and Groves, 2001; Eilu et al., 2011). Highest gold content was determined (up to 3.5 g/t in the quartzite-jasper with limonitization of the Ult occurrence cut by numerous small quartz veins and in arsenopyrites of this occurrence (238 ppm). It was suggested that ore mineralization from the three new gold occurrences has been formed via a gradual change in redox conditions. Because negative δ^{34} S isotope values are in oxidizing environments, common this environment destabilizes Au-bisulfide

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complexes and trigger gold deposition.

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