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Conversion of polycyclic aromatic hydrocarbons to graphite and diamond at high pressures

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Abstract

The polycyclic aromatic hydrocarbons (PAH): naphthalene, anthracene, pentacene, perylene, and coronene were submitted to temperatures up to 1500 °C at 8 GPa. To avoid catalytic action of metals on thermal conversion, graphite was used as container material. Moreover, graphite is very permeable to the gaseous products of thermal decomposition of PAH. The resulting thermal transformations and their evolution were studied by X-ray diffraction, Raman spectroscopy and scanning electron microscopy as a function of temperature for 60-s treatments. The nature of the initial compounds clearly affects the products of the different stages of carbonization and the first steps of graphitization. This becomes hardly discernible in the final stages of graphitization above 1000 °C. Above 1200 °C, graphite with high crystallinity forms in all cases. The temperature of the beginning of diamond formation does not seem to be influenced by the nature of the initial PAH and is equal to -1280 °C for all investigated compounds. Diamonds formed from the PAH are high-quality 5–40 μm single crystals. The $p$, $T$ values of diamond formation here obtained are significantly lower than those previously known for direct graphite–diamond transformation.

Keywords: A. Graphite, Diamond; B. High pressure, Carbonization, Graphitization

1. Introduction

Many investigations have been devoted to the study of the thermal transformations of organic compounds at atmospheric pressure. It was shown that these transformations represent a complicated sequence of physicochemical processes that are usually classified under two headings—carbonization and graphitization. Carbonization which occurs at temperatures up to 1000 °C includes processes such as polymerizations, polycondensations, cracking reactions, molecular rearrangements, hydrogen transfers etc. leading to pure carbon materials with various degrees of ordering by gradual loss of hydrogen and other chemical elements [1,2]. During the graphitization stage, these metastable carbon states evolve towards graphite, an equilibrium state of the carbon system, through gradual annealing of structural defects at temperatures up to 3000–3200 °C [3–5].

The action of pressure on the carbonization and graphitization of organic compounds has also been studied several times. Although most of these investigations were carried out at rather low pressures, not exceeding 1 GPa, they clearly proved that pressure considerably influences all steps of both carbonization and graphitization.

In the case of carbonization, pressure favours polymerizations and, as such, increases the yields of carbon residues. Pressure changes the conditions of the mesophase formation by increasing the viscosity of the liquid medium, thus acting upon the structure of the carbon residue [6,7]. Furthermore, pressure makes possible the solid-phase carbonization of organic compounds, so
that application of high pressures accelerates the
graphitization process and results in considerable
reduction of the temperature of graphite formation.

In the case of polycyclic aromatic hydrocarbons
(PAH), the pressure influence on carbonization and
graphitization of naphthalene and anthracene have been
studied most extensively [1,2,5–11]. The study of naph-
thalene carbonization [10] has shown that, at 8 GPa,
graphite formation occurs very rapidly (within seconds)
at only 1000 °C, to be compared with 2500–3000 °C at
atmospheric pressure. Increasing the treatment time at 8
GPa results in a further decrease of the graphite for-
mation temperature. But the main feature of the thermal
conversions of organic compounds under high pressures
is that they can be carried out in two different \( \rho, T \)
ranges, those of thermodynamic stability of graphite or
of diamond, respectively. In the latter range, diamond
should be the final product of the transformations. So, a
new stage—diamond formation—appears in the se-
quenCe of thermal transformations of organic com-
pounds under high pressure.

The first attempt to synthesise diamond through
treatment of various organic compounds at 12–13 GPa,
according to the current pressure scale, and 1300–3000
°C in the absence of a metal-catalyst was made by
Wentorf [12]. The work was carried out in a “Belt”
high-pressure apparatus using titanium as sample container
material. In the course of these studies, Wentorf showed
that the final product of the transformations of naph-
thalene, anthracene, and chrysene at 12 GPa and 2000
°C is graphite, whereas the treatment of adamantane,
camphene, pyrene, fluorene, and polyethylene in the
same conditions leads to diamond formation with a
—60% yield. The increase of the synthesis temperature
up to 2300 °C allowed diamond to form from
anthracene, but with a rather low yield not exceeding 30%.
Moreover, Wentorf pointed out that the diamonds thus
obtained were quite unusual: they were snow-white
“soft” diamonds, with the appearance and mechanical
properties of paraffin wax or chalk. Nevertheless, the X-
ray diffraction patterns of these products were those of
diamond, but with broadened diffraction lines. There-
fore, the transformation had produced very small dia-
mond crystallites. As a result, it was assumed that the
very nature of the starting carbonaceous material affects
decisively the nature of the products of pyrolysis under
pressure, even at 2000 °C or more. It was also concluded
that the synthesis of the usual “hard” diamonds from
organic compounds is not possible, at least at pressures
up to 12 GPa and temperatures up to 2300 °C. It is to be
stressed that the temperature of the direct graphite–
diamond transformation at 12 GPa found by Wentorf
was 1300 °C.

The failure of this first attempt to synthesise “hard”
diamond from organic compounds under high static
pressures stopped the investigations in this field for a
long time. Nineteen years elapsed before Yakovlev et al.
reported the possibility of “hard” diamond synthesis
from organic compounds: hydrocarbons [13] and car-
bohydrates [14], at static pressures and without metal-
catalysts. Furthermore, the authors [13,14] did not find a
well-defined influence of the pristine molecular struc-
tures on the final stages of the thermal conversions, al-
though diamond synthesis was performed at \( \rho, T \) values
lower than those used in Ref. [12]. A similar result was
obtained by Voronov and Rakhmanina on studying
nucleation of diamond crystals from \( sp^2 \) and \( sp^3 \)
hydrocarbons decomposed under high pressure [15].

Later, “direct” diamond synthesis from various pre-
cursors including terpene-family hydrocarbons, furfuryl
alcohol resin, phenolic resin, and mesophase pitch was
studied by Onodera and Suito [16]. They used an octa-
hedral anvil press with tantalum, molybdenum, and
Teflon as sample container materials. The authors paid
considerable attention to the influence of the content
of hydrogen in the starting substances on the \( \rho, T \) para-
meters of diamond formation and yield. They showed
that, when using Ta for the sample container and heater,
diamond formation is observed for all hydrocarbons
(camphene, adamantane, and fluorene) studied in the 6–
12 GPa pressure range at temperatures up to 1600 °C.
The lowest \( \rho, T \) values for diamond synthesis (6 GPa and
700 °C) were obtained for camphene. They also showed
that the threshold pressure for diamond synthesis in-
creases from camphene (6 GPa), to adamantane (8
GPa), and fluorene (9 GPa). The high threshold value
for fluorene was explained by the aromaticity of this
compound which necessitates more energy to break
the aromatic rings. Thus, it was concluded that the initial
hydrocarbon structure affects the \( \rho, T \) parameters of
diamond synthesis. The authors pointed out also that the
influence of the hydrogen content is not clear for the
model hydrocarbons (camphene, adamantane, and flu-
orene). But, in the cases of furfuryl alcohol resin and
mesophase pitch, there are optimal values for the H/C
atomic ratio which decrease the diamond formation
temperature and increase the diamond yield. When some
limiting value of the H/C ratio is exceeded, diamond
formation stops. It is worthy of note that the role of Ta in
diamond formation is not conclusively shown because no
diamond was formed under the pertinent conditions
when Ta, as the container material, was replaced by Mo
or by Teflon. Onodera and Suito [16] suggested that Ta
facilitates the dehydrogenation process of organic com-
pounds. But, taking into account that Ta is the sole metal
which has a catalytic action on the graphite–diamond
transformation without the formation of a solution
[17,18], it may be assumed that Ta also favours a de-
crease of the \( \rho, T \) parameters of diamond formation.

The aim of the present work was to get a better
understanding of the influence of the starting molecular
structure on the thermal conversions of organic com-
pounds at high pressures. PAH (naphthalene: C_{10}H_{8}, anthracene: C_{14}H_{10}, pentacene: C_{22}H_{14}, perylene: C_{20}H_{12} and coronene: C_{24}H_{16}, and adamantane: C_{20}H_{18}) were submitted to pressure treatments at 8 GPa and temperatures up to 1500 °C. In the above compounds, the H/C atomic ratio changes from 0.5 for coronene to 0.8 for naphthalene; it is 1.6 for adamantane. To avoid catalytic effects of metals on the conversion processes of these compounds, high-pressure high-temperature treatments were performed in containers made of special 99.9% pure graphite. The processes of graphite and diamond formations underwent careful scrutiny.

2. Experimental

Naphthalene and adamantane (Chemapol, impurity contents less than 0.5%) and anthracene, pentacene, perylene, and coronene (Aldrich Chemical Company, impurity contents less than 2%) were used without further purification. The work was performed in a "Toroid"-type high-pressure apparatus [19]. Cylindrical samples of the initial compounds (4 mm diameter and 5 mm height) previously obtained by cold pressing were put into graphite containers which were placed in the high-pressure apparatus. The experimental procedure consisted in loading the apparatus up to 8 GPa at room temperature, heating up to the desired temperature (with a heating rate of 15 °C per second) at which the samples were then maintained under constant load for 60 s. The resulting high-pressure states were quenched under pressure to room temperature. Temperature was measured during the high-pressure experiments by means of chromel-alumel and Pt-Pt/Rh thermocouples. The relative temperature accuracy was equal to ±2 °C. However, the absolute temperature accuracy above 1200 °C was about ±50 °C.

The recovered samples were analyzed at ambient conditions by means of X-ray diffraction, Raman spectroscopy and scanning electron microscopy (SEM). X-ray diffraction patterns were obtained with an INEL CPS 120 diffractometer using CuKα radiation. The Raman spectra were recorded on a Dilor XY spectrometer using the 514.5 nm emission line of an Ar+ laser as excitation source. Microscopic investigations were carried out by means of an electron scanning microscope DSM 982 Gemini.

3. Results

Fig. 1 shows the X-ray diffraction patterns of the pristine PAH and products of their treatments at 8 GPa and different temperatures.

According to this figure, the thermal transformations of PAH are characterized by common features which are also observed for adamantane. First of all, for treatment times not exceeding 60 s, the initial crystalline molecular phases of naphthalene, anthracene, pentacene, perylene, and coronene remain the basic components of the samples up to 400 °C. At 600 °C, in all cases, significant amorphization is observed. A further temperature increase leads to intensive departure of hydrogen and dramatic reconstruction of the treated compounds which begin to lose their initial individuality. At 800 °C, all patterns exhibit mainly two types of peaks: (0 0 2) resulting from stacks of parallel layer planes at 23–29°, and two-dimensional (01) peaks resulting from the regular structure within the individual layer plane segments at 41–47°. Peaks of the type (h k l) are absent, indicating that there is little or no stacking order in the arrangement of parallel layers. These patterns are known to be characteristics of carbon states described as clusters made of small fragments of graphene planes plus some amount of disorganized carbon [3,20,21].

The appearance of three-dimensional (10 0) and (101) peaks at 1000 °C shows that three-dimensional ordering of the graphene planes takes place and that graphite structure formation begins. The formation of diamond is observed at 1280 °C for all compounds. In consequence of the existence of a temperature gradient in the high-pressure apparatus, diamond forms first of all in the middle part of the surface of the sample cylinder, that is in the region of the highest temperature. In the lower temperature sample parts, graphite with a high degree of crystalline perfection is obtained. The corresponding diffraction pattern is shown in Fig. 2.

The increase of treatment temperature leads to increased diamond yield which reaches almost 100% at temperatures above 1350 °C. In all cases, diamond formation occurs within a few seconds. As the absolute accuracy of our temperature measurements above 1200 °C is ±50 °C, we cannot know whether the temperature of the beginning of diamond formation varies significantly from one compound to another.

The variations of the structural parameters vs. temperature for the carbon materials formed during graphitization were determined. Fig. 3 shows the temperature dependences of the distance between graphene planes (c_{002}) and the average sizes of the crystallites, L_{a} and L_{c}, where L_{a} and L_{c} are the crystallite sizes in the layer plane and in its normal direction, respectively. The c_{002} values were determined from the (0 0 2) diffraction peak positions. The dimensions of carbon crystallites have been determined from the analysis of X-ray diffraction line broadening. According to the Scherrer formula, L_{a} is equal to:

\[ L_{a} = \frac{\lambda}{2\theta} \cos \beta \]

where \( \lambda \) is the X-ray wavelength, \( \beta \) is the angular width (radians) of the (0 0 2) diffraction peak at half-maximum intensity (corrected for instrumental broadening) and \( \theta \) is...
is the Bragg angle for reflection (002) [21]. The layer dimension $L_a$ is calculated by use of the equation:

$$L_a \propto \frac{1}{84\beta\cos h_a D}$$

where $\beta$ and $h_a$ correspond to reflections (10) and (10 0) [21].

As shown in Fig. 3, evidences for two-dimensional carbon clusters formation appear at 700 °C. In the 700–1000 °C temperature range, the $d_{002}$ values for high-pressure states formed from perylene and coronene are systematically larger than those for states formed from naphthalene, anthracene, and pentacene at the same temperature. This demonstrates the influence
of the initial PAH structures on the thermal conversion products. Assuming that the $d_{002}$ value for graphitization onset is equal to 0.344 nm [3–5,21], the graphitization process for linear-condensed PAH begins at lower temperatures than for perylene and coronene (see Fig. 3a).

According to Fig. 3b and c, the crystallite sizes increase sharply above 1150 °C. The validity of the $L_a$, $L_c$ calculation methods is questionable for temperatures above 1150 °C where the crystallite sizes are larger than 100 nm. However, in Fig. 3, we presented also the crystallite sizes calculated from the X-ray data for graphite formed at the nominal temperature of 1280 °C which is the lowest temperature of diamond formation at 8 GPa. According to our results, graphite samples with $d_{002}$ ~ 0.3354 nm (which corresponds to the interplanar distance of perfect graphite) are formed at temperatures higher than 1150 °C from all PAH.

The SEM photographs of Fig. 4 show the evolution of the morphologies of the materials formed at different temperatures. Fig. 4a exhibits a characteristic example of the stratified structures resulting from PAH treatments (here perylene) at 8 GPa and 700 °C. When temperature increases, the layered character of the materials develops. Fig. 4b–d display typical views of graphite and diamond particles, respectively, obtained from PAH transformations (here naphthalene) at 8 GPa and the nominal temperature of 1280 °C.

From graphite views recorded at 1100 °C and above, we could not distinguish any influence of the initial substances (PAH and adamantane) on the products of the final stages of carbonization. Graphite obtained near the lowest diamond formation temperature is composed of platelets with rounded edges. Their sizes are equal to 3–30 μm for $L_a$ and 0.1–0.3 μm for $L_c$ (Fig. 4b and c). In this case, X-ray and SEM measurements are in good agreement for the $L_c$ values. But as expected, the X-ray $L_a$ values appear much smaller than those determined through SEM.

The diamond crystal sizes are equal to —5–35 μm (Fig. 4d). As a whole, the diamond crystals have greater sizes than the graphite particles from which they are formed. The SEM views of Figs. 4d and 5 show the sample regions with near total and partial diamond formation, respectively. The partial diamond formation is associated with the existence of the above-mentioned temperature gradient in the reaction zone. Fig. 5 allows to note an interesting feature of the diamond formation manifesting itself at “direct” synthesis of diamonds from hydrocarbons. At micron-scale, the untransformed graphite and the diamond crystals occur in distinct regions. Graphite platelets are frequently oriented perpendicular to the growing diamond facets.

Raman spectra were also studied in the same temperature range and those corresponding to materials obtained from naphthalene are displayed in Fig. 6.
Similar evolutions are observed for others PAH. But in the case of naphthalene, we can compare our data with the results of a recent study of carbon materials formed through carbonization of a naphthalene-derived meso-phase pitch at atmospheric pressure [22]. Comparison of the latter results with those obtained as a result of thermal conversions of naphthalene-derived pitches at 8 GPa confirms that high pressures reduce considerably both temperature and time of treatment necessary to reach any given carbonization or graphitization stage. This is especially true in the case of graphitization. So,

such states as observed at 900 °C and 8 GPa (Fig. 6) form only at 2080 °C at atmospheric pressure [22].
Furthermore, the structural quality of the graphite obtained at 8 GPa and 1000 °C is far better than that obtained at atmospheric pressure and 2700 °C.

The evolution of the Raman spectra exhibited in Fig. 6 can be interpreted in terms of a three-stage model recently suggested by Ferrari and Robertson for the interconversion between amorphous carbon states and graphite [23]. This model takes into account changes of positions and intensities of the G and D bands [24,25]. The changes of the G band positions (initial shift up to 1592 cm⁻¹ and subsequent reverse displacement down to 1580 cm⁻¹) and the values of the \( \gamma(D)/\gamma(G) \) ratio (initial increase almost up to 3 and subsequent decrease almost to 0 at 1280 °C) correspond perfectly to the final stages of "an ordering trajectory" of a carbon system in the course of its transformation from amorphous carbon to graphite [23]. According to this model, the first stage of spectrum evolution in the 600–800 °C temperature range (Fig. 6), related to the final steps of carbonization, corresponds the formation of nanocrystalline graphite. As this takes place, the fraction of sp³ carbon states in the system tends to 0. The second stage, associated with graphitization, corresponds to transformation of nanocrystalline graphite into microcrystalline graphite. The X-ray data illustrate this process very well.

4. Discussion

The results of this work constitute one more verification that high pressures affect radically all stages of the thermal conversion of hydrocarbons. In the following discussion, we will concentrate on three questions:

- The reasons of the discrepancies regarding the diamond synthesis parameters as determined in this study and in Refs. [12,16].
- The influence of the nature of the starting hydrocarbons on the diamond synthesis parameters and the limits of hydrogen content, above and below which diamond synthesis does not occur in the investigated \( p, T \) range.
- Mechanisms of diamond formation from hydrocarbons in the absence of metal-catalysts.

As noted above, the minimum \( p, T \) values for diamond synthesis from PAH and adamantane obtained in this work (8 GPa and 1280 °C) are much lower than those reported by Wentorf [12] for anthracene (12 GPa and 2300 °C), adamantane (12 GPa and 2000 °C) and other hydrocarbons; the pressure value (8 GPa) is even somewhat lower than that (9 GPa) obtained by Onodera and Suito for diamond synthesis from fluorine in the case of a Ta container [16]. In our opinion, these discrepancies arise from thermodynamic constraints. They are related to the choice of the container material and to its permeability for the gaseous products of hydrocarbon pyrolysis. The importance of this factor was already noted by Yakovlev in his analysis of thermodynamic conditions of diamond formation from hydrocarbons [26]. As a container material, graphite has a much higher permeability for the gaseous component and primarily for hydrogen than metals or Teflon. The greater issue of hydrogen from the reaction zone in the case of the carbon container ensures the greater degree of the PAH decomposition, the greater yield of the carbon residue, and consequently the greater yield of diamond. That is, thermal conversions in graphite capsules take place in the conditions of more open system. For this reason, higher \( p, T \) values are needed for an achievement of the same degrees of hydrocarbons decomposition and of the same diamond yield in the cases of the Ti, Mo or Teflon containers. According to our data, the total mass loss near 1280 °C, say for naphthalene, is equal to -25%. This mass loss in the dehydrogenation process coincides with the data of Ref. [10] and correlates formally with complete departure of hydrogen in the form of methane. Diamond synthesis in Ta containers [16] can probably be explained by the catalytic action of Ta and may be by the higher permeability to gaseous products of the Ta capsule in comparison with the Ti, Mo or Teflon capsules [12,16].

In our experiments, we did not observe any pronounced influence of the starting molecular structure of organic compounds on the parameters of diamond formation. Starting with H/C atomic ratios ranging from 0.5 to 1.6, we did not find an upper limit of the hydrogen content for diamond formation. For all investigated aromatic compounds and adamantane, diamond formation at 8 GPa begins practically at the same temperature of -1280 °C. The starting H/C ratio rather acts upon the diamond yield and on the sizes of the resulting diamond crystallites. However, from the results obtained by Onodera and Suito [16], it may be assumed that the existence of such an upper limit is possible at least for two reasons. The first one has a thermodynamic origin and is related to the choice of the container material as is discussed above. Materials with high gas-tightness induce confinement of the gaseous products of thermal conversion in the reaction zone. It interferes with an attainment of high degrees of the PAH decomposition and of a significant yield of the carbon residue—a necessary component for the diamond synthesis. On the other hand, when the gaseous products can leave the reaction zone, a very high content of hydrogen in the initial carbonaceous material can induce a very important decrease of the system volume at the stage of carbon residue formation. Thus, the pressure in the reaction zone can decrease below the one which is necessary for diamond synthesis in the conditions of fixed press load.
It is to be stressed that, during our experiments, it was observed that diamond is formed not only from the carbonaceous product of the hydrocarbons carbonization, but also from the graphite of the container walls in contact with this carbonization product. Thus, graphite–diamond transformation in the presence of products of the hydrocarbon carbonization is observed at $p$, $T$ values which are much lower than the known minimum parameters of direct graphite–diamond transformation, that are 12 GPa and 3000 °C [27], 12 GPa and 1300 °C [12], 11.5 GPa and 3000 °C [28], and 14 GPa and 800 °C [29]. In experiments with homogeneous graphite–hydrocarbon mixtures, we have tried to determine the minimum hydrogen content for which diamond formation at 8 GPa is still possible. It was shown that almost complete conversion of all carbon lying in the reaction zone, including graphite and carbon formed by carbonization of hydrocarbons, takes place if the hydrogen mass content in the starting mixture is equal to 1%. Fig. 7 shows diamond formed at 8 GPa and 1280 °C from binary mixtures of graphite with naphthalene and anthracene with this hydrogen content.

Decrease of hydrogen mass content below 1% reduces the diamond yield. When the hydrogen mass content is below 0.2%, no diamond is formed. Complete transformation of carbonaceous material into diamond is also observed when the initial hydrogen mass content increases above 1%. But in this case, the diamond produced becomes gradually looser and looser and easily breaks down into separate crystals and splinters when extracted from the high-pressure apparatus. The increase of porosity of the diamond product is obviously related to the increase of the density jump undergone by carbonaceous materials when diamond forms from the mixtures with high hydrogen contents.

These results suggest that hydrogen plays an important role in the processes of diamond formation from graphite and carbon residues resulting from hydrocarbon carbonization at high pressures. It is known that the hydrogen content decreases gradually in the course of the co-carbonization of hydrocarbons with graphite. During the final steps of carbonization, the chemically bound hydrogen atoms are mainly gathered in the defective graphite sheets [21]. An analogous promoting effect of hydrogen on the high-pressure synthesis of diamond was noted by Kindlein et al. [30] who showed that the implantation of hydrogen ions in graphite allows to obtain a 50% increase of diamond nucleation during the high-pressure graphite–diamond transformation. The interpretation of these results is not straightforward. Further investigation is currently being performed in our group by using high resolution transmission-electron microscopy and related techniques to provide insight into the understanding of the hydrogen action mechanism on the direct transformation of graphite to diamond.

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