

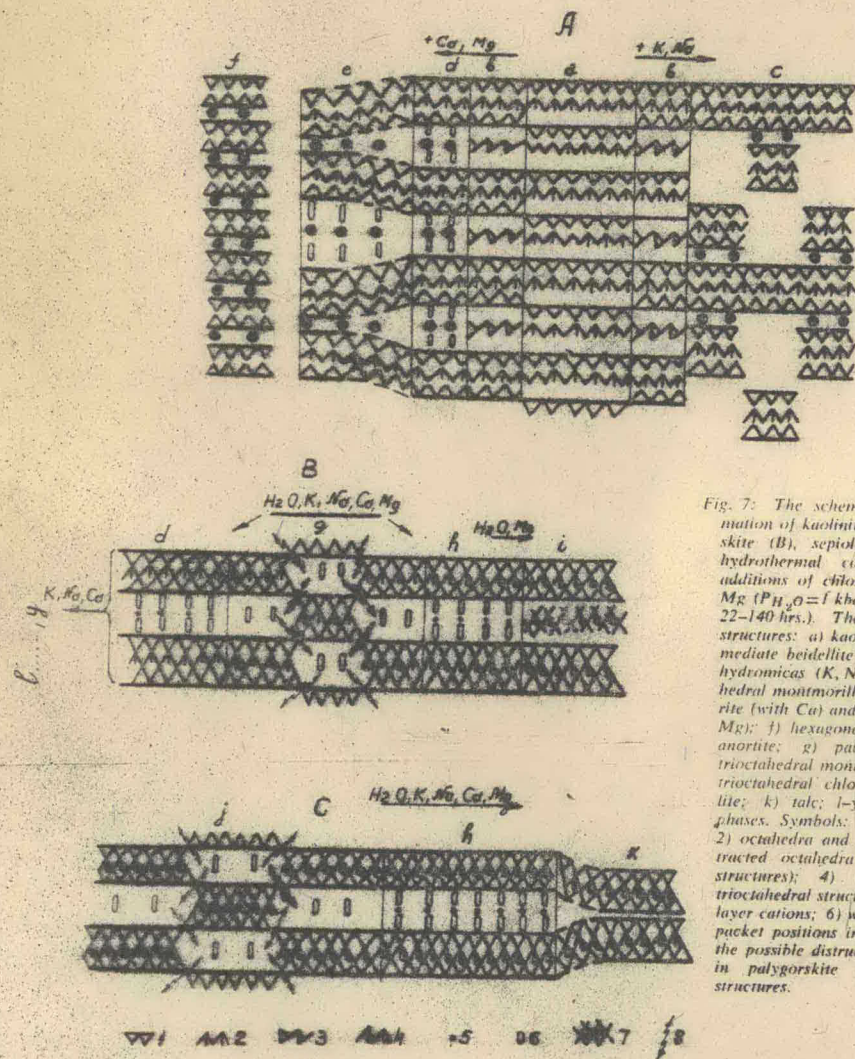


Fig. 7: The scheme of transformation of kaolinite (A), palygorskite (B), sepiolite (C) under hydrothermal conditions with additions of chlorides K, Na, Ca, Mg ($P_{H_2O} = 1$ kbar, 200–500°C, 22–140 hrs.). The schemes of structures: a) kaolinite; b) intermediate beidellite like phase; c) hydronicas (K, Na); d) diocahedral montmorillonite; e) rectorite (with Ca) and tosudite (with Mg); f) hexagonal analogue of anorite; g) palygorskite; h) triocahedral montmorillonite; i) triocahedral chlorite; j) sepiolite; k) talc; l-y) non-layered phases. Symbols: 1) tetrahedra; 2) octahedra and 3) partial disordered octahedra (diocahedral structures); 4) octahedra in triocahedral structures; 5) interlayer cations; 6) water; 7) interpacket positions in chlorites; 8) the possible dislocation positions in palygorskite and sepiolite structures.

montmorillonite type are at first formed, and finally micas of muscovite or paragonite composition are formed. In the presence of Ca and Mg-chlorides, kaolinite changes to an intermediate montmorillonite phase, followed by ordered mixed-layered phases, 1:1 of mica-montmorillonite (Ca) type and chlorite-montmorillonite (Mg).

By structural transformations of palygorskite under pressure of H_2O -vapour and also with $MgCl_2$ dis-

ordered mixed-layered phases of chlorite-montmorillonite type are formed. The process of transformation, described above and the process of which was discussed before [5, 7], can be schematically summarized as shown in Table 1.

The principal features of this process are:

1. The varieties of newly formed layered silicates keep their di- and tri-octahedral structural types.

2. Transformations, as a rule, have the intermediate smectite stage (montmorillonite-beidellite type) and their crystallochemical character determines the final phase.
3. Under the hydrothermal conditions and relatively low temperatures, depending on chemical compositions, ordered and disordered mixed-layered phases are formed as intermediate phases.
4. The positions of the fields of smectites, the mixed-layered and stable final phases (Table 1) probably show most common features of structural polymorphous conditions.

Conclusion

It can be said that the above mentioned results can be used for the estimation of some peculiarities of the processes of epigenesis and the first stage of metamorphism. Typical is the transformation mechanism of the process and a wide development of mixed-layered formations. Their nature is principally connected not with thermodynamical parameters (kinetics process), but with the chemistry of the surroundings. Disordered mica-montmorillonite structures are formed by the presence of K and Na in solutions, while Ca and Mg lead to the formation of ordered chlorite-montmorillonite (tosudite) and mica-montmorillonite (rectorite) formation. This process is characterized by the transformation mechanism and the formation of the intermediate smectite phase.

A number of data about the natural metamorphism of clay minerals [21, 22, 23] all allows to establish many common features with experimental data discussed above. New investigations of metamorphism of natural clay minerals in details and laboratory experimental investigations in this field will allow to define more accurately and concretely the conception of diagenesis and initial metamorphism of sedimentary rocks.

REFERENCES

1. Frank-Kamenetskij V.A. Strukturtypomorphismus und moderne Mineralogie. Berl. deutsch. Ges. geol. Wiss. B. Miner. Lagerstätten, 13, 3, 1968, s. 305, Berlin.
2. Tuttle O.F. A new hydrothermal quenching apparatus. Amer. Journ. Sci., 246, 1948, p. 628.
3. Бриджмен П.В. Физика высоких давлений. Гостехиздат, 1935.
4. Гойло Э.А., Котов Н.В., Франк-Каменецкий В.А. Экспериментальное исследование влияния стрессового и гидростатического давлений при различных температурах на кристаллическую структуру каолинита. Сб. Физические методы исследования минералов осадочных пород, "Наука", 1966, стр. 123.
5. Франк-Каменецкий В.А., Котов Н.В., Гойло Э.А. Гидротермальный синтез в системах галлуазит и каолинит III хлориды K, Na, Ca, Mg под давлением. Сб. Проблемы петрологии и генетической минералогии, "Наука", 1969, стр. 143.
6. Frank-Kamenetskij V.A., Kotov N.V., Goylo E.A. Strukturänderung von Tonmineralen unter verschiedenen thermodynamischen Bedingungen. 1. Kristall und Technik, Bd. 1, Hf. 3, 1966, s. 465.
7. Frank-Kamenetskij V.A., Kotov N.V., Goylo E.A. Strukturveränderungen von Tonmineralen bei verschiedenen thermodynamischen Bedingungen. 2. Kristall und Technik, Bd. 3, Hf. 4, 1968, s. 643.
8. Frank-Kamenetskij V.A., Kotov N.V., Schitov V.A., Goylo E.A. Strukturveränderungen von Tonmineralen bei verschiedenen thermodynamischen Bedingungen. 3. Kristall und Technik, Bd. 5, Hf. 1, 1970, s. 129.
9. Франк-Каменецкий В.А., Котов Н.В., Ключкова Г.Н. Фазовые превращения сепиолита и палыгорскита при разных давлениях в гидротермальных условиях. Геохимия, № 1, 1969, стр. 14.
10. Франк-Каменецкий В.А., Котов Н.В., Ключкова Г.Н. Фазовые превращения сепиолита и палыгорскита в гидротермальных условиях в присутствии хлоридов K, Na под давлением. Геохимия, № 1, 1970, стр. 14.
11. Barrer R.M., Denny P.I. Hydrothermal chemistry of the silicates. Part X. A partial study of the field $CaO-Al_2O_3-SiO_2$. Journ. chem. Soc., 3, 1961, p. 983.
12. Goldsmith J.R., Hiles E.G. The stability relations of anorite and hexagonal polymorph in the system $CaAl_2Si_2O_8-H_2O$. Journ. Geol., vol. 60, 4, 1952, p. 386.
13. Takeuchi Y., Donnay G. The crystal structure of hexagonal $CaAl_2Si_2O_8$. Acta Cryst., 12, 1959, p. 465.
14. Камаев А.А. О кристаллической структуре кимрита. Докл. АН СССР, сер. геол., т. 169, № 1, 1966, стр. 201.
15. Brauner K., Preisinger A. Structure of sepiolite. Miner. Petrogr. Mitt., vol. 6, 1956, s. 120.
16. Bradley W.F. Structure of attapulgite. Amer. Miner., vol. 25, 1940, p. 405.
17. Ключкова Г.Н. Сеполиит из Карамагара. Докл. АН Тадж. ССР, т. 11, № 9, 1968, стр. 45.
18. Ключкова Г.Н. К характеристике Памирского палыгорскита. Докл. АН Тадж. ССР, т. 11, № 8, 1968, стр. 36.
19. Green-Kelly R. The identification of montmorillonoids in clays. Journ. Soil. sci., vol. 4, 2, 1953, p. 233.
20. Sudo T., Kodama H. Aluminian mixed-layer montmorillonite-chlorite. Z. Krist., vol. 109, 1957, p. 379.
21. Sudo T. Mineralogical study on clays of Japan. Maruzen Co. Ltd., Tokyo, 1959.
22. Frank-Kamenetskij V.A., Logvinenko N.V., Drits V.A. Tosudite - a new mineral forming the mixed-layer phase 1:1 alusite. Int. clay conf., 1963, vol. 2, 1965, p. 181.
23. Франк-Каменецкий В.А., Логвиненко Н.В., Дриц В.А. О тосудите и алузите. Минер. сб. Львовск. гос. университета, вып. 1, № 22, 1968, стр. 70.
24. Милло Ж. Геология глин. "Недра", 1964 (русск. перевод 1968).
25. Коссовская А.П. Типизация и генетическое значение смешанных дольных минералов глин. Сб. Физические методы исследования минералов осадочных пород, "Наука", 1966, стр. 163.