

obviously be sought in the large displacements which take place during the transition of the shock wave from a medium with a given dynamic rigidity to another and which stimulate the process of phase transformation.

The crystalline structure of some samples was also analyzed with an electron microscope of the type UEMV-100, equipped with an electronograph. The results of these investigations are in full agreement with the x-ray measurements.

The interpretation of the diffractograms could be made on the basis of an orthorhombic lattice, which is more or less close to the one described in [8] where a new crystalline modification of  $\text{TiO}_2$  was synthesized in static conditions; the orthorhombic lattice of this modification belongs to the space group  $Pbcn$ , while the elementary cell contains in itself four molecules of  $\text{TiO}_2$ . The new phase of silica has the following parameters:  $a = 4.30$ ,  $b = 4.70$ ,  $c = 4.50 \text{ \AA}$ . The values of the interplanar distances are  $d_{ca1}$ , calculated from these parameters, satisfactorily coincide (see Table 1) with the observed values  $d_{obs}$ .

In the above-mentioned work, there is no detailed calculation of the intensities of the new-phase lines of  $\text{SiO}_2$ , but a comparative evaluation of the relative value of the diffraction maxima of the diffractograms and the relative intensities of the new phase of  $\text{TiO}_2$  reveals a

satisfactory agreement between the two. The crystal density, calculated from the lattice parameters, is equal to  $4.435 \text{ gm/cm}^3$ . We may note here that, due to the fairly strong broadening of the diffraction maxima and also due to the absence of diffraction maxima with large values of  $hkl$ , the accuracy of determination of the lattice parameters, did not exceed  $0.01\text{\AA}$ .

In conclusion, we must remark that the complex shock-wave process of sample loading and the subsequent removal of pressures from it, naturally, does not justify the assertion that the formation of the orthorhombic modification of silica takes place at the top of application of shock pressures to the  $\text{SiO}_2$  sample. As was already observed, especially in [15], one cannot rule out the possibility that the orthorhombic phase is a consequence of the transformation of silica from an unknown high-density phase (possibly from a fluorite type of structure), which forms after repeated compressions and which seems to be unstable in normal conditions.

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