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Fast collisional electron heating and relaxation in thin foils driven by a circularly polarized ultraintense short-pulse laser

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The creation of well-thermalized, hot and dense plasmas is attractive for warm dense matter studies. We investigate collisionally induced energy absorption of an ultraintense and ultrashort laser pulse in a solid copper target using particle-in-cell simulations. We find that, upon irradiation by a 2×10^{20} W cm⁻² intensity, 60 fs duration, circularly polarized laser pulse, the electrons in the collisional simulation rapidly reach a well-thermalized distribution with ~ 3.5 keV temperature, while in the collisionless simulation the absorption is several orders of magnitude weaker. Circular polarization inhibits the generation of suprathermal electrons, while ensuring efficient bulk heating through inverse bremsstrahlung, a mechanism usually overlooked at relativistic laser intensity. An additional simulation, taking account of both collisional and field ionization, yields similar results: the bulk electrons are heated to ~ 2.5 keV, but with a somewhat lower degree of thermalization than in the pre-set, fixed-ionization case. The collisional absorption mechanism is found to be robust against variations in the laser parameters. At fixed laser pulse energy, increasing the pulse duration rather than the intensity leads to a higher electron temperature.

PACS: plasma simulation, plasma heating, plasma dynamics

1. Introduction

The creation of warm dense matter (WDM) or hot dense matter (HDM) in a laboratory setting is of high interest for a broad field of research disciplines such as laboratory astrophysics (Remington 2005; Bailey *et al.* 2007; Fujioka *et al.* 2009), studies of planetary interiors (Ross 1981; Knudson, Desjarlais & Dolan 2008), inertial confinement fusion (Drake 2018; Le Pape *et al.* 2018), understanding the equations of state under such extreme conditions (Renaudin *et al.* 2003; Nettelmann *et al.*

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2008) and experimental verification of high energy density (HED) atomic physics models (Hoarty *et al.* 2013*b*; Faussurier & Blancard 2019). However, in order to benchmark atomic physics models against spectroscopic data, these must be obtained under well-controlled conditions. Since most such models assume Maxwellian plasma populations, this means that, when diagnosed, the heated samples should be as close to thermal equilibrium as possible.

The generation of WDM/HDM at uniform near-solid density requires that the sample be heated rapidly, i.e. before any significant hydrodynamic expansion. Such isochoric heating can be achieved using ultrahigh-intensity, short-pulse lasers, as has been done at various high-power systems (Evans *et al.* 2005; Gregori *et al.* 2005; Martinolli *et al.* 2006; Chen *et al.* 2007; Nilson *et al.* 2009; Pérez *et al.* 2010; Brown *et al.* 2011; Hoarty *et al.* 2013a). These experiments were conducted using laser pulses with 0.3–10 ps duration and energies in the range of 10–500 J, but there is a need for a wider access at lower-energy table-top facilities, typically delivering joule-level, femtosecond laser pulses. Promising results in this direction have recently been obtained by Purvis *et al.* (2013) and Bargsten *et al.* (2017) making use of nano-wire arrays to strongly enhance the laser-to-plasma coupling efficiency, thus creating keV temperature, sub-solid density plasmas. Yet, such structured targets are non-trivial to manufacture and are extremely sensitive to parasitic laser prepulses, which can destroy the nano-structures before the main pulse arrives.

Most laser-based isochoric heating experiments conducted so far have exploited the fast electrons driven by a linearly polarized laser pulse (Nilson *et al.* 2010; Santos *et al.* 2017; Sawada *et al.* 2019). Their energy dissipation through the plasma bulk enables heating to high temperatures (0.1–1 keV) at solid-range plasma densities, but usually at the expense of poor spatial uniformity (Dervieux *et al.* 2015) and relatively slow thermalization. Plasma heating in this case is caused by the interaction of the fast electrons with the bulk plasma via a combination of direct collisions (Robinson *et al.* 2014), ohmic dissipation of the colder return current (Lovelace & Sudan 1971; Guillory & Benford 1972; Bell & Kingham 2003; Robinson *et al.* 2014) or plasma waves driven by the fast electrons (Sherlock *et al.* 2014). Some experiments have been done with laser-accelerated ions to heat a secondary target (Patel *et al.* 2003; Dyer *et al.* 2008; Mančić *et al.* 2010). Yet, while this heating method can provide better spatial uniformity, it leads to much lower (~10 eV) temperatures.

At normal laser incidence and linear polarization (LP), and for sharp-gradient, highly overdense plasmas, the most commonly invoked mechanisms of laser energy conversion into fast electrons are $j \times B$ heating (Kruer & Estabrook 1985) and vacuum heating (Bauer & Mulser 2007; May et al. 2011). Both mechanisms hinge on the temporal modulation of the laser ponderomotive force around the target surface, and thus lead to periodic injection of MeV range electron bunches into the target at twice the laser frequency. Such suprathermal electrons thermalize relatively slowly (\sim ps), which may hinder those applications that require a closely Maxwellian dense plasma. In a work by Kemp & Divol (2016), it is shown that the fast electron bunches induce surface waves that can scatter the energized bulk electrons, thereby improving absorption. They also show the necessity of collisions to first heat up the target surface to keV temperatures, required for vacuum heating to commence.

By contrast, using laser pulses with circular polarization (CP), for which the ponderomotive force does not show high-frequency oscillations, the $j \times B$ and vacuum heating mechanisms are essentially suppressed in overdense targets, and so is the fast electron bunch production (and the surface waves induced by them). Still, some fast electrons can be produced with CP if the variation time scale of the laser envelope is not large compared to the laser cycle (Siminos *et al.* 2012, 2017).

In this paper, we study the effects of collisions on the energy absorption capability of the electrons in a thin, solid foil of a high-atomic-number element. Due to the high atomic number, it is not clear a priori what degree of ionization (Z^*) the ions have throughout the process and what influence the ionization history has on heating. While a high Z^* is desirable for the collisional heating process, the initially cold target will not be highly ionized in the beginning. Therefore, we have studied both different degrees of fixed ionization as well as the self-consistent ionization process including both field and impact ionization.

We demonstrate that the energy absorption of an intense short laser pulse in a high- Z^* solid-density target is mainly due to inverse bremsstrahlung electron heating within the plasma skin layer, and that this scenario holds in a broad range of experimentally relevant parameters. The front-layer electrons are collisionally scattered into the target body where they heat the plasma bulk to keV level temperatures, enough to reach Gbar range pressures, which is well in the regime of HDM. The scattered electrons have sufficiently low energies that they primarily heat the bulk via direct collisional thermalization. Since this mechanism relies on the scattering of the electrons accelerated by the laser field against the heavy ions, it is operative regardless the polarization. Inside the plasma, where the laser field is negligible, collisions cause fast relaxation of the electron distribution to a Maxwellian.

2. Simulation design

We have performed one- and two-dimensional (1-D and 2-D respectively) particle-incell (PIC) simulations of laser-solid interactions with and without collisions enabled. We have used the Smilei PIC code (Derouillat *et al.* 2018), which has a relativistic binary collision module (Pérez *et al.* 2012) based on the collisional algorithm by Nanbu (1997) and Nanbu & Yonemura (1998). In the case of a collisional plasma, we have considered either a fixed degree of ionization or self-consistent modelling of the ionization process – through field ionization and collisional impact ionization.

We ran 1-D simulations in a box of size 20 μ m with a resolution of Δx = 0.39 nm (51 200 cells). We considered both LP and CP laser pulses with wavelength λ = 800 nm, dimensionless amplitude¹ a_0 = 10 (intensity $I = \frac{1}{2}c\epsilon_0(m_e c\omega a_0/e)^2 \approx 2 \times 10^{20}$ W cm⁻², where ϵ_0 is the vacuum permittivity, m_e the electron mass, e elementary charge and ω the laser frequency) and a Gaussian temporal profile with $t_{\rm FWHM}$ = 60 fs full-width-at-half-maximum (FWHM) duration in the intensity. The plasma is 2.5 μ m thick, starting at x = 7.5 μ m with a linear density ramp over a distance of 20 nm. The plasma consists of electrons and copper ions at solid density, $n_{\rm Cu,0}$ = 48.4 $n_{\rm c} \approx 8.4 \times 10^{22}$ cm⁻³, with 400 macro-particles per cell for each species. Here, $n_{\rm c} = \epsilon_0 m_{\rm e} \omega^2/e^2$ is the critical density associated with the laser frequency ω . The particles are initialized from Maxwell–Jüttner distributions (in three momentum dimensions) with temperatures $T_{\rm e,0}$ = 1 eV for the electrons and $T_{\rm i,0}$ = 0.1 eV for the ions

In order to assess the influence of the plasma collisionality alone, we have first carried out simulations with fixed ionization degrees $Z^*=11$, 19, 24 and 27. Then, to ascertain the physical accuracy of these results, we have performed simulations describing both collisional and field ionization. The collisionless skin depth $l_{\rm s}=c/[\omega(n_{\rm e}/n_{\rm c})^{1/2}]$ is resolved, even for the highest ionization where

¹Note that the amplitude is normalized such that the intensity stays the same regardless of ellipticity, i.e. the field amplitude, at the same a_0 with circular polarization is $E_{\rm CP} = E_{\rm LP}/\sqrt{2}$ compared to that of LP, $E_{\rm LP}$.

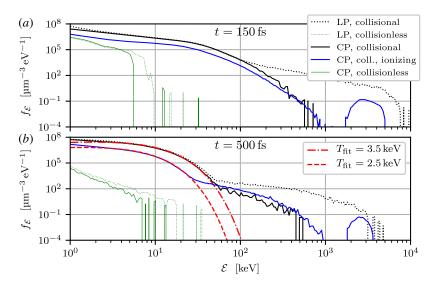


FIGURE 1. Electron energy spectra $f_{\mathcal{E}}$ at times t=150 fs (a) and t=500 fs (b), for LP (dotted lines) and CP (solid lines), with (black lines) and without collisions (thin, green lines); also showing CP with self-consistent field and collisional ionization (blue, solid line). Two Maxwell–Jüttner distributions are fitted to the bulk of the CP self-consistent and fixed-ionization electron spectra in panel (b) (red dashed and dash-dotted lines respectively).

 $l_{\rm s}^{(Z^*=27)} \approx 3.5$ nm. The values of $Z^*=11$, 19 and 27 correspond to full depletion of different electronic shells, $Z^*=27$ being the reference ionization used in other scans. An additional data point, $Z^*=24$, was chosen as an arbitrary value between 19 and 27. When modelling the ionization process self-consistently, the ions were initialized with $Z_0^*=5$, in accordance with the widely used Thomas–Fermi model (More 1983). Both field-tunnelling and electron–ion impact ionization were enabled. The self-consistent ionization simulation was only performed with CP.

We also performed one collisional 2-D simulation to check that our results are robust to multidimensional effects. This simulation uses the same CP laser and target parameters as our 1-D base case. In order to limit the computational cost at the increased dimensionality, it was performed at a reduced resolution of 640 cells per micron in both directions ($\Delta x = \Delta y = 1.56$ nm), and a simulation box size of 10 μ m longitudinally and 1.6 μ m (2 λ) transversely. Furthermore, the number of particles per cell per species was reduced to 50. A test of these resolution parameters in one dimension showed excellent agreement in electron kinetic energy spectrum of the main body of the electrons with the corresponding high-resolution, collisional 1-D simulation; however, the lower particle count led to a poorer statistics in the high-energy tail of the electron spectrum.

3. Results and discussion

Figure 1 compares the electron energy spectra as obtained at CP and LP ($Z^* = 27$), with or without collisions enabled and at CP with self-consistent ionization. The spectra are recorded at two successive times (t = 150 fs and t = 500 fs) – for reference, the peak laser intensity hits the target at $t \approx 110$ fs and the pulse FWHM duration is

60 fs. For both types of polarization, much higher electron energies are achieved when allowing for collisions. The collisionally enhanced absorption results in a bulk electron temperature of $T_{\rm e} \approx 3.5$ keV at t=500 fs for both LP and CP, determined by fitting Maxwell–Jüttner distributions to the bulk spectra (i.e. ignoring the tails). Meanwhile, the collisionless simulations only reach an electron temperature of $\sim 10-100$ eV; these electrons are, however, far from being thermalized and only their energetic tails are visible in the figure. The fact that both CP and LP reach very similar bulk electron temperatures when collisions are enabled indicates that the laser absorption mechanism is the same in both cases.

By comparing the electron spectra with the fitted Maxwellians, we conclude that the electrons have reached a degree of thermalization wherein less than 0.1% of the kinetic energy is in the high-energy tail for CP, and \sim 1% for LP. The time frame for this thermalization is consistent with a rough estimate for electron–electron collisional energy transfer rates. At t=500 fs, the deviation from Maxwellian starts at an electron energy $\mathcal{E}\simeq50$ keV. This is consistent with the \sim 300 fs collisional energy relaxation time of a 50 keV electron through a 3.5 keV plasma at 2.3×10^{24} cm⁻³ electron density (Huba 2016).

Note that these results compare LP and CP for the same on-target intensity, while in an experiment, the circularly polarized pulse would in practice be at a somewhat lower intensity due to losses in the conversion from LP to CP (through a quarter-wave plate). Another practical concern may be elliptical polarization, due to imperfections in the CP conversion. While the exact dependence of the fast electron generation on the ellipticity of the polarization is non-trivial, the bulk collisional absorption itself is not expected to be affected by the ellipticity, as demonstrated by the same bulk temperatures reached in the two extreme cases of CP and LP.

The electron temperatures we quote are technically calculated before the electrons have fully thermalized with the ions, which occurs over \sim ps time scales. However, due to the high degree of ionization, the electron–ion equilibrium temperature is $T_{\rm eq} \approx n_{\rm e}/(n_{\rm e}+n_{\rm i})T_{\rm e} \approx 0.96T_{\rm e}$. Thus, energy transfer from the electrons to the ions is insignificant.

When activating self-consistent (both field and collisional) ionization, the bulk electron temperature is slightly reduced (\sim 2.5 keV) compared to the fixed-ionization case. The front plasma is rapidly ionized, mostly through field ionization, so that collisional absorption quickly reaches an efficiency similar to that obtained with fixed $Z^*=27$ (see figure 2b showing that the average ionization $\langle Z^*\rangle\simeq 24$ at the plasma front already at t=100 fs). The lower T_e is mostly due to the energy spent on ionization – the average ionization energy from $Z^*=5$ to 27 is 0.9 keV.

Moreover, figure 1(b) shows that, for both CP and LP, collisions cause efficient bulk electron thermalization as early as t = 500 fs. High-energy tails are found to emerge above ~ 50 keV for the fixed ionization and ~ 30 keV for the self-consistent ionization. Note the large range of the logarithmic $f_{\mathcal{E}}$ scale, meaning that the tails are three to five orders of magnitude lower than the bulk spectra. The non-thermal tail is heavier in LP than in CP, due to the operative $j \times B$ and vacuum heating.

Also, the simulation with self-consistent ionization displays a larger tail, compared to the bulk spectrum, than its counterpart with fixed Z^* . The larger tail as well as an electron population at ~ 3 MeV can be explained by field-ionization events in the charge-separation layer, which is exposed to stronger laser fields. As the target front electrons are being pushed back by the ponderomotive force, the ions remaining in the charge-separation layer experience the less shielded laser field which quickly ionizes them further. Since these newly freed electrons are injected into regions of stronger

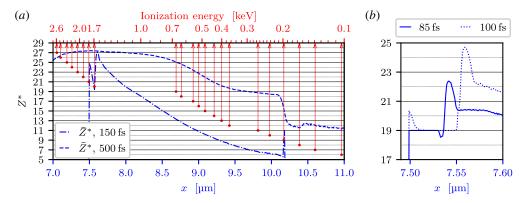


FIGURE 2. Average ionization level profiles \bar{Z}^* in the self-consistent field and collisional ionization simulation (blue lines, bottom axis) at times t=150 fs (dash-dotted line) and t=500 fs (dashed line), and ionization level as a function of ionization energies of copper (red dots and arrows, top axis) – ionization data obtained from the Atomic Spectra Database of the National Institute of Standards and Technology (Kramida *et al.* 2018). Panel (b) shows the average ionization level profiles \bar{Z}^* near the target front at t=85 fs (solid line) and t=100 fs (dotted line).

laser fields, they are energized similarly to vacuum heating in LP, thus resulting in a larger population of non-thermal electrons, which, as in LP, thermalize relatively slowly. Furthermore, the average ionization level is lower inside the target with self-consistent ionization, as seen in figure 2, thus reducing the efficacy of collisional thermalization. Both these effects act to give a larger high-energy tail.

The ionization level profiles (\bar{Z}^*) of the self-consistent ionization simulation shown in figure 2(a) represent the local ionization levels averaged over all macro-particles in each spatial cell. Comparing the t=150 fs (dashed-dotted line) and the t=500 fs (dashed line) average ionization curves, we see that the target front quickly reaches a high ionization degree while the bulk is ionized more gradually. Since there are no strong electric fields inside the plasma, the ionization of the bulk must be driven by impact ionization. The \bar{Z}^* curve at t=500 fs (dashed line) in figure 2(a) displays plateaus at $\bar{Z}^*=27$, 19 and to some extent 11. These plateaus result from the large jumps in ionization energies between the successive outermost electron shells (e.g. between $\bar{Z}^*=11$ and 12, or $\bar{Z}^*=19$ and 20; see figure 2a). The ionization energy to reach $\bar{Z}^*=28$ is ~ 11 keV

Figure 2(b) shows \bar{Z}^* at the target front surface ($x = 7.5 \, \mu \text{m}$). At time $t = 85 \, \text{fs}$ (solid line), the ionization level has saturated at $\bar{Z}^* = 19$ due to the jump in ionization energy after $Z^* = 19$. Later, at $t = 100 \, \text{fs}$ (dotted line), the laser field has become strong enough to sustain field ionization beyond $Z^* = 19$, yielding the peak in \bar{Z}^* near $x = 7.5 \, \mu \text{m}$. Apart from the laser field, the electrostatic field (E_x) induced by the laser ponderomotive force at the target front causes additional ionization. This results in the \bar{Z}^* peak seen around $x = 7.55 \, \mu \text{m}$, which moves into the plasma as the charge-separation layer is pushed forward by the laser ponderomotive force.

The difference between the simulations is made clearer when studying the electron phase spaces shown in figure 3.² The figure displays time sequences of the collisional

²The normalization f/f_{max} of the distribution functions in figures 3 and 5 are with respect to the initial maximum value of the distribution function in their respective planes of phase space f_{max} . The colour values of the plotted distributions can therefore be directly compared.

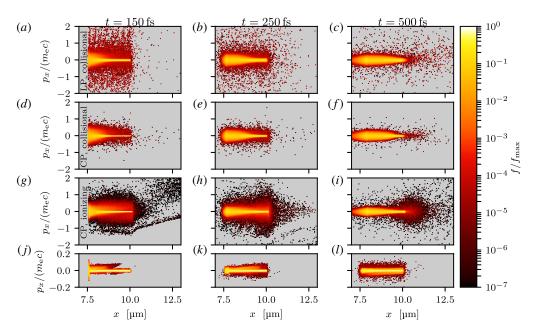


FIGURE 3. Electron phase-space distributions for collisional LP with fixed ionization (a-c); collisional CP with fixed ionization (d-f) and self-consistent field and collisional ionization (g-i) as well as for collisionless CP (j-l), at times t = 150 fs (a,d,g,j), 250 fs (b,e,h,k) and 500 fs (c,f,i,l). Note the different momentum scale for the collisionless CP.

distributions with LP in the top row and CP in the second row; the third row shows the self-consistent ionization CP simulation and the bottom row shows the collisionless CP distribution. In the LP simulation, high-energy electron bunches are produced at twice the laser frequency, as seen in the t=150 fs panel (top row), while CP with fixed ionization (second row) produces a more even distribution of hot electrons since $j \times B$ and vacuum heating mechanisms are inhibited. At t=500 fs, most of the fast electrons have thermalized in the fixed-ionization case, while there remains a significant population of high-energy electrons 'swarming' around the back of the target with self-consistent ionization.

With self-consistent ionization, two populations of relatively high-energy electrons are created during the rising phase of the laser pulse. These populations originate from two successive field-ionization phases. The first one occurs early in the interaction, when the ionization of the surface plasma momentarily saturates at $\bar{Z}^*=19$. In the t=150 fs panel of figure 3, this population accounts for the broad momentum distribution in the target bulk, and also for the beam (with momenta $p_x/(m_e c) \sim -1$) being reflected in the vacuum ($x>10~\mu m$) and refluxing into the target. The second phase starts at $t\simeq 90$ fs, when the laser pulse gets intense enough to ionize the surface plasma beyond $Z^*=19$ (compare the 85 fs and 100 fs curves in figure 2b). This yields fast electrons (visible in the upper right corner of the t=150 fs phase space) more energetic than those generated earlier, which correspond to the bump around ~ 3 MeV in the energy spectra of figure 1. A similar field-ionization injection of fast electrons from the surface ions was seen by Kawahito & Kishimoto (2017) in a carbon plasma, although they used LP which also caused bunching of the electrons at twice the laser frequency.

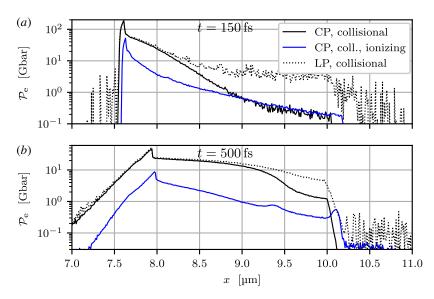


FIGURE 4. Electron kinetic energy density $\mathcal{P}_{\rm e}$ for collisional LP (dotted line) and CP (black solid line) and CP with self-consistent field and collisional ionization (blue solid line), at times t=150 fs (a) and t=500 fs (b). The peak laser intensity hits the target at $t\approx 110$ fs.

We now turn our attention to the energy density achieved in these scenarios. Since the heating process is fast compared to hydrodynamical time scales, the plasma bulk has not had time to expand, and hence the bulk electrons and ions remain at solid-range density. At the same time, the electrons reach keV temperatures, resulting in high energy densities of the order of $\sim 10~{\rm Gbar} = 10^9~{\rm J}~{\rm cm}^{-3}$. In figure 4, the electron kinetic energy density $\mathcal{P}_{\rm e}$ is displayed throughout the target at times $t=150~{\rm fs}$ (top panel) and $t=500~{\rm fs}$ (bottom panel). At the earlier time, the kinetic energy density is concentrated to the front of the target, while at the later stage the energy has spread out throughout most of the plasma. The energy density in the fixed-ionization simulations reaches approximately $\sim 10~{\rm Gbar}$ and is mostly homogeneous in the region $x=8-9~{\rm \mu m}$.

The high-energy electrons created with LP facilitate a better spatial homogenization of the energy density than with CP. Their slow thermalization results in a more spatially homogeneous target heating, since they can recirculate several times through the plasma. In a potential application, one should therefore make a compromise between good thermalization and homogenization. Another parameter that can be used to control homogenization is target thickness; decreasing it helps for a faster homogenization of the plasma heating. However, a thinner target will also explode faster hydrodynamically, which would give a HED application a shorter time frame to operate in.

Meanwhile, the self-consistent simulation gives an exponentially decreasing energy density profile throughout the target at t=500 fs, indicating that thermalization is taking longer. The lower temperature and electron density reached with self-consistent ionization result in approximately an order of magnitude lower energy density compared to the fixed-ionization results. However, there is still a significant region with $\mathcal{P}_{\rm e} > 1$ Gbar in the self-consistent ionization simulation at t=500 fs. In this

case, the energy density does not homogenize as efficiently, partly due to a decreased ability of the target to thermalize fast electrons (stemming from lower \bar{Z}^*), and partly due to the inhomogeneity of the ionization profile which affects the bulk electron density profile.

As a consequence of the strong gradients in \mathcal{P}_e around the target front side, a shock wave is launched. The shock wave presents itself as a sharp jump in electron pressure, most clearly seen close to $x = 8.0 \, \mu \text{m}$ in the $t = 500 \, \text{fs}$ panel in figure 4. The details of shock formation are sensitive to the laser and target parameters, and are more clearly seen from the ion phase space, as will be addressed by a paper in preparation (Sundström *et al.* 2020). However, no ion reflection occurs at the shock front, which means that the shock is hydrodynamic like in its nature.

At the high ionization levels discussed in this paper, such high temperatures and densities may result in significant energy losses due to bremsstrahlung. The total bremsstrahlung emission power density can be estimated as $S_{\rm BS}$ [W cm⁻³] $\approx 1.69 \times 10^{32} \times Z^{*3} (n_{\rm i} [{\rm cm}^{-3}])^2 (T_{\rm e} [{\rm eV}])^{1/2}$ (Huba 2016). By comparing the power density to the thermal energy density $\sim \frac{3}{2} n_{\rm e} T_{\rm e}$, we arrive at a radiative time scale of the order of several ps for keV range temperatures at a density of $n_{\rm i} = 8.4 \times 10^{22}$ cm⁻³. Hence, the radiative losses from bremsstrahlung will mostly be of concern at time scales longer than those studied in this paper. However, bremsstrahlung losses cannot be completely disregarded in a WDM/HDM experiments, where current spectroscopic temporal resolution is constrained to \sim ps time scales.

Besides bremsstrahlung, line emission from relaxation of excited states may be of concern. However, at electron energies above $\gtrsim \! 10$ eV, electron energy losses from excitations become subdominant compared to ionization (Joshipura, Vaishnav & Limbachiya 2006). Since energy loss from collisional ionization events is accounted for in the self-consistent ionization simulation, the temperature of $T_{\rm e} \approx 2.5$ keV is likely not affected much by losses through line emissions.

3.1. Illustration of the collisional absorption mechanism

To illustrate the mechanism by which collisions enhance absorption, we have performed a simplified set of simulations. These are designed to generate a quasi-steady state: the laser intensity is constant after a linear ramp-up over 10 laser cycles; the ions are stationary; the plasma is 2.5 μ m long and it terminates at a thermal boundary, meaning that particles which exit the boundary are reflected with momenta chosen randomly from a Maxwellian distribution at $T_{\rm e,0}=10$ eV for the electrons – the same at the initial temperatures. The other simulation parameters are: CP at $a_0=10$, $Z^*=27$ with and without collisions; resolution and other numerical parameters are as stated in § 2. The long ramp-up time has been chosen to reduce electron energization due to the laser amplitude envelope modulation (Siminos *et al.* 2012). Note that due to the steady state nature of this simplified simulation setting, it is hard to draw any quantitative conclusions that can be transferred to the time-varying situation.

We will now take a look at the interaction between the electrons and the laser electric field. The density of power S exerted on an electron population can be expressed as

$$S(x,t) = -e \int d^3 v \mathbf{E}_{\perp} \cdot \mathbf{v} f_e(\mathbf{v}) = -e n_e \mathbf{E}_{\perp} \cdot \mathbf{V}_{\perp}, \qquad (3.1)$$

where $E_{\perp} = E_{\perp}(x, t)$ is the laser electric field – which only lies in the transverse plane – and $V_{\perp} = V_{\perp}(x, t) \equiv [1/n_{\rm e}(x, t)] \int {\rm d}^3 v \, v_{\perp} f_{\rm e}(x, v; t)$ is the projection of electron velocity moment onto the transverse plane.

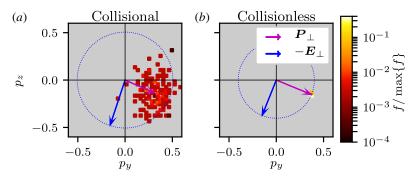


FIGURE 5. Electron transverse momentum distribution at x = 7.0-7.2 nm behind the ion front in the simplified simulations with fixed ions. The left and right panels correspond to collisional and collisionless simulations, respectively. The distributions are here recorded at t = 190 fs, which is well after the quasi-steady state has been reached, where E_{\perp} and P_{\perp} rotates (clockwise) in the transverse plane.

In a 1-D model, disregarding collisional effects, the transverse canonical momentum $\tilde{P}_{\perp} = P_{\perp} - eA_{\perp}$ is conserved, and $\tilde{P}_{\perp} = 0$. Hence $P_{\perp} = eA_{\perp}$, where A_{\perp} and P_{\perp} are the transverse component of the magnetic vector potential and the electron momentum moment, defined analogously to V_{\perp} . In quasi-steady state, A_{\perp} is just rotating in the transverse plane, so the electric field is $E_{\perp} \equiv -\partial A_{\perp}/\partial t = \omega A_{\perp} [\cos(\omega t)\hat{y} - \sin(\omega t)\hat{z}]/\sqrt{2}$, where $A_{\perp} = A_{\perp}(x)$ is the magnitude of the vector potential (necessarily transverse in one dimension). Importantly, the electric field vector is perpendicular to the vector potential and the magnitude of the electric field is $E_{\perp} = \omega A_{\perp}$. We therefore expect P_{\perp} and E_{\perp} to be perpendicular and their magnitudes – in normalized units – to be equal, $P_{\perp} = E_{\perp}$.

Figure 5 shows slices of the collisional (a) and collisionless (b) electron distributions in the transverse momentum plane at t=190 fs, well after quasi-steady state has been reached, and in a thin slice 7.0–7.2 nm (one cell length) behind the immobile ion front edge of the plasma. If we were to evolve this picture in time, we would see the (negative) electric field $-\mathbf{E}_{\perp}$ rotate clockwise, along the marked-out circular path in figure 5; the mean momentum \mathbf{P}_{\perp} would follow synchronously in this rotation. The most apparent difference between the collisional and collisionless distributions is the much larger momentum spread of the former, caused by collisional scattering of the electrons. In contrast to the collisionless case, $-\mathbf{E}_{\perp}$ and \mathbf{P}_{\perp} are not equal in magnitude nor are they perfectly perpendicular. The missing transverse canonical momentum has been collisionally transferred to the ions, where it disappears from the simulation due to the ions being static. Note that if $-\mathbf{E}_{\perp}$ and \mathbf{P}_{\perp} are not perfectly perpendicular in (3.1), then the absorbed power density S is non-vanishing.³ We can express (3.1) as

$$S = -neE_{\perp} \cdot V_{\perp} = neE_{\perp} V_{\perp} \sin(\alpha), \tag{3.2}$$

where the phase angle between $-E_{\perp}$ and V_{\perp} is $\pi/2 - \alpha$.

Figure 6 displays configuration space profiles of E_{\perp} , V_{\perp} and S – in dimensionless units – as well as $\sin(\alpha)$; the curves are produced from a time average over 21 time frames spanning 20 fs. In the collisionless case, we have $P_{\perp}(x) = E_{\perp}(x)$ and the phase

³For simplicity, we are ignoring relativistic effects in this discussion, which would otherwise complicate the relationship between P_{\perp} and V_{\perp} .

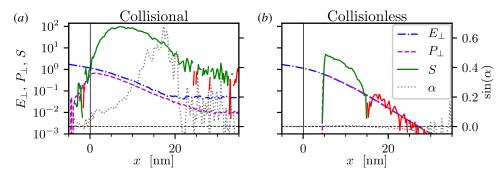


FIGURE 6. Magnitudes of the perpendicular electric field E_{\perp} (blue dash-dotted) and mean electron transverse momentum P_{\perp} (magenta dashed) as well as the absorbed power density S (solid line, green and red for S>0 and S<0 respectively). Also shown is the phase shift $\sin(\alpha)$ (grey dotted) between $-E_{\perp}$ and V_{\perp} , where V_{\perp} is the mean electron transverse velocity moment of the distribution. The vertical black line marks the location of the transverse momentum planes plotted in figure 5. All values are expressed in dimensionless units.

shift angle $\alpha \simeq 0$ throughout the first $\simeq 25$ nm $\approx 8l_s$. Due to a finite spread in the electron transverse velocities, there will be a continuous exchange of electrons in the longitudinal direction not accounted for in the fluid description above, which induces a small deviation from $\alpha=0$ and hence $S\neq 0$. However, S changes sign at S0 nm, beyond which the absorbed power is negative. In the collisional case, S1 is consistently smaller than S1 nm, Furthermore, the phase shift S1 is much larger, which is reflected in the about two orders of magnitude larger absorbed power S2 than in the collisionless case.

A final note on the collisional case in figure 6(a) is the numerical artefact that causes both $E_{\perp}(x)$ and $P_{\perp}(x)$ to level off near x=20 nm. As the Monte Carlo collisional algorithm used in Smilei (Pérez *et al.* 2012) only conserves momentum statistically, a P_{\perp} noise floor is generated which drives noise in E_{\perp} , i.e. the base level in figure 6(a). This effect could be alleviated by increasing the number of macro-particles. However, the absolute majority of the collisionally induced laser-energy absorption occurs in the region x=5-15 nm and is therefore not significantly affected by the collisional noise floor.

3.2. Parameter scans

We have also performed parameter scans in order to investigate the dependencies of the collisional heating mechanism. One such scan has been in ionization, with either fixed ionization ($Z^* = 11$, 19, 24 and 27) or self-consistent field and impact ionization. We have also conducted scans in laser intensity with a_0 ranging from 1 to 14, and pulse durations from $t_{\rm FWHM} = 15$ fs to 400 fs. The remaining parameters are as in § 2.

Let us first consider the absorbed energy from the laser. Figure 7(a) shows the kinetic energy gain by the electrons and ions (ΔU) following the laser irradiation, for a scan in a_0 (colour coded) and a scan in pulse duration (shape coded). The value displayed on the horizontal axis is the laser pulse energy $J = It_{\rm FWHM} [\pi/\log(4)]^{1/2} \propto a_0^2 t_{\rm FWHM}$, where I is the laser intensity.

In the case of a constant pulse duration, $t_{\rm FWHM} = 60$ fs (triangles), the trend scales like a power law with $\Delta U \propto J^{0.74}$ (dashed line) or $\Delta U \propto a_0^{1.48}$, since $J \propto a_0^2$. In

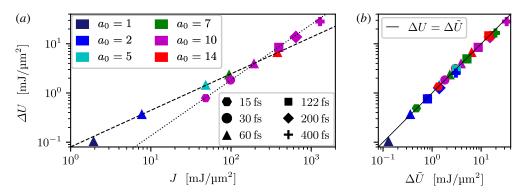


FIGURE 7. Total simulated kinetic energy gain ΔU against (a) the laser energy J and (b) the power law scaling $\Delta \tilde{U}$ for different combinations of laser parameter a_0 and duration $t_{\rm FWHM}$. Lines in panel (a) indicate power law scalings: $\Delta U \propto J^{0.74} \propto a_0^{1.48}$ at constant $t_{\rm FWHM} = 60$ fs (dashed) and $\Delta U \propto J^{1.13} \propto t_{\rm FWHM}^{1.13}$ at constant $a_0 = 10$ (dotted). These two power law scalings combine to give $\Delta \tilde{U} = 0.23$ mJ μ m⁻² $\times a_0^{1.48} \times (t_{\rm FWHM}/100 \text{ fs})^{1.13}$, which is shown in panel (b) to agree well with the full data set, also including parameter combinations not shown in (a).

other words, the absorption efficiency scales as $\Delta U/J \propto J^{-0.26} \propto a_0^{-0.52}$. The $a_0=1$ point seems to deviate from the above scaling, suggesting that it is mainly valid at relativistic intensities. Thus, $a_0=1$ was excluded from the fit. This scaling is similar to the $I^{-1/4}$ scaling of the normal skin effect, as described by Rozmus & Tikhonchuk (1990). Comparing the collisional mean free path, $\lambda_{\rm mfp} \sim 20$ nm, to the skin depth, $l_{\rm s} \approx 6$ nm,⁴ it is not completely clear that the normal skin effect ($\lambda_{\rm mfp} \lesssim l_{\rm s}$) can be ruled out. However, our simulation results do not support some other scaling laws derived by Rozmus & Tikhonchuk (1990) for normal skin effect, possibly due to the non-relativistic and simplified nature (constant intensity and semi-infinite plasma) of their analytic treatment.

The other scaling trend displayed in figure 7(a) is at constant $a_0 = 10$ (magenta). Here, the power law fit (dotted line) gives $\Delta U \propto J^{1.13} \propto t_{\rm FWHM}^{1.13}$. In this case the absorption efficiency still has a weak positive scaling of $\Delta U/J \propto t_{\rm FWHM}^{0.13}$. From the pulse duration scaling, we note that the 200 fs and 400 fs pulses begin to fall off below the scaling followed by the other data points, and they were thus also excluded from the fit. The decreasing trend with $t_{\rm FWHM}$ in the long-pulse limit may be a consequence of the recirculation of hot electrons, which could lower the effective plasma collisionality in the irradiated region.

Combining the two above scalings yields the approximate scaling

$$\Delta \tilde{U} = 0.23 \text{ mJ } \mu \text{m}^{-2} \times a_0^{1.48} \left(\frac{t_{\text{FWHM}}}{100 \text{ fs}}\right)^{1.13}$$
 (3.3)

in the (a_0, t_{FWHM}) plane. Figure 7(b) shows this scaling to agree well with the full set of data, including data points where both a_0 and t_{FWHM} are varied. The observed scaling does break down at the low- a_0 or long-duration limits. However, the range of validity stretches over two orders of magnitude in pulse energy and a similar range in absorbed energy, and the laser parameters captured by this power law are

⁴The skin depth adjusted for collisions has been inferred from figure 6(a).

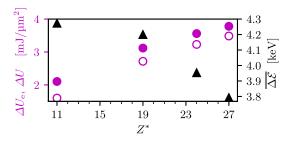


FIGURE 8. Collisional simulation with various (fixed) ionization degree Z^* . Left axis (magenta circles): gained kinetic energy by electrons and ions ΔU (filled circles) as well as only by electrons $\Delta U_{\rm e}$ (open circles). Right axis (black triangles): average kinetic energy gained by one electron $\overline{\Delta \mathcal{E}}$.

experimentally feasible and relevant to isochoric heating experiments. Collisional effects in general decrease at higher particle energies and the absorption happens through collisional scattering of the laser-driven electrons in the skin layer. Therefore, if the laser field (a_0) is increased, so that the electrons in the skin layer reach higher energies, then the efficiency of the collisional absorption should decrease.

We also report on a scan in (fixed) ionization degree Z^* . Although this parameter cannot be controlled independently in experiments, this scan aims to provide insight into the target collisionality, which scales as $(Z^*)^2$ – ignoring other effects, e.g. individual particle energy. However, by varying Z^* while keeping the ion density $n_{Cu,0}$ fixed, we inevitably also change the electron density $n_{e,0} = Z^* n_{Cu,0}$, which may introduce other density-related effects. Nevertheless, the electron density stays highly overcritical – the lowest electron density in this scan is $n_{e,0} = 532.4n_c$ for $Z^* = 11$. Figure 8 displays the energy absorbed by both electrons and ions ΔU (filled circles, left axis) and only by electrons ΔU_e (open circles, left axis) for the different ionization degrees. The absorbed energy increases with Z^* , while a smaller fraction of the absorbed energy goes into the ions at higher Z^* .

Due to the accompanying changes in electron density, the average absorbed energy per electron $\overline{\Delta \mathcal{E}}$, also shown in figure 8 (black triangles, right axis), happens to decrease by approximately 15% from $Z^*=11$ to 27. The decrease in $\overline{\Delta \mathcal{E}}$ with Z^* may seem surprising if the dominant heating mechanism is collisional. However, this might be due to other density effects, such as the increased skin depth at lower electron density which allows a deeper laser penetration and thus a stronger laser-to-electron coupling efficiency. When we examine the electron energy spectra (not shown) in this scan, the Maxwellian-fitted bulk electron temperatures are all $T_e=3.5\pm0.1$ keV.

Besides just the pure amount of energy absorbed from the laser, we are also interested in how well thermalized the plasma is. As a measure of that, figure 9 shows the fraction of electron kinetic energy in the non-thermal electrons. This is calculated by the fraction of the energy in the high-energy tail to the total electron kinetic energy, $\int_{2T_c}^{\infty} [f_{\mathcal{E}}(\mathcal{E}) - f_{\mathcal{E}}^{\text{MJ}}(\mathcal{E})] \mathcal{E} \, d\mathcal{E}$, where $f_{\mathcal{E}}^{\text{MJ}}$ is a Maxwell–Jüttner distribution fitted to the bulk of the electron energy spectrum $f_{\mathcal{E}}(\mathcal{E})$ (as shown in figure 1) and T_e is the temperature inferred from the fit. Figure 9 shows scans in fixed ionization degree Z^* (a), laser amplitude a_0 (b) and pulse duration t_{FWHM} (c). The displayed values are taken 200 fs after the end of the laser pulse. Due to a varying heat transport speed, the fraction of non-thermal electron energy is only taken in the

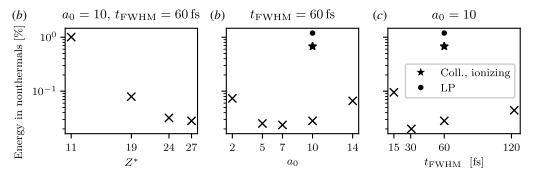


FIGURE 9. Fraction of the electron energy in non-thermal electrons, for collisional simulations, 200 fs after the end of the laser pulse, for scans in Z^* (a), a_0 (b) and t_{FWHM} (c) with CP, marked by crosses. The value marked with a dot is from LP, and self-consistent ionization is shown as a star.

region in which the bulk electron temperature is no longer increasing. This should still give a representative estimate of the non-thermal fraction, since the fast electrons have already recirculated by the chosen time, see the t=250 fs panels of figure 3. The precise values in figure 9 are sensitive to the choice of time and region to include, thus these results are only qualitative. Nevertheless, the general trends shown here are still representative of the observed situation – importantly, the relation between LP and CP is robust.

Figure 9(a) shows that there is a trend toward lower non-thermal fraction at higher ionization levels, which is consistent with the faster thermalization expected at high Z^* . This trend also suggests that the higher absorbed energy per electron at lower Z^* (figure 8) is linked to a relative increase in the non-thermal population.

Regarding the scans in the laser amplitude and duration in figures 9(b) and 9(c), respectively, no clear trend appears to emerge among the CP laser pulses (marked by \times). Then there are the self-consistent ionization (star) and LP (downward triangle) simulations: both have approximately one order of magnitude higher fraction of energy in non-thermal electrons than the equivalent (fixed-ionization, CP) counterpart. The higher fraction of non-thermal energy with LP stems from the $j \times B$ and vacuum heating mechanism. The higher non-thermal energy fraction with a self-consistent ionization process is discussed in conjunction with its phase-space distribution in figure 3.

Even a very small fraction of non-thermals may affect the interpretation of X-ray diagnostics (Rosmej 1997; Chen *et al.* 2009; Renner & Rosmej 2019), meaning that LP can be more intrusive than CP in WDM/HDM studies. We have also conducted simulations with a larger pre-expanded plasma (exponentially decaying density profile with a scale length of 80 nm). While not presented here, those simulations show that LP can result in up to $\sim 10\,\%$ of the electron kinetic energy in non-thermal electrons, which would of course be even more intrusive and significantly affect the X-ray diagnostics. With CP, the pre-plasma weakens the energy absorption by about a factor of two, but the fraction of energy in fast electrons stays $\lesssim 1\,\%$.

3.3. Two-dimensional simulation results

Up to this point, all the results presented have been produced in 1-D simulations. However, to investigate the applicability of these results in higher dimensions, where

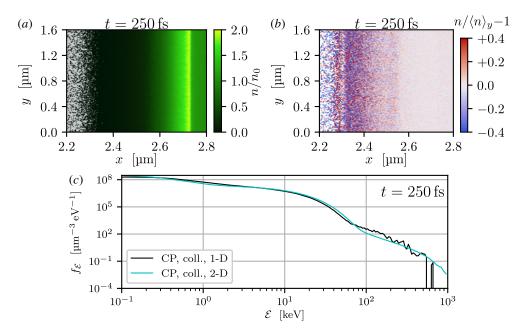


FIGURE 10. Map of the electron density (a) in units of background density n_0 and the relative transverse variation of the electron density (b) near the front target surface at 250 fs. Panel (c) shows a comparison of the electron energy spectra from the 1-D and 2-D simulations at the same time. The spectra are taken from the full simulation box. The transverse band of higher density at $x \approx 2.72~\mu m$ in the panel (a) corresponds to a shock front launched by the laser impact.

transverse plasma modulations can arise at the target boundary and affect the bulk heating and hot-electron generation (Kemp & Divol 2016), we have performed a 2-D simulation at our baseline laser parameters (see § 2 for details). A density map of the electron density near the illuminated target surface is shown in figure 10(a). The density is shown at t = 250 fs. There is a transverse band of higher density at $x \approx 2.72$ µm that represents a shock front propagating into the plasma. Notably, this shock front remains straight, with no evidence of substantial density modulations.

To more clearly demonstrate the absence of transverse instability effects, the relative transverse variation of the electron density $n/\langle n\rangle_y-1$ is displayed in figure 10(b), where $\langle \cdot \rangle_y$ denotes a transverse average over the full width of the simulation box in y. Any transverse density structure should therefore be clearly visible in this representation. The substantial deviations from the average density observed in the low-density $(n_e \lesssim 0.2n_0)$ region correspond to statistical noise due to a low number of computational particles in said region. Importantly, the deviations seen in figure 10(b) have no structure to them, and the same applies for the shock, suggesting that transverse effects are inoperative in the present highly collisional case (at least within the simulated time window).

Lastly, to confirm that the collisional heating behaves similarly in the 2-D and 1-D simulations, figure 10(c) shows the electron kinetic energy spectra of the corresponding 1-D and 2-D simulations at t = 250 fs.⁵ The spectra of the 2-D

⁵Due to the 2-D simulation having a smaller longitudinal box size, and thus the target front being located at $x = 2.5 \mu m$ instead of $x = 7.5 \mu m$, the times of comparison for the 1-D simulation are shifted by 15 fs

and 1-D simulations are essentially the same. Although not shown here, the two spectra agree similarly well also at earlier times. Since the following thermalization process is almost entirely collisional, and thus independent of dimensionality, it is safe to conclude that the energy absorption is not affected by going from one to two dimensions, under the interaction conditions considered.

4. Conclusions

We have performed collisional and collisionless 1-D and 2-D PIC simulations and shown that a collisional, inverse bremsstrahlung, absorption can be used for strong plasma heating in a solid-density, high-Z* material, such as copper, with ultrahigh intensity, short-pulse lasers. Using CP, the electron population quickly thermalizes to well-formed Maxwellian distributions suitable for experimental verification of HED physics models. The collisional simulations show that the target electrons are quickly heated to $T_c \sim 3.5$ keV bulk temperature on a ~ 300 fs time scale. The target energy density reaches ~10 Gbar, which is within the realm of ultrahigh energy density. The use of CP provides faster collisional thermalization of the electron population compared to LP, something which is valuable for experimental tests of HED atomic physics models. A test of the collisional absorption using a 2-D simulation, demonstrates the transferability of the 1-D results to higher dimensions. In contrast to previous work conducted with lower-Z* targets (Kemp & Divol 2016), the high collisionality is not favourable for driving transverse plasma modulations, resulting in the same absorption levels in two dimensions as in one dimension, at least for CP.

We have carried out scans over laser parameters and ionization. The scans over laser settings show that the mechanism is robust to changes in the laser, over two orders of magnitude in laser energy, with lower intensity and longer pulses at the same laser pulse energy yielding better energy absorption. Also, the higher collisionality incurred from a higher ionization level improves energy absorption and electron thermalization. A more realistic simulation run with self-consistent ionization, including both impact and field ionization, reached $T_{\rm e} \sim 2.5$ keV, confirming that collisional heating is still operational in a self-consistently ionized plasma, although its thermalization is then less complete than at fixed ionization, due to high-energy electrons generated through ionization events in strong-field regions.

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later compared to the 2-D simulation, due to increased travel time for the laser pulse. The simulation time at which the 1-D spectrum is plotted is thus t = 265 fs.

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