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The direct growth of planar and vertical graphene on Si(100) *via* microwave plasma chemical vapor deposition: synthesis conditions effects†

 Š. Meškiniš, *^a A. Vasiliauskas,^a A. Guobienė, ^a M. Talaikis, ^b G. Niaura^b and R. Gudaitis^a

In the present research, graphene was synthesized directly on a Si(100) substrate *via* combining direct microwave plasma-enhanced chemical vapor deposition and protective enclosure. The graphene flake orientation was controlled using suitable synthesis conditions. We revealed that high processing temperatures and plasma powers promote vertical graphene growth. The main related physical mechanisms were raised temperature gradients, thermal stress, ion bombardment, and elevated electric field effects. Lowering the synthesis temperature and plasma power resulted in planar graphene growth. An elevated synthesis temperature and long deposition time decreased the graphene layer number as the carbon desorption rate increased with temperature. Dominating defect types and their relationships to the graphene growth conditions were revealed. Planar graphene n-type self-doping was found due to substrate-based charge transfer. In the case of vertical graphene, the increased contact area between graphene and air resulted in the adsorption of more molecules, resulting in no doping or p-type doping.

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1 Introduction

The 2D nanomaterial graphene is of very significant interest due to its exciting combination of electrical, optical, and mechanical properties. High levels of charge carrier mobility and multiplication, optical transparency, flexibility, very high Young's modulus and failure strength, high ductility, and chemical inertness are all present.^{1–4} Graphene can be grown as “classic” planar sheets, as vertical graphene flakes and nanowalls, and as a filler in nanocomposite materials. Among the numerous possible graphene applications, it has been considered for use as a new transparent conductor, a monolayer alternative to Schottky contact metals, and even as an active layer in semiconductor devices and different sensors. In particular, vertical graphene nanowalls provide a high surface area that is beneficial for the fabrication of biosensors and energy-storage devices.⁵ Planar graphene is beneficial for transparent electronics and transparent conductor applications, such as in various photoelectric and microelectronics devices.⁵

Usually, planar graphene is fabricated *via* graphite exfoliation or chemical vapor deposition (CVD) on catalytic metal (Cu, Ni, Co) foil, with subsequent transfer to the target substrate.⁶

Graphene transfer onto the target semiconductor or dielectric substrate is a long process. This can result in graphene contamination by different adsorbents⁷ or the formation of wrinkles or ripples on graphene.⁸

Graphene can be synthesized directly on semiconducting or dielectric substrates using plasma-enhanced chemical vapor deposition (PECVD). Usually, remote plasma systems are used for direct graphene synthesis to avoid radiation-based defect formation or even the total etching-out of the growing graphene film.⁹ In such a way, planar graphene layers are grown.^{9,10} Vertical graphene sheets and nanowalls are synthesized in direct plasma systems.⁹

Some studies have shown the growth of vertical and planar graphene in the same PECVD system on dielectric or semiconducting substrates.^{5,11–16}

Switching between the growth of planar and vertical graphene on a glass substrate can be achieved using a copper-foam-based Faraday cage with different pore sizes.⁵ However, this method has some disadvantages. Copper atoms can be deposited on the graphene.¹⁷ It was found that the density of graphene nanowalls directly grown on SiO₂ and Al₂O₃ substrates *via* microwave PECVD can be reduced *via* combining a protective metallic shield with lowered acetylene/hydrogen gas flow ratios.¹¹ However, planar graphene was grown on the entire substrate surface only when using additional NH₃ gas flow.¹¹ Thus, only n-type doped graphene could be achieved. In that study, no details regarding the geometric parameters of the protective shield were provided. Planar and non-planar graphene flakes were synthesized on SiO₂ film using a microwave

^aInstitute of Materials Science, Kaunas University of Technology, K. Baršausko St. 59, LT51423 Kaunas, Lithuania. E-mail: sarunas.meskinis@ktu.lt

^bDepartment of Organic Chemistry, Center for Physical Sciences and Technology, Saulėtekio av. 3, LT-10257 Vilnius, Lithuania

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plasma torch.¹² The graphene orientation was changed upon choosing an appropriate ethanol gas flow and plasma power in that study. However, the separate planar graphene flakes covered only a tiny part of the substrate.¹²

In some studies, the graphene flake orientation was changed *via* controlling the methane and etching gas flow ratios without reporting the use of Faraday-cage-type protective shields. Notably, vertical graphene was grown on silicon *via* microwave PECVD using CH₄/H₂ gas flow ratios higher than 5.5, and horizontal graphene was synthesized using lower methane/hydrogen gas flow ratios.¹³ However, in other studies, vertical graphene was grown using very different CH₄/H₂ gas flow ratios, and no planar graphene was reported.^{18–20}

The orientation of graphene flakes directly synthesized on SiO₂ film was changed from planar to vertical *via* varying the CH₄/Ar gas flow ratio in a CH₄ : Ar:H₂ gas mixture in an ECR PECVD system.^{14,15} However, in numerous studies, vertical graphene was grown on SiO₂ and other substrates from a CH₄ : H₂ mixture without any argon gas (see reviews^{18,21–26}). On the other hand, planar graphene was synthesized on SiO₂ and glass substrates *via* ECR PECVD using a C₂H₄ : Ar gas mixture.²⁷ The addition of CO₂ gas to the methane, hydrogen, and argon gas mixture to grow graphene on glass and quartz substrates using magnetron microwave PECVD resulted in planar graphene rather than vertical graphene.¹⁶ However, other authors grew vertical graphene from a CH₄/CO₂ gas mixture²⁸ or even CO₂ gas alone.²⁹ Thus, changing the graphene sheet orientation *via* adjusting the synthesis gas flow ratio or composition seems to be very system specific.

In the present research, we achieved control of the orientation of PECVD-synthesized graphene flakes *via* setting appropriate growth process conditions. The main technological parameters affecting the graphene structure and flake orientation were considered. Those parameters are plasma power, pressure, the gas mixture composition, temperature, and time.^{18,21,26,30} Technological processing conditions beneficial for the growth of planar graphene and vertical graphene on Si(100) substrates were found. High processing temperatures and plasma powers promoted vertical graphene growth due to increased temperature gradients, thermal stress, ion bombardment, and electric field effects. The suppression of these effects using lower synthesis temperatures and plasma powers promoted planar graphene growth.

2 Experimental

Vertical and planar graphene was synthesized on Si(100) using a Cyrannus microwave-plasma-enhanced chemical vapor deposition (PECVD) system (Innovative Plasma Systems (Iplas) GmbH, Germany). More information on the Cyrannus microwave plasma source can be found elsewhere.^{31–35} A methane and hydrogen gas mixture was applied. Prime grade n-type double-sided polished monocrystalline Si(100) (Sil'tronix, France) was used. The selected graphene samples were grown on p-Si. According to the manufacturer data, the silicon wafer resistivity was in the range of 1–10 Ω cm. The size of the silicon substrates was 1 × 1 cm.

First, the substrate was pretreated with hydrogen plasma without any wet chemical cleaning of the silicon. The plasma power was 1.7 kW, the processing temperature was 900 °C, the hydrogen flow was 200 sccm, the pressure was 22 mBar, and the time was 15 min. After plasma treatment, methane gas was introduced. A special enclosure, consisting of a rectangular non-magnetic steel sheet bent perpendicularly in two places, was used to protect the sample from direct plasma action and prevent excessive etching of the growing graphene. The steel enclosure can withstand much higher processing temperatures and powers without being damaged or without graphene poisoning due to evaporation or sputtering products. We were not able to grow graphene inside the plasma discharge zone when no protective enclosure was used.³⁶ Furthermore, a graphene sample placed inside CH₄/H₂ plasma without the protective enclosure was fully etched-out.

The graphene synthesis process parameters (plasma power, pressure, methane and hydrogen gas flows, temperature, and time) were varied. The sample direct synthesis conditions are presented in Table 1.

Raman scattering spectra (further Raman spectra) of synthesized samples were acquired using an inVia Raman spectrometer (Renishaw, Wotton-under-Edge, UK). The exciting laser-beam wavelength was 532 nm, and the power was 1.5 mW. The Lorentz function was used to fit the D, G, D', and 2D peaks. The 2D/G peak intensity ratio ($I(2D)/I(G)$ ratio) was estimated to evaluate the graphene layer number.³⁷ The $I(D)/I(G)$ peak

Table 1 The graphene growth processing conditions used in the study

Sample no.	P, kW	H ₂ , sccm	CH ₄ , sccm	p, mBar	T, °C	t, min
1	1.4	150	50	22	900	50
2	1.1	150	50	22	900	50
3	0.9	150	50	22	900	50
4	0.7	150	50	22	900	50
5	0.7	150	50	10	900	50
6	0.7	150	50	10	800	50
7	0.7	150	50	10	700	50
8	0.7	75	25	10	900	50
9	0.7	75	25	10	800	50
10	0.7	75	25	10	700	50
11	0.7	75	35	22	700	50
12 ^a	0.7	75	35	10	700	50
13	1.7	150	50	22	800	50
14	1.7	150	50	22	700	50
15	1.7	150	50	22	600	50
16	1.7	150	50	22	900	50
17	1.7	150	50	22	900	30
18	1.7	150	50	22	900	70
19	1.7	150	50	22	900	10
20	1.7	150	50	22	900	90
21	1.7	150	50	10	900	90
22	0.7	150	50	10	900	90
23	0.7	150	50	10	800	90
24	0.7	75	25	10	800	90
25	0.7	75	25	10	700	90
26	0.7	75	25	10	700	150

^a The graphene was grown on both n-Si and p-Si.

intensity ratio was calculated to estimate the defect density of graphene.^{38,39} It should be mentioned that the $I(2D)/I(G)$ ratio can decrease depending on the defect density.³⁸ However, a different dependence was found for most samples in our case (Fig. S1†). Thus, we used the $I(2D)/I(G)$ ratio for graphene layer number evaluation, despite the presence of defects. The D/D' peak intensity ratio revealed the types of dominating defects.⁴³ The full width at half maximum of the G peak ($\text{FWHM}(G)$)^{40–42} and positions of the G and 2D peaks ($\text{Pos}(G)$ and $\text{Pos}(2D)$)^{36,41,42,44,45} were also taken into account as parameters sensitive to graphene doping and stress. The spectra were measured in several different places on the sample. Average values and standard deviations of different Raman spectra parameters were calculated for each sample.

Raman mapping was accomplished for selected samples using an inVia Raman microscope (Renishaw, Wotton-under Edge, United Kingdom) equipped with a thermoelectrically cooled ($-70\text{ }^{\circ}\text{C}$) CCD camera. Spectra were excited using 532 nm laser radiation and dispersed using a grating with 1800 lines per mm. The laser power was restricted to 0.45 mW and focused on a spot on the surface with a diameter of approximately $1\text{ }\mu\text{m}$ by using a $50\times/0.75\text{ NA}$ objective lens (Leica). The Raman maps were collected from a surface area of $50\times 50\text{ }\mu\text{m}^2$ with a $1.5\text{ }\mu\text{m}$ step size. The acquisition time for each spectrum was set to 4 s. To plot the $I(2D)/I(G)$ maps, experimental spectra were fitted with a single Gaussian–Lorentzian shape component for each of the spectral modes (G and 2D) using software written in our lab based on MATLAB R2013 (The MathWorks Ins.).

Atomic force microscopy (AFM) experiments were carried out at room temperature using a NanoWizardIII atomic force microscope (JPK Instruments, Bruker Nano GmbH, Berlin, Germany). The data were analyzed using JPKSPM data processing software (Version spm-4.3.13, JPK Instruments, Bruker Nano GmbH). The AFM images were collected using an ACTA (Applied NanoStructures, Inc., Mountain View, USA) probe. A pyramidal cantilever was used. The radius of curvature (ROC) was $<10.0\text{ nm}$ and the cone angle was 20° . The reflex side coating was Al with a thickness of $50\pm 5\text{ nm}$. The probe force constant was set at 40 N m^{-1} , and the resonance frequency was in the 300 kHz range. Height, amplitude, and phase diagrams were recorded in steps with a scan size of $2\text{ }\mu\text{m}$ and a scan speed of 1 Hz . The integral gain was set to 2, while the proportional gain was set to 5. Pixels for samples and lines were 516×516 , operating in tapping mode.

Some samples were additionally investigated using a Helios Nanolab 650 scanning electron microscope (SEM) (FEI, Eindhoven, The Netherlands).

3 Results and discussion

3.1 Synthesis of planar and vertical graphene

The structures of the graphene layers synthesized using different growth process conditions were investigated *via* Raman spectroscopy. Typical Raman spectra are presented in Fig. 1 and S2.† The G and 2D peaks typical of graphene can be seen at $\sim 1600\text{ cm}^{-1}$ and $\sim 2700\text{ cm}^{-1}$. Defect-related D ($\sim 1350\text{ cm}^{-1}$), D + D'' ($\sim 2500\text{ cm}^{-1}$), and D + D' ($\sim 2900\text{ cm}^{-1}$)

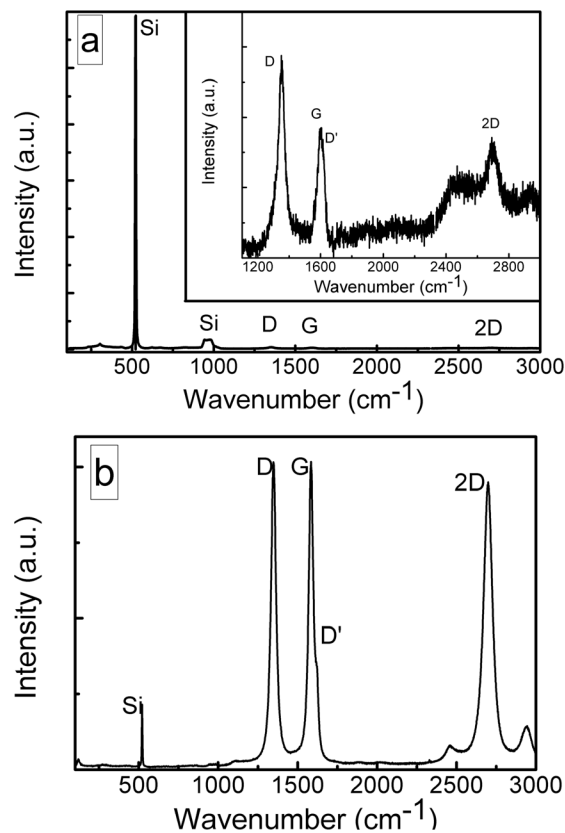


Fig. 1 Characteristic Raman spectra of graphene grown on Si(100): (a) the plasma power was 0.7 kW and the synthesis temperature was $700\text{ }^{\circ}\text{C}$; (b) the plasma power was 1.7 kW and the synthesis temperature was $900\text{ }^{\circ}\text{C}$.

peaks³⁹ were observed. The D' peak at $\sim 1620\text{ cm}^{-1}$ is present as a shoulder of the G peak. The presence of defect-related Raman bands is typical for graphene films directly synthesized on silicon and silicon-based dielectrics, such as SiO_2 and Si_3N_4 , due to the nanocrystalline nature of these layers.^{9,10,43} The Si-substrate-related peak at 520 cm^{-1} (ref. ¹⁹ and ⁴⁶) is visible in the Raman spectra of our samples; the broad feature in the vicinity of $950\text{--}1000\text{ cm}^{-1}$ corresponds to the silicon first overtone mode.

Significant differences between the synthesized graphene Raman spectra were found. The Si(100) substrate-related peak at 520 cm^{-1} was much stronger than the graphene-related peaks for some samples. In other cases, the intensity of the G peak was comparable with the intensity of the Si-related peak or it was even higher.

It should be noted that the number of graphene layers can be evaluated based both on the relative intensities of the Si and G bands and the $I(2D)/I(G)$ ratio.^{46,47} The $I(G)/I(\text{Si})$ ratio should increase and the $I(2D)/I(G)$ ratio should decrease with an increased graphene layer number. It should be mentioned that the $I(2D)/I(G)$ intensity ratio can decrease not only due to an increase in the number of graphene layers but also because of an increase in the disorder/defective nature of a sample.³⁸ However, as discussed above, in our case this ratio can be used to study the graphene layer number changes in different

samples (Fig. S1†). The plot of $I(2D)/I(G)$ vs. $I(G)/I(Si)$ revealed contradictory results (Fig. 2a). Three groups of samples were found. For the first group, the $I(2D)/I(G)$ ratio decreased with an increase in the $I(G)/I(Si)$ ratio. In numerous studies, the presence of this behavior for Raman spectra of planar graphene samples indicates a rising graphene layer number.^{46,47} For the second group, the tendency of the $I(2D)/I(G)$ ratio to rise with the $I(G)/I(Si)$ ratio is seen. In the case of samples with the highest $I(G)/I(Si)$ ratios, no clear relationship is found between the $I(2D)/I(G)$ ratio and $I(G)/I(Si)$ ratio. These contradictory results can be explained *via* considering different mechanisms affecting the dependence of the $I(2D)/I(G)$ and $I(G)/I(Si)$ ratios on the graphene layer number.^{46,47} The $I(2D)/I(G)$ ratio is related to changes inside graphene.^{46,47} At the same time, the $I(G)/I(Si)$ ratio depends on silicon substrate signal attenuation.⁴⁶ In such a case, the orientation of the graphene flakes can be crucial because vertical graphene nanowalls can absorb several tens of times more light than planar graphene with the same layer number.⁴⁸ Thus, one can suppose that for vertical graphene nanowalls the $I(G)/I(Si)$ ratio does not depend on the graphene layer number. The selected samples were studied *via* atomic force microscopy (AFM) and scanning electron microscopy (SEM) to test this assumption (Fig. 2b, 3, and 4, and S4–S21†).

The $I(G)/I(Si)$ and $I(2D)/I(G)$ ratios increased with the graphene roughness value (R_q) (Fig. 2b). Thus, based on the $I(2D)/I(G)$ ratio, the sample roughness increases despite a decrease in the graphene layer number. This means that a graphene roughness increase is not related to a graphene layer number change. Taking into account R_q values above 5 nm, in this case, growth of non-planar graphene flakes can be supposed (Fig. 3 and 4). A thorough study of the graphene height histograms and AFM images of selected samples revealed that the samples with the lowest R_q values are planar graphene (Fig. 2b and 3a, S13–S15, S18, and S20†). In this case, the maximum feature height is 2–3 nm. Significant phase contrast was found for these low-roughness graphene layers (see, *e.g.*, Fig. S12 and S13†), indicating the presence of different phases on the surfaces of the samples.⁴⁹

Thus, some through-holes are present in the planar graphene layers. According to ref.⁵⁰ and ⁵¹, the graphene layer thickness is 0.41 nm. Thus, the maximum planar graphene thickness is 5–6 layers. No surface features can be seen in a typical SEM image measured for a sample from this group, indicating the growth of planar graphene (Fig. 4a). The Raman microscopy mapping study confirmed the deposition of graphene over the entire sample area (see the insets of Fig. 4a and b). Thus, the suppression of the Si-related peak is in good accordance with the increased sample roughness and non-planar graphene growth.

Samples with larger R_q values and with G/Si ratio values similar to planar graphene represent a mixture of planar and vertical graphene (Fig. 2). In this case, according to the height histograms, large proportions of the feature heights are within the range typical for planar few-layer graphene (Fig. 3c and d, and S11–S13, S16, and S18†). However, significant proportions of the heights are well beyond the thickness of ten graphene layers, despite the Raman spectra of the samples still being typical of graphene (Fig. 1b).

The last group of samples involves vertical graphene films. Their height histogram values are between several nm and several tens of nanometers (Fig. 3b, S4, S9, and S18–S20†). Vertical graphene flakes can be clearly observed in a typical SEM image (Fig. 4b). In this case, the presence of a mixture of bilayer and single-layer graphene can be supposed according to the $I(2D)/I(G)$ ratio values of close to one.

Clarifying the analysis of the $I(2D)/I(G)$ vs. $I(G)/I(Si)$ graph, one can see that the first group of samples is made up of planar graphene and some planar graphene samples with the inclusion of non-planar graphene flakes (Fig. 2). The second group of samples is made up of mixed vertical and planar graphene samples and some vertical graphene samples. The third group is made up of vertical graphene samples only. This is related to the different graphene synthesis conditions. Most of the samples in group I are grown at a temperature of 700 °C using plasma power of 0.7 kW. However, all the samples in group III were synthesized at a temperature of 900 °C, plasma power of

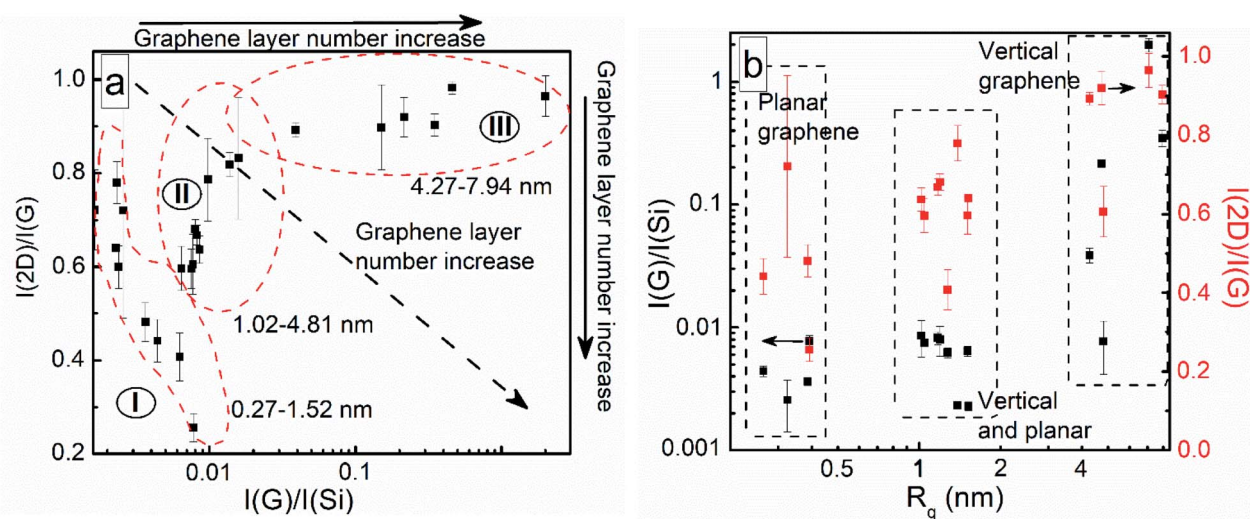


Fig. 2 (a) A plot of $I(2D)/I(G)$ vs. $I(G)/I(Si)$; and (b) plots of $I(G)/I(Si)$ vs. R_q and $I(2D)/I(G)$ vs. R_q .

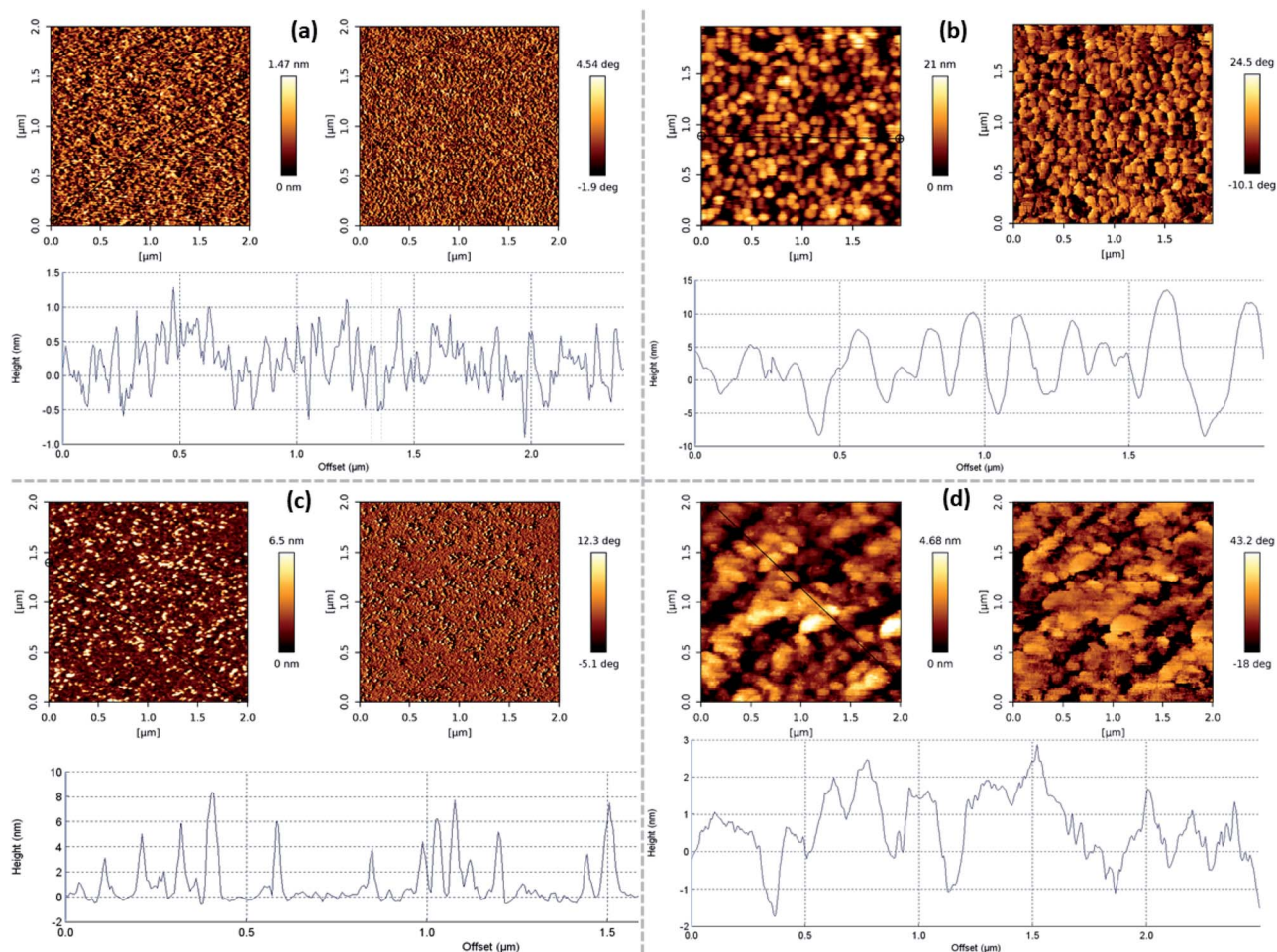


Fig. 3 Typical graphene AFM images. The sample synthesis conditions: (a) plasma power, 0.7 kW; hydrogen flow, 150 sccm; methane flow, 50 sccm; pressure, 10 sccm; synthesis temperature, 700 °C; growth time, 50 min; (b) plasma power, 1.7 kW; hydrogen flow, 150 sccm; methane flow, 50 sccm; pressure, 22 mBar; synthesis temperature, 900 °C; growth time, 50 min; (c) plasma power, 0.7 kW; hydrogen flow, 75 sccm; methane flow, 25 sccm; pressure, 10 mBar; temperature, 900 °C; time, 50 min; (d) plasma power, 0.7 kW; hydrogen flow, 150 sccm; methane flow, 50 sccm; pressure, 22 mBar; temperature, 900 °C; time 50 min.

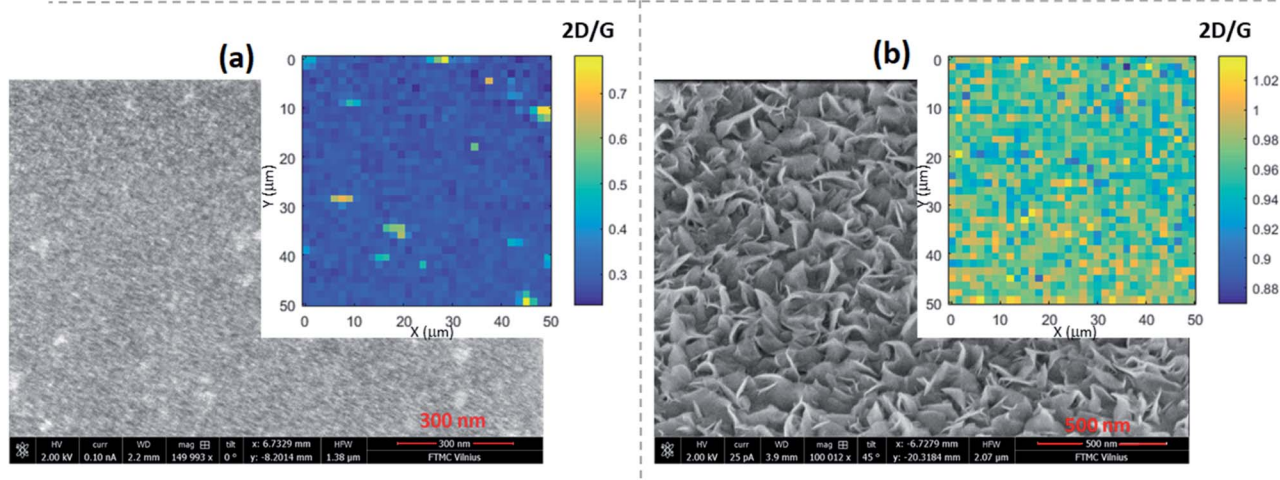


Fig. 4 Typical graphene SEM images and $I(2D)/I(G)$ mapping. The sample synthesis conditions: (a) plasma power, 0.7 kW; hydrogen flow, 75 sccm; methane flow, 35 sccm; pressure, 22 mBar; temperature, 700 °C; (b) plasma power, 1.7 kW; hydrogen flow, 150 sccm; methane flow, 50 sccm; pressure, 22 mBar; temperature, 900 °C; time, 90 min.

1.7 kW, and with a synthesis time of at least 30 min. Therefore, the effects of the graphene synthesis conditions were then analyzed more thoroughly.

3.2 Synthesis condition effects

The technological parameters studied were plasma power, pressure, gas flows, temperature, and time. The electric field strength at the substrate surface increases with plasma power, promoting vertical graphene growth and potentially increasing the processing temperature.³⁰ Lower pressure helps to achieve higher electron mean free paths and higher electron temperatures, subsequently increasing the gas ionization rate.¹⁸

The carbon-containing active species flux at the surface promotes graphene growth.⁵² The graphene nucleus density decreases with an increased H₂ content due to suppressed hydrogen etching, resulting in a smaller graphene grain size.⁵³ A higher substrate temperature increases the temperature gradient and thermal stress, promoting the upward curling of graphene flakes and subsequent vertical graphene growth.^{18,54} Finally, the deposition time is important because the growth of vertical graphene begins with planar graphene growth.⁵⁴ Then, due to stress, ion bombardment, and lattice mismatch between the substrate and graphene, vertical graphene growth begins.^{18,21,55}

Via changing the gas flows, one can adjust the fluxes of active species, controlling the graphene growth⁵² and etching⁵³ processes. Therefore, the gas-flow effects were also studied for necessary control of the graphene structure.

An analysis of AFM images revealed that for graphene synthesized at a temperature of 900 °C, a mixture of planar and vertical graphene was grown using plasma power in the range of 0.7–1.4 kW (Fig. S6–S9†). For all synthesized samples, graphene flakes with a width of 70–90 nm, length of 300 nm, and height of up to 4 nm are seen (Fig. S6–S9†). An increase in the plasma power to 1.7 kW resulted in the growth of vertical graphene flakes with a width of 60–80 nm (Fig. S5†). At the same time, the $I(G)/I(Si)$ ratio and R_q value significantly increased (Fig. 5a and c). This can be explained because the electric field strength and ion bombardment increased with an increase in the plasma power, promoting vertical graphene growth.³⁰ In contrast, possible plasma-induced temperature increase effects are somewhat limited (see Fig. S3†). At the same time, the $I(2D)/I(G)$ ratio increased and the $I(D)/I(G)$ ratio decreased with plasma power, indicating a lowering of the graphene layer number and defect density (Fig. 5e and g). The methane and hydrogen dissociation rates increase with plasma power.⁵² Thus, graphene etching by hydrogen prevailed over graphene growth, promoting increased C and CH_x species densities.^{52,56} The decrease in defect density despite an increase in plasma power can be explained considering the prevailing defect types. For graphene synthesized using plasma power from 0.7 to 1.4 kW, boundary defects prevailed, and a further power increase resulted in the domination of boundary defects and vacancies (Fig. 5i). Thus, the $I(D)/I(G)$ ratio decrease can be attributed to the increased graphene crystallite size. Increased ion irradiation can explain vacancy formation when using plasma power of 1.7 kW, as shown in numerous studies.^{57–62}

Doubling the processing pressure did not influence the graphene synthesized using a power of 1.7 kW (Fig. S4†). However, we achieved some control of the graphene layer number, defect density, and defect type for samples grown at 700 °C using power of 0.7 kW. Some on-site defects appeared along with boundary defects when the gas flows were increased (Fig. 5i). The presence of a more significant number of hydrocarbon species with a lower dissociation degree can be supposed.⁶³ Synthesis pressure doubling slightly decreased the defect density and increased the layer number (Fig. 5g and h, S4†) due to the reduced gas ionization,¹⁸ resulting in lower ion etching and the formation of fewer radiative defects.

The synthesis temperature effects are shown in Fig. 6. It can be seen that no graphene was grown at 600 °C. In the synthesis temperature range of 700–900 °C, the $I(G)/I(Si)$ and $I(2D)/I(G)$ ratios increased with temperature when using a power of 1.7 kW for synthesis (Fig. 6a and e). When using power of 0.7 kW, the lowest graphene layer number was obtained at 800 °C. The graphene layer number changes are related to competition between decreased graphene etching by hydrogen⁵³ and increased carbon atom desorption.⁶⁴ The defect density slightly increased with temperature for graphene synthesized using power of 0.7 kW, while it non-monotonically depended on the temperature for graphene grown using power of 1.7 kW. The graphene R_q value increased with the synthesis temperature for all studied samples (Fig. 6c). The increased temperature promoted vertical graphene growth, activating the formation of non-planar graphene flakes along with planar ones, and the coverage of the entire sample surface with vertical graphene flakes occurred upon a further temperature increase (see Fig. S4, S13, S10, S14, S17, and S19–S22†). This can be explained based on the presence of higher temperature gradients and thermal stress, which promote the upward curling of graphene flakes and subsequent vertical graphene growth.^{18,54} The effects of temperature on the types of prevailing defects depended on the plasma power used for synthesis (Fig. 6i). For graphene synthesized using power of 1.7 kW, an increase in temperature resulted in the formation of vacancies along with boundary defects. Thus, additional thermal activation is necessary for ion-irradiation-induced vacancy formation.⁶⁵

The $I(G)/I(Si)$ ratio and surface roughness significantly increased with the graphene synthesis time when a power of 1.7 kW and temperature of 900 °C were used for synthesis (Fig. 6b and d). After 10 min of growth, a planar and non-planar graphene mixture was formed. The $I(G)/I(Si)$ ratio of that sample places it in group I, which mainly contains planar graphene. For the planar graphene areas, flakes with a height of 0.2–0.8 nm and a width of 15–25 nm were observed, and for non-planar graphene, flakes with a height of up to 5.5 nm and a width of 40–75 nm were seen (Fig. S22†). A further increase in the deposition time resulted in the growth of vertical graphene (Fig. S5 and S18–S20†). This follows the vertical graphene growth mechanism suggested in ref. ⁵⁴. At first, planar graphene grows,⁵⁴ and it is necessary for some time to elapse before vertical graphene growth begins due to thermal stress release.^{9,18,21,66} The $I(2D)/I(G)$ ratio increased and the $I(D)/I(G)$ ratio decreased with growth time for graphene deposited using power of 1.7 kW (Fig. 6f and h). The prevailing defect type changed from vacancy defects for graphene grown for

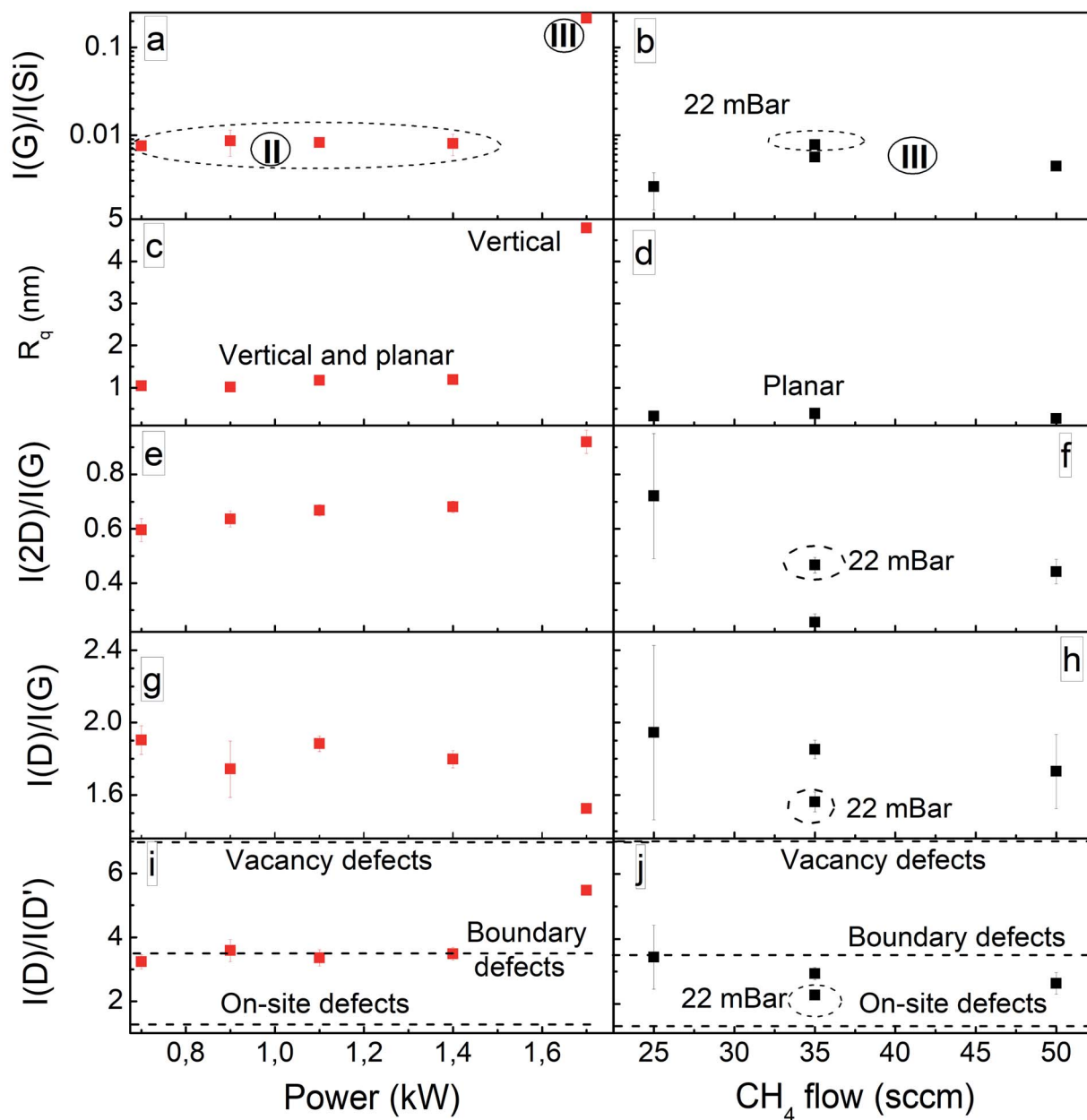


Fig. 5 The effects of synthesis power (a, c, e, g, and i) and methane gas flow (b, d, f, h, and j) on the graphene structure: $I(G)/I(Si)$ ratio (a and b); R_q (c and d); $I(2D)/I(G)$ (e and f) and $I(D)/I(G)$ ratios (g, h); and $I(D)/I(D')$ ratio (i and j).

10 min to boundary defects for the sample synthesized for 90 min (Fig. 6j). The decrease in the graphene layer number with an increase in processing time can be explained based on the domination of hydrogen etching over additional graphene layer growth.⁶⁴ The defect density decrease over time can be explained based on the self-annealing of vacancy defects⁵⁹ and an increased graphene crystallite size.⁶⁷

3.3 Graphene self-doping

The graphene FWHM(G), Pos(G), and Pos(2D) were analyzed to reveal possible doping and stress effects (Fig. 7). One can see the

tendency of FWHM(G) to increase with an upshift in Pos(G) for the group III samples (Fig. 7a). This can be explained based on the increased defect density⁴¹ due to the decreased graphene crystallite size.⁶⁷ Another tendency is for FWHM(G) to decrease with an upshift in Pos(G) for group I samples (Fig. 7a). That is typical of graphene doping effects.^{40–42} Thus, graphene self-doping can be suggested for group I samples.

Analysis of the Pos(2D) vs. Pos(G) plot revealed that for all group I samples, Pos(2D) is downshifted with an upshift in Pos(G) (Fig. 7b). This is typical of n-type doped graphene.^{36,40,42,44,45} For some samples, the Pos(2D) downshift with an upshift in Pos(G) was much more significant, and this could

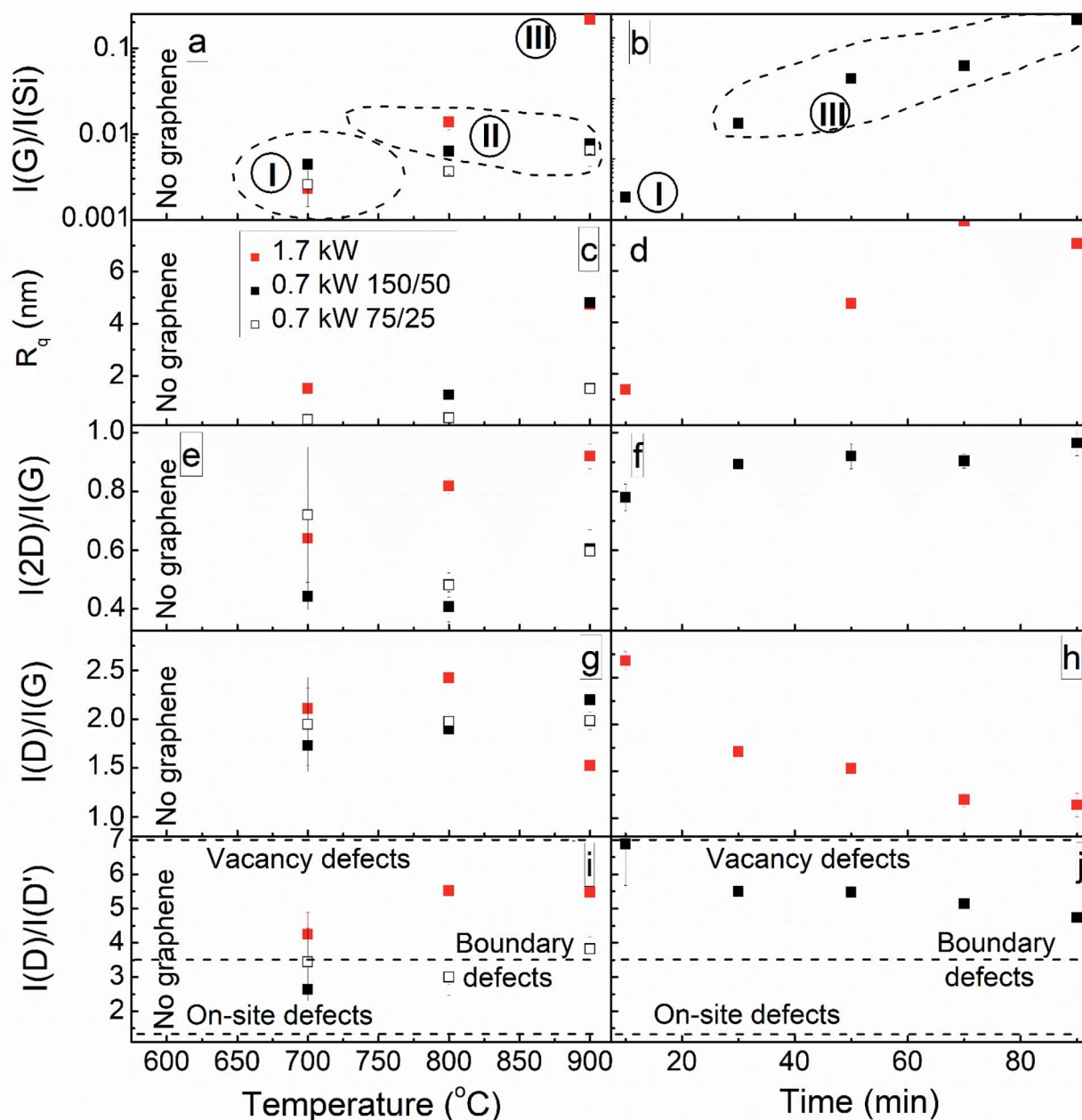


Fig. 6 The dependence of the graphene structure on temperature (a, c, e, g and i) and time (b, d, f, h and j).

be explained only based on n-type self-doping (Fig. 7). The Pos(2D) downshift can occur due to a graphene layer number increase.⁴⁷ However, in our case, no clear correlation between the large Pos(2D) downshift and $I(2D)/I(G)$ ratio changes was found (Fig. S23†). Thus, a combination of n-type self-doping and stress effects can be suggested. Pos(2D) slightly upshifted with Pos(G) for other graphene samples (Fig. 7b). Thus, weak p-type doping can be supposed.^{40–42,44,45}

Graphene self-doping has already been reported in numerous studies.^{36,68–74} In our case, n-type self-doping was found for all planar graphene samples and some samples consisting of a planar and non-planar graphene mixture. This is in accordance with the substrate-induced n-type self-doping hypothesis.^{71–74}

Substrate-induced graphene n-type self-doping was reported for graphene placed on a SiO₂ substrate due to positive silanol groups.^{75,76} It is noteworthy that a native oxide layer can be present at the interface between directly synthesized graphene and silicon due to silicon surface reoxidation.⁶² Thus, one can suppose that, in our case, electron transfer from the native silicon oxide inter-layer to directly synthesized graphene takes place. The Pos(2D) vs. Pos(G) data for graphene synthesized on p-type Si(100) are in good accordance with this assumption (Fig. 7b and S24†).

For vertical graphene, most of the graphene surface is substrate-free.

Thus, charge transfer from the substrate to graphene is limited. However, a much larger graphene surface area is

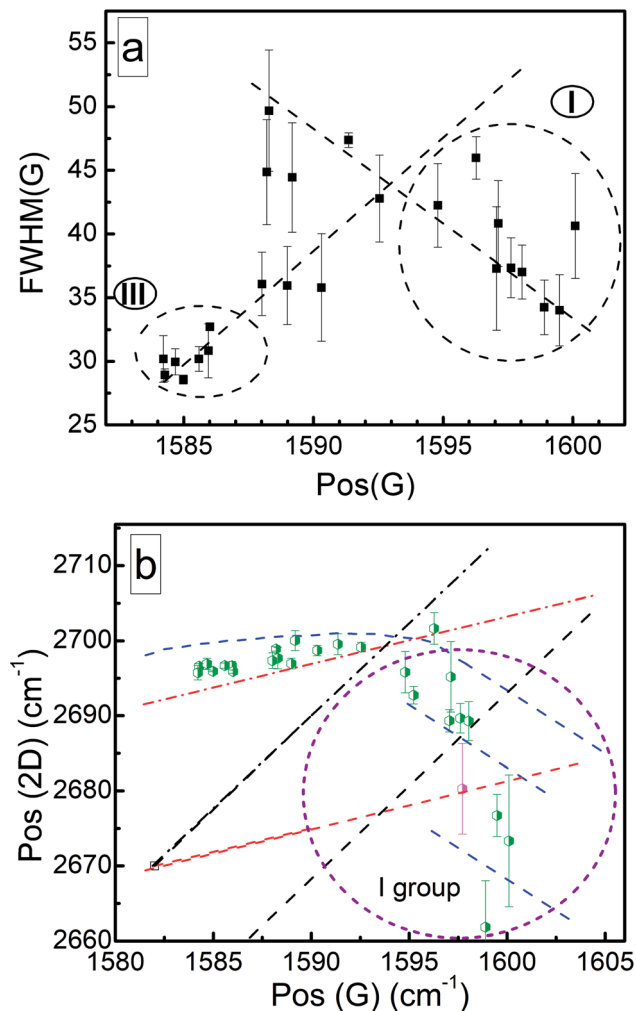


Fig. 7 Plots of (a) FWHM(G) vs. Pos(G) and (b) Pos(2D) vs. Pos(G). In the Pos(2D) vs. Pos(G) plot, the black dashed-dotted line refers to undoped strained graphene (plotted according to the method in ref. 44). The black dotted line refers to p-type doped strained graphene (constant hole concentration and different stress levels) (plotted according to ref. 44). The red dotted line refers to unstrained p-type graphene (plotted according to ref. 44). The red dashed-dotted line refers to p-type doped strained graphene (constant stress level and different hole concentrations) (plotted according to the method in ref. 44). The blue dotted lines refer to n-type doped graphene (plotted according to ref. 44). The open square symbols refer to unstrained and undoped graphene.⁴⁴ The green diamonds refer to the experimental points of graphene synthesized on n-type silicon substrates. The magenta-colored diamonds refer to the experimental points of graphene synthesized on p-type Si(100) (the synthesis conditions are the same as those for sample no. 12).

exposed to the air, and the adsorption of different ambient molecules is possible. This can induce p-type doping.^{68–70}

4 Discussion and conclusions

The mechanisms causing the observed results can be generalized considering the effects of the technological conditions on the graphene structure and morphology and related physical

and chemical processes. Vertical graphene growth is promoted by two main processes (Fig. 8).

The temperature gradients and thermal stress increase with an increase in the synthesis temperature. At elevated temperatures, thermal stress is released by the partial detachment of graphene flakes from the substrate and curling. Some time is necessary for stress accumulation when the critical temperature is reached. In addition, it seems that the diffusion of radicals to the tops of vertical graphene flakes is more critical than the density of radicals on the surface.⁷⁷ Another factor is ion bombardment and electric field effects,⁷⁷ which are enhanced with an increase in plasma power.³⁰ Ion bombardment generates surface defects and CH_x radicals.⁷⁷ If radicals are adsorbed on a defected flake edge, they can diffuse along the edge with a low probability of evaporation or transition to a side surface. The nanoflake sharp edges enhance the electric field, intensifying reactions due to ion focusing.⁷⁷ Without the activation of these mechanisms, planar graphene grows.⁷⁷ A decrease in pressure promotes vertical graphene growth due to increased gas ionization.¹⁸ This is, again, more important than the CH_x radical density on the surface.⁷⁷ The growth of planar graphene due to increased methane flow shows that an increased hydrocarbon species flux promotes planar graphene growth.

A decrease in the graphene layer number due to elevated temperature and a long synthesis time can be explained based on an increase in the carbon desorption rate with temperature.⁶⁴ Thus, the vertical graphene flakes were thinner than the planar ones in most cases. Defect density changes should be analyzed, considering the presence of different defects. Notably, in most samples, boundary defects dominated. In the planar graphene samples, on-site defects can be formed. A low synthesis temperature, low power, and elevated methane flux may benefit their creation. Vacancies were found in samples containing vertical/planar graphene and vertical graphene. A combination of high power and increased temperature or a combination of low power with increased temperature and a longer growth time activated vacancy creation. Thus, carbon desorption and ion irradiation are essential processes.

In the case of planar graphene, n-type self-doping was revealed. This was explained based on charge transfer from the substrate. The growth of vertical graphene flakes resulted in a decreased contact area between the graphene flakes and the substrate. Therefore, charge transfer from the substrate was suppressed. In the case of vertical graphene, no doping or p-type doping was found. In this case, p-type doping can be explained based on an increased contact area between graphene and air, and more molecules can be adsorbed.

In conclusion, we were able to control the orientation of growing graphene flakes *via* adjusting the technological deposition process conditions. High temperature, high plasma power, and a prolonged synthesis time promoted vertical graphene growth due to increased temperature gradients, thermal stress, ion bombardment, and elevated electric field effects. Without these mechanisms, planar graphene grows at lower synthesis temperatures and lower plasma powers. The graphene layer number decreased with elevated temperatures and longer synthesis times because the carbon desorption rate increased

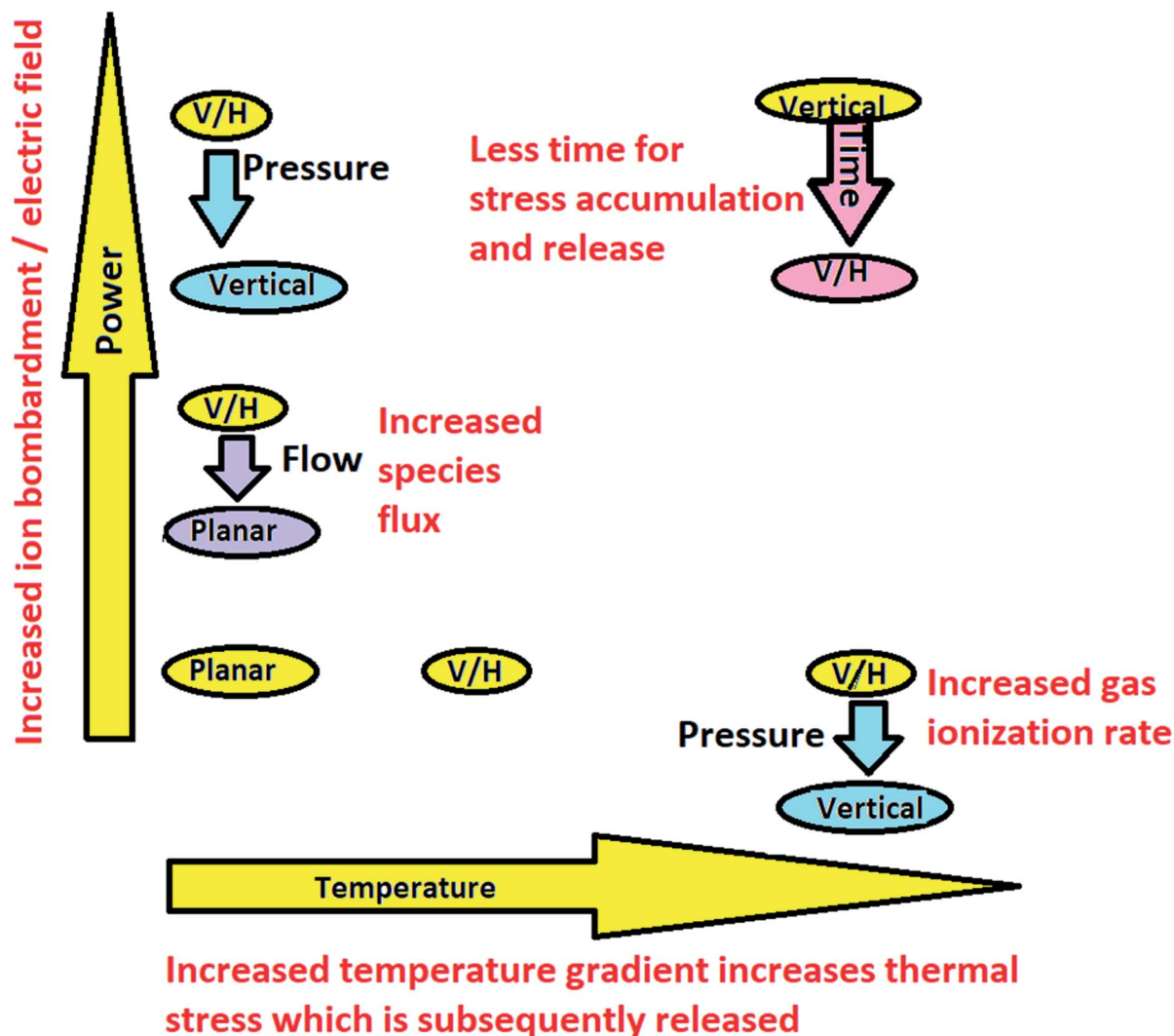


Fig. 8 The dependence of the synthesized graphene flake orientation on the synthesis conditions. V/H refers to a vertical and planar graphene mixture.

with temperature. In most samples, boundary defects prevailed. A low synthesis temperature, low power, and elevated methane gas flow were beneficial for the formation of some on-site defects. Vacancy creation was activated by high plasma power, an increased growth temperature, and a shorter synthesis time. Substrate-based charge transfer resulted in planar graphene n-type self-doping. In the case of vertical graphene, no doping or p-type doping was found. The adsorption of more molecules can explain this, due to the increased contact area between graphene and air.

Author contributions

Š. Meškiniš: conceptualization; visualization; writing – original draft; writing – review and editing; project administration; funding acquisition; A. Vasiliauskas: investigation; A. Guobienė: investigation; visualization; M. Talaikis: investigation; visualization; G. Niaura: investigation; writing – review and editing; R. Gudaitis: conceptualization; investigation.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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