Dynamical scattering in ice-embedded proteins in conventional and scanning transmission electron microscopy

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Structure determination of biological macromolecules using cryogenic electron microscopy is based on applying the phase object (PO) assumption and the weak phase object (WPO) approximation to reconstruct the 3D potential density of the molecule. To enhance the understanding of image formation of protein complexes embedded in glass-like ice in a transmission electron microscope, this study addresses multiple scattering in tobacco mosaic virus (TMV) specimens. This includes the propagation inside the molecule while also accounting for the effect of structural noise. The atoms in biological macromolecules are light but are distributed over several nanometres. Commonly, PO and WPO approximations are used in most simulations and reconstruction models. Therefore, dynamical multislice simulations of TMV specimens embedded in glass-like ice were performed based on fully atomistic molecular-dynamics simulations. In the first part, the impact of multiple scattering is studied using different numbers of slices. In the second part, different sample thicknesses of the ice-embedded TMV are considered in terms of additional ice layers. It is found that single-slice models yield full frequency transfer up to a resolution of 2.5 Å, followed by attenuation up to 1.4 Å. Three slices are sufficient to reach an information transfer up to 1.0 Å. In the third part, ptychographic reconstructions based on scanning transmission electron microscopy (STEM) and single-slice models are compared with conventional TEM simulations. The ptychographic reconstructions do not need the deliberate introduction of aberrations, are capable of post-acquisition aberration correction and promise benefits for information transfer, especially at resolutions beyond 1.8 Å.

1. Introduction

Understanding structural details down to the atomic scale is essential to decipher the chemical interactions of protein complexes and ultimately their function. In recent years, conventional transmission electron microscopy at cryogenic temperatures (cryo-CTEM) has reached near-atomic to atomic resolution for proteins in their natural state using single-particle structure determination (Herzik, 2020; Yip et al., 2020; Nakane et al., 2020; Zhang et al., 2020; Kato et al., 2019) by employing TEM imaging under parallel illumination conditions with the specimen embedded in glass-like amorphous ice. The direct structural interpretation of individual molecular images, however, remains challenging, primarily due to three aspects. First, only the projection of a rather extended 3D object along the electron-beam direction is observable. Second, CTEM phase contrast emerges from a
complex entanglement of the specimen and electron-optical parameters. Third, dynamical scattering and propagation occur within the specimen, hampering the direct inversion of the recorded data to obtain the structure of interest when multiple scattering becomes significant. The first aspect is resolved in contemporary single-particle cryo-CTEM by recording large numbers of particles in different orientations, followed by their 3D alignment and reconstruction (Penczek et al., 1992). Similarly, correcting schemes are employed to account for the contrast transfer function (CTF) governing the second aspect, so as to reverse the distortion effects of the native spatial frequencies of the object in the imaging process, caused mainly by spherical aberration and an intentionally applied defocus to transfer phase to amplitude contrast (Crowther & Böttcher, 1996). Importantly, most of these correction schemes are inherently based on the assumption that the interaction of the electrons with the specimen can be treated within the (weak) phase object \( (W)PO \) approximation, which implies that dynamical scattering is negligible (Wade, 1992). Indeed, the dominant presence of light elements such as hydrogen, oxygen, carbon, nitrogen and sulfur often justifies the projection assumption in both forward simulations and object retrieval, despite the complexity of biological specimens with \( 10^4 \) to \( 10^6 \) atoms, although limitations of this simplification have been discussed (Hall et al., 2011). Moreover, different continuum and atomistic concepts have been used to include structural noise arising from the glass-like ice embedding (Himes & Grigorieff, 2021; Vulović et al., 2013; Shang & Sigworth, 2012) in forward simulations.

A detailed and comprehensive treatment of the effect of multiple electron scattering in vitrified proteins via multislice simulations is required to shed light on the ultimate resolution limits, accuracy and precision of contemporary single-particle cryo-CTEM. These simulations will also form the foundation for the applicability of novel acquisition and reconstruction schemes that employ cryo-scanning TEM (cryo-STEM). In particular, the feasibility of differential phase contrast (Dekkers & de Lang, 1974) cryo-STEM utilizing segmented ring detectors acquired under in-focus conditions has recently been demonstrated for biological specimens leading to near-atomic resolution (Lazić et al., 2022). When employing momentum-resolved STEM techniques that use ultrafast cameras to collect a 4D dataset, the obtainable information can still be further extended. In analogy to cryo-CTEM simulations, the interpretability of centre of mass (COM) images (Müller et al., 2014), as well as highly dose-efficient direct ptychographic phase-retrieval schemes such as single sideband (SSB) (Rodenburg et al., 1993; Yang et al., 2015; O’Leary et al., 2021) and Wigner distribution deconvolution (WDD) (Rodenburg & Bates, 1992) ptychography, is equally governed by the validity of the weak phase approximation (for SSB) or by a multiplicative object (for WDD).

A coarse figure of merit for the breakdown of the projection assumption is provided by the Fresnel number (Vulović et al., 2014), which depends on the targeted resolution, the acceleration voltage and sample thickness. The Fresnel number is well known in light optics and helps to separate the validity of the Fresnel, Fraunhofer and ray optics concepts, respectively. Gureyev et al. (2020) presented a multislice study with biological molecules to compare the contributions of multiple scattering and free space propagation. For small molecules such as the amino acid aspartate (141 Da) and lysozyme (14.3 kDa), the conclusion was drawn that propagation is the dominant factor. However, the chosen proteins were rather small and had been simulated in a vacuum, reducing the scattering power and the likelihood of dynamical scattering as well as neglecting the impact of glass-like ice surrounding the protein.

Baxter et al. (2009) emphasized the importance of including structural noise, e.g. owing to the ice background, for the simulation of realistic images. To account for improved modelling of the surrounding ice, Himes & Grigorieff (2021) used coarse grained water pseudomolecules represented by randomly seeded single isotropic scattering centres of the water molecules with a density of low-density amorphous ice. To account for the hydration radius of the protein, they calculated hydration shells in the periphery of the protein derived by the protocol of Shang & Sigworth (2012). Vulović et al. (2013) employed molecular dynamics (MD) simulations to model the glass-like ice embedding using empirical potentials.

In the present study, we treated both the protein and the ice fully atomistically. MD simulations were conducted to generate the atomic structure of the ice up to a thickness of 95.5 nm, followed by a hydration procedure to embed the protein. Subsequently, to obtain the electron wavefunction at the exit face of the specimen, dynamical multislice simulations were performed. The main focus of this study is to investigate the effect of dynamical scattering and propagation inside vitrified proteins using the tobacco mosaic virus (TMV) as a well known test sample in cryo-EM (Sachse et al., 2007; Fromm et al., 2015; Weis et al., 2019). In particular, the deviations of treating the scattering problem as a single-scattering approach are quantified for cryo-CTEM, as well as in-focus cryo-STEM techniques such as integrated COM (iCOM) and ptychographic imaging.

2. Methods

2.1. Complete atomic model of specimen

In order to generate a simulation of ice-embedded TMV, the most recent atomic model of TMV was retrieved from the Protein Data Bank [PDB code 6sag (Weis et al., 2019)] and helically expanded 147 times to generate an atomic model of three complete helical turns along the y axis. MD simulations were performed to model the glass-like ice background (Thompson et al., 2022) using the four-site rigid water model TIP4P/2005 (Abascal & Vega, 2005), as schematically shown in Fig. 1. The fourth site M is massless and contains the charge associated with the oxygen atom. Starting from a water crystal with a size of 251.1 × 251.1 × 15.5 Å with periodic boundary conditions, the MD simulations were initiated by applying a velocity distribution corresponding to a temperature of 280 K.
2.2. Single and multislice simulations

The dynamical interaction between the specimen and the electron wave, including multiple scattering and propagation, was modelled by the multislice algorithm following the theories of Cowley & Moodie (1957, 1959a,b). We briefly recall the central aspects of the multislice theory here to introduce single scattering, then dynamical scattering in general, and to define the terminology. The reader is referred to the comprehensive literature for detailed derivations (e.g., De Graef, 2003; Kirkland, 1998).

The multislice method provides a numerical solution to the relativistic Schrödinger equation by dividing the 3D electrostatic potential of the specimen into slices of thickness $\Delta z$. The potentials within the slices are projected along the electron-beam direction and placed in the slice centres, at which the interaction is taken into account. Between the slice centres, the electron wave is propagated along the distance $\Delta z$ through potential-free space. For $\Delta z \rightarrow 0$, the multislice algorithm is an accurate solution to the relativistic scattering problem. In this work, isolated-atom scattering factors for electrons as tabulated by Lobato & Van Dyck (2014) were used.

2.2.1. Single scattering

Assuming a specimen thickness $\Delta z$ and an incident wavefunction $\psi_{in}(r)$, single scattering is formally described by

$$\psi^{(1)}(r) = F_{\Delta z/2} \otimes \left[ \exp[i\sigma V_{1,\Delta z}(r)] \cdot \left[ F_{\Delta z/2} \otimes \psi_{in}(r) \right] \right], \quad (1)$$

with $F_{\Delta z/2}$ being the Fresnel propagator for propagating a distance $\Delta z/2$, $\sigma$ being the relativistic interaction constant, $V_{1,\Delta z}$ being the projected potential of the first slice, $r$ being a position vector perpendicular to the optical axis and $\otimes$ being the convolution. $\psi_{in}$ is the incoming plane wave in CTEM and a focused probe in STEM. In practice, the convolution theorem is used, which translates the convolution in real space to a multiplication in Fourier space. There, the Fresnel propagator reads

$$\mathcal{F}[F_{\Delta z/2}] = \exp \left( -i\pi \frac{\Delta z}{2} |k|^2 \right), \quad (2)$$

with the spatial frequency vector $k$ and the Fourier transform $\mathcal{F}$. Equation (2) is identical to the effect of defocus, i.e. a phase factor applied in the diffraction plane. The first propagation in CTEM leaves the plane wave unchanged, such that $\psi^{(1)}(r) = F_{\Delta z/2} \otimes \exp[i\sigma V_{1,\Delta z}(r)]$. The exponential term is often termed ‘phase grating’. Equation (1) consists of imprinting a phase modulation on $\psi_{in}$ such that it is termed a PO model. In any single-slice model, the interaction is fully described by the multiplication of the incident electron wave with an object transmission function (OTF).

If, additionally, the phase $\phi(r) = \sigma V_{1,\Delta z}(r)$ is small, Taylor expansion of the phase grating yields $1 + i\sigma V_{1,\Delta z}(r)$, known as the WPO model. Only in this case, the image intensity can be given analytically in closed form. The transfer of spatial frequencies is best studied in Fourier space and the Fourier transformed image intensity $I$ of a WPO is

$$\mathcal{F}[I](k) = \delta(k) - 2A(k)\mathcal{F}[\delta(k) \cdot k_s \cdot k_C \cdot \sin[\chi(k)]], \quad (3)$$

where $A(k)$ is a potentially present objective aperture, $k_s$ and $k_C$ are spatial and temporal coherence envelopes, $\chi(k)$ is the aberration function, and $\delta(k)$ is Dirac’s delta function. From this equation, the established Scherzer conditions for the defocus and the aperture can be derived to maximize contrast transfer within the Scherzer passband for spatial frequencies below the point resolution limit of the microscope.
2.2.2. Dynamical scattering. In the following, we refer to the term ‘dynamical scattering’ as the consecutive action of phase grating and Fresnel propagation. To simulate dynamical scattering, the multislice scheme yields for the wavefunction $\psi^{(n)}$ after interacting with the $n$th slice:

$$\psi^{(n)}(r) = \exp[i\sigma V_{n,L}(r)] \left[ F_{L, n} \otimes \psi^{(n-1)}(r) \right].$$

(4)

The exit wave results from a final propagation by $\Delta z/2$. According to the iterative application of equation (4), multiple scattering and propagation cannot be separated anymore for a given exit wave $\psi^{\text{exit}}(r)$.

The small lattice vibrations at liquid nitrogen temperature were accounted for by multiplying the atomic scattering amplitudes with an isotropic Debye–Waller factor according to a Debye parameter of $B = 0.3$ Å$^2$. For those simulations where the surrounding glass-like ice was to be approximated by its mean inner potential (MIP) (e.g. Himes & Grigorieff, 2021), we used an artificially high Debye–Waller factor of 1809 Å$^2$ for the ice regions, being five times the distance of the bond length between oxygen and hydrogen in the TIP4P/2005 water model. As a result, the potential of the ice becomes practically constant within the embedding, but without introducing abrupt potential steps where the hydration shell of the protein was applied.

2.3. Conventional transmission electron microscopy image simulations

The CTEM simulations were performed using the exit waves formed under plane wave illumination. Aberrations, as well as partial spatial ($k_s$) and temporal coherence ($k_c$), were taken into account within the framework of linear imaging theory by multiplication of the diffraction pattern wavefunction with a phase defined by the aberration function $\chi$ and by partial coherence envelopes $[k_s k_c \exp(i\chi)]$. Additionally, an objective aperture with a radius of 20 mrad was applied if not stated otherwise. An inverse Fourier transform of equation (4) yields the image wavefunction, whose squared modulus corresponds to the recorded image. The simulation parameters can be found in Table 1. Dose considerations as well as detection performance are not included in the current CTEM image simulations.

2.4. Scanning transmission electron microscopy simulations

For STEM, multislice simulations with a focused probe and 24 slices were performed, and the diffraction patterns for each scan pixel have been stored, resembling a recording with a pixelated STEM detector. This 4D dataset was used to calculate the centre of mass (Müller et al., 2014), from which the iCOM signal was obtained by an integration in Fourier space (Section S3 of the supporting information) (Lazić et al., 2016), the SSB reconstruction (Rodenburg et al., 1993) and the WDD (Rodenburg & Bates, 1992; McCallum & Rodenburg, 1992). Both iCOM and WDD and SSB yield the projected electrostatic potential of the specimen in PO and WPO approximation, respectively. WDD will only be discussed very briefly because, for low-dose applications, SSB is advantageous compared with WDD since no deconvolution is needed (O’Leary et al., 2021). Importantly, these methods allow one to reconstruct frequencies up to twice the radius of the probe-forming aperture (Rodenburg et al., 1993). Therefore, we used 10 mrad for the probe-forming aperture, being half as large as the CTEM counterpart. Because SSB and iCOM show the highest contrast in the absence of aberrations, the defocus was chosen as $C_1 = -97.5$ Å, being the centre of the specimen, and no other aberrations were used. It turned out that 24 slices were by far sufficient and were chosen as the maximum number of slices in the STEM simulations.

For iCOM, diffracted intensities up to 1.4 times the bright field disc were used. To deconvolve the probe and $\sigma V$, an aberration-free probe was assumed, which only has consequences for the comparison in real space because it is a radially symmetric rescaling in Fourier space and ignored in the Fourier ring correlation (FRC) curve introduced below. For SSB, the complex value of each pixel was calculated from the average of the double-overlap regions in the 4D dataset, Fourier transformed with respect to the scan position, resulting in a constant transfer function up to a maximum spatial frequency equal to twice the aperture radius (O’Leary et al., 2021).

2.5. Comparison in real and Fourier space

We used two concepts to compare the reconstructions of the proteins for different slicings and imaging methods. An assessment of the impact of dynamical scattering in CTEM is best performed on the basis of the complex specimen exit wavefunctions simulated with different numbers of slices. This poses the challenge of introducing an error metric for a given complex exit wave regarding modulus and phase with respect to the ground truth; here a simulation with 120 slices. To this end, the complex normalized difference is used for comparison in real space and FRC is used in reciprocal space. The complex difference for two matrices $X, Y \in \mathbb{C}$ is calculated by

$$d(X', Y) = \frac{X' - Y}{|Y|},$$

(5)

where $X' = X \cdot \exp(i\phi)$, with $\phi \in (0, 2\pi)$, which contains a constant phase offset such that $||Y_{\text{ice}} - X_{\text{ice}} \cdot \exp(i\phi)||_F$ is minimal in the glass-like ice region, with $||X||_F$ being the

<table>
<thead>
<tr>
<th>CTEM</th>
<th>STEM</th>
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<tbody>
<tr>
<td>Acceleration voltage (kV)</td>
<td>300</td>
</tr>
<tr>
<td>Objective aperture radius (mrad)</td>
<td>20</td>
</tr>
<tr>
<td>Source size (mrad)</td>
<td>0.1</td>
</tr>
<tr>
<td>Energy spread (eV)</td>
<td>0.7</td>
</tr>
<tr>
<td>Pixel size (Å)</td>
<td>0.487±0.12</td>
</tr>
<tr>
<td>Chromatic aberration (Å)</td>
<td>2.7</td>
</tr>
</tbody>
</table>

† The larger pixel size was only used in comparisons with STEM simulations.
Frobenius norm of $X$. The division and the calculation of the absolute value are performed pointwise. The complex difference is invariant against uniform rescaling by a real factor $a > 0$, $d(X, Y) = d(aX, aY)$.

The FRC curve measures the normalized cross-correlation between resolution rings of two matrices ($X$ and $Y$) in Fourier space. It is given by (van Heel et al., 1982; Saxton & Bau-meister, 1982; van Heel & Schatz, 2005)

$$\text{FRC}(k) = \frac{\sum_{k_i=|k|+\Delta k} |\mathcal{F}[X](k_i)|^2 \sum_{k_j=|k|+\Delta k} |\mathcal{F}[Y](k_j)|^2}{\sum_{k_i=|k|+\Delta k} |\mathcal{F}[X](k_i)|^2 \sum_{k_j=|k|+\Delta k} |\mathcal{F}[Y](k_j)|^2},$$

where $\overline{X}$ denotes the complex conjugate of $Y$ and $\Delta k$ denotes the width of the resolution shell, which was chosen as the smallest present spatial frequency determined by the size of the field of view. A consequence of the normalization is that, except for the sign, the FRC is invariant against uniform rescaling with an isotropic function $f(k) = f(k)$ in Fourier space.

3. Results

In order to understand the effects of dynamical scattering, we use a simulation with 120 slices as the ground truth. When only single scattering is considered, two quantities are of interest. Firstly, the OTF itself, whose phase or imaginary part are directly proportional to the projected potential of the specimen in PO and WPO approximation, respectively. These OTFs represent the electron wavefunction in the slice centre. However, multislice simulations yield the electron wavefunction at the exit face of the specimen. Secondly, we thus Fresnel propagated the PO and WPO results from the slice centre to the exit face and termed the results as $PO_{exit}$ and $WPO_{exit}$, respectively. Initially, the impact of dynamical scattering is mainly studied at the level of exit wavefunctions, then at the basis of the image formed by the optical system, and different ice thicknesses are considered. Finally, a comparison between the performance of CTEM and recent phase-contrast STEM techniques is elaborated.

3.1. Dynamical scattering effects on exit waves under parallel illumination

The comparison of the phases of the OTF as well as the phases of the exit waves using single- and 120-slice interaction models are shown in Fig. 2. Figs. 2(a)–2(e) show the phases of the wavefunctions using different interaction models and Figs. 2(f)–2(l) depict the corresponding moduli. Interestingly, the phases of the PO and WPO approximation in Figs. 2(a) and 2(b), respectively, resemble the 120-slice simulation best, indicating that a propagation to the exit face of the specimen leads to less accurate phase distributions of $PO_{exit}$ and $WPO_{exit}$ in Figs. 2(c) and 2(d), respectively. Nevertheless, all phase results show the glass-like ice on the left side and the typical ladder-like structure of TMV at the right side limited by the central TMV symmetry axis. A comparison of the moduli with the ground truth in Fig. 2(i) reveals systematic differences for both WPO approaches in Figs. 2(f) and 2(h), and a similar pattern is shown for $PO_{exit}$ in Fig. 2(g). The dynamic range of the modulus of the 120-slice simulation ranges from 0.34 to 1.65 and is significant, suggesting that one should not conclude here that TMV is a PO. Clearly, the spatial information about the virus assembly is dominantly contained in the phase. In essence, this first overview means that none of the OTF models, defocused to the exit plane or not, yield suitable visual agreement with the multislice ground truth.

Figure 2
Comparison of different simulations of the specimen exit wave of TMV protein embedded in ice. The total specimen thickness $\Delta z_{\text{specimen}} = 19.5$ nm (with no water molecules on the top or bottom surfaces). Only a part of the simulation is displayed due to the symmetry of TMV, as depicted by the red square in the top-right schematic drawing. The phase and modulus of different exit waves are shown in grey scale (min black and max white, as indicated in the bottom-left panels of each part) for (a)–(e) and (f)–(l), respectively: (a) Exit wave of the PO transmission function, (b) Exit wave of the WPO approximation, (c) Exit wave of the PO after an additional propagation of half the specimen thickness, 0.5$\Delta z_{\text{TMV}}$ (d), (h) Exit wave of the WPO after an additional propagation of 0.5$\Delta z_{\text{TMV}}$ (e), (i) Exit wave of a multislice simulation with 120 slices. The scale bar is 25 Å.
truth for phase and amplitude simultaneously. In that respect, PO_{exit} exhibits qualitative agreement in the phase and modulus for low and medium spatial frequencies, whereas high frequencies arise from Fresnel propagation by half the specimen thickness.

The obtained simulations raise two questions for further investigation. First, how many slices are actually needed to take into account dynamical scattering effects reliably? Second, do the observed differences between the single- and multi-slice models have any impact on the results of single-particle cryo-CTEM?

First, the complex normalized real-space differences of exit waves for several models with respect to the 120-slice simulation are depicted colour-coded in Fig. 3. Large errors for the moduli are found for single-slice OTF approaches according to Figs. 3(a)–3(d), decreasing by a factor of 2–4 for 2 slices and by a factor of 6–14 for 20 slices in Figs. 3(e) and 3(f), respectively. Interestingly, the PO approaches in Fig. 3(b) do not show distinct colour features distinguishing ice and TMV. Recalling that the difference to the ground truth is plotted here, this means that the PO approach yields a phase of a wave in the specimen centre that is in very good agreement with a full dynamical simulation, at least by visual inspection. This notion is also supported by the similarity of the PO phase with that of the 120-slice simulation in Fig. 2. Larger and more systematic phase differences for the PO_{exit} and WPO_{exit} simulations in Figs. 3(c) and 3(d), i.e. the Fresnel propagation of the OTF from the slice centre to its exit face, are observed. At least for the phase, a closer match to the ground truth is found at the slice centre as opposed to the exit wave. In conclusion, the phase error decreases with the number of slices and becomes negligible at 20 slices. As the described properties represent real-space data of Figs. 2 and 3, they cannot be easily interpreted with respect to attainable resolution. Therefore, we turn to resolution-based correlations included in FRC curves.

A comparison of the exit waves in Fourier space is provided in Fig. 4 for single-slice OTF-based models and multislice simulations with increasing number of slices (2, 3, 6 and 20). Considering an exit wave simulated with 120 slices as the ground truth again, with an increasing number of slices the FRCs gradually approach 1 across the full spatial frequency band up to 1 Å resolution. In fact, already three slices are sufficient to attain an FRC threshold of 0.4 at 1 Å⁻¹. Taking only two slices into account yields an FRC of 0 at 1 Å⁻¹, and still 0.25 at k = 0.9 Å⁻¹, corresponding to distances of 1.1 Å in the specimen. As the single-slice simulation is typically used for cryo-CTEM, it warrants more careful interpretation. None of the (W)PO FRCs show a significant correct spatial frequency transfer beyond 0.7 Å⁻¹. From the FRC comparison considering the complex-valued waves, the propagation to the exit face of the specimen is favoured over the central slices, as the WPO and PO curves in Fig. 4 have nulls at smaller spatial frequencies than their PO_{exit} and WPO_{exit} counterparts. Among the exit-wave approaches, the WPO_{exit} is the slightly better model. An FRC analysis separated into modulus and phase shows the same trend and is shown in Fig. S1 of the supporting information.

One might argue critically that the similarity of the wave-function to the OTFs is the ultimate criterion, since the retrieval of the structure in terms of the single projection...
might be seen as the ultimate goal. In that case, the FRC of all simulations could have been performed with respect to the PO as ground truth, after propagating back to the centre of the specimen. Even in practice, one could move the objective-lens focus to this location; and indeed, the oscillatory behaviour of the FRC(PO) in Fig. 4 originates from the fact that two functions in different planes along the optical axis are compared here, introducing effects of the CTFs with different foci. In any case, one can revert this argumentation to the equivalent statement that if the PO approach was sufficient in the sense that the 120-slice simulation yielded the same wavefunction in the slice centre after Fresnel propagation back to this plane then the forward propagation of the PO to the exit face of the specimen, POexit in Fig. 4, would yield the same result as the 120-slice simulation. This is obviously not the case for higher spatial frequencies. Therefore, we conclude that a few-slice approach with slice thicknesses below \( \frac{\lambda}{24n_m} \) is needed to describe the interaction between specimen and incident electrons reliably.

The single-slice projection into an OTF can be considered physically relevant only in a limited manner, in contrast to the wavefunction created by dynamical scattering, as obtained by multislice simulation with a converged number of slices. This wavefunction must be considered to reflect the wave present in the object plane of the objective lens in experiments and therefore needs to be chosen as the ground truth. A significant deviation between OTFs and multislice simulation suggests using reconstruction techniques that either account for multiple scattering, introduce prior knowledge or do both.

3.2. Dynamical scattering in CTEM imaging

To simulate the experimentally observable CTEM image intensities, the WPOexit, POexit and 120-slice exit waves from Section 3.1 were subjected to the imaging process with a defocus of \(-500\) nm and a spherical aberration of \(1.5\) mm. The results are compiled in Fig. 5, where the left column represents the image intensities as simulated. The right column depicts the result after correcting the sign of the CTF, also known as phase flipping, to mitigate contrast inversions, assuming equation (3) to be valid.

All three simulations show the outline of the helical assembly of TMV including features of the simulated glass-like ice matching the characteristic appearance of experimental cryo-EM images. However, the WPOexit case does not show internal features, in contrast to the POexit and 120-slice cases, which are both very similar and agree in fine structure. The main difference between the POexit and 120-slice cases is seen in the absolute contrast and the contrast between TMV and the glass-like ice, both being smaller in magnitude for 120 slices. The same effect is observed in the phase-flipped images on the right.

According to Fig. 2, the structural information of TMV is dominantly encoded in the phase of the exit wave, meaning that the recorded image should ideally be similar to Fig. 2(e). Therefore, Fig. 6(a) depicts the FRCs between this signal and Figs. 5(a)–5(c). On the other hand, the image intensity is the experimentally observable quantity, such that FRCs taking the image corresponding to the 120-slice simulation as reference are shown in Fig. 6(b). As expected due to the focus setting, an oscillating CTF can be seen in both comparisons, while the single-slice models (W)POexit exhibit an additional beating of the envelope in Fig. 6(a). Interestingly, the null of the
Nevertheless, the (W)POexit cases yield suitable agreement with the same envelopes and aberrations. This is due to the fact that the reference was subjected to an all models is observed on the level of image intensities, which is rarely taken in the spatial frequency range above 0.2 Å⁻¹, suggesting that a continuum model does not match a fully atomistic treatment over a large range of the spatial frequency spectrum. Furthermore, the FRCs show that using a vacuum or the MIP for amorphous ice yield the same results because the effect of a constant potential offset leads to a rather small global modification of the electron wavelength in the specimen.

3.4. Influence of ice thickness in cryo-CTEM imaging

Fig. 8 shows both the specimen exit-wave phases and the CTEM image intensities for different specimen thicknesses. In particular, the ideal case with no ice embedding on top or below the TMV resulting in 19.5 nm thickness is depicted in Figs. 8(a) and 8(c), whereas Figs. 8(b) and 8(d) show the maximal case of 94.5 nm thickness with two additional ice layers of Δz_{ice} = 37.5 nm thickness encapsulating the TMV. Both results were obtained by fully atomistic MD. In both cases, the contrast is dramatically reduced for the thicker ice, concerning the internal structure of the TMV, as well as the contrast between the TMV and the adjacent ice embedding.

To mimic the averaging process for 2D class averages in single-particle cryo-EM, a total of 500 simulated CTEM images were added for thin, intermediate and thick specimens (34.5, 64.5 and 94.5 nm, respectively). Contrast inversions were accounted for by phase flipping according to the respective CTFs, again assuming equation (3) to be valid. In analogy to Fig. 6(a), FRCs with the exit-wave phase for the minimal specimen thickness of 19.5 nm from Fig. 8(a) have been calculated and plotted in Fig. 8(c). The same curves can be observed when the averaged CTEM simulations are compared with the averaged CTEM simulation with no additional ice on top and bottom, as seen in Fig. S4. As expected from the visual impression in Figs. 8(c) and 8(d), increasing ice thickness leads to higher noise, and decreasing FRCs over the whole range of spatial frequencies are observed. In particular, FRCs drop linearly up to a frequency of 0.4 Å⁻¹ and show a plateau with some modulations above. Interestingly, a threefold increase of the ice layer from 34.5 to 94.5 nm only leads to an FRC drop from ~0.75 to 0.5. Nearly 40 nm of ice on both top and bottom face causes a marginal further FRC decrease by 0.1. To conclude, image quality deteriorates when specimen thickness increases, but more quickly as soon as ice layers of a few nanometres are present, whereas the additional effect imposed by growing ice with tens of nanometres thickness is noticeably smaller.

3.5. Comparison of cryo-CTEM and phase-contrast cryo-STEM

The above analysis has shown several limitations of single-slice models. Nevertheless, it was also found that the (W)PO
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Figure 9
Comparison of conventional and scanning TEM simulations. The total specimen thickness $\Delta z_{\text{specimen}} = 19.5$ nm (with no water molecules on the top or bottom surfaces). (a)–(c) Integrated COM, and phase and modulus of an SSB reconstruction. For the iCOM reconstruction, a deconvolution with the magnitude squared of an aberration-free probe was performed to show the reconstructed $\sigma V$. (d)–(f) Complete power spectra of CTEM, iCOM and SSB on the left, and up to spatial frequencies of 0.25 Å$^{-1}$ on the right. (g) FRC curves with respect to the phase of the OTF. For SSB and WDD, only the phase was used, and for the CTEM simulation a defocus of $-500$ nm and no other aberrations were used. The CTEM simulation was back-propagated to the centre of the specimen. The scale bar is 25 Å.

Figure 8
Comparison of simulations with different specimen thicknesses. (a), (b) Phase of the exit wave for the minimal and maximal ice thicknesses of 19.5 and 94.5 nm, respectively. (c), (d) CTEM simulations for the minimal and maximal ice thicknesses. The total specimen thickness is given in the captions. (e) FRC curves of 2D class averages from 500 CTEM simulations with defocus values randomly distributed over 0.1 to 1.0 μm for different specimen thicknesses of 34.5, 64.5 and 94.5 nm. The exit wave of the central slices that contain TMV is used as the reference. The average is performed with CTEM images that are phase flapped and the additional propagation is considered as an additional defocus. The scale bar is 25 Å.

4. Discussion
From a very general viewpoint, dynamical scattering impacts the direct interpretability of TEM data twofold. Firstly, the electron wave exiting the specimen differs significantly from the projected structure according to the Coulomb potential. Therefore, Fig. 9(g) depicts the FRCs with the projected potential according to the PO approximation. The smooth and quasi-identical frequency transfers of iCOM, SSB and WDD reveal high structural fidelity of the reconstructed signals across the whole frequency range, with an FRC of 0.5 at 0.75 Å$^{-1}$ and 0.25 at 1 Å$^{-1}$. The defocus-related oscillation of the CTF for CTEM, however, is bound by an envelope with a null at 0.7 Å$^{-1}$, suggesting that a larger number of CTEM images with different foci over STEM images is needed to retrieve, in particular, high spatial frequencies faithfully. In this context, phase-contrast STEM appears promising, although a robust and comprehensive comparison with cryo-CTEM will need to account for dose efficiency and noise-transfer characteristics in a future study.
intensity being the pivotal question in quantitative CTEM, the impact of dynamical scattering has naturally been addressed in cryo-CTEM literature before (Vulovic et al., 2014; Gureyev et al., 2020; Himes & Grigorieff, 2021). The outcome, however, has been largely focused on including the principal theory and simulation, but to a smaller extent on the impact on the resulting information transfer in single-particle structure determination in comparison with the state-of-the-art WPO approximation. In that respect, the present study sheds light on the quantitative analysis of how many slices are needed to obtain a certain accuracy in describing interaction and imaging, on the necessity of the MD-based atomistic modeling of the ice embedding, on the impact of various ice thicknesses on the specimen in the electron-beam direction, and on the perspective of recent cryo-STEM techniques for protein structure determination.

For TMV, a large protein assembly of high molecular weight, one can clearly draw the conclusion that single-slice approximations are an inaccurate way to describe the scattering in the specimen in comparison with multislice simulations, as was demonstrated at the level of specimen exit waves. However, we also found that this interpretation concerns mostly high spatial frequencies, since the WPO models show partly acceptable FRCs with a full multislice result up to 0.6 Å⁻¹. Beyond, at least 2–3-slice interaction models are required to achieve 1 Å⁻¹.

In accordance with the exit-wave analysis, CTEM images simulated with single-slice approaches differ substantially at high spatial frequencies from the image formed by the wavefunction from full dynamical scattering imaged with the same optical parameters (Fig. 6). In other words, interpreting high-resolution features within recorded images in terms of single-slice models can be inaccurate, which includes the commonly applied WPO-based CTF correction. At first sight, the observed relationship points to an inherent limit of the achievable spatial resolution independent of the quality of the imaging system used. Interestingly, to date, TMV structures have been experimentally resolved up to 1.9 Å (Weis et al., 2020). This observation for elastic scattering suggests that thick ice coverage reduces the FRC over a minimally embedded protein much less than expected after adding the first thin ice layers. However, contrast is also affected by inelastic scattering (Angert et al., 1996), which needs to be considered for recordings without zero-energy-loss filter or correction of chromatic aberration. Regardless, the simulation results emphasize the requirement of experimentally controlling and minimizing ice thickness for obtaining optimal protein contrast.

As multislice approximations are currently not included in cryo-EM structure determination workflows, the computed 3D density maps will result in lower-resolution maps as well as other systematic density inaccuracies. When atomic models are built, chemical building blocks of amino acids are commonly placed in the obtained isosurface Coulomb potential (Henderson et al., 1990). Such prior information may mitigate the violation of projection assumptions when generating atomic models. In cases when multiple scattering becomes relevant, it is possible to include multislice simulations of the 3D density in the iterative projection matching workflow, which could increase the precision orientation parameter determination for each experimental particle image. Moreover, by improving the simulations of CTEM images of atomic models through multislice approximations, it may ultimately become realisable to include the simulation based on the atomic protein model in the orientation-determination iterative optimization until the atomic model matches the experimental images. The maximization of agreement between experimental diffraction data and the atomic protein model is commonly being performed in X-ray protein crystallography (Brünger et al., 1998).

Formerly predicted (Rose, 1976) and recently implemented (Lazić et al., 2022) STEM phase-contrast imaging motivated the comparison of STEM techniques with cryo-CTEM. Although iCOM, SSB and WDD ptychography involve the projection assumption, the FRC extends up to 1 Å⁻¹ and is
smooth across the whole frequency range. Note that the STEM FRCs still reach a value of 0.5 at 0.7 Å⁻¹, where the FRC between plane-wave PO and the multislice result is already 0 (Fig. 9). In general, the monotony of iCOM, SSB and WDD will be advantageous when small numbers of particles are studied, as the CTF correction in CTEM is an additional processing step enhancing noise that can only be overcome by averaging larger numbers of molecular views. While it was assumed here that all CTF oscillations can be corrected perfectly, a final comparison between CTEM and STEM needs to account for the detection stage, namely Poissonian noise and modulation transfer functions of the cameras. In this respect, SSB ptychography has performed highly efficient and very well with regards to spatial resolution (Maiden et al., 2012) and developments for inverse multislice ptychography (Bangun et al., 2022; Chen et al., 2021; Maiden et al., 2012) provide solutions beyond the projection assumption. Overcoming the (W)PO limitations to CTEM by 2–3-slice phase enhancement in 4D STEM is thus a promising way to explore the opportunity of resolution and contrast enhancement in future cryo-EM.

5. Data availability statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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