# Epithermal Mineralization in the Okhotsk–Chukchi Volcano-Plutonic Belt

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**Abstract**—This paper considers the geochemistry and conditions of generation for the Mesozoic Au–Ag epithermal deposits in the Okhotsk—Chukchi volcanic—plutonic belt (OChVB) in Northeast Russia. We provide new data on the composition and concentrations of trace elements, including REEs, in the ores of epithermal Au–Ag deposits. The ores were found to be enriched in a wide range of trace elements. The REE distributions of these ores are dominated by light "hydrophile" lanthanoids of the "cerium" group. The Eu anomalies were found to vary between high negative to low and high positive levels. Comparative analysis over the classes of gold concentration showed a synchronous enrichment of the ores in similar sets of trace elements. A study of fluid inclusions revealed that the ore-forming solutions had hydrocarbonate potassium or hydrocarbonate sodium compositions. The fluids had high concentrations of sulfate ions for most deposits. The salinity of the fluids was frequently found to increase toward later low-temperature mineralization phases. We identified the tendency of increasing K<sup>+</sup> percentage in the fluid from the earlier oreless quartz to productive quartz with increasing depth, as well as some decrease in the percentages of Na<sup>+</sup>, Ca<sup>++</sup>, and Cl<sup>-</sup>. The results indicate magma chambers of andesite magmas and meteoric waters as the most likely sources of the fluids that generated the epithermal Au–Ag ores in the OChVB deposits.

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## **INTRODUCTION**

The older placer areas are sufficiently well known in Northeast Russia that it became obvious many years ago that rapid growth in gold mining should primarily be expected from exploration to detect epithermal Au– Ag deposits in volcano-plutonic belts (the OChVB and others). Over 140 tons of gold and 2000 tons of silver have been extracted in the short timespan of 10 years in the unique Kupol Au–Ag epithermal deposit.

The OChVB is the longest among all volcano-plutonic belts in Northeast Russia (over 3000 km) (Fig. 1). Most of the known OChVB Au–Ag epithermal deposits are being mined today. Several deposits have been exhausted: Karamken, Vetvistoe (Magadan Region), Sopka Rudnaya, and Severo-Vostok (Chukotka). A number of deposits (Moroshka, Sentyabr'skoe, Zhil'noe, and Gornoe) are being developed. Most OChVB deposits are of the low-sulfidation type (Kupol, Dvoinoe, Dzhul'etta, and others); the second most numerous group of deposits have intermediate (medium) sulfidation (Dukat, Lunnoe, Gol'tsovoe, and others), while only two deposits (Svetloe and Perekatnoe) can be classified as belonging to the highsulfidation subtype. The OChVB was found to contain several tens of promising ore occurrences, over 2000 mineralization sites, and several thousand geochemical anomalies. The mining companies reported extracting 30 tons of Au and over 960 tons Ag from the OChVB epithermal deposits in 2017.

The main goal of the present study is to sum the data, both that available in the literature and new data obtained by these authors, in order to identify geochemical and thermal, pressure, and geochemical features in the Mesozoic OChVB Au–Ag epithermal mineralization, to derive new information concerning the conditions of mineralization, to use the sum of this knowledge in order to design new criteria and refine known ones for assessment of industrial significance and prediction of deposits. The method used in the analytical studies was described in detail in our previous paper (Volkov et al., 2017b).

## THE MESOZOIC VOLCANISM AND EPITHERMAL Au–Ag MINERALIZATION IN THE OCHVB

The OChVB is 3000 km long; its average width is 200 km. The OChVB was forming during 25 myr



**Fig. 1.** A map showing the regionalization of the OChVB and the location of the known epithermal Au–Ag deposits. (1) a generalized boundary of the OChVB igneous formations: Albian–Santonian: on land (a) and in sea (b); (2) boundary between the outer and inner zones of the OChVB; (3) boundary of the subzones of depression grabens and of that of magmatogenic uplifts; (4) inferred boundaries in water area; (5) boundaries of sectors in the outer zone and in flank zones; (6) volcanic rocks in outer zone; (7) OChVB magmatogenic uplifts; (8–11) OChVB outer zone, sectors: (8) Okhotsk, (9) Penzhina, (10) Anadyr, (11) Central Chukchi; (12, 13) flank zones: (12) West Okhotsk, (13) East Chukchi); (14) Late orogenic (late Barremian to early Albian) Mesozoic structures: O Omsukchan graben, P Verkhne-Penzhina, U Umkuveem, A Ainakhkurgen, T Tytyl'veem basins, KH Khurchan–Ortukan TMA zone; (15) gold–silver deposits: (1) Nyavlenga, (2) Dzhul'etta, (3) Tikhoe, (4) Dukat, (5) Agan, (6) Oira, (7) Dal'nee, (8) Kupol, (9) Moroshka, (10) Televeem, (10) Dvoinoe, (11) Sentyabr'skoe, (13) Pepenveem, (14) Pechal'noe.

(Belyi, 1994) at the boundary between the continental Verkhoyansk–Chukchi and Koryak–Kamchatka terrane (see Fig. 1). The OChVB evolution proceeded in two phases: (1) the earlier (Albian–Cenomanian) phase in which approximately 90% of all volcanic rocks were produced, and (2) the later (Turonian–Santonian) phase when the Koryak–Kamchatka structures evolved. The formation of the OChVB was probably caused by the Pacific plate being subducted under the collage of the terranes accreted to the Siberian continent. The main sources of acid magma were in the continental crust during this volcanic episode; the crust seems to have been Neoproterozoic (Tikhomirov et al., 2008).

The OChVB consists of three elements (see Fig. 1): the main arcuate part that extends from the lower reaches of the Ul'ya River to the coast of the East Siberian Sea (approximately 2000 km); this part is called the Tauisk–Chaun arcocline, with the other elements being the western Sea-of-Okhotsk and East Chukchi flank zones (Belyi, 1994). In fact, these elements are independent structures that are joined together by a left lateral end-part juncture that were formed during the same span of geologic time.

The outer and flank zones of the OChVB typically contain various isometric negative volcanic structures: subsidence structures with a caldera form (depressions and subsidences) and calderas; circular intrusive– effusive edifices; and volcanic grabens and semi-grabens that have inherited the fault directions in the basement. Dome-like intrusive–effusive and cryptointrusive structures are abundant in all zones, but their role is relatively small, although the mineralization location is not infrequently controlled by these structures.

One important element in the OChVB structure consists in magmatogenic uplifts where the principal parts of the larger intrusive bodies are concentrated. In the inner zone, most of these structures were evolving inheriting the analogous uplifts in the Koni—P'yagin zone of the Taigonos volcanic arc. These were forming in the outer zone during the earlier phase in the OChVB evolution and their relative area is considerably below that in the inner zone. Some of these are elongate parallel to the overall OChVB trend, while others are transverse. Magmatogenic uplifts are infrequent in the flank zones.

The Verkhovansk-Chukchi Mesozoic features experienced the greatest activation in the late Cretaceous to early Paleogene: the activation seems to have been synchronous with a phase in the OChVB evolution (Sidorov et al., 2013). The area of influence exerted by the Late Mesozoic tectono-magmatic activation (TMA) is identified as a perivolcanic zone of a hazy boundary (as wide as 500 km) that covers the southern part of the Yana-Kolyma system, the Oloi zone, and the Chukchi system. One feature that is noticeable in the deep structure of the perivolcanic zone is the presence of large low-density locations in the lithosphere. These locations are frequently adjacent to the OChVB volcanogenic depressions and basins that show some inheritance of the Mesozoic structural plan. The linear TMA (tectono-magmatic activation) zones are treated here as branches away from the OChVB; chains of volcanic fields and intrusive bodies are also frequently located on extensions of tongues of the belt blanket formations of varying lengths. At the same time, they are different from the belt formations in having different tectono-magmatic features, which are largely controlled by isolated local occurrences of magmatism and by metallogenic features.

All the OChVB structures and the feather TMA zones described above have ore potential. However, the ore potential of volcanogenic structures and of the feather TMA zones are known fragmentarily for the southern part of the Okhotsk sector of the OChVB, for the outer part of the Central Chukchi sector, and very poorly for the inner part of the Anadyr sector.

The complex structure of the basement terranes gave rise to a great diversity of ore assemblages in the OChVB (Sidorov et al., 2009). The inner zone of the belt more frequently contains deposits of the Cu-Moporphyry formation containing Au, Ag, and platinoids. The outer and perivolcanic TMA zones typically contain Au-Ag, Ag-base-metal and Sn-Ag mineralization (Sidorov et al., 2011). Epithermal Au-Ag deposits proper (with Au/Ag 1/1-1/10) are more abundant in the outer zone. Silver-dominated deposits (Au/Ag = 1/10 - 1/1000) are confined to the Omsukchan riftogenic volcanic depression, which introduces complications into the OChVB between the Yana-Kolyma and Omolon terranes (Sidorov et al., 2009). It should also be remarked that the OChVB does not contain any significant Au-Ag-Te deposits and occurrences, which were thought by Nolan (1933) to be related to small volcanic bodies in the Cenozoic volcanic belts of the western United States.

The generation of epithermal deposits of dominantly Ag and Au–Ag types in the volcanic belts is largely due to the fact that they are underlain by potentially gold- and silver-bearing rock sequences of the Verkhoyansk and Chukchi terrigenous units, as well as by older metamorphic units of median massifs (cratons), which were extra sources of metals for volcanogenic–plutonogenic deposits (Volkov et al., 2006).

Comparison among different OChVB sectors by the intensity of epithermal mineralization shows that the well-known Okhotsk sector (the Karamken, Nyavlenga, Dzhul'etta, and Dukat deposits, the Evensk group, etc.) is inferior to the poorly known Anadyr sector in terms of proved Au reserves in epithermal deposits. The Anadyr deposits are Kupol, Moroshka, Sentyabr'skoe, Valunistyi, and Dvoinoi.

This fact seems to be due to the rather complex geological structure of the Okhotsk sector. In this sector one finds abundant rejuvenated epithermal deposits (Goncharov and Sidorov, 1979). In addition, the Okhotsk sector is dominated by acidic igneous units of the Ag–Sn–Mo type (Sidorov et al., 2009). At the same time, one discerns a pronounced relationship between epithermal Au–Ag mineralization and the Cu-porphyric ore-forming system on the Kupol deposit in the Anadyr sector (Volkov et al., 2012).

The geochemical features of Au–Ag ores in epithermal deposits that will be discussed below show many similarities to the ore assemblage patterns pointed out above.

# THE GEOCHEMICAL FEATURES OF THE OCHVB EPITHERMAL ORES

The ores sampled from the Mesozoic epithermal Au–Ag deposits are dominated by  $SiO_2$  (67.6–92.8%) along with noticeable concentrations of  $Al_2O_3$  (3.2– 14.0%), K<sub>2</sub>O (0.44–5.5), and Fe<sub>2</sub>O<sub>3</sub> (0.48–5.1) (Table 1). The ores of the Tikhoe deposit are found to contain noticeable concentrations of Na<sub>2</sub>O (2.27%) and CaO (4.83%) (see Table 1), which is caused by mineralization in an argillite zone (Volkov et al., 2015). The Dukat ores typically show high concentrations of MnO (3.9) and the Nyavlenga ores have high concentrations of MgO (3.46) (see Table 1), which is due to the special type of mineralization in these deposits (Volkov et al., 2014). The metasomatic ores of the Agan deposit (Volkov et al., 2015) show the highest total  $Na_2O + K_2O$  (see Table 1). Judging from the data in this table, the concentration of sulfides in the ore samples studied here varies in a wide range (S<sub>total</sub> is between 0.3% and 3.2%).

The results of trace-element analyses for the OChVB ores of volcanogenic epithermal Au-Ag deposits and host rocks are presented in Table 2 and in Fig. 2 where the values have been normalized by the means for the upper crust (Taylor and McLennan, 1985). It follows from Table 2 and Fig. 2 that the ores are clearly enriched in a wide range of elements (Li, P, Rb, V, Cr, Sc, Ba, Mn, Au, Ag, As, Sb, Te, Cu, Mo, Zn, Pb, In, Cd, Ga, Co, Mo, Bi, Tl, Cs, Be, Se, W, and U) compared with the upper crustal means (Tayler and McLennan, 1985). The enrichment ratios vary between a few times (Li, P, Sc, Rb, Cs, Cr, V, Tl, Ba, Se, W, and U) and a few tens (Mo, Mn, and Sn), hundreds (Pb, Cu, Zn, Be, and Te), and thousands (As, Sb, In, Cd, and Bi), reaching a few tens or hundreds of thousands of times (Au and Ag), thus providing evidence of a geochemical affinity of trace elements and their synchronous participation in the mineralization.

Comparative analysis of mean trace-element concentrations in the ores of the OChVB epithermal Au– Ag deposits shows great similarities in composition and distribution (see Table 2, Fig. 2). This shows that the mineralization of these deposits occurred under similar conditions. At the same time, the Dukat ores are characterized by an appreciable synchronous enrichment in Be and Mn (see Fig. 2), which is due to a wide abundance in the ores of minerals that contain these admixtures (Konstantinov et al., 1998). The Pechal'noe ores show an obvious synchronous enrichment in Li, Be, Rb, Cs, Mo, W, and Tl, which corresponds with the composition of the host volcanic rocks (Egorov et al., 2005; Volkov et al., 2017b). The

Components,		o nonicodi						DEPOSITS						
%	Dvoinoi	Dzhul'etta	Dukat	Kupol	Moroshka	Oira	Dal'nee	Pechal'noe	Tikhoe	Nyavlenga	Sentyabr'skoe	Pepenveem	Televeem	Agan
и	9	6	5	6	5	9	4	9	5	7	9	5	2	5
SiO <sub>2</sub>	87.015	84.17	75.186	92.838	80.788	75.023	76.94	85.923	76.834	67.589	68.267	93.254	82.48	84.41
TiO <sub>2</sub>	0.057	0.03	0.046	0.118	0.098	0.153	0.328	0.255	0.224	0.487	0.28	0.046	0.07	0.07
$Al_2O_3$	5.788	4.463	6.06	3.845	6.552	7.928	10.898	7.17	6.228	14	9.618	2.754	7.71	7.92
${\rm Fe_2O_{3total}}$	0.487	2.44	5.122	0.645	2.818	2.053	5.023	2.232	2.91	5.159	4.538	1.794	1.345	1.03
MnO	0.039	0.049	3.925	0.107	0.112	0.063	0.113	0.022	0.126	0.082	0.104	0.013	0.01	0.01
MgO	1.203	0.293	1.08	0.535	1.482	2.085	1.253	0.392	1.534	1.241	3.468	0.084	0.12	<0.10
CaO	0.758	0.613	1.044	0.785	3.134	0.94	0.39	0.072	7.736	4.83	2.417	<0.10	<0.10	0.06
$Na_2O$	0.128	0.142	0.306	0.048	0.078	0.635	0.328	0.145	0.348	2.277	0.433	0.052	0.16	0.33
$K_2O$	3.615	2.682	1.97	0.442	2.304	3.35	3.24	2.858	1.406	3.647	2.88	0.812	5.57	5.51
$P_2O_5$	0.022	0.015	0.008	0.027	0.02	0.057	0.1	0.103	0.072	0.214	0.082	0.028	0.02	0.03
$S_{total}$	0.303	3.21	0.746	0.418	1.326	0.273	1.243	0.542	1.248	0.421	2.987	0.356	1.315	0.56
L.O.I.	Ι	Ι	I	0.923	2.052	0.99	I	I	I	1.226	0.378	I	Ι	I
Ω	99.415	98.107	95.493	100.731	100.764	93.55	99.856	99.714	98.666	101.173	95.452	99.193	98.8	99.93
X-ray fluorescen	ce analysis	, at IGEM R	AN labor	atory, Ana	lyst A.I. Yak	ushev; L.	O.I. is short	t for loss on ig	gnition; her	e and below	n denotes the nun	nber of sampl	es.	

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Pepenveem     Televeem       178.056     48.108       1.45     4.569       n.d.     11.823       1.45     4.569       n.d.     11.823       149.345     343.755       149.345     343.755       149.345     343.755       149.345     343.755       12.157     9.548       12.157     9.548       12.157     9.548       12.157     9.548       12.157     9.548       12.157     9.548       11.943     248.27       11.943     248.27       318.184     0.506       12.153     11.943       255.582     243.508       21.144     84.575       255.582     243.508       21.144     84.575       2.687     1.926       12.158     1.926       12.158     1.926       12.158     1.926       12.158     1.926       12.158     43768       0.48.07	0.075     3.95     0.297     72.186       5.61     13.033     3.512     4.729       348.857     421.5     151.308     216.842       11.886     7.552     5.941     4.426
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<b>EPOOSITTS Pecchal'noe</b> 152.047     21.773     21.773     n.d.     1.687     1.687     1.687     1.687     1.687     1.687     1.687     1.687     1.687     1.687     1.687     13.39.038     20.06     50.386     113.516     113.516     113.516     113.516     1.977     8.701     59.599     18.635     13.088     25.929     6.118     44.386     3.341     44.386     3.341     44.44     106.698     0.144     0.647     0.647	0.279 0.279 21.176 147.48 8.231
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Kupol       108.667       108.667       0.902       833.667       19.985       652.5       30.167       35.55       30.167       35.55       30.167       35.55       30.167       35.55       30.167       35.55       30.167       35.55       31.85       5.958       5.958       5.958       31.85       5.958       5.958       31.85       5.958       5.958       5.958       5.958       5.958       5.958       5.958       5.958       5.958       5.958       5.958       5.958       18.417       0.215       10.215       10.817       1.333       395.333       395.333	0.235 5.4 27 3.798
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S S S S S $\mathcal{A}_{\mathcal{A}}$ S S S S S S S S S S S S S S S S S S	Te Cs Ba La

Table 2. The element composition (g/t) of ore samples from the OChVB epithermal Au–Ag deposits

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	Elements,							Ω	EPOSITS						
	g/t	Dvoinoe	Dzhul'etta	Dukat	Kupol	Moroshka	Oira	Dal'nee	Pechal'noe	Tikhoe	Nyavlenga	Sentyabr'skoe	Pepenveem	Televeem	Agan
	Ce	3.21	1.64	6.592	6.75	6.98	8.462	22.225	16.379	12.729	24.143	14.738	11.561	8.126	16.280
	Pr	0.373	0.188	0.824	0.747	0.892	1.193	2.775	1.996	1.525	3.129	1.912	1.344	0.877	1.954
	Nd	1.528	0.822	3.444	2.835	3.4	5.267	11.225	7.291	5.735	12.529	7.513	4.87	2.93	8.140
	Sm	0.3	0.178	0.85	0.635	0.78	1.248	2.208	1.482	1.147	2.429	1.364	1.026	0.541	1.630
	Eu	0.059	0.171	0.105	0.253	0.378	0.338	0.7	0.311	0.727	0.799	0.422	0.159	0.145	0.322
	Gd	0.39	0.237	0.89	0.611	0.912	1.25	2.158	1.17	1.265	2.357	1.181	0.896	0.301	1.592
J	Tb	0.034	0.017	0.151	0.136	0.129	0.187	0.358	0.196	0.167	0.356	0.188	0.132	0.054	0.284
OU	Dy	0.243	0.153	1.037	0.838	0.872	1.185	1.923	1.105	0.971	2.057	1.065	0.711	0.346	1.820
RN/	Но	0.037	0.022	0.202	0.187	0.143	0.23	0.37	0.237	0.189	0.411	0.22	0.114	0.065	0.356
AL (	Er	0.139	0.071	0.616	0.503	0.382	0.72	0.978	0.703	0.551	1.15	0.627	0.266	0.218	1.024
OF '	Tm	0.01	0.002	0.077	0.061	0.048	0.094	0.142	0.103	0.071	0.168	0.093	0.033	0.031	0.130
VOL	Yb	0.149	0.068	0.606	0.449	0.372	0.719	0.808	0.689	0.485	1.093	0.63	0.212	0.261	0.988
CAN	Lu	0.01	0.001	0.072	0.073	0.044	0.098	0.122	0.096	0.065	0.161	0.093	0.031	0.044	0.132
IOLO	Hf	0.184	0.004	0.308	0.733	0.44	1.127	1.37	1.244	0.858	1.8	1.552	0.005	0.448	1.194
GY A	Та	0.00	n.d.	0.032	0.112	0.026	0.075	0.145	0.184	0.083	0.487	0.217	0.038	0.088	0.111
AND	W	0.202	0.147	0.203	12.3	0.88	0.707	2.15	17.861	0.029	1.613	1.687	3.616	n.d.	0.106
SEIS	IL	1.977	0.387	1.222	1.453	1.414	1.678	1.038	6.86	3.995	1.824	1.017	0.192	3.106	3.830
MOL	Pb	438.667	11389.167	3479	75.483	1038.2	305.5	16.75	13.534	5582.222	26.629	25243.167	2570.633	374.958	27.940
.0GY	Bi	0.54	0.125	0.167	0.034	0.18	0.112	1.304	0.798	0.188	0.27	0.077	152.29	0.468	0.205
v	Th	1.043	0.125	1.14	0.962	0.427	1.233	1.025	2.162	1.877	3.129	4.143	2.134	1.756	2.480
ol. 12	U	0.544	0.067	0.469	0.499	0.305	0.51	0.324	0.702	0.549	1.171	1.021	1.44	0.74	3.732
2 1	Au	51.218	1805.721	32.132	67.965	84.434	25.937	360.349	2.919	1714.051	7.449	461.729	1.377	158.005	3.202
No. 6	Cu	328.333	768.333	1424	33.333	226	138.333	82.5	15.318	632.084	30	2096.667	43.844	6587.089	4.000
20	u	9	9	5	9	5	9	4	9	5	7	6	5	7	5
18	Inductively with electro	coupled plass thermal atom	ma mass specta ization using a	roscopy (IC Spectr AA	(The State of the	GEM RAN ctrometer (/	laboratory Analyst V.A	(Analyst) A. Sychkov	Ya.V. Bychko 'a); n.d. mea	ova). Gold ii ns not detec	n the samples ted.	was determined	by atomic abs	orption spec	troscopy

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Table 2. (Contd.)

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Fig. 2. Trace elements in the ores of the OChVB epithermal Au–Ag deposits as normalized by the mean upper crustal values (Taylor and McLennan, 1985).

epithermal ores in the Agan deposit have a distinguishing feature, viz., a high enrichment in Sn (see Fig. 2), which classifies the ores as belonging to the tin-porphyritic mineralizing system (Volkov et al., 2015). Similar distributions but different intensities of individual elements occur in the Kupol, Moroshka, Dvoinoi, Sentyabr'skoe, Televeem, Dzhul'etta, Oira, Dal'nee, Nyavlenga, and Tikhoe deposits (see Fig. 2), which may have been because they belong to the Cu-Mo-porphyritic mineralizing system (Volkov et al., 2012, 2014). The low intensity of elements in the distribution of the Nyaylenga deposit compared with the ores of the other known deposits seems to be due to thermal metamorphism of the ores (Savva et al., 2007). The noticeable enrichment in Bi and W observed in the Pepenveem ores provides evidence that this deposit belongs to a mineralizing system related to granitoid intrusions (Lang et al., 2000).

Comparative analysis of the trace-element distributions in the ores over gold concentration class (Fig. 3) emphasizes a synchronous enrichment of the ores in a similar set of trace elements from low (1-5 g/t) to high class (>100 g/t).

The OChVB host volcanic rocks are also enriched in a wide range of elements (Li, Sc, Ti, V, Mn, Sr, Co, Ba, REEs, etc.) compared with the upper crustal values (Taylor and McLennan, 1985), but the enrichment ratios for most of them are low (2 to 5 times). The highest enrichment ratios ranging between 5 and 10 or higher (As, Ag, Pb, Sb, and Mo) were found for host rocks within the epithermal ore fields (Sakhno et al., 2015; Kravtsova, 2010).

According to V.A. Zharikov et al. (1999), light REEs pass into aqueous fluid as the pressure increases, while heavy ones are retained in the magma, so that one can treat the former as "hydrophile" elements and the latter as "magmaphile" ones. Three REE sets can be identified (Mineev, 1974): the cerium set (La, Ce, Pr, and Nd), the yttrium set (Sm, Eu, Gd, Dy, and Ho), and the scandium set (Er, Yb, and Lu). Judging from the data in Table 3, the REE distributions of the ores are dominated by light "hydrophile" lanthanoids of the "cerium" set. A similar REE composition is also typical of the OChVB host volcanic rocks (Kravtsova et al., 2005; Kravtsova and Almaz, 2006; Kravtsova, 2010; Tikhomirov et al., 2008, 2016; Sakhno et al., 2015).

We know that Cl-bearing hydrothermal fluids can effectively concentrate light REEs, but are depleted in heavy REEs (Oreskes and Einaudi, 1990); in that case the Hf/Sm, Nb/La, and Th/La ratios in the ores are generally below 1, while F-rich fluids synchronously concentrate light and heavy REEs, since the Hf/Sm, Nb/La, and Th/La ratios are usually greater than 1 (Oreskes and Einaudi, 1990). The ores in most of the OChVB epithermal Au–Ag deposits are evidently enriched in light REEs and depleted in heavy REEs, and have Hf/Sm, Nb/La, and Th/La ratios below 1 (see Table 3); one exception is the Sentyabr'skoe deposit where Hf/Sm = 1.13. It follows that the oreVOLKOV et al.



**Fig. 3.** The trace-element distribution over classes of Au concentration in the ores of the OChVB epithermal Au–Ag deposits as normalized by the mean upper crustal values (Taylor and McLennan, 1985). Classes of Au concentration (g/t): >100 (a), 10–20 (b), and 1–5 (c).

ş							DI	EPOSITS						
Dvoin	oe	Dzhul'etta	Dukat	Kupol	Moroshka	Oira	Dal'nee	Pechal'noe	Tikhoe	Nyavlenga	Sentyabr'skoe	Pepenveem	Televeem	Agan
7.86	-	4.3	18.355	17.873	18.958	24.497	56.714	39.99	32.057	62.666	37.598	27.296	18.365	43.252
6.84	<u>8</u>	3.73	14.703	15.018	16.056	20.014	49.858	35.691	28.293	54.913	33.501	24.901	17.045	36.926
1.01	Э	0.57	3.652	2.855	2.902	4.483	6.857	4.3	3.764	7.753	4.097	2.396	1.32	6.326
6.7	61	6.541	4.026	5.26	5.533	4.465	7.272	8.3	7.517	7.083	8.178	10.395	12.911	5.837
0.6	613	0.021	0.362	1.156	0.564	0.903	0.621	0.839	0.747	0.741	1.137	0.005	0.828	0.733
0.2	248	0.046	0.238	0.233	0.149	0.324	0.246	0.406	0.198	0.726	0.373	0.092	0.327	0.240
0.0	757	0.171	0.395	0.253	0.118	0.352	0.096	0.263	0.292	0.263	0.549	0.359	0.397	0.288
39.	91	42.481	30.495	31.891	33.38	25.399	28.514	25.789	28.81	28.715	27.473	23.538	29.777	33.034
0.	521	0.537	0.411	0.519	0.715	0.414	0.316	0.325	0.293	0.374	0.247	0.675	0.422	1.505
ω.	432	1.344	15.265	0.578	1.024	1.157	0.618	8.376	0.346	0.459	0.811	1.21	2.879	7.685
0	439	0.02	0.026	1.18	0.308	0.361	0.407	0.176	1.117	0.721	0.273	0.14	0.39	0.062
70.	952	1355.909	36.753	31.514	39.605	39.822	37.226	35.688	43.169	34.444	36.042	2382.513	35.747	39.983
37.	273	Ι	21.5	7.899	20.769	15.133	18.141	18.173	15.292	17.713	12.986	14.211	16.475	18.559
0.	889	1.713	9.286	1.728	1.614	3.597	6.108	0.227	0.787	1.677	2.363	0.584	0.24	0.065
0.	811	0.158	0	0.008	I	4.576	0.469	0.021	2.524	0.034	0.62	Ι	1.485	0.609
0.	643	3.495	0.125	0.441	0.543	0.637	1.907	0.027	32.483	4.574	11.126	0.007	0.147	0.015
0.	773	2.635	0.488	1.12	1.382	0.916	1.011	0.779	1.746	1.066	1.033	0.627	1.004	0.696
Ξ.	15	1.089	1.071	0.909	0.952	1.089	1.032	1.009	1.005	1.012	0.989	0.992	0.97	0.953
6.	271	7.314	3.237	5.753	6.625	3.313	9.023	8.111	8.998	7.388	8.143	18.994	L11.517	5.913
6	871	2.557	2.122	3.738	2.903	1.754	3.034	3.468	3.499	3.056	3.457	3.617	5.11	3.295
	114	2.827	1.189	1.101	1.985	1.407	2.162	1.372	2.108	1.745	1.517	3.413	0.933	1.304
14.	074	90.968	4.152	5.413	8.593	3.708	9.125	8.855	10.225	7.656	8.428	20.179	10.502	6.763
6.	489	3.381	13.748	14.13	14.898	18.428	46.95	33.898	26.418	51.686	31.715	23.716	16.359	34.974
÷	063	0.778	3.236	2.659	3.214	4.438	7.715	4.501	4.466	8.409	4.44	3.038	1.452	6.004
0.	298	0.139	1.294	1.024	0.798	1.537	1.907	1.488	1.101	2.404	1.35	0.509	0.523	2.144
0.	197	0.956	0.124	0.399	0.485	0.271	0.317	0.209	0.634	0.329	0.309	0.155	0.268	0.197
21.	496	24.177	10.878	15.05	18.774	11.769	27.523	23.757	26.227	22.092	23.394	54.412	31.127	16.478
0	.018	0.104	0.016	0.038	0.054	0.04	0.031	0.019	0.057	0.033	0.029	0.014	0.018	0.020

Table 3. The indicator parameters of ore sampled from the OChVB epithermal Au–Ag deposits

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 $Eu/Eu^* = Eu_N/(Sm_N^*(Tb_N^*Eu_N)^{1/2}); Ce/Ce^* = Ce_N/((2La_N + Sm_N)/3), REE , LREE stands for light REE, HREE for heavy REE.$ 

forming fluids of the deposits belonged to a NaCl– $H_2O$  hydrothermal system that was enriched in Cl relative to F; this agrees with studies of fluid inclusions in ore quartz (Volkov et al., 2012; Kolova et al., 2015; Prokof'ev et al., 2015, among others).

The U/Th ratio for an ore reflects the redox conditions of the host rocks (Jones and Manning, 1994): U/Th  $\leq 0.75$  occurs in an oxidizing environment; U/Th 0/75–1.25 is characteristic of an environment without oxygen, and U/Th > 1.25 for a reducing environment. Judging from the data in Table 3, the U/Th ratios in the ores of the deposits studied here are below 0.75 (they vary between 0.24 and 0.71), providing evidence of an oxidizing environment during their mineralization (Jones and Manning, 1994). The high values of that ratio (1.5) in the Agan ores (see Table 3) indicate a reducing environment of mineralization that is characteristic for the shallow level of the epithermal system where the deposit was formed (Volkov et al., 2015).

The Co/Ni ratios in the Dvoinoe, Pechal'noe, Tikhoe, Pepenveem, Televeem, and Agan ores (see Table 3) are considerably below 1.0 (they vary between 0.06 and 0.88), which is characteristic for medium- and lowtemperature hydrothermal fluids of meteoric origin (Kun et al., 2014). Values above 1.5 in the Dzhul'etta, Dukat, Kupol, Moroshka and other ores (see Table 3) provide evidence of an obvious role played by a magmatic fluid in the mineralization (Kun et al., 2014). The high Co/Ni ratio in the ores shows the superposition of a later magmatic fluid on the earlier mineralization and is a consequence of a rejuvenated mineralization (Goncharov and Sidorov, 1979).

Effective use of the Y/Ho ratio for assessing the origin of mineralizing fluids was shown in (Bau, 1991; Jones and Manning, 1994; Monecke et al., 2002). According to Table 3, the Y/Ho ratios in the OChVB epithermal Au–Ag ores studied here vary between 23.53 and 42.48, which corresponds with the range of the ratios that are characteristic of the present-day hydrothermal fluids in backarc basins (Bau, 1991; Jones and Manning, 1994; Monecke et al., 2002).

The REE composition in the OChVB epithermal Au–Ag ores studied here is shown in Table 2, while the chondrite-normalized REE distributions are plotted in Fig. 4. Abnormally low values of  $\Sigma REE$  (between 4.3 and 18.95 g/t) were found in the Dvoinoe, Dzhul'etta, Dukat, Kupol, Moroshka, and Televeem ores (see Table 3). Lower values of  $\Sigma REE$  (24.49– 39.99 g/t) are characteristic for typical epithermal ores in the Oira, Pechal'noe, Tikhoe, Pepenveem, and Sentyabr'skoe deposits. Higher values (43.25-62.66) of  $\Sigma REE$  were found in the Agan, Nyavlenga, and Dal'nee ores. Low total REE concentrations, the same as in the OChVB deposits, were found in productive quartz in the ores of the epithermal deposits on the Kuramin Range, Uzbekistan and of the Banská Štiavnica area, Slovakia (Vinokurov et al., 1999). The total REE concentration in the epithermal Au–Ag ores of the deposits studied here (see Table 3) is appreciably lower than those in the OChVB igneous and volcanic rocks and in island-arc andesites (Kravtsova et al., 2005; Kravtsova and Almaz, 2006; Kravtsova, 2010; Tikhomirov et al., 2008, 2016).

The chondrite-normalized REEs of the ores studied here make slightly dipping near-chondrite distributions (see Fig. 4a) that exhibit many similarities in their configurations to those of the REE distributions for the OChVB rocks (see Fig. 4c), as well as for island-arc andesites (Kravtsova et al., 2005; Kravtsova and Almaz, 2006; Kravtsova, 2010, among others).

The REE distributions clearly distinguish three sets of ore (see Fig. 4a). The distributions of the first set (Dzhul'etta, Moroshka, Tikhoe, and Kupol) show well-pronounced Eu maxima (see Figs. 4a, 5). Pronounced Eu minima were found for the distributions of the Dvoinoe, Dukat, Pepenveem, and Agan ores (see Figs. 4a, 5). The slightly dipping near-chondrite distributions that involve no pronounced Eu maxima and minima are characteristic for the ores of the third set (Nyavlenga, Televeem, Pechal'noe, Oira, and Sentyabr'skoe). Differently directed distributions were found for the Kupol ores (see Fig. 5), which supports the identification of two types, viz., high- and low-sulfidation epithermal mineralization, on this deposit (Volkov et al., 2012).

The REE distributions clearly separate the OChVB host volcanic rocks into two sets. Acidic rocks typically show distributions with pronounced Eu minima, while intermediate rocks make slightly dipping nearchondrite distributions. We note that the epithermal Au–Ag Cenozoic deposits of Kamchatka that are localized in intermediate (andesitic to dacitic) rocks were found to have near-chondrite REE distribution configurations (Takahashi et al., 2007; Andreeva, 2013).

To summarize, we can state that the different ratios in the ore-containing section that is composed of volcanic rocks with different compositions seems to control the REE distribution configurations in the ores. These results suggest that the OChVB host rocks, as well as the host rocks in the other volcano-plutonic belts in Northeast Russia (Volkov et al., 2017a) might be the source of the REEs and possibly of other trace elements for the ore-forming fluids.

According to S.V. Vinokurov et al. (1999), positive Eu anomalies constitute a characteristic feature of productive quartz in epithermal Au–Ag deposits, while oreless quartz in such deposits exhibits negative Eu anomalies (Vinokurov et al., 1999). This inference is also supported by our results (see Fig. 4b). Judging from this figure, the distributions of rich ores differ from those of poor ores in having pronounced Eu maxima. It may be hypothesized that the Eu maxima in the ores of the first set (see Fig. 4a) are due to their richness.



**Fig. 4.** The distribution of chondrite-normalized mean REE values ((a) over deposits, (b) over classes of Au concentration) in the ores of epithermal Au–Ag deposits and host rocks (c) in the OChVB. Numerals in circles (1–14) refer to known deposits: (1) Nyavlenga, (2) Dal'nee, (3) Agan, (4) Pechal'noe, (5) Oira, (6) Sentyabr'skoe, (7) Dukat, (8) Tikhoe, (9) Kupol, (10) Moroshka, (11) Pepenveem, (12) Televeem, (13) Dvoinoe, (14) Dzhul'etta. OChVB rocks: (1) rhyolite lavas; (2) rhyolite ignimbrites; (3) andesites; (4) dacites; (5) dacitic tuffs; (6) rhyolite dikes (Tikhomirov et al., 2008; Sakhno et al., 2015).

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**Fig. 5.** The chondrite-normalized REE distribution in the ores of the OChVB epithermal Au–Ag deposits. Ag.37 etc. are sample numbers.

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Densit	Physi	co-chemical p	arameters of fluid	ds	Defense
Deposit	<i>T</i> , °C	<i>C</i> , wt %	d, g/cm <sup>3</sup>	P, bars	References
Dukat	201-357 (6)	2.5-2.8	0.63-0.88	90 (1)	Berman et al., 1993
Dal'nee	220-355 (22)	0.6-2.4	0.57-0.86	_	Kravtsova, 2010
Dukat	185-435 (46)	1.5-37.0	0.64-1.11	_	Kravtsova, 2010
Kupol	222-267 (4)	2.1	0.78-0.86	_	Volkov et al., 2006
Kupol	225 (1)	1.9	0.85	_	Sidorov et al., 2007
Kupol	222–276 (12)	0.5-3.2	0.75-0.86	_	Volkov et al., 2012
Dvoinoe	154-251 (7)	0.2-5.0	0.80-0.95	_	Volkov et al., 2012
Arykvaam	234–267 (10)	0.4-1.2	0.79-0.82	_	Volkov et al., 2012
Kupol	224–276 (12)	0.5-3.2	0.75-0.86	_	Prokof'ev et al., 2012
Dvoinoe	154–251 (7)	0.2-5.0	0.80-0.95	_	Prokof'ev et al., 2012
Dvoinoe	133–254 (9)	0.5-3.9	0.82-0.94	_	Nikolaev et al., 2013
Sentyabr'skoe	155-360 (32)	0.9-8.1	0.24-0.94	80-570 (8)	Nikolaev et al., 2013
Nyavlenga	157-359 (27)	0.3-8.2	0.56-0.93	_	Volkov et al., 2014
Dzhul'etta	126-222 (20)	1.2-5.6	0.86-0.97	_	Prokof'ev et al., 2015
Tikhoe	105-260 (11)	1.0-9.2	0.82-1.01	_	Kolova et al., 2015
Sentyabr'skoe	137–296 (34)	0.0–7.4	0.69-0.96	_	Savva et al., 2016

Table 4. The physico-chemical parameters of mineral-forming fluids at the OChVB Au-Ag deposits

The number of determinations is enclosed in parentheses, a dash means no data.

Eu and Ce anomalies are commonly treated as marking the redox potential of a mineralization environment (Bortnikov et al., 2007; Goryachev et al., 2008; Jones and Manning, 1994). The ores sampled from the OChVB deposits (see Table 3) show  $\delta$ Ce and  $\delta$ Eu varying between negative or weakly negative and moderately positive values ( $\delta$ Ce = between 0.90 and 1.15) and ( $\delta$ Eu = between 0.48 and 2.63). This combination of  $\delta$ Ce and  $\delta$ Eu indicates low-oxidation and oxidation conditions that existed during the deposition of the OChVB epithermal ores. The low Eu/Sm ratios (<1) in the ores studied here (see Table 3) suggest the inference that the OChVB mineralization occurred in the upper crust under similar physical and chemical conditions (Vinokurov, 1996).

Comparison between the data obtained here and known published examples (Vinokurov et al., 1999; Kravtsova, 2010, among others) indicates that the patterns identified here, viz., the depletion of ores in rareearth elements, the prevalence of light lanthanoids

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over heavy ones, low Eu/Sm ratios (<1), and positive Eu anomalies, all constitute, in all probability, typical features for the epithermal mineralization system.

### THE DISTINGUISHING FEATURES OF THE ORE-FORMING FLUIDS

Many results have been reported concerning physico-chemical parameters and composition of oreforming fluids for the OChVB Au–Ag deposits (Table 4, Fig. 6). Papers exist concerning the conditions of mineralization for the Dukat (Goncharov and Sidorov, 1979; Berman et al., 1993; Kravtsova et al., 2003; Konstantinov et al., 1998, among others), Dal'nee (Kravtsova et al., 2003; Kravtsova, 2010), Kupol (Volkov et al., 2006, 2012; Sidorov et al., 2007; Prokof'ev et al., 2012), Dvoinoe (Volkov et al., 2012; Prokof'ev et al., 2012; Nikolaev et al., 2013; Kolova et al., 2018), Arykvaam (Volkov et al., 2012), Sentyabr'skoe (Nikolaev et al., 2013; Savva et al., 2016),



**Fig. 6.** The temperature–salinity diagram for mineral-forming fluids in the OChVB epithermal gold–silver deposits. (1) Kupol, (2) Dvoinoe, (3) Dzhul'etta, (4) Nyavlenga, (5) Tikhoe, (6) Sentyabr'skoe.

Nyavlenga (Volkov et al., 2014), Dzhul'etta (Prokof'ev et al., 2015), and Tikhoe deposits (Kolova et al., 2015).

The ore-forming fluids of most deposits had temperatures of  $105-359^{\circ}$ C and salinities of 0-9.2 wt %equi NaCl. Higher temperatures (up to  $435^{\circ}$ C) and salinities (up to 37.0 wt %-equi NaCl) have only been recorded for fluids of the late mineralization phase on the Dukat deposit (Goncharov and Sidorov, 1979; Berman et al., 1993; Kravtsova, 2010). This phase is thought to have been related to the influence of a granitoid massif that was emplaced near the deposit. Some deposits were found to involve the action of heterogeneous fluids (Dukat and Sentyabr'skoe); this enables us to estimate the pressure that existed during the mineralization to be in the range 80-570 bars. These pressures are relevant to shallow and medium (subvolcanic) depths of mineralization.

Bulk analysis of the composition of fluid inclusions in quartz monofractions furnished data on fluid compositions for ten OChVB Au–Ag deposits (Table 5, Fig. 7): Dvoinoe, Kupol, Nyavlenga, Pepenveem, Pechal'noe, Dukat, Moroshka, Dal'nee, Dzhul'etta, and Tikhoe. The main components in the fluid include (g/kg H<sub>2</sub>O): carbon dioxide (1.3–58.1), methane (0.02–3.18), chlorine (<0.1–5.15), sulfate ion (<0.3–21.65), hydrocarbonate ion (0–116.51), Na (0.79–23.90), K (0.16–18.79), Ca (0–4.70), and Mg (0–7.02). The following trace elements have been determined (mg/kg H<sub>2</sub>O): Br (0–972.8), As (6.52– 1921.4), Li (2.39–184.42), Be (0–0.19), B (38.85– 7603.0), Rb (0–61.98), Cs (0–26.42), Sr (0–144.36), Mo (0–41.18), Ag (0–15.98), Sb (0.88–903.82), Cu (0–641.75), Zn (0–654.57), Cd (0–0.66), Pb (0– 122.51), Bi (0–0.08), Th (0–0.20), U (0–0.10), Ga (0–0.59), Ge (0–1.01), Ti (0–12.74), Mn (0–1599.9), Fe (0–1772.2), Co (0–10.15), Ni (0–5.46), V (0– 3.84), Cr (0–5.46), Y (0–4.50), Zr (0–1.20), Sn (0– 3.09), Ba (0–33.65), W (0–184.64), Au (0–0.669), Hg (0–1.35), Tl (0–0.85) and REE (0.003–0.49). The ratios of the most typical components vary as follows: Na/K from 0.2 to 15.4, CO<sub>2</sub>/CH<sub>4</sub> from 3.2 to 216.0, and K/Rb from 122.8 to 2725.5.

The analyses for the composition of mineral-forming fluids at the OChVB deposits are consistent with the results of cryometric studies: the fluids have low salinity and low concentrations of chlorine, carbon dioxide, and methane. The anions are dominated by the hydrocarbonate ion and, occasionally, by the sulfate ion, while the cations are dominated by sodium and potassium. Among the trace elements, one notes higher concentrations of Br, As, Li, B, Sb, Zn, and W.

The concentrations of cations and anions show a tendency of increasing K+ from earlier oreless quartz to productive quartz with increasing depth, as well as a slow decrease in the percentage of Na<sup>+</sup>, Ca<sup>++</sup>, and Cl. One observes a direct Ag–K correlation and an inverse

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						Deposits								
lents I	Dukat	Mor	oshka	Dal'	nee	Ι	Dzhul'etta	T			Tikl	hoe		
	Duk-4	KW12-85	KW11-049	MSDL 43531004	MSDL 14481003	EV-1	EV-1	v-2c	Ti-1	304	290	Tikhii	Ti-1	Ti-7
					Macrocomponent	s, g/kg w	ater							
	2.91	3.80	5.74	2.98	4.27	11.23	6.15	6.89	11.84	31.10	24.24	9.24	25.99	19.46
	0.05	0.27	0.17	0.13	0.05	0.31	0.13	0.09	0.26	0.52	0.17	0.34	0.23	0.23
	0.86	<0.5	0.58	2.14	6.13	2.69	0.57	2.52	3.37	2.53	5.15	0.54	3.60	4.17
	<0.2	$\stackrel{\scriptstyle \sim}{\sim}$	<0.5	<0.5	<0.5	<0.3	0.31	<0.2	<0.3	<3	$\leq 1$	<0.5	<0.5	9.22
$\mathbf{D}_3^-$	6.84	18.30	6.57	6.77	10.50	4.11	1.35	0.53	0	84.19	28.42	13.29	12.19	10.20
	2.84	5.27	1.81	1.26	1.52	3.19	0.79	1.20	1.58	25.46	9.86	4.48	69.9	6.81
	0.39	1.24	0.60	0.90	0.89	0.21	0.16	1.09	0.68	14.37	1.66	0.70	0.47	5.32
	0.04	0.24	0.13	1.65	4.70	0	0.002	0	0	0	1.95	0.41	0	0.57
	0.03	0.33	0.29	0.14	0.28	0	0	0.01	0.03	0	0.54	0	0	0.21
				N	ficrocomponents,	10 <sup>-3</sup> g/kg	water							
	0	513.44	152.82	323.75	47.53	125.04	8.02	104.28	10.40	36.53	112.91	322.89	33.45	234.58
	33.15	1072.7	66.55	111.98	183.62	28.32	16.18	26.80	105.15	1216.8	135.80	343.29	880.26	1807.3
	5.18	5.07	6.41	3.02	3.40	4.43	3.03	2.59	2.39	184.42	3.08	4.47	2.62	4.94
	0	0.09	0.10	0	0.05	0.06	0.05	0	0.08	0	0	0.19	0	0
1	43.59	4559.4	2223.0	853.55	896.08	67.18	240.28	68.72	38.85	9676.5	2233.2	4676.7	432.31	434.29
	1.69	3.64	1.46	3.20	3.32	0.16	0	2.20	2.13	17.56	5.20	0	1.15	8.55
	0.77	0.56	0.62	2.47	1.03	0.08	0	0.28	0.08	1.27	4.67	0	0.42	0.27
	1.23	0.26	1.93	36.09	86.07	0	0	0	0.74	0	23.52	0	1.41	56.54
	0.85	26.20	2.62	1.60	1.92	0	0.08	0	0.12	0	41.18	2.52	0.22	2.50
	0	15.98	0	0	0.27	0.13	0	0.89	0.01	0	0	0	0.15	0
	24.38	903.82	32.94	72.36	45.48	168.34	17.02	201.78	11.55	53.48	29.59	38.17	41.73	272.76
	0	0.34	0.03	0	0	1.71	0	0	0.47	0	0	0.02	0	0.34
	0	20.03	0	21.15	0	2.29	0	0	2.33	0	0	0	0	2.49
	0	0.28	0	0	0.08	0.003	0	0.01	0	0	0.26	0.02	0	0.14

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# EPITHERMAL MINERALIZATION

						Г	eposits								
	Elements	Dukat	Mor	oshka	Dal'i	nee	П	Zhul'etta	t			Tikh	106		
		Duk-4	KW12-85	KW11-049	MSDL 43531004	MSDL 14481003	EV-1	EV-1	v-2c	Ti-1	304	290	Tikhii	Ti-1	Ti-7
	Pb	0.01	2.31	0	0	0	0	0.03	3.67	0.23	1.11	0	0	3.53	122.51
	Bi	0	0	0	0	0	0.003	0	0	0.02	0	0	0	0	0
	Th	0	0	0	0	0	0	0	0.05	0.03	0.20	0.09	0	0	0
	Ŋ	0	0	0	0	0	0	0	0.01	0.01	0.10	0	0	0	0
	Ga	0	0.12	0.70	0	0.59	0.10	0	0.19	0.22	0.23	0	0	0.20	0
JOU	Ge	0.05	0.10	0.19	0.13	0.06	0.09	0.18	0	0	0	0.19	0.37	0.17	0.83
RN	Ti	0	0	0	3.50	12.74	0	0	4.29	0	0	0.60	0	0	5.95
AL (	Mn	8.64	1.64	0	29.48	86.02	0	0	0	1.55	0	118.63	0	0.54	10.66
کر DF	Fe	4.30	82.39	0	0	30.10	0	0	8.98	32.55	0	0	0	0	19.36
VOL	Co	0	0	0	0.02	0.10	0	0	0	0.06	0	0.31	0	0.03	0.04
CAN	ïZ	0	0	0.27	0.24	0.66	0.37	0.31	0	0	5.46	0	0.03	0.23	0.30
JOL	>	0	4.50	0.65	0	0	0.10	0.55	0.63	1.55	0.05	0.87	0	0	0
0GY	Cr	0.07	0.61	1.68	0.14	0.16	0.25	0	0.71	0.40	5.46	0	0	0.07	0
AN	Y	0.01	4.50	0.65	0	0	0.01	0	0.06	0.02	0	0.87	0	0	0
ID S	Zr	0	0.04	0.29	0	0	0.03	0	1.20	1.07	0.04	0	0	0.05	0
EIS	Sn	0	0.10	0	0.03	0	0	0	0	1.36	3.09	0	0.04	0.02	0.003
MOI	Ba	0.13	3.43	0.34	1.40	19.12	0.07	0.17	1.17	1.95	33.65	0	0	0	5.89
LOG	M	9.08	11.03	9.35	4.44	1.78	0.01	0	0.25	2.94	33.65	11.48	4.02	0.24	2.44
Y	Au	0	0.191	0.035	0	0	0.052	0.028	0.031	0.203	0.304	0.295	0.054	0	090.0
Vol	Hg	0	0.76	0.19	0.09	0.19	0	0	0.03	0	0	0	0	0	0.01
. 12	IT	0	0.02	0.02	0.10	0.13	0.003	0	0.03	0	0.15	0	0	0.01	0.11
Ν	REE	0.03	0.05	0.12	0.07	0.04	0.003	0.03	0.17	0.05	0.06	0.24	0.14	0.01	0.07
o. 6	Na/K	7.4	4.3	3.0	1.4	1.7	15.4	3.7	1.1	2.3	1.8	5.9	6.4	14.2	1.3
20	$CO_2/CH_4$	63.3	14.3	34.2	22.9	86.7	36.7	46.7	80.1	46.1	60.0	146.7	27.4	113.4	83.6
18	K/Rb	227.7	341.1	408.4	281.9	268.0	1281.9	I	494.7	317.5	818.6	319.6		409.4	621.8

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Table 5. (Contd.)

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**Fig. 7.** The compositions of mineral-forming fluids in the OChVB gold—silver deposits. (a) Dvoinoe, (b) Kupol, (c) Nyavlenga, (d) Pepenveem, (e) Pechal'noe, (f) Dukat, (g) Moroshka, (h) Dal'nee, (i) Dzhul'etta, (j) Tikhoe.

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Ag–Na correlation for productive quartz. The K/Rb ratio for solutions is generally high (occasionally reaching 2725), which may be evidence of an interaction between the ore-forming fluid and the host rocks. One occasionally encounters values below 250, which must most likely have resulted from a magmatic source of the fluid.

The most-pronounced feature that distinguishes the deposits studied here consists in sulfate concentrations in the fluid. One notes deposits with appreciable sulfate concentrations in the fluid (Dvoinoe, Kupol, Nyavlenga, Pepenveem, and Pechal'noe) and deposits with low sulfate concentrations (Dukat, Moroshka, Dal'nee, Dzhul'etta, and Tikhoe).

# CONCLUSIONS

We conclude by noting that the ores of the OChVB epithermal Au–Ag deposits are characterized by an obvious enrichment in a wide range of elements compared with the mean upper crustal values.

Comparative analysis of the trace-element distributions in the ores studied here over classes of gold concentration shows a synchronous enrichment of the ores in similar sets of trace elements and allows indicator elements to be used to determine the type of oreforming fluid.

The patterns identified here, viz., the prevalence of light lanthanoids over heavy ones and positive Eu anomalies, seem to be typical of the epithermal oreforming system. The presence of Eu maxima in the REE distributions of epithermal ores is a characteristic feature of ore columns. The REE distributions of the ores studied here, as well as of the host rocks, are dominated by light "hydrophile" lanthanoids of the "cerium" set.

The epithermal ores studied here were mostly formed by fluids coming from depth (in the lower crust), as well as by solutions that were produced by deep fluids mixed with shallow infiltration waters. The fluid salinity increased toward later, low-temperature phases during the mineralization process. The oreforming solutions are of the hydrocarbonate potassium type, having 60-80% hydrocarbonate and potassium of the total amount of cations and anions. The concentration of anions and cations shows a tendency of the potassium percentage to increase from the earlier oreless quartz toward productive quartz with increasing depth, as well as a slow decrease in the percentages of Na<sup>+</sup>, Ca<sup>++</sup>, and Cl<sup>-</sup>. One observes a direct Ag-K correlation and an inverse Ag-Na correlation for productive quartz. The most-pronounced feature of the deposits studied here consists in sulfate concentrations in the fluid.

The results indicate magma chambers of andesite magmas and meteoric waters as the most likely sources

of the fluids that went to form the epithermal Au–Ag ores in the OChVB deposits.

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