Monte Carlo simulation of nanoscale material focused ion beam gas-assisted etching: Ga\(^+\) and Ne\(^+\) etching of SiO\(_2\) in the presence of a XeF\(_2\) precursor gas†

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Elucidating energetic particle-precursor gas–solid interactions is critical to many atomic and nanoscale synthesis approaches. Focused ion beam sputtering and gas-assisted etching are among the more commonly used direct-write nanomachining techniques that have been developed. Here, we demonstrate a method to simulate gas-assisted focused ion beam (FIB) induced etching for editing/machining materials at the nanoscale. The method consists of an ion–solid Monte Carlo simulation, to which we have added additional routines to emulate detailed gas precursor–solid interactions, including the gas flux, adsorption, and desorption. Furthermore, for the reactive etching component, a model is presented by which energetic ions/target atoms, and secondary electrons, transfer energy to adsorbed gas molecules. The simulation is described in detail, and is validated using analytical and experimental data for surface gas adsorption, and etching yields. The method is used to study XeF\(_2\) assisted FIB induced etching of nanoscale vias, using both a 35 keV Ga\(^+\), and a 10 keV Ne\(^+\) beam. Remarkable agreement between experimental and simulated nanoscale vias is demonstrated over a range of experimental conditions. Importantly, we demonstrate that the resolution depends strongly on the XeF\(_2\) gas flux, with optimal resolution obtained for either pure sputtering, or saturated gas coverage; saturated gas coverage has the clear advantage of lower overall dose, and thus lower implant damage, and much faster processing.

1 Introduction

Controlling/editing materials at the nanoscale is critical for the realization of many emerging technologies. One synthesis technique for sculpting materials at the nanoscale is focused ion beam (FIB) nanomachining. FIBs are used in many applications, ranging from imaging with the helium ion microscope (HIM),\(^1–^3\) ion beam induced deposition,\(^4–^6\) and, nanomachining.\(^7\) Owing to the high achievable beam resolutions using the gas field ionization source (GFIS), FIBs using light ions, such as Ne\(^+\) and He\(^+\), have attracted significant attention in recent years. However, a draw back of nanomachining performed using FIB sputtering is that it requires sufficiently high ion doses to sputter material from the target surface, which may produce significant subsurface damage.\(^8\) Subsurface damage may be mitigated through the use of focused ion beam induced etching (FIBIE); in FIBIE, a reactive precursor gas, such as XeF\(_2\), is used to enhance the material removal rate, which therefore reduces the required dose. However, the factors determining resolution for FIB sputtering are not perfectly known, and the addition of a precursor gas further influences achievable nanomachining resolutions in a way which is dependent on the interaction of the gas flux, and scan parameters. In this paper, we describe a Monte Carlo simulation for FIBIE with XeF\(_2\), and use it to study the resolution of etched features in SiO\(_2\).

FIBIE reduces the subsurface damage associated with nanomachining by enhancing material removal rate, and thereby reducing the total ion dose required. Damage to the substrate may result from high doses of energetic ions, in the form of implanted ions in the subsurface (which may result in, for example, bubbling\(^9\)), as well as implanted energy, which may cause such effects as amorphization of a crystalline target.\(^8,^{10}\) FIBIE works similarly to electron beam induced etching: a reactive precursor gas is injected near the substrate, which then adsors to the surface as a monolayer. In the case of SiO\(_2\) with a XeF\(_2\) precursor, the gas does not spontaneously etch; however, the input of energy from the ion beam leads to
dissociation of adsorbed molecules. The volatile compounds resulting from the dissociation of the XeF₂ molecule, and reaction with the substrate, may be removed from the substrate spontaneously, or by additional interactions resulting from the ion beam.

While atomic layer control via FIB sputtering and etching has not been achieved (see ref. 11 for a recent perspective/review on atomic scale fabrication), the reduced sub-surface scattering that occurs in 2D materials provide avenues for enhanced resolution. In this regard, we have explored controlled defect introduction via scanning He⁺ irradiation in multilayer and single layer WSe₂. Additionally, subsurface nanoscale nanomachining via laser-assisted He⁺ focused ion beam sputtering was shown to enhance the etching rate in single layer graphene nanochannels.

Monte Carlo method simulations are widely used to study interactions of energetic ions with solid targets, and FIB processing specifically. Such applications include studying subsurface damage, imaging with the HIM, and ion beam induced deposition. One of the most popular Monte Carlo programs is SRIM, which simulates the interaction of numerous ion species and target compositions. However, despite its versatility and widespread use, SRIM is unable to simulate the effects of cumulative material removal due to either sputtering, or gas assisted etching. While several Monte Carlo methods have been developed which incorporate target evolution due to sputtering, we are unaware of any simulation that fully accounts for both sputtering, and material removal due to FIBIE. For this reason, we have developed the EnvizION Monte Carlo code, which has been previously used to simulate FIB milling and secondary electron (SE) emission. While chemically assisted etching was estimated previously using EnvizION, this was emulated by reducing the surface binding energy of the target material, and was not sufficient to understand all factors related to material removal rate, as well as etching resolution. In this paper, we introduce our method for simulating gas assisted etching using EnvizION; this method incorporates details about the incoming gas flux, population of irregular surface features with gas, and depletion of gas due to reaction with substrate surface atoms driven by energy added from the ion beam. While specifically developed for high-energy focused ion beam applications, low-energy variants to the energy transfer could be implemented for atomic layer etching, as well as the inverse plasma enhanced atomic layer deposition.

The ion solid interaction of the EnvizION simulation method is based closely on SRIM. Unlike SRIM, EnvizION employs a voxel based representation of the computational domain, which permits the target to evolve as material is sputtered, and redeposited elsewhere. Bulk relaxation, and surface smoothing routines are employed to reduce porosity, and other simulation artifacts, during sputtering. The excitation and emission of SEs by the primary ion may additionally be simulated, according to the type of substrate. The FIBIE method, presented in this paper, consists of a method for simulating both the incoming gas flux, as well as the interactions by which the primary ion beam may cause dissociation of adsorbed XeF₂, and the associated etching of material. The mechanism by which the ions trigger dissociation of adsorbed XeF₂ is modeled similarly to the atomistic mechanism of ion beam induced deposition described by Dubner. In this model, the interaction between the primary ion and the adsorbed surface molecules is mediated by the substrate. The primary ion generates large numbers of displaced atoms in the near surface region of the target; some of these displaced atoms may reach the surface of the target, where they are permitted to interact with adsorbed gas through a binary collision. In contrast to Dubner’s model, we additionally allow secondary electrons to cause dissociation with an energy dependent cross section.

In Section 2, we describe the simulation method in detail. Section 3 compares simulations with experiments for five sets of scan parameters for Ga⁺ etching, and three sets of scan parameters for Ne⁺ etching. We study the etching resolution with respect to gas flux in Section 4, and our results are summarized in Section 5.

2 Simulation method

In this section, we survey the novel aspects of the method for simulating FIBIE of SiO₂ with a XeF₂ precursor. Details of the sputtering method have been presented previously, with studies focusing on the validation of the code, and sputtering in monatomic substrates, and the use of simulations to understand sputtering in SiO₂ targets. Monte Carlo modeling of SE emission was previously implemented only for conducting targets; in the present work, we use a similar method, but with the electron scattering determined by interaction with optical phonons, using the Monte Carlo method of. In contrast, the method described in our previous work for conducting targets was based on the inelastic interactions of excited electrons with other electrons. The SiO₂ Monte Carlo method for SE emission that we employ does not account for charging of the substrate; this subject will be studied in the near future.

In this paper, we study line scans, which consist of a discrete set of “pixels”, with length , and with pixel spacing or “pitch” ; the ion beam dwells at each pixel center for a time . The number of pixels in a single line, , defines a “frame”. After a frame is scanned, the experiment may pause or blank the beam for some time to allow the gas flux to repopulate the surface with XeF₂; this time is referred to as the “refresh time”, denoted . After refreshing, another frame is etched by scanning the beam back along the line (i.e., we study “serpentine” scans, though any patterning option is possible). This process continues until the full dose has been reached (generally on the order of thousands of frames for the experiments presented in this paper).

The computational domain consists of the region

\[ \{(x, y, z) : x_{\text{min}} < x < x_{\text{max}}, y_{\text{min}} < y < y_{\text{max}}, z \in (-h_{\text{max}}, h_{\text{pad}})\} \]

All substrates that we consider here consist of SiO₂, initially occupying the region \( z < 0 \). The empty space at the top of the
domain is taken to be $h_{\text{pad}} = 25$ nm, and $x_{\text{min}}, x_{\text{max}}, y_{\text{min}}, y_{\text{max}},$ and $h_{\text{max}}$, vary by simulation. The domain is represented by a 3D cubic lattice of voxels, with the centers of each voxel separated by a distance $\delta \approx 0.25$ nm, with $\delta$ being the average distance between atomic centers in SiO$_2$. Each voxel is represented by a flag which corresponds to the contents of the voxel, either empty, or occupied by an atom of Si, O, or an implanted neutral atom of the impinging ion type; the initial substrate is populated at random with 1/3 Si and 2/3 O voxels. Ga$^+$ ions are assumed to occupy lattice sites a distance of travel $V$, reaches the $x$ or $y$ boundaries, its direction of travel is set to

$$V' \leftarrow V - 2(V \cdot \hat{n})\hat{n}$$

where $\hat{n}$ is the outward pointing normal of the boundary. In previous versions of EnvizION, the simulation had open boundary conditions so that moving ions and target atoms which leave the simulation domain are no longer tracked. Open boundary conditions can lead to significant artifacts in via depth if the domain is not large enough, and are problematic for the gas simulation, as discussed below.

The method that we employ for gas assisted etching (GAE) is based on that described by Dubner$^{38}$ for modeling ion beam induced deposition. In this model, the precursor gas (in our case, XeF$_2$) is assumed to follow a Langmuir isotherm on the SiO$_2$ surface. Once adsorbed, the XeF$_2$ does not etch spontaneously, but remains adsorbed for its residence time (100 ms), or until it is activated, either directly or indirectly, by the ion beam. Once etching occurs, the simulation emulates the chemical reaction as follows:

$$\text{SiO}_2 + 2\text{XeF}_2 \rightarrow \text{SiF}_4 + \text{O}_2 + 2\text{Xe}$$

The etch products are assumed to immediately volatilize, and are removed from the substrate. The method for simulating gas assisted etching (GAE) consists of two components:

1. A method for the population of the surface with gas molecules.
2. A method for the ion beam to activate the adsorbed gas to etch.

Diffusion of the adsorbed XeF$_2$ along the surface may be important for both the etch rate and the resolution of etched vias; however, it is computationally costly, and we ignore diffusion in the present study.

In simulations, the surface is populated by an isotropic gas flux, with gas molecules adsorbing to the surface following a Langmuir isotherm model. Each gas molecule may occupy exactly one surface site, which in simulations is defined as an empty voxel which has at least one nearest or next-nearest neighbor voxel which contains a substrate atom. While surface sites and adsorption orientations are not distinguished in this version of the simulation, for systems where atom specific and molecular orientation specific orientations are known, we can accommodate these specifics with modified residence times and dissociation energies.

We illustrate the isotropic gas population routine in Fig. 1. The population of the surface by gas molecules is simulated as an isotropic flux. For each gas molecule, an initial position $P$ is generated uniformly at $z = h_{\text{pad}}$ and an initial trajectory $V$ is generated from the half sphere of trajectories with $z$ component less than zero. The gas molecule then travels according to the following steps:

function gas recoil ($P$, $V$)
  $it \leftarrow 0$
  while ($it < it_{\text{max}}$) do
    (1) Trace a ray from $P$ along $V$ until either a surface site is reached, or we reach the boundary $z = h_{\text{pad}}$ or $z = -h_{\text{max}}$. If a boundary is reached, the routine terminates. Otherwise, let the position of the surface site be given by $P'$.
    (2) If the surface site is empty, generate a uniform random number $r \in (0, 1)$. If $r < \alpha$, the gas molecule adsorbs to the surface site at $P'$.
    (3) For all other cases, generate a new random trajectory $V'$, and update the molecule’s position and trajectory with $P \leftarrow P'$, $V \leftarrow V'$.
  (4) $it \leftarrow it + 1$.
end while

end function

The parameter $\alpha$ is referred to as the sticking coefficient, and expressed the probability that a gas molecule will adsorb to a site on which it is incident. During the ray trace step of gas recoil, it is important that open boundary domains are not used, as this will prevent gas trajectories with small $z$ component from reaching the surface. Gas molecules that have adsorbed onto the surface may spontaneously desorb with a mean residency time $\tau$, which we take to be 100 ms, following Harriott (1993). A gas molecule which desorbs at position $P_{\text{desorb}}$ is assigned a randomized trajectory $V_{\text{desorb}}$, and we call gas recoil $\{P_{\text{desorb}}, V_{\text{desorb}}\}$, i.e., desorbed gas molecules may recoil throughout the domain until either escaping, or re-adsorbing to an empty site. In simulations, the gas population routine is called between each pixel dwell, and during the gas refresh time. In the first case, after a pixel dwell of length $\tau_{\text{dwell}}$, the gas flux is simulated for $\tau_{\text{dwell}}$ as well, before the next pixel is simulated. Dwell times are typically a few $\mu$s, or sub-$\mu$s, while the timescales characterizing gas population are on the order of thousands of $\mu$s, so that this simplification is expected to introduce little error.

To understand the population of a surface due to the setup we have described, it is helpful to compare the gas population of an irregular surface with that of a perfectly flat surface. For a flat surface, in the absence of diffusion or ion beam induced dissociation, our gas population scheme is described by the following equation:

$$\rho(t) = \rho_0 + (\rho_0 - \rho_\infty)\exp(-t\tau_{\alpha}^{-1})$$

(2)
Here $\rho(x,y,t)$ is the number of adsorbed molecules per unit area on the surface, $\rho_\infty$ is the equilibrium surface coverage density, $\rho_0$ is the initial surface coverage density, and $\tau_a$ is the time scale for gas population. We sometimes find it more convenient to refer to the gas fraction, $C(x,y,t) = \rho(x,y,t)\phi^2$. The effective gas flux, in molecules per area per unit time, is given by $\phi$. Fig. 2(a) plots the gas fraction as a function of time for an initial flat substrate with zero gas population, for varying gas flux. When the gas flux is as high as $10^{18}$ molecules per cm$^2$ per s, the gas fraction rapidly approaches full coverage. Importantly, in the experiments we consider here, we are interested in the formation of high aspect ratio valleys, which populate quite differently than a flat surface. Fig. 2(b) compares the population of the flat surface from eqn (2), to the simulated result using the described gas recoil routine in a rotationally symmetric Gaussian via exposed to a gas flux of $10^{17}$ molecules per cm$^2$ per s; this Gaussian via has a 15 nm full width at half maximum (FWHM), and a depth of 100 nm. While the simulated population at the top of the via and the surrounding flat surface ($z = 0$) is the same as the analytical theory, deeper in the via we see that gas population occurs much more slowly; near the via bottom ($z = -99$ nm) gas population occurs on a timescale that is more than two orders of magnitude slower than the flat surface.

An important component of the gas recoil routine is that a gas molecule which fails to adsorb to the surface spontaneously desorbs with a random trajectory; this emulates transient adsorption to an already adsorbed gas molecule, or a failure to adsorb due to a sticking coefficient less than one. After desorbing, a gas molecule recoils throughout the domain, which, for brevity, we refer to as gas ricochet. We consider the effects of ignoring ricochet in Fig. 3. Fig. 3(a) shows a top down view of the simulated gas fraction at $t = 10^6$ ms, while Fig. 3(b) shows the same simulation, except not permitting ricochet (i.e., if a gas molecule reaches an already occupied gas site, the gas recoil routine for that molecule stops, and the gas molecule disappears); in this manner gas sites are only populated in proportion to the solid angle exposed to the top of the domain. Ignoring ricochet results in a significantly lower gas fraction. Fig. 3(c) plots the gas fraction from Fig. 3(a) and (b), at various times and depths in the via; we can see in particular that the gas population with ricochet becomes higher relative to the simulation without ricochet as the gas fraction increases. This can be understood as follows: as the gas fraction inside of a via increases, this leads to more incident gas molecules landing on...
already full surface sites, and therefore subsequent desorption allows gas molecules to find non-occupied sites; this contribution is a critical component to populating high-aspect ratio vias, where direct adsorption is minimal due to the low solid angle relative to the top of the domain.

The primary ion Monte Carlo which depends on the ion type (here Ne\(^+\) and Ga\(^+\)) and the target substrate (here SiO\(_2\)) proceeds as described in our previous work\(^{29}\), which is in turn closely based on SRIM.\(^{20}\) Interested readers should refer to our previous work\(^{29}\) for the relevant SiO\(_2\) parameters that affect the ion–solid interactions, as we focus here on the precursor–solid interactions that are operative in the new simulation. Summarily, moving atoms (i.e., primary ions and recoiling target atoms) exhaust their energy through a combination of elastic collisions with atoms in the substrate, and inelastic interactions with the electrons of the substrate. Inelastic interactions may give rise to secondary electron emissions. Elastic collisions give rise to the scattering of moving atoms; if an elastic collision transfers more energy than a displacement threshold \(E_{\text{disp}}\) to the atom in the target, the target atom becomes a recoil, and travels through the substrate similarly to the incident ion. In the near surface region, \(E_{\text{disp}} = 0\), while \(E_{\text{disp}} = 8\) eV elsewhere. If a moving atom reaches the surface with energy greater than the surface binding energy \(E_{\text{surf}}\), then it may be sputtered. Atoms which are not sputtered, and whose initial energy is less than 8 eV, are returned to their initial position after their energy drops to zero.

In the EnvizION model, gas-assisted etching is driven by interactions between moving atoms, excited electrons, and adsorbed XeF\(_2\). Any moving atom which reaches an occupied surface site is permitted to interact with the adsorbed XeF\(_2\) molecule – importantly, even if an atom with energy insufficient to sputter reaches the surface, it is permitted to interact with an adsorbed XeF\(_2\). If an electron reaches an occupied surface site, it etches with a probability \(P_e\). Secondary electron dissociation cross sections have been studied in detail for electron beam induced deposition\(^{36-38}\) and electron beam induced etching studies\(^{39,40}\). For simplicity, in what follows we set \(P_e = 1\); notably, even with a unit etching probability, we find that electron etching is a very minor component in ion beam induced etching.

Moving atoms interact with adsorbed XeF\(_2\) through a binary collision model (see Dubner\(^{35}\)). The interaction between a moving atom A and an adsorbed XeF\(_2\) molecule is calculated as a binary collision between the moving atom, and one of the component atoms of the XeF\(_2\). There is a 1/3 chance that the collision is with the Xe, and a 2/3 chance that the collision is with a F atom; for convenience, we refer to the selected knock-
on atom in the precursor gas molecule as B. The maximum impact parameter \( p_{\text{max}} \) is set to be \( r_A + r_B \) where \( r_A \) and \( r_B \) are the van der Waals radii of atoms A and B respectively. The impact parameter of the binary collision is then set to be \( p = \sqrt{\text{rand} p_{\text{max}}} \), where rand is uniformly distributed in \((0, 1)\). If the amount of energy transferred from A to B exceeds an activation energy \( E_{\text{act}} \), then dissociation and subsequent etching occurs. Based on eqn (1), if etching occurs, the simulation flags the nearest occupied voxel to empty, and then flags another nearest neighbor, or next-nearest neighbor, voxel to empty with probability 0.5. In this way, each gas molecule removes on average 1.5 atoms when it dissociates, in accordance with eqn (1). Note that the etching routine in simulations does not distinguish Si and O atoms, nor does it allow for two \( \text{XeF}_2 \) molecules to participate in a reaction, and instead removes, on average, the correct number of substrate atoms per dissociated \( \text{XeF}_2 \).

Based on the above discussion, there are two parameters whose values are unknown: the activation energy \( E_{\text{act}} \), and the sticking coefficient \( \alpha \). By comparison with the experiments described in Section 3, and further experimental benchmarks described in the ESI,† we use an activation energy of \( E_{\text{act}} = 0.05 \) eV. We use a value of \( \alpha = 1 \) for simulations; however the comparison with previously published experiments, presented in the ESI,† requires a value of \( \alpha = 0.025 \) to account for the extremely high gas flux values in the experiments. The gas etching parameters used for simulations in this paper are summarized in Table 1.

Again, for computational efficiency, we ignore diffusion in all simulations. When \( \text{XeF}_2 \) adsors to the surface, it may diffuse, similar to the case of IBID.41 In the present paper, we will ignore diffusion in all simulations. We will consider the effects of diffusion in a future paper, which, for 3D electron beam induced deposition studies, even short range diffusion has been shown to have interesting effects when transitioning from electron to mass transport limited regimes.42

### 3 Results

In Fig. 4, we illustrate etching due to 10 keV spot dwells of varying ion species. For these simulations, the gas fraction, or coverage, on the surface is kept at 1 during each simulation.

| Recipe | Energy (keV) | \( L_{\text{scan}} \) (nm) | \( \Delta x \) (nm) | \( t_{\text{dwell}} \) (\( \mu \)s) | Current (pA) | \( t_{\text{refresh}} \) (\( \mu \)s) | Frames | Dose (nC cm\(^{-1}\)) |
|--------|-------------|-----------------|----------------|----------------|-------------|----------------|-------|----------------|
| Ga\(^+\) 1 | 35 | 128 | 4 | 0.05 \( \mu \)s | 1 pA | 2400 \( \mu \)s | 15700 | 2.0 |
| Ga\(^+\) 2 | 35 | 128 | 4 | 0.1 \( \mu \)s | 1 pA | 2400 \( \mu \)s | 9500 | 2.4 |
| Ga\(^+\) 3 | 35 | 128 | 4 | 0.05 \( \mu \)s | 1 pA | 1200 \( \mu \)s | 17349 | 2.1 |
| Ga\(^+\) 4 | 35 | 128 | 4 | 0.2 \( \mu \)s | 1 pA | 1200 \( \mu \)s | 8000 | 4.0 |
| Ga\(^+\) 5 | 35 | 128 | 4 | 0.5 \( \mu \)s | 1 pA | 1200 \( \mu \)s | 4000 | 5.0 |
| Ne\(^+\) 1 | 10 | 128 | 2 | 1 \( \mu \)s | 0.3 pA | 1000 \( \mu \)s | 20105 | 62 |
| Ne\(^+\) 2 | 10 | 128 | 2 | 0.5 \( \mu \)s | 0.3 pA | 500 \( \mu \)s | 34849 | 54 |
| Ne\(^+\) 3 | 10 | 128 | 2 | 0.2 \( \mu \)s | 0.3 pA | 300 \( \mu \)s | 67017 | 41 |

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Fig. 4  Comparison of etching from 10 keV He\(^+\), Ne\(^+\), and Ga\(^+\) ion point dwells. Histograms of the radial location of all etch events are shown in (a)–(c), computed from 10 000 incident ions. Ion paths are shown in red, and recoil atom paths are shown in blue, for 50 incident ions, in (d)–(f); the surface of the target is indicated by a dashed line. The etch yields for each ion are as follows: 4 atoms per ion for He\(^+\), 29 atoms per ion for Ne\(^+\), and 34 atoms per ion for Ga\(^+\). Note that (d)–(f) have projected all data along the \( y \)-axis, while simulations were three dimensional.
These simulations were run without voxel updating, so that the surface remains smooth even if sufficient energy is transferred to a gas molecule to cause dissociation. He⁺ has a very low etch yield (5 atoms per ion) compared to Ne⁺ and Ga⁺ (21 and 35 atoms per ion, respectively). This is due to the fact that He⁺ travels deeper into the target, and generates few recoil atoms in the near surface region. Ne⁺ and Ga⁺ have similar etch yields, with Ga⁺ being significantly higher at this energy, due to the fact that it generates more recoils in general, as well as having an interaction volume nearer to the surface. However, the sputter yield varies greatly among the ion species, so that He⁺ ions show significantly narrower than the effective beam profiles reported in Mahady et al. (2018) for pure sputtering, suggesting that the lower doses associated with etching may reduce artifacts which broaden the sputtered/etched profile. We consider a comparison between the effective beam profiles here with those measured using the teardrop method of Tan et al. in the ESL.
We study XeF₂ assisted FIB etching using five scan parameter “recipes” for the Ga⁺ ion beam, and three for the Ne⁺ ion beam. The parameter setups are described in Table 2. The simulated scan lengths are given in Table 2, while in experiments, the scan lengths are 500 nm, and 250 nm, for Ga⁺ recipes and the Ne⁺ recipes, respectively. For the Ne⁺ experiments, each frame consists of two parallel lines, separated by 1 nm, which are scanned in a serpentine pattern; Ga⁺ line scans consist only of a single line. In simulations, we take advantage of the mirror boundary conditions to reduce the total number of simulated ions by a factor of four. The computational domain is defined by \( \{ x, y, z \} : -256\delta < x < 256\delta, 0 < y < 512\delta, -1024\delta + h_{\text{pad}} < z < h_{\text{pad}} \}; \) the beam is scanned from \( x = 0 \) to \( x = 256\delta \), and the scan is centered on \( y = 0 \). We simulate 1/2 of the beam current of each recipe given in Table 2, and 1/2 of the scan length, since for each simulated ion, the boundary condition effectively gives us a simulation of three additional reflected ions. The use of this symmetry, to reduce the number of ions simulated, has been studied in detail; for very high dwell times artifacts emerge, however the mirror conditions are appropriate in the parameter regime we study here.

We approach the experiments performed with each ion species in the following manner. The Ne⁺ and Ga⁺ experiments are performed on two different machines, with different gas injection systems, so it is expected that the Ne⁺ recipes (respectively, the Ga⁺ recipes) should all have similar XeF₂ fluxes to one another, but this flux may differ from that of the Ga⁺ recipes (respectively, the Ne⁺ recipes). However, the localized gas flux for each recipe set is not well known, so it must be treated as a fitting parameter. Therefore, we first simulate one recipe with a variable gas flux in order to identify the flux that results in best agreement with experimental via depth and shape; we call the simulated gas flux for the Ga⁺ (respectively, the Ne⁺) experiments \( \phi_{Ga} \) (respectively, \( \phi_{Ne} \)). Once the gas flux for each recipe set is identified, the remaining recipes are simulated.

### Table 3  Material removal yields for each recipe (in atoms per ion)

|       | Ga⁺1 | Ga⁺2 | Ga⁺3 | Ga⁺4 | Ga⁺5 | Ne⁺1 | Ne⁺2 | Ne⁺3 |
|-------|------|------|------|------|------|------|------|------|
| Yield | 74.2 | 73.7 | 73.3 | 75.0 | 64.8 | 8.2  | 8.4  | 9.5  |

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#### 3.1 Ga⁺ etching

We first study XeF₂ assisted etching using the Ga⁺ beam recipes described in Table 2. Experiments for each of the five Ga⁺ recipes were performed twice; these experiments demonstrate non-trivial variability in the etch rate, with via depths differing by as much as 25% between experiments. The experimental results are the most consistent for recipe 5, with the difference in via depth being less than 4 nm, so we use this via to calibrate...
simulation parameters; the results are shown in Fig. 5(a), where a gas flux of $\phi_{Ga} = 4 \times 10^{17}$ molecules per cm$^2$ per s gives close agreement between the simulated via profile, and experiment. Fig. 5(b) shows a surface plot of the simulated Ga$^+$ etched via, with a gas flux $\phi_{Ga}$; note that only 1/4 of the actual line scan is simulated, since we use mirror boundary conditions. Fig. 5(c) compares the derivative of the via depth with respect to dose, and the gas coverage in the bottom of the via; the derivative and gas coverage tend to decrease with increasing dose, however they both exhibit significant noise, likely due to the small variations in gas coverage in this dose range.

We simulate Ga$^+$ recipes 1–4 using a XeF$_2$ flux of $\phi_{Ga}$; Fig. 6(a) and (b) shows vias resulting from simulations and experiments with a refresh time of 2400 $\mu$s, while those in Fig. 6(d)–(f) have a refresh time of 1200 $\mu$s. For the purposes of comparison, we plot the simulated via cross-sections with via profiles extracted from the experimental TEM images, shown in Fig. 6 (see ESI† for a comparison between experiments and the actual TEM images). While there is variability in the via depths between the experiments, overall, the experimental etch volumes are reproduced well by simulations. The differences between simulations and experiments are within the variability of the experiments,

Fig. 7 Ne$^+$ etching using the Ne$^+$ Effective Beam Profile, using Ne$^+$ etching recipe 1. (a) Comparison of Ne$^+$ etched vias with experimental SEM image as the gas flux is varied. Gas flux is given in units of molecules per cm$^2$ per s. (b) Surface profile of the etched via with flux $7 \times 10^{16}$ molecules per cm$^2$ per s. The entire simulation domain is shown, and there is a mirror boundary condition at the $y = 0$ plane, and the $x = 64$ nm plane. The initial position of the beam in the $x$–$y$ plane is indicated by $P_0$. (c) Comparison of the fractional gas coverage at the via bottom (left axis, square markers) and the absolute value of the change in via depth per ion (right axis, circular markers), for the etching simulation with gas flux $7 \times 10^{16}$ molecules per cm$^2$ per s. The bottom coverage is calculated for surface sites with $x \in [0, 64]$ nm and $y \in [0, 15]$ nm.

Fig. 8 Comparison of simulated etched profile to experimental via profiles for the Ne$^+$ etched via with recipes 1 through 3.
and the trends in \textit{via} shape and material removal yield are replicated by the simulations. Additionally, in Fig. 6, we plot the simulated total depth/dose for each recipe. As the dwell time is increased, while the refresh time is held constant, the depth etched per ion decreases; the greatest variability in etch rate is seen between recipes 3 and 5, where the change in depth per ion decreases by a third as the dwell time is increased by a factor of 10. The total removal yields for all Ga$^+$ line scan recipes are summarized in Table 3. Interestingly, the removal rates of Table 3 do not vary as much as the ratio of the etched \textit{via} depth to ions plotted in Fig. 6(f); this difference can be understood by the fact that the \textit{via} FWHM varies between recipes, even when etching to the same depth. Ga$^+$ \textit{via} 4, for example, is 15\% wider at the same depth as Ga$^+$ \textit{via} 3.

### 3.2 Ne$^+$ etching

The results of simulating Ne$^+$ etching recipe 1 of Table 2, are shown in Fig. 7. Again, we treat the effective gas flux as a fitting parameter, and the best agreement with experiments is obtained for $\phi_{\text{Ne}} = 7 \times 10^{16}$ molecules per cm$^2$ per s, which is lower than that observed in the Ga$^+$ etching experiments. A plot of the surface of the etched \textit{via} is also shown in Fig. 7; compared to the Ga$^+$ recipes, the Ne$^+$ etch leads to a deeper and higher aspect ratio \textit{via}. Note that the Ne$^+$ dose is higher compared to the Ga$^+$ experiments; the lower gas flux, greater \textit{via} depth, and lower mass of Ne$^+$, lead to a correspondingly lower cumulative etch yield, and sputter yield (6.8 and 0.5 atoms per ion, respectively). The higher aspect ratio \textit{via} also leads to lower gas

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**Fig. 9** Comparison of the simulated spatial distributions of sputtering and etching for the formation of the Ne$^+$ \textit{via} with recipe 1. The parameters correspond to the first (a) 121 frames, and (b) 1218 frames of the dose. The blue shaded region indicates the cross section of the volume of material removed by etching, and the orange to the volume removed by sputtering. In the shallow \textit{via}, sputtering is more significant near the beam center, where gas is exhausted, than near the tails where etching dominates.

**Fig. 10** Comparison of the beam shape (dashed lines) and the corresponding etched \textit{via} profile for the 35 keV Ga$^+$ with recipe 5 with gas flux $\phi_{\text{Ga}}$ (a), and the 10 keV Ne$^+$ beam with recipe 1 with gas flux $\phi_{\text{Ne}}$ (b).
coverages, as we would expect from Fig. 2, as shown in Fig. 7(c) on the left axis. Similar to the Ga⁺ recipes, gas etching is the dominant mechanism by which material is removed from the surface, so the change in via depth with dose is approximately proportional to the coverage of gas in the via bottom, as shown in Fig. 7(c).

Fig. 8 shows the via profiles resulting from simulations for all Ne⁺ recipes in Table 2, compared with experimental images. All simulations in Fig. 8 were performed with a gas flux equal to $\phi_{Ne}$. Both the dwell time and refresh time are varied for the Ne⁺ recipes, however we can see that the overall trend is for the etch rate to increase as the dwell time is reduced: via 3, with a dwell time of 0.2 μs has a 34% higher total removal rate per incident ion, than does via 1, which has a dwell time of 1 μs. When compared to the experimental images, simulations are slightly shallower in recipes 2 and 3. All Ne⁺ recipes result in vias with a similar FWHM. The material removal rates for all Ne⁺ recipes are given in Table 3.

### 4 Discussion

We consider the relative contributions to via shape formation from sputtering and etching in Fig. 9; here we plot the profile of the simulated etched via using Ne⁺ recipe 1, with a XeF₂ flux of $\phi_{Ne}$, after 121 frames (a), and 1218 frames (b). The shaded regions indicate the cross sections of volume removed by etching, and by sputtering, over this dose range. As expected, etching accounts for the majority of the material removed, and largely determines the via shape. Sputtering is negligible away from the via center, however it contributes a significant fraction of the material removal near the via center. The relative importance of sputtering in the via center can be attributed to lower gas coverage, caused by depletion from the ion beam, and, in deeper vias, a reduced incoming gas flux due to shadowing; this combination of factors reduces the overall etch rate, while the sputter rate is unaffected by gas coverage. Additionally, we see that as a via increases in depth, the via depth is not completely explained by the influence of etching and sputtering, but instead, there is an additional contribution from atoms being sputtered/displaced and redeposited elsewhere in the via (see Fig. 9(b)).

We compare the beam profile with the associated simulated etched via shape, for both Ga⁺ recipe 5, and Ne⁺ recipe 1, in Fig. 10. The Ga⁺ etched via shows a close similarity between the via shape and the beam shape; however, for the Ne⁺ etched via, we see that there is a significant difference between the beam and via shapes, with the etched via being wider than the beam profile. This is in contrast to the case for pure sputtering without XeF₂, where simulations show that the sputtered via shape is similar to the beam profile for both Ne⁺ and Ga⁺ sputtering.

To understand why there is such a difference between the beam profile and the via profile for the simulated Ne⁺ etched via in Fig. 10, we plot the FWHM as a function of via depth for Ne⁺ recipe 1 with varying gas flux in Fig. 11(a). When the gas flux is $1 \times 10^{14}$ molecules per cm² per s, there is zero chemical etching and the line scan is in a pure sputtering regime. As the gas flux increases, and etching becomes significant, the FWHM increases as a function of depth, reaching a peak around $1 \times 10^{16}$ molecules per cm² per s, after which it begins to decrease with increased gas flux. When the gas flux reaches a sufficiently high value, the FWHM of the etched via becomes comparable to the pure sputtered via. Importantly, we see that $\phi_{Ne}$ falls in the intermediate regime where the FWHM is near maximal as a function of depth. If the flux in the Ne⁺ etch simulation is increased to $5 \times 10^{17}$ molecules per cm² per s, we see that the etched via profile is narrower, just as in the Ga⁺ case (see Fig. 11(b)). Therefore, simulations predict that etching resolution is optimal for pure sputtering, or very high gas fluxes. Furthermore, the optimal resolution resulting from XeF₂ assisted etching is similar to the effective beam profile, just as in sputtering.

### 5 Conclusion

We present a computational method for simulating gas-assisted ion beam etching; in particular, we study XeF₂ assisted FIB
induced etching of SiO$_2$. This method is incorporated into the EnvizION simulation, which uses a Monte Carlo simulation of focused ion beam processing. The presented method accounts for an isotropic flux of XeF$_2$ molecules incident on a surface with arbitrary geometry, allowing population of high aspect ratio features such as sputtered valleys. The adsorbed gas can react with the SiO$_2$ molecules due to added energy from the focused ion beam. The etching reactions are dominated by interactions between the recoiling atoms in the target which are displaced by the primary ion; consequently, the etching is driven by energy which is transferred from the primary ion to the substrate by the elastic nuclear energy loss. Diffusion of adsorbed gas molecules is neglected for the present work.

We apply the simulation method to study the formation of nanoscale vias with reference to experiments; five recipes are studied for FIB etching using a 35 keV Ga$^+$ beam, and three recipes are studied for FIB etching using a 10 keV Ne$^+$ beam. Effective ion beam profiles are identified for the Ne$^+$ and Ga$^+$ beams, which reproduce the shape of experimental etched vias by combining the innate ion beam profile with possible machining artifacts. Using these beam profiles, we simulate experimental vias using a variable gas flux to identify the experimental gas flux. After identifying the experimental gas flux, the Ga$^+$ and Ne$^+$ etching simulations reproduce experimental vias for all parameter recipes we consider.

Unlike sputtering simulations, the resolution of the simulated etched vias is shown to be strongly dependent on the incoming gas flux. The FWHM of etched vias is found to increase with $vi$ depth for the Ne$^+$ etched vias, and the FWHM can vary by as much as a factor of two depending on gas flux. Intermediate gas fluxes are shown to lead to the lowest resolution etching, while pure sputtering, and very high gas flux, produce comparable FWHM as a function of depth.

In a future work, we will analyze the factors determining resolution in detail. In particular, we expect that surface diffusion of adsorbed gas molecules plays a significant role in both the etch rate, and the etching resolution. Additionally, it remains an important open problem to incorporate a simulation of the gas flux, to remove the need for fitting the flux to experimental data.

Conflicts of interest

There are no conflicts to declare.

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