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Prospects for Electron Imaging with Ultrafast Time Resolution

Michael R. Armstrong, Bryan W. Reed, and Ben R. Torralva

Materials Science and Technology Division, Chemistry and Materials Science Directorate, Lawrence Livermore National Laboratory, P.O. Box 808; L-356, Livermore, CA 94550

Nigel D. Browning

Department of Chemical Engineering and Materials Science, University of California-Davis, One Shields Ave, Davis, CA 95616

Abstract

Many pivotal aspects of material science, biomechanics, and chemistry would benefit from nanometer imaging with ultrafast time resolution. Here we demonstrate the feasibility of short-pulse electron imaging with 10 nanometer/10 picosecond spatio-temporal resolution, sufficient to characterize phenomena that propagate at the speed of sound in materials (1-10 kilometer/second) without smearing. We outline resolution-degrading effects that occur at high current density followed by strategies to mitigate these effects. Finally, we present a model electron imaging system that achieves 10 nanometer/10 picosecond spatio-temporal resolution.

Due to the speed limitations of electronics, typically ultrafast time domain experiments (including imaging) require "flash" illumination of the sample in analogy with time-resolved pump-probe optical spectroscopy. The time resolution for this technique is determined by the duration of the pump and probe pulses – the pulse duration must be shorter than the time scale of dynamics of interest. The analogous experiment with electron imaging (or diffraction) involves initiation of sample dynamics followed by sample illumination with an electron probe pulse. Scattered electrons from the probe pulse may be projected via electron optics onto a 2D detector to obtain a magnified image (or diffraction pattern) of the sample, as shown in Fig. 1. For single shot imaging, the image must be obtained with a single electron probe pulse after a single pump pulse.

Pump-probe x-ray and electron diffraction techniques have recently been employed in the investigation of phenomena ranging from biomolecule structural dynamics to ultrafast melting [1-9]. Diffraction experiments have a great advantage over spectroscopy, as diffraction directly probes atomic positions. Typically, diffraction experiments can achieve atomic (Angstrom scale) spatial resolution even with ultrafast time resolution, while static imaging with electrons can achieve sub-Angstrom resolution.[10,11] Although such results provide invaluable insight into ultrafast processes on atomic length scales, diffraction is significantly limited in field of view, requires a sample with a periodic character, and is inherently an ensemble measurement [6, 12]. Yet, material dynamics often occur in contexts which are inherently non-periodic with a larger field of view [13]. Diffractive imaging techniques are being developed to address these issues [6,14-16], but direct imaging is ideally suited to characterize such systems and electron imaging has played a pivotal role in the discovery of phenomena that are difficult to observe with diffraction [17,18]. Also, the

dynamics of phenomena at the 10 nanometer scale are often unique and non-reversible. At ultrafast time scales, such data may only be obtained with a single shot probe. Here we define ultrafast as sufficient time resolution to characterize effects that propagate at the speed of sound in the sample without smearing at the achievable spatial resolution.

The primary problem with obtaining high temporal and spatial resolution with electrons in this scheme involves the degradation of resolution through Coulomb interactions within the probe pulse. Although ultrafast stroboscopic imaging avoids space charge issues [19], in general imaging is most useful for the observation of stochastic, irreversible processes which are difficult to consistently prepare. Yet, ultrafast stroboscopic imaging requires a sample which can rapidly, identically refresh for up to 10⁷ experiments to built up an image. Here we outline the relevant issues for single shot ultrafast imaging and present a model electron optical system which obtains high spatial resolution ultrafast images.

To obtain a high quality image, a minimum scattered particle fluence (i.e. detected particles scattered per unit sample area) is required to achieve a given spatial resolution via shot noise statistics. A rough constraint for distinguishing features in an image is given by requiring the shot noise in a single pixel to be less than the contrast in the image by a quality factor – the Rose criterion [20],

$$Shot \ noise \le \frac{contrast}{\# \ of \ gray \ scale \ levels} \tag{1}$$

Typically, the number of gray scale levels (the contrast divided by the noise) must be around 3-5 for minimum acceptable image quality. Shot noise is given by

$$Shot \ noise = \frac{1}{\sqrt{N}} \tag{2}$$

where N = the number of detected particles per pixel. So, for 10^6 detected particles scattered from a sample with 50% contrast in a shot noise limited imaging system, the noise would have to be below 10% by eq. 1 (assuming 5 gray scale levels). By eq. 2, this requires a pixel size that is sufficiently large to detect an average of 100 particles per pixel, giving 10^4 pixels for 10^6 total detected electrons across the field of view – a 100x100 image. In the case of a $1 \mu m^2$ illuminated sample region, this implies a shot noise limited resolution of 10 nm. For imaging systems which use direct current to illuminate the sample, the peak current can be arbitrarily small (i.e. low current with long integration time), effectively eliminating Coulomb interactions between electrons. For ultrafast imaging, the added requirement to obtain high temporal resolution in a single shot constrains the peak current to be much higher. In particular, an electron pulse of 10 ps duration containing 10^6 electrons has a peak current of approximately 20 mA, many orders of magnitude larger than the current found in standard transmission electron microscopy (~ nA). In this regime Coulomb effects are significant and must be considered in the electron optical design of the imaging system.

The effects of Coulomb interactions in an electron pulse can be broadly divided into two categories: large scale, or "global" space charge (GSC) effects, and stochastic scattering between individual electrons. GSC effects are relatively straightforward: on large scales a uniform distribution of charge remains uniform as it propagates [21, 22]. From an electron optical perspective, uniform global space charge behaves precisely as a negative lens and the effects of global space charge can be corrected to first order by electron optics. In a real imaging system, the electron pulse will not be uniform, and global space charge may result in higher order aberrated distortions of the electron pulse, potentially requiring the use of aberration corrected electron lenses [23]. Global space charge will not significantly degrade the resolution with proper design of the

imaging system, but random smaller scale electron interactions still limit the achievable resolution at high current density.

Stochastic scattering fundamentally limits the achievable resolution through random electronelectron scattering, which results in irreversible loss (blurring) of image information as the pulse
propagates [24, 25]. Essentially, image information encoded in electron trajectories (after being
scattered from the sample) is lost via thermal equilibration. This effect has a significant impact on
high current charged particle optical systems and has been modeled within this context. [24-30]
Typically, the degree of resolution degradation is characterized by a blur profile – a histogram of
the deviation (in the image plane) between image forming electrons which are ray traced through
the optical system and electrons whose trajectories are perturbed by Coulomb interactions. This
procedure gives a blur profile, which is similar to the point spread function used as a transfer
function in traditional optics and may be used to estimate the optimum resolution of the imaging
system. Since blur cannot be completely eliminated, it must be mitigated by reducing the current
density through instrument design.

We explored two strategies to reduce the effects of stochastic blur: the use of relativistic pulses and annular dark field imaging. For picosecond duration pulses of sufficient charge to image in a single shot, the use of relativistic electron energy is necessary to mitigate stochastic blur and maintain an ultrashort pulse duration over relatively long propagation lengths [31]. Although blur is significantly reduced at high electron energy, our simulations indicate that even for short propagation lengths (~1 cm) at 5 MeV electron energy, 10^6 electrons in 10 ps gives roughly ~10 nm of blur. An actual imaging system would involve much longer propagation lengths (resulting in

higher blur), so this strategy alone is insufficient to achieve 10 nm/10 ps spatial-temporal resolution. In principle, arbitrarily high energy pulses could be used to reduce blur indefinitely, but this is practically limited by the strength of electron lenses.

To achieve high resolution, it is also necessary to reduce the average current density. One way to accomplish this is annular dark field imaging (ADF). In ADF, the image is formed by scattered electrons, as shown in Fig. 1. Light regions in the image correspond to regions of strong scattering in the sample [32]. ADF is advantageous because it removes electrons that would only contribute to a bright background without carrying image information about the sample. Also, by only accepting electrons that are scattered to large angles, the average current density in the optical system can be significantly reduced. Finally, since only scattered electrons form the image in ADF, the total image contrast is 100%, which decreases the minimum required fluence from the Rose criterion.

To demonstrate the feasibility of high spatio-temporal electron imaging, we modeled an imaging system with a 5 MeV illumination electron pulse focused to a 1 micron diameter spot on a copper sample with sufficiently small divergence for unscattered electrons to be blocked after the sample, allowing scattered electrons to continue through the imaging system (as shown in Fig. 1). We used electron differential scattering cross sections calculated by the ELSEPA computer program [33] assuming a Fermi-Thomas model of the nucleus for a solid copper sample. The simulated electron optical system consists of thin lenses with focal lengths of 1.0 cm (objective), 0.74 cm (1st intermediate), and 1.6 cm (2nd intermediate), with magnifications of 6.3x, 47x, and 18x, respectively, for a total magnification of ~5300x. The dynamics of the electron pulse were solved by computation of Coulomb forces without approximation and propagation of point particles via a

symplectic leap frog algorithm, typically performed on between 10-40 nodes of a Unix cluster at LLNL for 2 x 10⁴ electrons at a density consistent with 10⁶ electrons in a 10 ps duration pulse. There was not a strong dependence on the number of electrons at the same density for $> 2 \times 10^4$ electrons, indicating that edge effects are negligible in this model. The stochastic blur was calculated for three stages of magnification since blur did not increase significantly thereafter, where the current density is smaller and blur is less significant on the scale of the magnified image. The blur profile of this imaging system with a Gaussian profile fit is shown in Fig. 2. As illustrated in the figure, these simulations indicate it is possible to obtain less than 10 nm of stochastic blur with a 10 ps duration pulse, assuming 10⁶ electrons scattered to greater than 5 mrad half angle. Assuming single electron sensitive detection and 10⁶ image forming electrons, the illumination pulse need not have more than a few pC of charge to obtain 10 nm spatial resolution with a pulse that is short enough to obtain 10 ps time resolution. A more intuitive illustration of the spatial resolution is shown in Fig. 3, which shows a CdTe tetrapod, CdTe nanoparticle and gold nanoparticle imaged using high angle ADF imaging [32] (figure 3a) and a simulation of this image assuming the blur and noise limitations of the model (figure 3b). The object image was used as presented in ref. 32. Figure 3b was constructed by assuming a random, spatially uniform distribution of electrons are scattered in proportion to the image intensity of figure 3a, with a scattered, detected fluence sufficient to satisfy the Rose criterion (10⁶ e/µm²) where the intensity of figure 3a is 100%. To generate figure 3b, these "scattered" electrons are then binned into 5x5 nm² pixels (on the scale of figure 3a) after randomly smearing their positions using the distribution of figure 2 to account for stochastic scattering. Fine structure at the 10 nm scale (such as the boundary of the gold nanoparticle) is captured in this image, demonstrating the possibility of high resolution imaging using an ultrashort pulse of electrons. Without relativistic electrons and ADF imaging with

10⁶ scattered electrons, stochastic scattering would randomly distribute these image electrons over a region that is larger than the field of view, completely washing out image information on the scale of fig. 3a.

Given the feasibility of this method in principle, two technologies require further development to realize such an instrument in practice: high beam quality short-pulse electron guns and powerful magnetic lenses systems capable of operating in the several-MeV range. To obtain a sufficiently small illumination area and beam divergence, the transverse geometric beam emittance must be on the order of some nm. Furthermore, standard transmission electron microscopy is limited by chromatic aberration, which constrains the variation in electron energy to less than 10⁻⁵ for Angstrom scale resolution. Although typical radio frequency (RF) accelerators can achieve transverse normalized emittance around 1 µm with 0.1-1.0% energy variation, more work is required to achieve the generation of high quality ultrashort electron beams [21, 34, 35]. Furthermore, we have found that stochastic blur is mitigated by reducing the distance to achieve high magnification, which implies very strong lenses with focal lengths of order 1 cm or less for 5 MeV electrons. Such short focal lengths at MeV energy will require powerful, possibly superconducting lenses [36, 37].

In conclusion, the optimum resolution in electron imaging is constrained at low current density by signal to noise requirements and at high density by stochastic scattering of electrons. Simulations of stochastic scattering in a model electron imaging system indicate that it is possible to obtain 10 nm spatial and 10 ps temporal resolution by imaging with 5 MeV electrons and the use of annular dark field imaging. Such an imaging system would require improvements in the beam quality available

from short-pulse MeV electron sources and the development of high power electron lenses, and would open a horizon in the study of ultrafast material dynamics.

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Figure captions

Figure 1: Schematic of annular dark field (ADF) imaging. Scattered "dark field" electrons are collected to form the image. An image may be formed with "bright field" electrons, but this image includes unscattered background electrons which increase the current density without carrying image information.

Figure 2: Blur profile for the modeled optical system with 20k electrons at the equivalent density of 10⁶ electrons in 10 ps, fit to a Gaussian profile. The FWHM is 6.6 nm, which is within the resolution limit given by shot noise.

Figure 3: (a) ADF image of a CdTe nanoparticle, tetrapod, and gold nanoparticle [32] and (b) a simulation of the same image assuming a scattered, detected fluence sufficient to satisfy the Rose criterion, and spatial smearing corresponding to the distribution shown in Figure 2. The image would be washed out on the scale of fig. 3(a) without ADF and relativistic electrons.

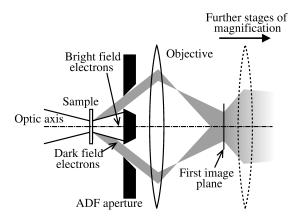


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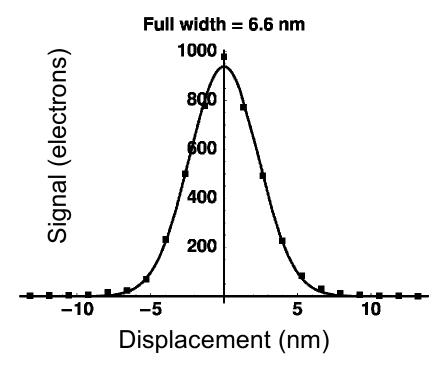


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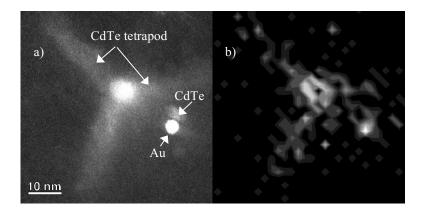


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