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Tailoring the stoichiometry of C_3N_4 nanosheets under electron beam irradiation†

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Two-dimensional polymeric graphitic carbon nitride ($\text{g-C}_3\text{N}_4$) is a low-cost material with versatile properties that can be enhanced by the introduction of dopant atoms and by changing the degree of polymerization/stoichiometry, which offers significant benefits for numerous applications. Herein, we investigate the stability of $\text{g-C}_3\text{N}_4$ under electron beam irradiation inside a transmission electron microscope operating at different electron acceleration voltages. Our findings indicate that the degradation of $\text{g-C}_3\text{N}_4$ occurs with N species preferentially removed over C species. However, the precise nitrogen group from which N is removed from $\text{g-C}_3\text{N}_4$ ($\text{C}-\text{N}-\text{C}$, $=\text{NH}$ or $-\text{NH}_2$) is unclear. Moreover, the rate of degradation increases with decreasing electron acceleration voltage, suggesting that inelastic scattering events (radiolysis) dominate over elastic events (knock-on damage). The rate of degradation by removing N atoms is also sensitive to the current density. Hence, we demonstrate that both the electron acceleration voltage and the current density are parameters with which one can use to control the stoichiometry. Moreover, as N species were preferentially removed, the *d*-spacing of the carbon nitride structure increased. These findings provide a deeper understanding of $\text{g-C}_3\text{N}_4$.

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Introduction

Graphitic carbon nitride ($\text{g-C}_3\text{N}_4$) is regarded as the most stable polymeric allotrope of the carbon nitride family under ambient conditions.¹ Due to the semiconducting and exceptional photocatalytic properties of $\text{g-C}_3\text{N}_4$,² it holds great promise for use in several applications,^{3,4} including catalysis,⁵ energy conversion,^{6,7}

biosensing,⁸ and bioimaging.⁹ Moreover, $\text{g-C}_3\text{N}_4$ can be used for the preparation of metal nitrides nanoparticles, for example, GaN and VN,¹⁰ as well as metal carbide nanoparticles, for instance, NbC, TaC, and WC.¹¹

A bandgap of 2.7 eV renders $\text{g-C}_3\text{N}_4$ powder photoluminescent when dissolved in solvents and irradiated by a light source with a wavelength of ~ 450 nm.¹² The photoluminescence of $\text{g-C}_3\text{N}_4$ nanosheets and quantum dots can be quenched when in contact with certain ions, thus making it a promising material for chemical sensing.^{13–15} It has also been reported that $\text{g-C}_3\text{N}_4$ can absorb near-infrared light¹⁶ generating heat when exposed to near-infrared light for an extended period. This suggests the potential for $\text{g-C}_3\text{N}_4$ application in photodynamic therapies to treat tumors by selectively overheating specific tumorous tissue regions. Because of its semiconducting properties,¹⁷ $\text{g-C}_3\text{N}_4$ also shows promise in nanoelectronics, energy conversion and storage, photocatalysis, among other potential applications.^{18–20}

In an ideal case, $\text{g-C}_3\text{N}_4$ nanosheets are comprised solely of covalent bonds between carbon and nitrogen atoms in an sp^2 -hybridized configuration. In reality, however, it has also been reported that hydrogen is present, which is associated with primary and/or secondary amine groups at termination edges and on the surface.¹² The presence of these functional groups on nanosheet surfaces is mostly linked to incomplete polymerization produced during the synthesis, which produces $\text{g-C}_3\text{N}_4$ nanosheets with defects distributed across the lattice.

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These defects can either be an advantage, *e.g.*, to facilitate further functionalization and tailor the desired properties for bespoke applications, or they can be a source of instability within the g-C₃N₄ crystalline structure, which could reduce their performance for certain applications. For example, it has been reported that the morphology and surface area of g-C₃N₄ are important parameters for controlling photocatalytic performance.²¹ Thus, understanding the actual structure of g-C₃N₄ and aspects regarding its stability is of great importance.

Moreover, its potential for industrial use can only be realized once an in-depth understanding of the stability of g-C₃N₄ under different conditions is obtained. Typically, material stability is investigated through mechanical, chemical, and thermal stress testing, which is usually conducted with bulk quantities. The rise in diverse applications that require extreme precision and reliability, such as in biomedicine and space exploration, demands characterization and quality controls under specific conditions at smaller scales, even down to the atomic level. For example, the biomedical use of g-C₃N₄ nanosheets requires that the material interacts with specific proteins and enzymes to develop its functional ability, whereas employing g-C₃N₄ in nanoelectronic devices for use in space exploration may necessitate tolerance of the material to certain doses of radiation. Although it is known that g-C₃N₄ has high chemical and thermal stability (up to 500 °C at pH from acidic to alkaline),^{22,23} its stability under electron beam irradiation is, to the best of our knowledge, little known.

Transmission electron microscopy (TEM) is a structural and spectroscopic characterization tool that allows the investigation of materials at the atomic level. Moreover, it permits control of the electron irradiation level (dose) that the specimen receives. This can be achieved by varying the electron acceleration voltage and by tuning the electron current density. This is important because the mechanisms through which electrons interact with the specimen vary dramatically by changing the conditions of the incident electron beam. The electron/specimen interaction can be of an inelastic or elastic nature, and both scattering processes are the result of Coulomb interactions between the incident electron beam with either the atomic nuclei of the specimen (elastic interaction) or the electron cloud surrounding the atom nuclei of the specimen (inelastic interaction). These physical processes are important for imaging and characterizing samples, but they can simultaneously damage the specimen, which in many cases modifies or even degrades the crystalline structure of the material.^{24,25}

In the case of inelastic scattering, the energy of incident electrons is transferred to atoms in the specimen. This energy is converted into kinetic energy and momentum, leading to the loss of mass or atom displacement, known as radiolysis. In elastic interactions, there is no change in the incident beam energy but only a deflection of incident electrons due to electrostatic interactions with the nuclei of atoms. To maintain constant energy and momentum, incident electrons and atoms in the specimen deflect. If deflection from atoms of a specimen is above a threshold energy (which is dependent on the material in question), then atoms can be displaced in a process known

as knock-on damage.^{25,26} Whether inelastic or elastic interactions dominate is not only highly dependent on the energy and current density of the incident electron beam but also on the nature of the specimen being investigated.²⁷

The use of high acceleration voltages leads to knock-on damage dominating because electrons carrying more energy have a higher probability of interacting elastically with the specimen. This effect can be even more prominent when exposing thin materials to high-energy beams²⁸ and can be easily observed by imaging graphene with acceleration voltages above its knock-on damage threshold energy (86 kV).^{25,29} The use of low acceleration voltages reduces knock-on damage but may increase damage due to radiolysis. This is because low electron voltages increase the interaction cross-section of the incident electrons, which increases the probability of inelastic interactions taking place. Therefore, there is no single set of optimum operating conditions for TEM measurements and a balance of the operating conditions must be determined for each type of sample.^{28,30,31}

Herein, we investigate the stability of g-C₃N₄ nanosheets under different electron irradiation conditions for different TEM acceleration voltages (300, 80, 60, and 30 kV) and current densities (1.0×10^{-11} – 6.5×10^{-10} nA nm⁻²). Our results suggest that the stability of g-C₃N₄ is highly dependent on the operating conditions of electron beam exposure, which provides an exciting approach for tailoring the nitrogen content and *d*-spacing of g-C₃N₄, and ultimately its properties.

Experimental section and methods

Synthesis of g-C₃N₄

Crystalline g-C₃N₄ nanosheets were synthesized by a two-step annealing of the NH₄SCN precursor under deaerated conditions. NH₄SCN (5 g) was heated at 230 °C for 12 h in a tubular furnace (ramping rate: 1 °C min⁻¹) under an Ar atmosphere (flow rate: 50 mL min⁻¹). The obtained white powder was then washed with deionized (DI) water to remove residual ions. The washed powder was dried at 100 °C for 12 h, loaded into a silica glass tube, evacuated to 10⁻⁵ mbar, and then sealed. The ampule was eventually heated to 630 °C for 12 h in an oven at a ramping rate of 1 °C min⁻¹ to obtain g-C₃N₄.^{32,33}

TEM sample preparation

A nominal amount of g-C₃N₄ (20 µg) was dispersed in 0.5 mL of isopropanol in a bath sonicator at room temperature to improve the dispersibility and to exfoliate the g-C₃N₄ flakes into thin layers. Sonication separates and exfoliates the flakes, allowing the preparation of individual, freestanding g-C₃N₄ nanosheets (a few atom layers thick). To explore the efficiency of the procedure, a mixture of isopropanol containing the g-C₃N₄ flakes was sonicated for different time intervals (180, 600, 900, and 1800 s) and shown in Table S1 of the ESI.† The sonicated samples were left to stand for 600 s to allow the precipitation of clusters and larger agglomerations. For each sonication time, a small volume (3 µL) from the supernatant

was collected with a pipette and drop-coated onto a TEM grid of lacey carbon film. The samples were then left to dry freestanding (on the tip of cross-locking tweezers) on a hot plate at 40 °C for 1 h prior to TEM measurements.

TEM imaging

The samples were imaged using a double-aberration-corrected FEI Titan microscope operating at 300 and 80 kV. The results obtained at acceleration voltages of 60 and 30 kV were acquired using a spherical-aberration-corrected Titan Themis microscope. For each acceleration voltage, the stability of the g-C₃N₄ flakes was investigated by varying the current densities from high (10^{-3} nA nm⁻²) to low (10^{-5} nA nm⁻²). The electron energy loss spectroscopy (EELS) spectra were acquired locally with the microscope operating in the scanning transmission electron microscopy (STEM) mode at each corresponding acceleration voltage with a camera length of 91 mm and a convergence angle of 21.4 mrad.

Results and discussion

As-synthesized g-C₃N₄ nanosheets were isolated as bright yellow powders and shown in Fig. S1A in the ESI.† Careful analysis of the results indicated that even a short sonication time was sufficient to disperse the flakes (Fig. S1B–E, ESI†), as no significant differences between the samples sonicated for 180 s and 1800 s were identified (Fig. S2, ESI†). Since sonication is known to damage two-dimensional (2D) materials, we used samples sonicated for 180 s for the electron irradiation experiments to minimize the possibility of such damage.

After sonication, the g-C₃N₄ dispersion was left to settle for 600 s to allow larger agglomerates to precipitate. A small aliquot (3 µL) of the upper portion of the suspension was then removed with a pipette and drop-coated onto a standard TEM grid with a lacey carbon film. TEM grids containing the g-C₃N₄ dispersion were dried in air on the tip of cross-locking tweezers left on top of a hot plate at 40 °C for 1 h.

Initial characterization of the material begins with determining whether the as-synthesized g-C₃N₄ sample is crystalline and then the identification of its crystal parameters. The lattice parameters of g-C₃N₄ are highly dependent on the C:N ratio.³⁴ For example, Fina *et al.*³⁴ measured the *d*-spacing of the (100) plane for their g-C₃N₄ sample with a C:N ratio of 0.68 as 0.68 nm, whereas the *d*-spacing of the same (100) plane for fully polymerized triazine and tri-*s*-triazine (C:N ratio 0.75) was smaller (0.41 nm and 0.615 nm, respectively) in other studies,^{22,35,36} suggesting that lower g-C₃N₄ C:N ratios lead to increased *d*-spacings. This is vitally important because it indicates that varying the nitrogen content affects the *d*-spacing, *viz.*, it provides a pathway to tailor the properties of g-C₃N₄.

TEM images of our g-C₃N₄ specimen (at 300 kV) are shown in Fig. 1. The g-C₃N₄ nanosheets are indeed crystalline (Fig. 1A), with lattice fringes easily identified (see inset). The Fourier filtered image shows two crystal spacings at high resolution (Fig. 1B). The corresponding fast Fourier transform (FFT) image is shown in Fig. 1C. To minimize the exposure of the sample to the electron beam, all of the imaging parameters were determined on a section of the specimen near the region to be evaluated. Once the TEM acquisition parameters were set, the stage was then moved to the nearby non-irradiated region, and an image was immediately acquired. The *d*-spacing values not only provide feedback on the different C:N ratios of g-C₃N₄ but also indicate the presence of defects in the lattice. In this case, the obtained value (attributed to the (100) orientation) is taken as the basis of the “non-irradiated” sample.

Specimen stability studies at an acceleration voltage of 300 kV reveal that the g-C₃N₄ nanosheets were sensitive to electron irradiation. At low magnification (<165 000 \times), it was possible to image the sample without visible changes to the flakes. This was achieved with a current density of up to 3.5×10^{-10} nA nm⁻². However, when the magnification was increased to 210 000 \times , the g-C₃N₄ nanosheets rapidly lost their crystalline structure during the same acquisition time (1 s). This happens because the current density is approximately two times larger

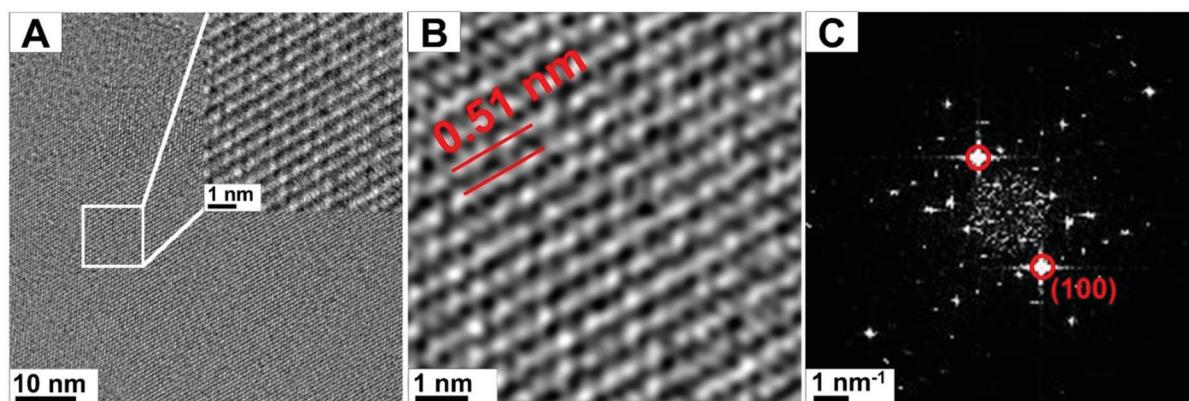


Fig. 1 Low- and high-resolution TEM images of C₃N₄ nanosheets. (A) TEM crystalline image of the g-C₃N₄ lattice. The inset is of a cropped region showing the lattice in more detail. (B) Fourier filtered image of the inset presented in A. The lattice parameters were measured directly from the TEM image. (C) FFT of the image in B showing the lattice reflex attributed to the (100) plane. TEM images were acquired using an acceleration voltage of 300 kV and a current density of 4.0×10^{-11} nA nm⁻².

(6.5×10^{-10} nA nm $^{-2}$) for a magnification increase from 165 000 \times to 210 000 \times . Successful imaging of g-C₃N₄ with a magnification of 210 000 \times was only possible by decreasing the current density to 4.0×10^{-11} nA nm $^{-2}$ (which was achieved by increasing the illumination area). At this low current density, the g-C₃N₄ lattice remained significantly more stable and could be irradiated for up to two minutes, which is sufficient for image capture with longer acquisition times. These results indicate that the current density is an important electron beam parameter concerning the stability of g-C₃N₄ nanosheets. Examples of the differences observed for different magnifications (and therefore current densities) are presented in Fig. 2A and B, which shows high-resolution TEM images of a g-C₃N₄ flake and its corresponding FFT, respectively. The image shown in Fig. 2A was acquired using a current density of 6.0×10^{-11} nA nm $^{-2}$. Suitable images with current densities up to 8.5×10^{-11} nA nm $^{-2}$ could be obtained, while for higher current densities, degradation was readily observed. An example of this degradation can be seen in Fig. 2D and E; panel D clearly shows the amorphous nature of the material after electron irradiation, as does the corresponding FFT in panel E, which lacks any reflexes and confirms the amorphous nature of the heavily irradiated samples. Typically, the transition to an amorphous state occurred after ~ 20 s irradiation for current densities in the range of 4.0×10^{-10} to 6.0×10^{-10} nA nm $^{-2}$ (one order of magnitude larger than that used in Fig. 2A). We also examined the stoichiometric changes between C and N in the material when transitioning from its crystalline state to an amorphous state through EELS probing studies. A representative example from a crystalline g-C₃N₄ state is shown in Fig. 2C, which displayed stoichiometric values of 62% C and 38% N. For the amorphous material obtained after extended electron irradiation, the stoichiometric values

change to 96% C and 4% N (Fig. 2F). The EELS data, therefore, suggest a preferential loss of nitrogen from the g-C₃N₄ nanosheets during the transition process, ultimately yielding amorphous carbon.

To further investigate the amorphization process and the stability of the g-C₃N₄ nanosheets, studies using an electron acceleration voltage of 80 kV were conducted. Reducing the acceleration voltage should, in principle, reduce the role of knock-on damage.^{25,28} The same experimental imaging conditions were used as were implemented at an electron acceleration voltage of 300 kV. Interestingly, the degradation of g-C₃N₄ was similar for both electron acceleration voltages (Fig. 3). Fig. 3A–C shows low magnification (31 000 \times –99 000 \times) TEM images for current densities in the range of 9.5×10^{-11} to 3.5×10^{-10} nA nm $^{-2}$, during which the g-C₃N₄ nanoflakes remained crystalline for up to 180 s. However, when the magnification was increased (165 000–210 000 \times), the total current density increased by one order of magnitude and the nanosheet transition to an amorphous state was again rapid and occurred within 20 s. EELS studies showed that this is related to the loss of nitrogen species in the material (for example shown in Fig. 1F).

To study the electron-beam-driven transition process, we also examined specimen stability at two lower acceleration voltages, namely, 60 and 30 kV. At lower acceleration voltages in the same current range explored at 80 and 300 kV (9.5×10^{-11} to 6.0×10^{-10} nA nm $^{-2}$), the crystalline g-C₃N₄ nanosheets became amorphous after exposure for 15 s with a current density of 6.0×10^{-10} nA nm $^{-2}$. This suggests that the degradation of the g-C₃N₄ material occurs more readily at electron acceleration voltages below 80 kV.

To better comprehend the irradiation susceptibility of g-C₃N₄ at low electron acceleration voltages, a series of

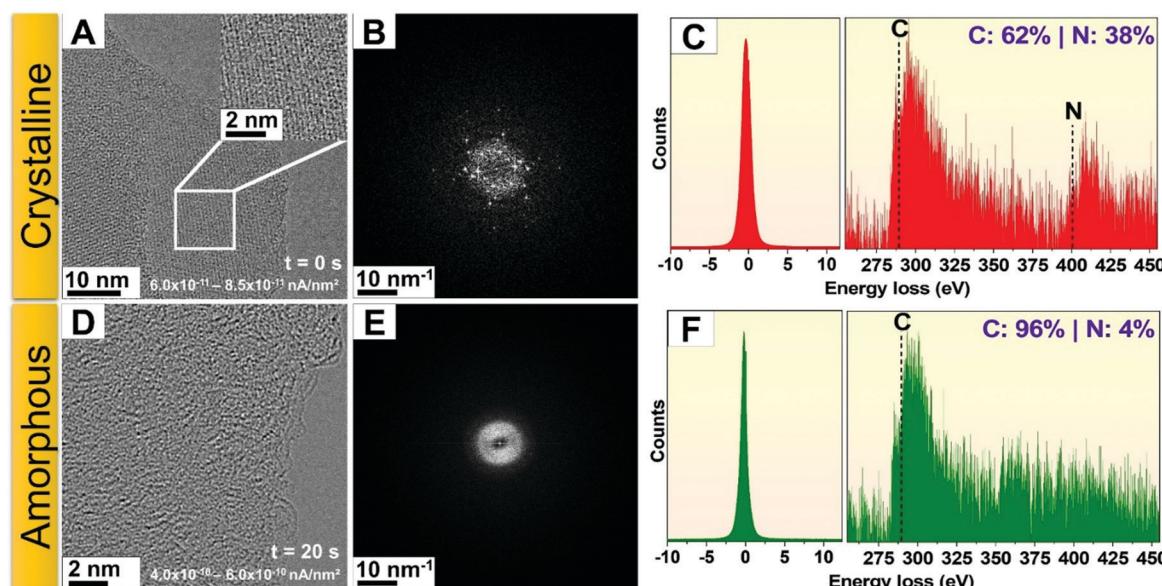


Fig. 2 Sample of g-C₃N₄ sonicated for 180 s in isopropanol and imaged at 300 kV. (A) TEM image of the g-C₃N₄ nanosheet indicating that it is crystalline, as highlighted in the inset and in the FFT shown in panel (B). (C) EELS data showing the presence of carbon and nitrogen. (D) TEM image showing that after 20 s of electron irradiation, the g-C₃N₄ nanosheet quickly loses its crystallinity and becomes amorphous. (E) FFT showing the absence of reflexes in the amorphous state. (F) EELS data indicating that nitrogen is lost after electron irradiation.

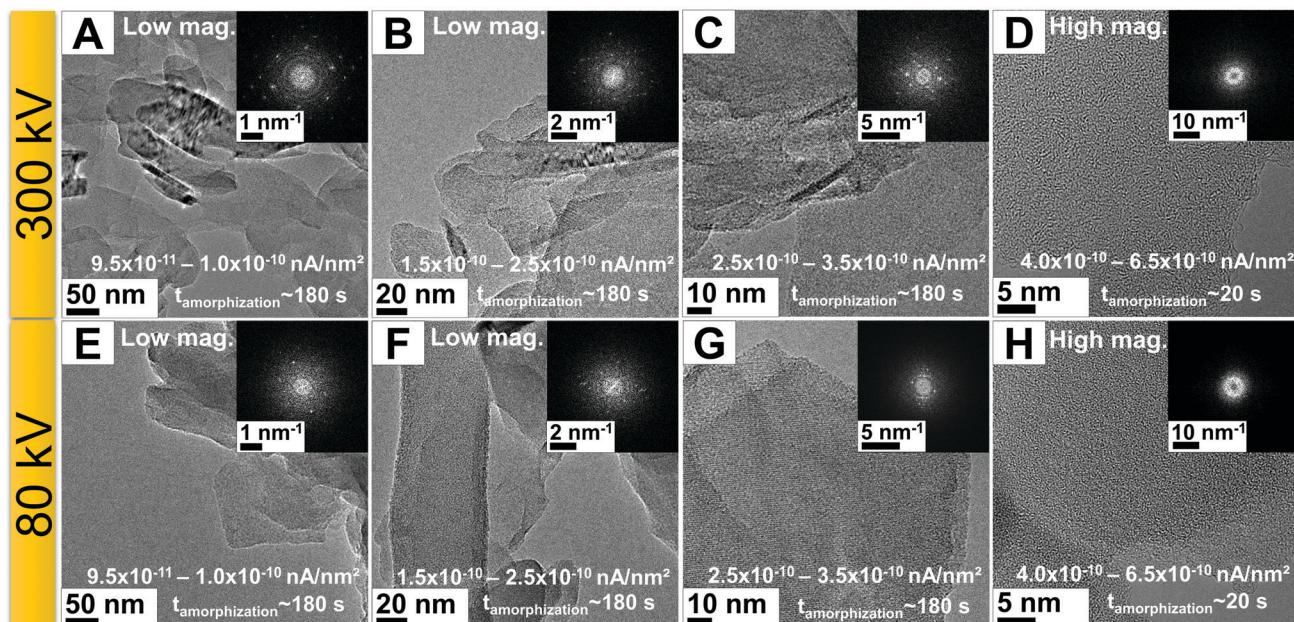


Fig. 3 TEM images of $\text{g-C}_3\text{N}_4$ under different electron acceleration voltages. (A)–(D) TEM images of $\text{g-C}_3\text{N}_4$ at 300 kV. (A)–(C) Low magnification images show the presence of crystalline $\text{g-C}_3\text{N}_4$ nanosheets, as can be seen in the FFT insets. (D) Amorphization of the $\text{g-C}_3\text{N}_4$ nanosheet within 20 s after increasing current density. (E)–(H) TEM images of $\text{g-C}_3\text{N}_4$ imaged at 80 kV. (E)–(G) Low magnification images also show that the $\text{g-C}_3\text{N}_4$ nanosheets are crystalline. (H) Amorphous $\text{g-C}_3\text{N}_4$ after 20 s of electron irradiation at higher current density.

experiments using low current densities (1.0×10^{-11} to $4.0 \times 10^{-11} \text{ nA nm}^{-2}$) were conducted at acceleration voltages of 300, 80, 60, and 30 kV. At these lower current densities, successful image acquisition was only possible with longer exposure times (four times higher, 4 s). At low acceleration voltages, the image resolution decreased owing to increased electron scattering in the specimen and the predominance of lens aberrations.³⁷ The resolution in TEM is limited by spherical (Cs) and chromatic (Cc) aberrations of the objective lenses.³⁸ While the Cs occurs because electrons reach the optical axis at points slightly shifted from the image plane, Cc arises from energy spread in the electron beam, which causes electrons to focus at different image planes.³⁸ At lower acceleration voltages, the Cc predominates. To overcome these image quality limitations at low acceleration voltages, it is essential to use a transmission electron microscope equipped with a Cc correction lens or an electron-beam monochromator.^{37,39,40} To obtain high-resolution images (Fig. 4), two imaging parameters (acceleration voltage and current density) were used. When a lower acceleration voltage was used, a lower current density was needed. Since electrons with low acceleration have a higher probability of interacting with the specimen (leading to an increase in electron-driven damage), one needs to reduce the current density to reduce the damage due to electron irradiation. However, this strategy can lead to poor image quality in addition to the lower resolution caused by the change in acceleration voltage. One can easily see that the image obtained at 300 kV (Fig. 4A) is of higher quality (is clearer) compared to that collected at 30 kV (Fig. 4D).

The microscopes used in this study are equipped with a Cs corrector at the objective lens as well as an electron

monochromator. The electron monochromator reduces the electron energy spread and consequently reduces the Cc.^{37,38} Good-quality imaging at 60 kV (Fig. 4B) and 30 kV (Fig. 4D) was only possible by reducing the energy spread of the electron beam (also known as electron beam monochromatization). With an electron acceleration voltage of 60 kV, it was possible to acquire images with reasonable image quality, whereas for an acceleration voltage of 30 kV, the image quality was visually limited, even when using a monochromated beam. Even with only a slight increase in current density (from 1.0×10^{-11} to $4.0 \times 10^{-11} \text{ nA nm}^{-2}$), the image quality deteriorates for the lowest acceleration voltages tested (60 and 30 kV) (Fig. S3, ESI†). For example, imaging with a current density of $6.0 \times 10^{-11} \text{ nA nm}^{-2}$ (Fig. S3C and D, ESI†) led to image qualities significantly worse than when using a current density of $1.0 \times 10^{-11} \text{ nA nm}^{-2}$ (Fig. S3C' and D', ESI†).

As mentioned in the introduction, there is an intrinsic correlation between the beam energy and the type of interaction with the specimen (either elastic or inelastic). Electron beams with high energies will favor elastic interactions (knock-on damage), whereas low-energy beams favor inelastic interactions (radiolysis). Therefore, the hypothesis to explain the observed phenomena is that with a decrease in the acceleration voltage, albeit with beam energy below the knock-on threshold energy, the energy transfer (inelastic interaction) from the incident beam to the specimen is high. At higher current densities, the greater the irradiation dose of the specimen, the more rapid the degradation of the lattice. As mentioned previously, this can produce defects on the $\text{g-C}_3\text{N}_4$ surface that are regions of instability in the lattice, which contribute to faster degradation of the material.

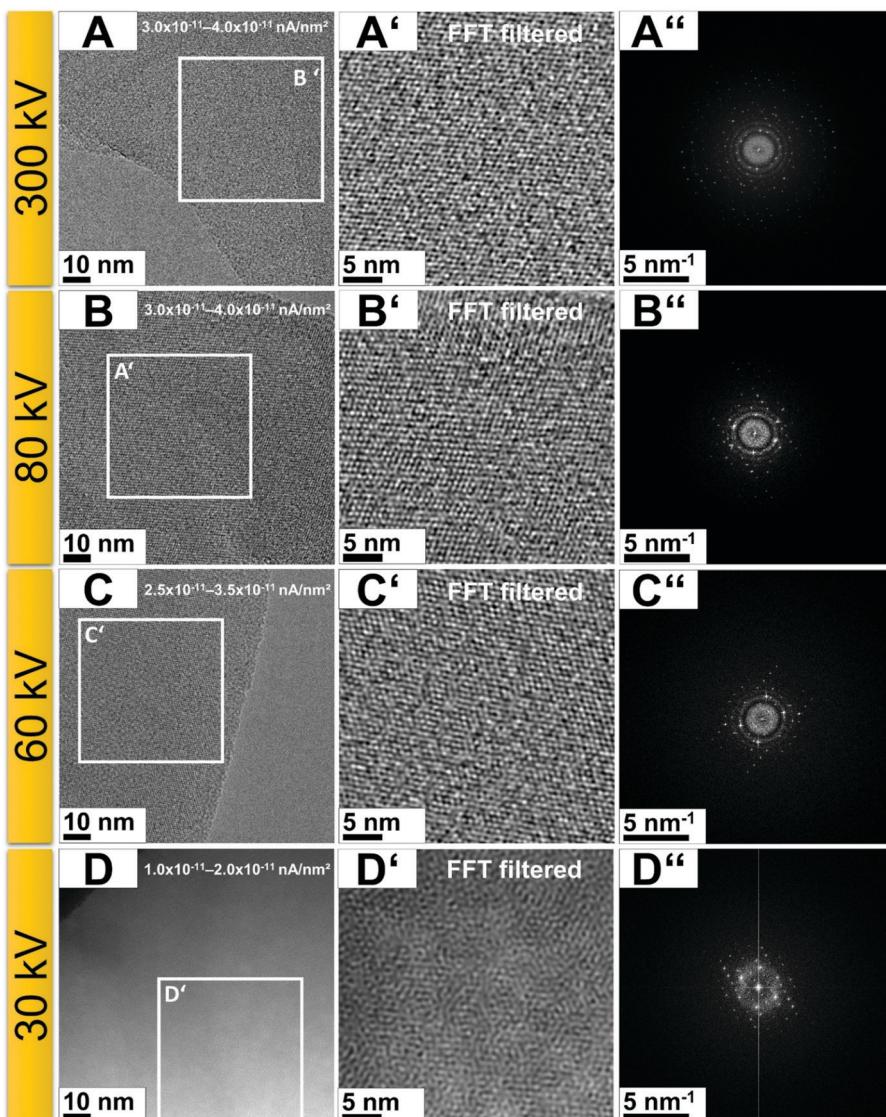


Fig. 4 Differences in imaging C_3N_4 nanosheets with decreasing electron beam energy. TEM images of $\text{g-C}_3\text{N}_4$ at (A–A'') 300 kV, (B–B'') 80 kV, (C–C'') 60 kV, and (D–D'') 30 kV. (A'), (B'), (C'), and (D') Fourier filtered images of the boxed regions in the TEM images, for 300, 80, 60, and 30 kV, respectively. The corresponding FFT images are shown in panels A'', B'', C'' and D''. In all cases, TEM imaging of $\text{g-C}_3\text{N}_4$ was performed using a low current density (1.0×10^{-11} to $4.0 \times 10^{-11} \text{ nA nm}^{-2}$).

To better illustrate our observations, plots of the degradation time (amorphization time) for different electron acceleration voltages and currents are provided in Fig. 5. In Fig. 5A, amorphization of the $\text{g-C}_3\text{N}_4$ nanosheets is plotted against the acceleration voltage for two low current densities (on the order of $10^{-11} \text{ nA nm}^{-2}$). It can be seen that for current densities between 1.0×10^{-11} and $3.0 \times 10^{-11} \text{ nA nm}^{-2}$, the nanosheets become amorphous over a time frame of approximately 120 s. With just a small increase in the current density (from 3.0×10^{-11} to $4.0 \times 10^{-11} \text{ nA nm}^{-2}$), the amorphization time decreases from 120 s to 30 s (reduction of 75%) for an acceleration voltage of 30 kV. Fig. 5B displays the amorphization time plotted as a function of the current density, which indicates that low-energy electron beams (30 and 60 kV) are more efficient at amorphizing the crystalline $\text{g-C}_3\text{N}_4$ nanosheets to amorphous carbon (due to the

preferential release of N species). The preferential release of N can be thought of as akin to N defects in graphene. In graphene, the presence of dopants changes bond lengths as well as the local chemical properties, and are thus a source of instability that can facilitate degradation during electron irradiation.³⁸ This is also observed with hexagonal boron nitride (h-BN), where B is preferentially removed over N from the graphene-like lattice.^{39–41}

Moreover, since the C:N ratio (stoichiometry) may play an important role in the nature of the $\text{g-C}_3\text{N}_4$ lattice, and given that N appears to be ejected preferentially, we decided to investigate the loss of N in further detail. EELS data were collected for acceleration voltages of 300, 80, and 60 kV, as shown in Fig. 6A–C. Unfortunately, measurements for 30 kV were not possible owing to hardware limitations. The EELS spectra show that in all cases, the carbon peak (284 eV) and the nitrogen peak



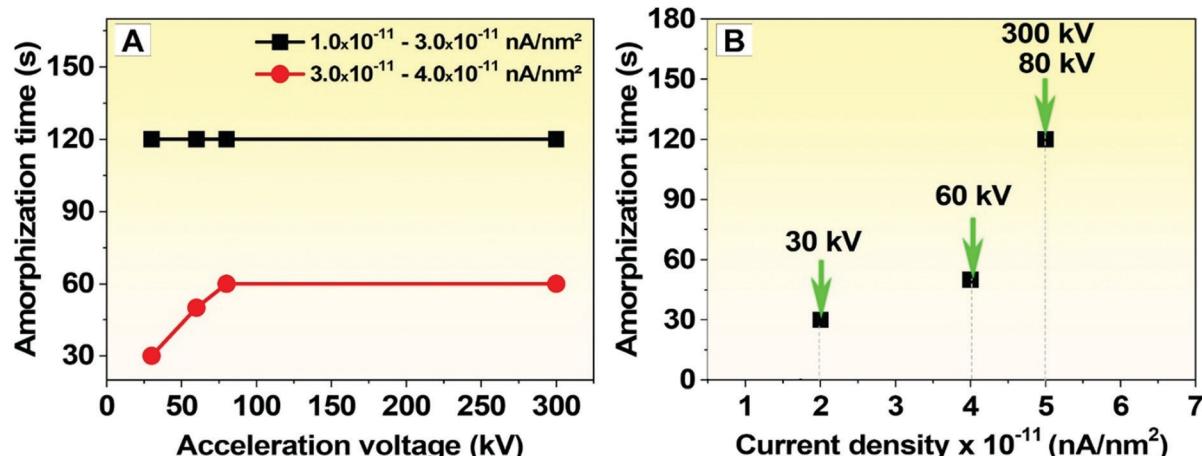


Fig. 5 Comparison of the amorphization time, acceleration voltage, and current density. (A) Typical amorphization time plotted with respect to the microscope acceleration voltage for current densities above and below $3.0 \times 10^{-11} \text{ nA nm}^{-2}$. (B) Typical amorphization time plotted according to microscope screen currents below $3.0 \times 10^{-11} \text{ nA nm}^{-2}$, indicating that g-C₃N₄ is more stable at higher electron acceleration voltages.

(401 eV) are present. However, the intensity of each peak varies for the different acceleration voltages used. Because the relative intensity of the EELS peaks is related to the elemental content in the specimen, the atomic content of carbon and nitrogen can be determined. Quantification of the EELS data suggests that the carbon content increases as the electron acceleration voltage decreases (Fig. 6D). The carbon and nitrogen content

(given in molar %) measured at 300 kV was found to be 58% ($\pm 1\%$) and 42% ($\pm 1\%$), respectively. At 60 kV, the atomic content of the investigated elements was 79% ($\pm 1\%$) for carbon and 21% ($\pm 1\%$) for nitrogen.

Since the interaction of the electron beam with the g-C₃N₄ nanosheets induces their transformation into amorphous carbon for all of the electron acceleration voltages tested – even

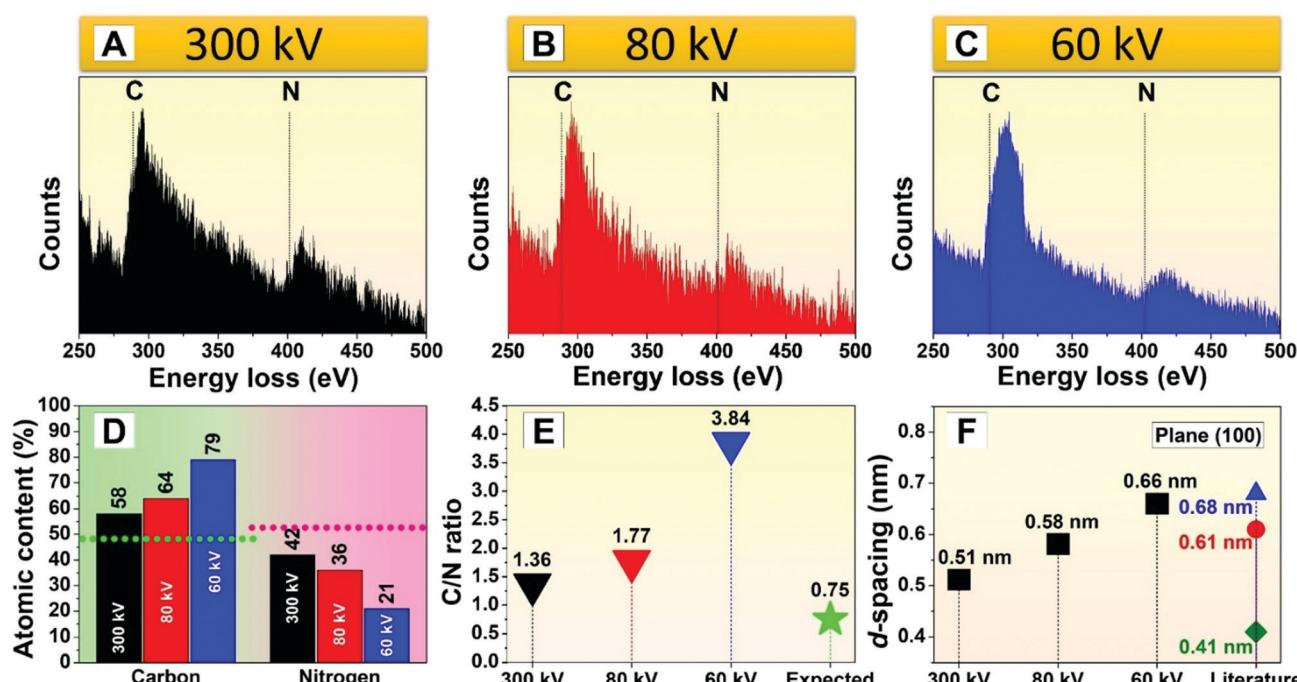


Fig. 6 Chemical analysis of C₃N₄ nanosheets after irradiation with different electron beam energies. (A)–(C) EELS spectra of g-C₃N₄ measured using different acceleration voltages. Carbon and nitrogen were detected in all measurements. The atomic content for each measurement was quantified, with the relative amount of carbon increasing with decreasing acceleration voltage, whereas the opposite occurs for nitrogen, which decreases to a greater extent than carbon, as shown in panel (D). The theoretical atomic percentage for carbon and nitrogen for fully polymerized g-C₃N₄ are displayed as green and pink horizontal dotted lines, respectively. (E) The C : N ratio increases as the acceleration voltage decreases for 300, 80, and 60 kV. The green star indicates the C : N ratio described in the literature for fully polymerized g-C₃N₄. (F) The *d*-spacing of the (100) plane measured for 300, 80, and 60 kV compared to values published in the literature (see text).

with short exposure times – the accuracy of EELS quantification is therefore limited. This should be kept in mind when interpreting the data in comparison to the theoretical values of a completely non-irradiated flake. Since higher electron acceleration voltages interact less with the specimen, EELS data collected at 300 kV will be more accurate; however, in practice, we find that the values still differ from the theoretical values for fully polymerized $g\text{-C}_3\text{N}_4$ nanosheets (*i.e.*, 48.48% for carbon, green dotted line; and 51.52% for nitrogen, pink dotted line in Fig. 6D). These values are relative to the atomic mass percentage).

Electron beam-induced stoichiometric changes are related to structural changes. As mentioned in the introduction, $g\text{-C}_3\text{N}_4$ can be obtained with various degrees of polymerization^{41–43} depending on the precursor materials and synthetic conditions, which play a crucial role in the polymerization of the final product and its subsequent properties.⁴⁴ Our data show that under electron irradiation, there is a clear increase in the C:N ratio. The difference between the measured C:N³⁴ changes significantly with the selected electron acceleration voltage, as shown in Fig. 6E. Our estimated C:N ratio when using 300, 80, and 60 kV acceleration voltages is much higher than the value reported for the ideal fully polymerized $g\text{-C}_3\text{N}_4$ (C:N ratio of 0.75), especially when the 60 kV electron beam was used. Because lower acceleration voltages increase the interaction cross-section of the electron beam with the specimen, the probability of radiolysis-induced damage increases.^{28,30,31} This confirms our observations that lower acceleration voltages remove nitrogen atoms more easily and faster (<20 s) from $g\text{-C}_3\text{N}_4$. However, it does not indicate which nitrogen atom is removed from $g\text{-C}_3\text{N}_4$ (C–N–C, $\equiv\text{NH}$ or $-\text{NH}_2$). This would require detailed mass spectroscopy analysis of C, H, and N before and after irradiation of a large amount of $g\text{-C}_3\text{N}_4$ (25–50 mg), which is not feasible in this instance.

Our data also show changes in the lattice *d*-spacings as N is preferentially removed from the $g\text{-C}_3\text{N}_4$ specimen under irradiation. The literature suggests that fully polymerized $g\text{-C}_3\text{N}_4$ exhibits smaller *d*-spacings when compared to partially polymerized materials. Our systematic data confirmed this trend. In our study, the *d*-spacing corresponding to the (100) plane measured with an acceleration voltage of 60 kV (0.66 nm) is larger than that at 80 kV (0.58 nm) and 300 kV (0.51 nm), as shown in Fig. 6F. It should be noted that there is a lack of clarity for the assignment of X-ray diffraction (XRD) based diffraction patterns for $g\text{-C}_3\text{N}_4$.⁴⁵ For example, Wang *et al.*⁴⁶ calculated the XRD results from the different phases of $g\text{-C}_3\text{N}_4$ and assigned the *d*-spacing for the (100) plane (space group $P\bar{6}m2$) as 0.411 nm (Fig. 6F, green diamond). Tyborski *et al.*⁴⁷ assigned the same (100) plane of a heptazine-based $g\text{-C}_3\text{N}_4$ (space group $P\bar{6}_3cm$) with a *d*-spacing of 0.61 nm (Fig. 6F, red circle). Although both references point to the existence of different phases of $g\text{-C}_3\text{N}_4$, no investigation of the C:N ratio was considered. Experimental data by Fina *et al.*³⁴ assigned the (100) peak to a *d*-spacing of 0.68 nm (Fig. 6F, blue triangle), which is larger than the calculated values mentioned above. They attributed the larger *d*-spacing to their sample being a partially polymerized $g\text{-C}_3\text{N}_4$ structure (C:N ratio of 0.68).

Our results confirm that there is a correlation between the measured *d*-spacing and the C:N ratio. The electron beam produces structural damage in $g\text{-C}_3\text{N}_4$ across all examined acceleration voltages and current densities, changing the C:N ratio of $g\text{-C}_3\text{N}_4$ (degree of polymerization). This opens up new possibilities for tailoring the structure, and hence the properties, of $g\text{-C}_3\text{N}_4$ by selectively altering the *d*-spacing and C:N ratios with the use of electron beams. The ability to controllably engineer $g\text{-C}_3\text{N}_4$ to exhibit the desired function is attractive for specialized applications. For example, Gilbert *et al.*⁴⁸ demonstrated the possibility of creating defined pore sizes in multi-layer h-BN by controlling the vacancies introduced using an electron beam. They claim that such h-BN nanopores can be used for water desalination, molecular separation, and sensing. Similar possibilities can be expected with $g\text{-C}_3\text{N}_4$.

The enhancement of the catalytic properties of $g\text{-C}_3\text{N}_4$ has been demonstrated by introducing dopants,⁴⁹ which is a strategy that can also improve its potential in sensor development.⁵⁰ The synthesis of porous $g\text{-C}_3\text{N}_4$ using urea as a precursor showed promising results for the photocatalytic production of hydrogen by water splitting using sunlight.⁵¹ Tailoring the nitrogen content of $g\text{-C}_3\text{N}_4$ also opens up the possibility of improving energy storage technologies.⁵² Chen *et al.*⁵³ demonstrated that nitrogen-deficient $g\text{-C}_3\text{N}_4$ exhibits high electrochemical performance for lithium-ion battery applications. The increase in *d*-spacing due to changes in the C:N ratio of $g\text{-C}_3\text{N}_4$ as the result of bond breakage also creates new active sites, which could, for example, enhance the photocatalytic properties of $g\text{-C}_3\text{N}_4$. We also envision the possibility of incorporating tailored $g\text{-C}_3\text{N}_4$ in devices to detect radiation doses, for diverse applications in space exploration, biomedical radiation treatment, or nuclear power plants.

Conclusions

The work at hand presents an approach to selective change the stoichiometry of $g\text{-C}_3\text{N}_4$ nanosheets using an electron beam with different energies and current densities. In this way, our approach provides a facile method to control the C:N ratio of $g\text{-C}_3\text{N}_4$ and, as a result, the physical properties of the material. The $g\text{-C}_3\text{N}_4$ nanosheets degrade over time under electron irradiation for all explored electron acceleration voltages and current densities. Systematic EELS studies indicate that N was preferentially lost, such that amorphous carbon eventually remained. Through the careful temporal tracking of the process, the structural changes in-between crystalline and amorphous phases revealed that as N is lost, the lattice *d*-spacing increases. Corresponding EELS studies allowed us to correlate the *d*-spacing changes with the stoichiometry of the $g\text{-C}_3\text{N}_4$. The material with stoichiometry could be altered by changing both the electron acceleration voltage and current density. Faster deterioration was clearly observed for the lower acceleration voltages studied (≤ 60 kV), suggesting that inelastic scattering (radiolysis) is the predominant material loss mechanism. The rate of change (material loss) also increased with increasing



current density, increasing rapidly above a certain threshold. These findings indicate that with a careful adjustment of the electron acceleration voltage or current density, the nitrogen content of the crystalline g-C₃N₄ nanosheets can be modified. This work not only provides fundamental insight into the stability of g-C₃N₄ under electron irradiation, but also allows the tailoring of the stoichiometry of g-C₃N₄, and thus broadens its potential for applications such as photocatalysis, energy storage, and sensing.

Author contributions

The experiment was devised and managed by MHR. RGM conducted the TEM experiments and primary data analysis. AM and BBI prepared the g-C₃N₄ material. RGM, HQT, XY, AB, PP, AM, BBI, RS, TG and MHR contributed to the data analysis and preparation as well manuscript preparation.

Conflicts of interest

There are no conflicts to declare.

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