

HIGH PRESSURE EXPERIMENTAL STUDY ON GROWTH AND RESORPTION OF DIAMOND IN KIMBERLITE MELT

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Studies of inclusions in natural diamonds suggest a genetic link between diamond formation and volatile- and alkali-rich mantle melts or fluids, broadly similar to kimberlite composition (Navon et al., 1988). Arima et al. (1993) demonstrated that diamond crystallized and grew in a volatile-rich silicate melt of kimberlite composition in high-pressure and high-temperature experiments (1800-2200°C and 7-7.7 GPa). This paper reports detailed morphological features of newly crystallized diamonds in the experiments by Arima et al. (1993). In addition, resorption (dissolution) of diamond in kimberlitic melts was studied experimentally in the graphite stability field (1300-1500°C and 2.5 GPa) under the W-I condition. This paper also reports a variety of resorption features observed on the seed diamonds.

Crystallization and growth of diamond at 7 - 7.7 GPa

Detailed experimental procedures were given in Arima et al. (1993). The kimberlite used in the experiments is an aphanitic kimberlite from Wesselton Mine, South Africa. In the runs at 1800-2200° C and 7-7.7 GPa, the kimberlite powder was loaded in a graphite capsule with and without a seed diamond (about 1 mm in diameter), which was emplaced in a Mo capsule. Diamond crystallized and grew from kimberlite melt in all the run. Newly formed diamonds occur in the interface between the kimberlite and graphite capsule, indicating the kimberlite acts as the solvent-catalysts. No systematic difference exists between the runs with seed diamond and those without seed diamond at the same P-T condition. No diamond was observed in the "dummy" run in which graphite was encapsulated in a Mo capsule without kimberlite powder. The kimberlite part of the run products consists of a fine-grained crystal aggregate. No glass was observed. Dendritic or acicular morphology of the crystals indicates that they are quench phases crystallized from the kimberlitic melt during quenching at high pressure. Overall, the results indicate that the diamond crystallized from kimberlitic melt and was not formed by solid-state transformation of graphite.

The newly formed diamond crystals, up to 100 μm in diameter, are colorless and transparent under optical microscope. They are isolated well faceted octahedron, hopper crystals, or dendritic crystals. The newly crystallized diamonds generally have well developed {111} face. In a rare case they have small rugged {100} face. Octahedron twinned on {111} twin plane is common. In a rare case, cyclic twins are present. The morphology of newly formed diamond in the present experiment indicates that the growth rate of {111} is several times lower than those of {100}, being resemble those of natural diamond but contrast to those of synthetic diamonds grown from metal solvent-catalysts. In addition, in the runs at 2000-2200°C, a depression at the center of the {111} face is locally present, and growth steps indicative of layer-growth are clearly observed in it.

No apparent change was noted in the original habits of the seed diamonds. However, all the seed diamonds had newly formed triangular patterns on the {111} faces. They correspond to growth hillocks that have rounded triangular, truncated triangular, or triangular shape, and have the same orientation as the original triangular {111} faces of the seed crystal. In a rare case, growth pyramids (~0.74 μm high) of triangular morphology clearly show growth steps similar to those observed on natural diamonds (Sunagawa, 1984; van Enckevort, 1992). These features suggest a spiral growth mechanism of the hillocks. It is likely that carbon atoms diffused from the graphite capsule through the kimberlite melt and deposited as diamond on the seed crystal.

Experiment on resorption of diamond at 2.5 GPa

Resorption of diamond in kimberlitic silicate melts was investigated at 2.5 GPa and 1300-1500° C for 10 to 120 min. using a piston cylinder high pressure apparatus. A natural diamond crystal and a resorption agent (kimberlite powder) were emplaced in a sealed inner Pt capsule. The inner sample capsule was then loaded with the W-I buffer powder in a sealed outer Pt capsule. The natural diamonds used are smooth-faced octahedral crystals (1 mm in diameter) with slightly resorbed rounded corner and edge. To examine the effect of carbon (or CO₂) concentration in the melt on the diamond resorption, kimberlite compositions with variable carbon (or CO₂) concentrations were prepared. They were (1) the natural group 1 kimberlite composition from Wesselton Mine, South Africa (4.77 wt.% CO₂ and 6.2 wt.% H₂O), (2) the graphite-doped compositions in which graphite powder (20 wt.%) was added to the Wesselton kimberlite, (3) the carbonate-doped compositions in which CaCO₃ (30 and 50 wt.%) was added to the natural kimberlite, and (4) the synthetic kimberlite compositions in which SiO₂, Al₂O₃, TiO₂, MnO, MgCO₃, CaCO₃, Na₂CO₃, and K₂CO₃ were mixed with Fe, FeO, or Fe₂O₃. No H₂O was added to these synthetic compositions.

All diamonds in the run-products exhibited a variety of resorption features on {111} face including both negatively and positively oriented trigons, hexagonal deep pit, and/or shallow depression with hexagonal morphology. In addition, in most of the run-products, except for those with graphite-doped kimberlite, the diamonds showed preferential resorption of corners and edges of octahedron. In the runs with the natural kimberlite and carbonate-doped natural-kimberlite compositions, extensive resorption occurred and no octahedral faces were preserved after 120 min. In those run, knob-like asperities were noted on corners and edges. Kimberlite portions of the run-products were composed of dendritic quench-crystal aggregates. No graphite was present in all the runs, except for those with graphite-doped kimberlite composition.

All diamonds in the runs-products with natural-kimberlite and carbonate-doped natural-kimberlite compositions showed negatively oriented trigon. A majority of the pits are sharply edged trigons but some shallow pits are in the rounded triangular shape with rounded bottom surface. The trigons include the types of pyramidal-shallow, pyramidal-deep, flat-bottomed-shallow, and flat-bottomed-deep, as classified

by Frank and Lang (1965). Small trigons ($< 1 \mu\text{m}$) of both flat-bottomed and pyramidal exists within the relatively large trigons. The size of trigon clearly increases with run

temperature and run duration. Addition of carbonate to the kimberlite composition suppressed the resorption of diamond.

In the runs with graphite-doped kimberlite composition, the diamonds exhibited no noticeable change in the original habit. No deep-trigon was observed in those runs. The diamonds were covered by graphite and showed shallow depressions with hexagonal morphology on $\{111\}$ face but their corner and edge of octahedron were well preserved.

In the runs with synthetic-kimberlite compositions, both negatively and positively oriented trigons were formed on $\{111\}$ face. In some runs, sharply edged hexagonal deep pits were present on $\{111\}$ face. The run with a melt containing iron of higher oxidation state showed higher degree of resorption features. For example, in the run with Fe_2O_3 -doped synthetic composition at 1500°C and 2.5 GPa for 10 min, corners and edges of octahedron were extensively resorbed and only 50 % of area of $\{111\}$ face was preserved. To the contrary, in the run with FeO -doped synthetic composition at the same run condition, only 25 % of area of $\{111\}$ face was resorbed, and no apparent change was noted in the original habit of the diamond of the run with Fe-doped synthetic composition.

Resorption of diamond is thought to occur within transporting kimberlite magma (Robinson et al., 1989). The present experiments clearly demonstrate that diamond resorbed in the kimberlite melts in the graphite stability field. The resorption features observed are similar to those reported from natural diamonds (Frank and Lang, 1965; Robinson et al., 1989). The results suggest that the resorption of diamond to kimberlitic melt is a process controlled by temperature, time, carbon concentration and oxidation state of a resorption agent. Higher temperature, higher oxidation state, and lower CO_2 concentration of the kimberlitic melts seem to be favorite conditions for the dissolution of diamond to kimberlite melt. It is likely that carbon was dissolved in the melt as CO_2 .

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