

FORMATION OF SULFATES AT THE THIAPHES AREA OF MILOS ISLAND: POSSIBLE PRECURSORS OF KAOLINITE MINERALIZATION

AKIS E. KELEPERTSIS

Department of Geology, National University of Athens, Panepistimiopolis, Ano Ilisia, GR-15784 Athens, Greece

ABSTRACT

The physicochemical conditions at the Thiaphes area of Milos Island are favorable for the recent formation of alunogen, alunite, natroalunite, melanterite, gypsum, sulfur, sylvite, quartz, cristobalite and kaolinite. The local environment is characterized by gas emissions rich in H₂S and CO₂, thermal waters enriched in SO₄ and Na by mixing with seawater, porous and permeable alluvial aluminosilicate-rich soils, and the presence of atmospheric O₂. Alterations started in the Quaternary in an acid environment and continue up to the present day in areas of H₂S-rich fluids. The hydrated sulfates represent precursors of the very common kaolinite mineralization on the island. Kaolinite was also formed by the hydrolysis of feldspars.

Keywords: hydrothermal alteration, sulfates, kaolinite, acid environment, Milos, Greece.

SOMMAIRE

Le milieu physico-chimique dans la région de Thiaphes, sur l'île de Milos, en Grèce, est propice à la formation récente de l'association de alunogène, alunite, natroalunite, mélanterite, gypse, soufre, sylvite, quartz, cristobalite et kaolinite. Des émissions gazeuses riches en H₂S et CO₂ caractérisent ce milieu. Sont aussi importants les eaux thermales, enrichies en SO₄ et Na suite au mélange avec l'eau de mer, un sol alluvionnaire poreux, perméable, riche en aluminosilicates, et la présence de l'oxygène atmosphérique. Les phénomènes d'altération ont débuté au Quaternaire dans un milieu acide, et continuent aujourd'hui dans les endroits enrichis en H₂S. Les sulfates hydratés seraient les précurseurs de la kaolinite, qui est répandue sur l'île. L'hydrolyse des feldspaths a aussi produit de la kaolinite.

(Traduit par la Rédaction)

Mots-clés: altération hydrothermale, sulfates, kaolinite, milieu acide, Milos, Grèce.

INTRODUCTION

The island of Milos is located in the central part of

the South Aegean active volcanic arc (Fig. 1). Only a small exposure of metamorphic rocks occurs at the southern part of the island; these rocks belong to a flysch formation (phyllites) associated with ophiolites and allochthonous blocks of limestones (Fytikas 1977). Most of the island consists of volcanic rocks (Fig. 2), which are part of the southern Aegean island arc, and which formed during the Pliocene as a consequence of the northward subduction of the African plate beneath the Aegean plate (Fytikas *et al.* 1984). The volcanic rocks belong to a calc-alkaline series of andesites and rhyolites accompanied by tuffs, ignimbrites and pyroclastic rocks.

On Milos, there is a high-enthalpy geothermal field (Fytikas & Marinelli 1976). The heat flow was responsible for intense hydrothermal activity, which caused widespread bentonitization and alunization, kaolinization and the formation of various hydrothermal mineral deposits (kaolinite, bentonite, sulfur, barite, galena, alunite and manganese oxides). Kaolinite deposits are especially large; most of them are being exploited. Kaolinite is used as raw material in the paper and cement industries.

Fumaroles (up to 102°C), submarine gas leakage (>50°C), thermal springs (up to 75°C) and hot grounds (up to 100°C) are some surface thermal manifestations. At the sites of altered volcanic rocks, active mud volcanos, lignitic deposits, and volcanic emanations (Velinov *et al.* 1970, Baltatzis *et al.* 1986), hydrous sulfate (*e.g.*, rozenite, melanterite, römerite, halotrichite, epsomite and alunite) are formed.

This paper reports on the mineralogy and the conditions of formation of sulfate minerals and other secondary products, especially kaolinite, in the Thiaphes area near the village Adamanda (Fig. 2).

GEOLOGY OF THE THIAPHES AREA

The area in which humid-hot soils and gas exhalations occur is approximately 1 km northeast of Adamanda; it is known as Thiaphes, meaning "sulfur concentrations". The soils are unconsolidated alluvial deposits consisting of pebbles, sands, silt and clay derived by weathering of the surrounding geological formations, such as tuffs, rhyolitic and

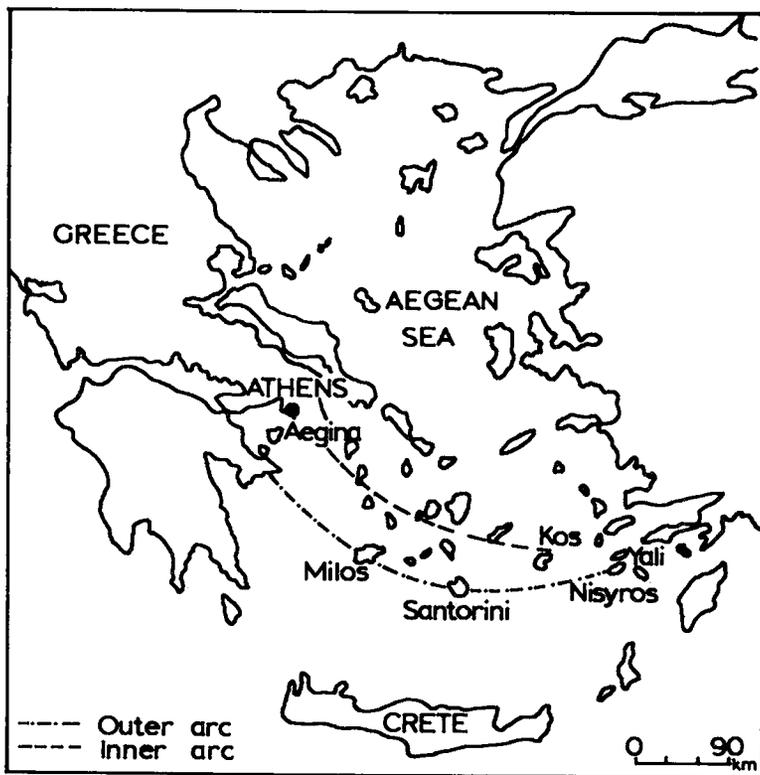


FIG. 1. Map of Greece showing Milos Island and the active volcanic arc.

rhyodacitic lavas, and lahars, which are younger than Upper Pliocene (Fytikas 1977).

Where gas emanations take place, the soils are altered and covered with thin layers of secondary minerals that make the soils yellowish, greenish, white and brownish black. The soils are damp, probably as a result of the effect of the underlying geothermal field, whose presence has been confirmed by drilling of the surrounding alluvial deposits.

ANALYTICAL METHODS

Powder X-ray diffractometry was used to identify the hydrated sulfate aggregates and the associated material. A Philips PW1010 diffractometer was operated with Ni-filtered $\text{CuK}\alpha$ radiation at 36 kV and 24 mA, at the Department of Geology, National University of Athens. The samples (10) were also analyzed by atomic absorption spectrometry for K, Na, Al, Fe, Ca, Co, Ni, Pb, Zn, Cu and Sr. X-ray-diffraction data and chemical results are shown in Tables 1 and 2, respectively.

RESULTS

Mineralogy

The minerals identified by X-ray diffractometry include alunogen, melanterite, gypsum, natroalunite, alunite (all sulfates), quartz, cristobalite, sulfur; halite, and sylvite. Of the two forms of silica, cristobalite is the newly formed mineral present, whereas part of the quartz may be of detrital origin. The aluminosilicates are represented by kaolinite; halite and sylvite also are present.

Chemistry

The chemical composition of the sulfate-coated aggregates analyzed is quite simple and is as expected from the mineral determinations; concentrations of the major elements (Al, Ca, K, Na, Fe) correlate well with the mineral assemblages. For example, the sample with high gypsum content (M5) is characterized by a high Ca content; the presence of alu-

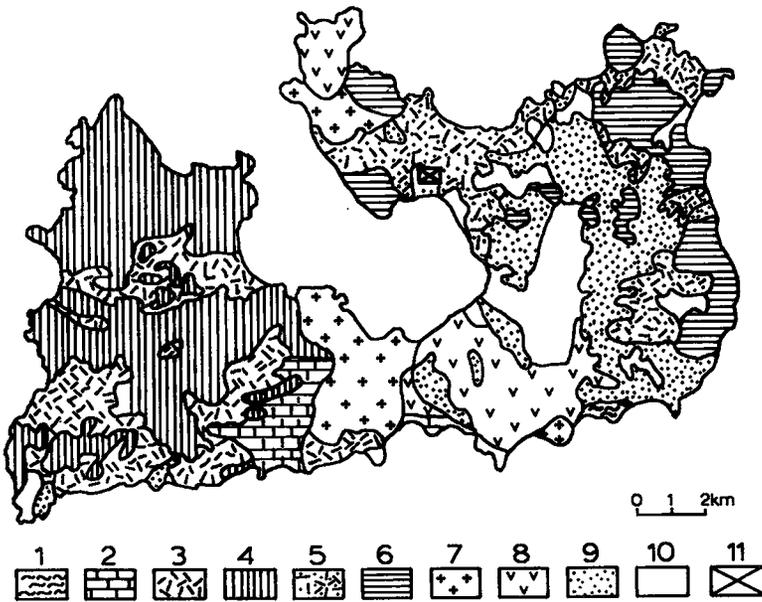


FIG. 2. Geological sketch map of Milos Island (from Fytikas *et al.* 1986), with location of the studied area. Symbols: 1 metamorphic basement, 2 Neogene sediments, 3 basal pyroclastic series (Middle-Upper Pliocene), 4 complex of domes and lava flows (Upper Pliocene), 5 pyroclastic rocks, 6 lava domes, 7 Halepa and Plakes domes, 8 rhyolitic complexes (Upper Pleistocene), 9 products of phreatic activity, 10 Quaternary sediments, 11 study area.

nogen $\text{Al}_2(\text{SO}_4)_3 \cdot 17\text{H}_2\text{O}$, alunite $\text{KAl}_3(\text{SO}_4)_2(\text{OH})_6$, natroalunite $\text{NaAl}(\text{SO}_4)_2(\text{OH})_6$ and kaolinite $\text{Al}_2\text{Si}_2\text{O}_5(\text{OH})_4$ is reflected by the high Al contents in the samples. Sodium and potassium are present in alunite, natroalunite, halite and sylvite. The high Fe-contents in the samples reflect the presence of melanterite ($\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$), the only iron sulfate detected. Ni, Pb, Zn concentrations are never anomalous, suggesting that other sulfate minerals are absent. Cu, Co and Sr were not detected in the samples, except for a Sr value of 230 ppm in a gypsum-rich specimen. All these results suggest that the altered parent soils, which were affected by the hydrothermal solutions and H_2S gases, were low in trace elements (Cu, Pb, Zn, Ni), and this is reflected by their mineralogical composition. The alluvial soils consist of particles derived from the alteration of the surrounding geological formations, which are mainly white tuffs and hydrothermally altered (mainly kaolinized) volcanic rocks. In general, the soils are composed of Si-, Al-, Fe-, K- and Na- bearing silicates. Part of the Na is attributed to contamination of the hydrothermal solutions by seawater.

DISCUSSION: FORMATION OF ALTERATION MINERALS

Hydrous sulfate minerals typically form in the oxidation zone of sulfide ores (Velinov *et al.* 1970, Good 1971, Zodrow *et al.* 1979), but also have been found in areas of volcanic activity (Stamatakis *et al.* 1987). From field observations and published data (Fytikas 1977), the chemical environment at the Thiaphes area is favorable for the formation of hydrated sulfate minerals. This environment is characterized by H_2O , H_2S and CO_2 emanations and by high humidity from the geothermal field. The SO_4 contents of thermal waters that apparently penetrate the permeable alluvial soils are high and hence pH is low (4.0).

The hydrous sulfates at the Thiaphes area of Milos must be the alteration products formed by the reaction of H_2S with alluvial soils in a humid microenvironment of low pH. The abundances of Al-sulfate minerals (alunogen, natroalunite, alunite) reflect the existence of Al-rich minerals in the soils. The occurrence of sulfur in thin layers in the surface around

TABLE 1. ASSEMBLAGES OF MINERALS, AS DEDUCED FROM XRD ANALYSES

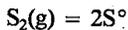
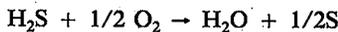
Sample	Mineralogy
M1	sulfur, alunogen, cristobalite
M2	alunogen, melanterite, halite, kaolinite, natroalunite
M3	sulfur, halite, sylvite, quartz, alunite
M4	alunogen, melanterite, halite, kaolinite, natroalunite, sylvite
M5	cristobalite, quartz, gypsum
M6	alunogen, cristobalite, quartz, gypsum
M7	alunogen, cristobalite, quartz
M1-1	sulfur, cristobalite
M4-1	alunogen, cristobalite, melanterite, natroalunite, sylvite, quartz
M5-1	sulfur, cristobalite, quartz

TABLE 2. CHEMICAL COMPOSITION OF THE MINERAL AGGREGATES

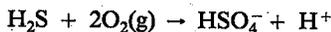
Sample	Na	K	Al	Fe	Ca	Ni	Pb	Zn	Sr	Cu
M1	0.062	0.03	0.10	0.07	0.01	467	0	124	0	0
M2	0.18	0.54	7.94	0.70	0.01	37	0	65	0	0
M3	0.13	3.38	3.73	0.38	0	29	87	53	0	0
M4	0.14	0.77	8.76	0.54	0	43	0	82	0	0
M5	0.70	0.73	1.91	0.20	15.26	49	34	49	230	0
M6	0.97	0.99	3.59	0.96	0.09	46	46	87	0	trace
M7	1.47	1.58	5.00	1.34	0.17	49	35	84	0	0
M1-1	0.58	0.79	1.81	0.29	0.02	467	55	89	0	0
M4-1	1.17	1.51	7.84	1.53	0.01	63	78	122	0	0
M5-1	0.50	0.61	1.30	0.30	0.02	24	53	92	0	0

Concentrations of Na, K, Al, Fe and Ca are expressed in Wt.%; those of Ni, Pb, Zn, Sr and Cu are expressed in ppm.

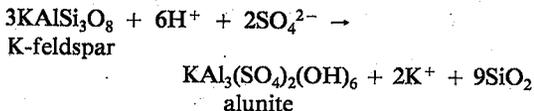
the H₂S emanations is due to the direct oxidation of S²⁻ to S⁰ in the atmosphere:



Where the solutions are subjected to a higher oxygen fugacity, the oxidation of H₂S leads to the formation of HSO₄⁻ according to the reaction:

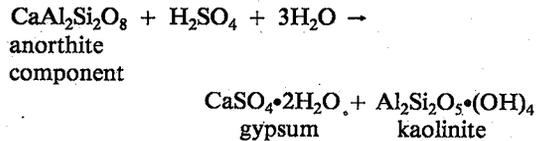


This process causes an increase of the acidity in the moist alluvial soil, which favors the release of metals such as Al, Si, K, Na, Ca from the aluminosilicate minerals and the subsequent formation of sulfates. For example, the formation of alunite from K-feldspar is represented by the following reaction:



The presence of alunite in the mineral aggregates suggests the alteration of acid volcanic material in an environment of high H⁺ and SO₄²⁻ activity (Hemley & Jones 1964). This proposal is consistent with the composition of the thermal waters on Milos (Fytikas 1977).

The formation of gypsum occurs by the transformation of the anorthite component of plagioclase according to the reaction:



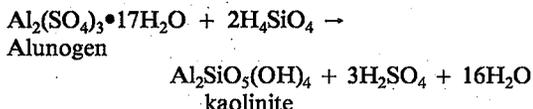
As mentioned before, on Milos there are widespread gaseous emanations, thermal springs, barite and sulfur deposits, etc., as well as many kaolinite deposits. Most of these deposits are aligned along fractures that formed during recent tectonic activity and that favored the surfacing of gases and hydrothermal solutions. These solutions consist of resurgent heated meteoric water mixed in some way with seawater.

Kaolinization, alunization and silicification phenomena had their maximum development in proximity of the high-thermal-gradient sites, as is shown on maps of thermal anomalies (Fytikas 1977). These anomalies, together with the widespread recent volcanic activity on the island, suggest the existence at depth of magmatic bodies, which represent the source of energy for the processes of alteration and mineralization.

The alteration processes are still active today near Thiaphes, giving rise to silica minerals, sulfates, sulfur, chlorides and aluminosilicate minerals (kaolinite). The mineralogy of the kaolinite mineralization includes amorphous phases of silica, cristobalite, quartz, alunite and natroalunite. Gypsum, pyrite and sulfur also are present in a few deposits. A comparison in the mineral associations between the present-day aggregates and the kaolinite deposits shows that sulfate minerals like alunogen and melanterite are absent in the deposits. This absence is reflected in their color: the kaolinized masses have a whitish color, but the recently altered masses of the Thiaphes area are impure, owing to silica minerals, sulfates and iron oxides. In addition, the relative proportions of sulfate minerals to kaolinite are higher in the present-day aggregates than in the kaolinite deposits. In the Kastriani kaolinite mine (Kelepertsis *et al.* 1988), the proportions of minerals are: 41% kaolinite, 44% cristobalite and amorphous phases, 6% quartz, 2% undetermined minerals and 7% alunite.

Based on the field observations and the above data, two processes of alteration seem to be responsible for the formation of kaolinite: (a) hydrolysis of feldspars and muscovite under conditions of low pH and humidity, and (b) development of early hydrated sulfate facies into sulfates having a higher structural order and a lower water content. These eventually were transformed into kaolinite and other minerals

according to the following reaction:



CONCLUSIONS

Fluids at the surface at the Thiaphes area of Milos Island are represented by gases (CO_2 and H_2S) and hydrothermal solutions enriched in SO_4^{2-} and Na^+ , possibly mixed with seawater. Deep magmatic bodies represent the source of energy for the upward migration of the fluids.

The formation of the sulfate minerals is favored by H_2S -bearing gases emanating at the surface, the low pH of the thermal waters and the high porosity and permeability of the alluvial soils composed of aluminosilicate minerals.

The present-day aggregates consist of alunogen, melanterite, gypsum, alunite, natroalunite, halite, sylvite, sulfur, quartz, cristobalite and kaolinite. According to these studies, the early hydrated sulfate phases may represent the precursors of the minerals in the paragenesis of the late Milos kaolinite deposits. The precursors eventually developed into sulfates with a higher structural order and kaolinite. Kaolinite was also formed by the alteration of feldspars.

ACKNOWLEDGEMENTS

The author thanks Mr. G. Samiotakis, chemist of the Silver-Barytine Ores Company, Greece, for the chemical analyses, Prof. S. Augoustidis for reading the final manuscript, Prof. R.D. Schilling and an anonymous referee for their helpful comments on the manuscript of this paper.

REFERENCES

BALTATZIS, E., STAMATAKIS, M.G. & KYRIAKOPOULOS, K.G., (1986): Rozenite and melanterite in lignitic layers from the Vóras mountain, western Macedonia, Greece. *Mineral. Mag.* **50**, 737-739.

FYTIKAS, M. (1977): *Geology and Geothermics of Milos Island*. Ph.D. thesis, Thessaloniki Univ., Thessaloniki (in Greek with English summary).

_____, INNOCENTI, F., KOLIOS, N., MANETTI, P., MAZZUOLI, R., POLI, G., RIFA, F. & VILLARI, L. (1986): Volcanology and petrology of volcanic products from the island of Milos and neighbouring islets. *J. Volcanol. Geotherm. Res.* **28**, 297-317.

_____, _____, MANETTI, P., MAZZUOLI, R., PEC-CERILLO, A. & VILLARI, L. (1984): Tertiary to Quaternary evolution of volcanism in the Aegean region. In *The Geological Evolution of the Eastern Mediterranean* (J.E. Dixon & A.H.F. Robertson, eds.). *Geol. Soc. London, Spec. Publ.* **17**, 687-699.

_____, & MARINELLI, G. (1976): Geology and geothermics of the island of Milos (Greece). *Proc. Geothermal Energy* **1**, 516-524.

GOOD, R. (1971): Melanterite - rozenite - pyrrhotite occurrence in Carroll Country, Virginia. *Virginia Minerals* **17**, 10.

HEMLEY, J.J. & JONES, W.R. (1964): Chemical aspects of hydrothermal alteration with emphasis on hydrogen metasomatism. *Econ. Geol.* **59**, 538-569.

KELEPERTSIS, A.E., SKOUNAKIS, S. & ECONOMOU, K. (1988): Kaolins of Milos island - their industrial use. Report submitted to the Ministry of Research and Technology, Greece.

STAMATAKIS, M.G., BALTATZIS, E. G. & SKOUNAKIS, S.B. (1987): Sulfate minerals from a mud volcano in the Katakolo area, western Peloponnesus, Greece. *Am. Mineral.* **72**, 839-841.

VELINOV, I., ASLANYAN, S., PUNEV, L. & VELINOVA, M. (1970): Ferratosulfates halotrichite and alunogen from the oxidized zones of the hydrothermally altered volcanic rocks, near the village of Krusha, district of Sofia. *Bull. Geol. Inst. Bulgarian Acad. Sci., Ser. Geochem. Mineral. Petrogr.* **19**, 243-264.

ZODROW, E.L., WILTSHIRE, J. & MCCANDLISH, K. (1979): Hydrated sulfates in the Sydney coal field of Cape Breton, Nova Scotia. II. Pyrite and its alteration products. *Can. Mineral.* **17**, 63-70.

Received April 5, 1988, revised manuscript accepted July 31, 1988.