Parametric study of cycle modulation in laser driven ion beams and acceleration field retrieval at femtosecond timescale

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High-frequency modulations appearing in the kinetic energy distribution of laser-accelerated ions are proposed for retrieving the acceleration field dynamics at the femtosecond timescale. Such an approach becomes possible if the laser-cycling field modulates the particle density in the ion spectra and produces quasitime stamps for analysis. We investigate target and laser parameters determining this effect and discuss the dependencies of the observed modulation. Our findings refine a basic mechanism, the target normal sheath acceleration, where an intense and ultrafast laser pulse produces a very strong electrical field at a plasma-vacuum interface. The field decays rapidly due to energy dissipation and forms a characteristic spectrum of fast ions streaming away from the interface. We show that the derived decay function of the field is in accordance with model predictions of the accelerating field structure. Our findings are supported by two-dimensional particle-in-cell simulations. The knowledge of the femtosecond field dynamics helps to rerate optimization strategies for laser ion acceleration.

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I. INTRODUCTION

Possible applications of laser-accelerated ion beams (cf., e.g., [1-5], and references therein) need tailored energy distribution functions (EDFs) of the ions. This EDF is coupled to the dynamics of the field which accelerates the ions. The fundamental mechanisms of laser-driven ion acceleration have been studied for nearly two decades. Special attention was drawn to the target normal sheath acceleration (TNSA) [6] (and references therein) and the radiation pressure acceleration (RPA) [7,8]. TNSA is, to some extent, a robust process, while RPA yields the highest energies of accelerated ions. The underlying dynamics of the acceleration field has been successfully probed at a timescale of picoseconds [9] down to about 100 fs [10,11]. This knowledge is important for benchmarking theoretical models which allow the extrapolation of required laser parameters for desired ion beam parameters.

In this work, we suggest to use a modulation phenomenon in laser-driven proton beams for tracing the acceleration process at a femtosecond timescale. This allows comparing experimental EDF results with model functions of the acceleration field dynamics. The knowledge of these

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fast-evolving processes is important to understand the limitations of optimization efforts in laser ion acceleration. For our analysis, we use and study in more detail faint density modulations in the kinetic energy spectrum of TNSA protons [12] which become visible if the laser intensity contrast between the pulse background and the peak is at a level of about 10^{10} . At such conditions, even the cycling action of the laser field within the temporal envelope of the laser pulse imprints onto the acceleration field and the velocity distribution of the accelerated ions. Our interpretation is based on numerical and analytical model calculations [12], which showed that the laser-cycledriven variation of the oscillatory part of the Lorenz force at twice the laser frequency releases hot-electron bunches such that the sheath field in the TNSA process is modulated, which translates to density variation in the accelerated ion bunch. The lightest ion, the proton, is most sensitive to such a kind of field variation. In combination with proton beam modulation, we have also observed at some conditions the same modulation effect in the carbon spectra, which is not further discussed here. An exclusive appearance of beam density modulation in low-energy, low-charge-state carbon ions has been discussed very recently [13]. This finding as well as other ion beam modulations [14,15] has been observed with different characteristics at specific interaction parameters.

First, we will discuss how the observed density modulation in the proton spectrum depends on laser parameters (polarization, temporal contrast) and on target parameters (thickness of the target foil, surface modification via structuring). Second, we use our interpretation of the modulation effect to retrieve the acceleration field strength at the femtosecond timescale. This result we compare to model predictions and 2D particle-in-cell (PIC) simulations describing the decline of the acceleration field. We conclude that the process of energy dissipation and the corresponding decline of the acceleration field strength are related to the gain of kinetic energy during the ion acceleration in the field.

II. EXPERIMENT

Ion acceleration experiments were performed with the 70 TW Ti:sapphire laser arm in the former High Field Laboratory at Max Born Institute. The central wavelength of the laser was about 800 nm, the pulse duration was ~35 fs, and the maximum laser energy on the target reached up to ~2 J. Focusing the laser pulse with an f/2.5 off-axis parabolic (OAP) mirror to spots with a FWHM of $\sim 4 \mu m$ realized peak intensities of up to $\sim 10^{20}$ W/cm². The incident angle of the laser beam is between 0.1 and 1 degree in order to avoid exact backreflection. In comparison with our first work [12], the second arm of the former two-beam High Field Laser system (other preamplifier architecture—multipass instead of regenerative) was used, allowing higher laser intensities and the possibility of switching to ultrahigh temporal contrast with a plasma mirror. Furthermore, a different Thomson-spectrometer configuration with an electric field to separate different ion species was employed. The temporal intensity contrast of the laser pulse was realized

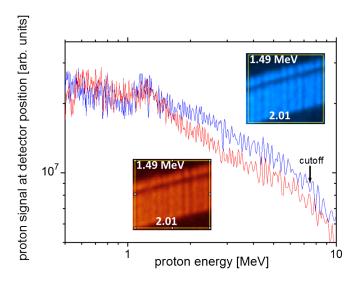


FIG. 1. Fast protons produced with laser-irradiated 5 μ m Ti foils using linear (blue line) and circular (red line) laser polarization (HC, parameters; cf. text). The proton signals are pixel counts at the pixel position (at the calculated energy position) of the detection system. Inset: Contrast-enhanced parts of the recorded proton traces.

by a cross-polarized wave generation (XPW) front end yielding a peak-to-amplified spontaneous emission (ASE) contrast of better than 10^{10} (high contrast, HC). For driving ultrathin targets, the contrast was further be enhanced by a double plasma mirror (DPM) and reaches values better than 10^{14} (ultrahigh contrast, UHC). Targets are 5- μ m-thick titanium foils or 30-nm-thick plastic (poly-vinyl-formal PVF, $C_5H_7O_2$, density 1.23 g/cm³) foils [16]. Kinetic energies of accelerated ions are measured with a Thomson mass spectrometer using a slit [(50–340) μ m width] as an entrance aperture. Recording the momentum dispersed ion traces with such an entrance slit allows discriminating faint density modulations against noise. This is a mandatory prerequisite.

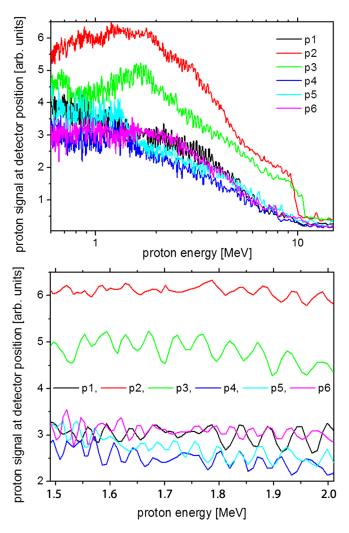


FIG. 2. Fast protons produced with 7×10^{19} W/cm² laser-irradiated 5 μ m Ti foils (the lower graphic shows parts of the traces with an enlarged energy interval similar to insets in Fig. 1): linear polarization applied for p1 (black), p5 (cyan), and p6 (purple) and circular polarization applied for p2 (red), p3 (green), and p4 (blue); spectrometer slit widths were 340 μ m for p1 and p2, 240 μ m for p3, and 50 μ m for p4-p6.

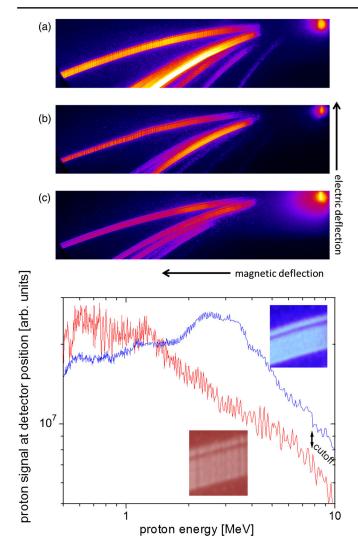


FIG. 3. Upper part: Recorded detector images of the ion spectra (parabolic traces) in the same false color scaling: (a) 5 μ m Ti foil, HC, circular polarization, spectrometer slit 240 μ m, (b) 5 μ m Ti foil, HC, linear polarization, spectrometer slit 50 μ m, and (c) 30 nm plastic foil (UHC, linear polarization, parameters cf. the text), spectrometer slit 50 μ m; lower part: scans of proton traces from (a) and (c): 5 μ m Ti foil (HC, circular polarization, red line; cf. Fig. 1), 30 nm plastic foil (UHC, linear polarization, parameters cf. the text) (blue line); inset: magnified part of signal traces. The proton signals are pixel counts at the pixel position (at the calculated energy position) of the detection system.

The entrance slit for the Thomson mass spectrometer is oriented parallel to the magnetic and the electric fields (both fields have the same orientation). Therefore, ions with the same kinetic energy and the same charge-to-mass ratio appear at the detector plane as a line being perpendicular to the magnetic deflection direction and in line with the electric field direction. The bending of the traces (photo of the parabolic traces in Fig. 3) is caused by the electric field. The width of the traces is determined by the length of the slit. Therefore, our observation is a proof that the

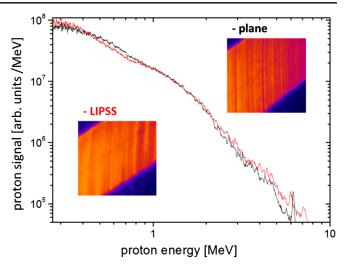


FIG. 4. Appearance of density modulation in fast proton bunches produced with HC laser pulses irradiating plane or LIPSS Ti foils (parameter cf. the text). The noncalibrated proton signal is evaluated with energy binning. Inset: Contrast-enhanced parts of the recorded proton traces.

density maxima of the signal (lines) and their orientation correspond to ions (protons) with the same energy.

An imaging Hamamatsu MCP, 100 mm in diameter, detected the ions, and the visible Thomson parabolas on the fluorescence screen were recorded with a CCD camera system. The detector response was cross-checked with alpha particles from a radioactive Am²⁴¹ source. In most experiments, *p*-polarized laser pulses irradiated the target foils close to normal incidence.

For some experiments, in order to generate circularly polarized pulses, a 200- μ m-thick mica $\lambda/4$ plate was inserted in front of the second-to-last turning mirror delivering the beam to the focusing OAP mirror. The polarization setting with the $\lambda/4$ wave plate was aligned with a Glan-Thompson prism. In order to visualize the density modulations, Figs. 1 and 2 display proton raw signals of a detector pixel without considering the binned energy interval of the pixel, and in Fig. 4 the proton number is calculated per binned energy interval for comparison.

III. PARAMETRIC STUDY OF MODULATION IN THE ENERGY DISTRIBUTION OF LASER-ACCELERATED PROTONS

In past proton imaging experiments [17], using a slit spectrometer, the temporal pulse contrast was at a level of 10^6 – 10^7 , and this did not allow observing the modulation effect. For the first time [12], we could detect high-frequency modulation of the proton EDF when an XPW front end was set into operation and, consequently, the temporal intensity contrast was enhanced by nearly 3 orders of magnitude (HC).

Within the experiments described in Ref. [12], we inserted thin Al foils into the ion beam path in front of

the spectrometer slit and repeated the experiment without any other changes. The scattering in the filter foil changed the longitudinal emittance of the proton beam, and the weak modulation feature disappeared. This is a strong proof to exclude any instrumental artifact as, e.g., structures of the imaging detector or high-frequency modulation of the deflecting fields inside the spectrometer. Within the course of this work, we describe parametric target and laser dependencies of the modulation effect.

In order to substantiate the modulation in laser-driven ion acceleration further, it is useful to identify additional conditions which suppress the effect. Changing the laser polarization from linear to circular for a normally incident light wave at a plane surface is one obvious strategy, because the electron trajectories in the light field are different and the forces accelerating the electrons are different, too. (In the case of circular polarization, the vector product between the electron velocity and the B field of the electromagnetic wave vanishes; at linear polarization, it does not.) This in turn changes the oscillatory motion of the electrons and in specific also the hot-electron population which is responsible for the TNSA mechanism. But, as is visible in Fig. 1, the experiment with a 5- μ m-thick titanium foil at HC condition does not show any significant difference between linear and circular polarization.

In order to substantiate this observation, we picked six successive laser shots (p1-p6) from one experimental run with the same laser energy irradiation and focusing (laser intensity at FWHM $\sim 7 \times 10^{19}$ W/cm², high contrast) and target conditions (5- μ m-thick titanium foil) in Fig. 2. We applied different spectrometer slit widths and a laser polarization as described in the caption of Fig. 2.

In all shots, the proton cutoff energy reaches 10 MeV. Although our available data are not backed with a statistics built on hundreds of shots, the presented successive ones in Fig. 2 demonstrate that the observed modulation phenomenon is reproducible if laser and target parameters allow the phenomenon to occur. Shot-to-shot variation of the signal strength and spectral shape are apparent. This becomes visible if the signals are recorded with different slit widths. While smaller slits lead to lower signals, the signal reduction did not follow exactly the reduced slit width. The used imaging system for the phosphorous screen had a resolution of 0.108 mm per CCD-camera pixel. The MCP has a channel diameter and channel pitch of about 25 and 31 μ m, respectively. The width of the density maxima is about 2–3 CCD pixels with the 50 μ m slit and 3–4 CCD pixels in the case of the 240 μ m slit. The slit itself was at a distance of 40 cm from the target, and the MCP detector was 100 cm located from the slit. The center of the magnet and the electric field plates was about 35 cm behind the slit.

For the smallest slit width, the signal fades already around the cutoff. The modulation phenomenon is clearly reproduced in all shots and does not show a qualitative difference between linear (LP) and circular (CP)

polarization. A similar number of peaks (11–14) appear in the selected energy interval. The peak position fluctuates as expected for a non-phase-locked multioptical-cycle femtosecond laser pulse. In different experiments with ultrathin target foils, we applied also this switching technique between LP and *CP* pulses. We investigated high harmonic generation in the laser transmission direction and observed emission only in the case of LP pulses. This clearly shows that our technique of *CP* pulse generation does not leave a strong LP background.

Therefore, we exclude a systematic experimental error. Instead, we will argue in the following that at the given laser intensities the plane of critical plasma density is bent. The magnitude of the bending angle Θ_{sf} can be approximated with the analytical expression $\langle \tan(\Theta_{sf}) \rangle \approx$ $c(I^{18}_{L})^{0.5}/(\omega_{pe}w_{L})$, where c is the speed of light, I^{18}_{L} the laser intensity as a multiple of 10^{18} W/cm², ω_{pe} the plasma frequency of electrons, and w_L the laser beam waist. The equation is derived for a Gaussian laser beam profile assuming balance of the ponderomotive pressure by the created ambipolar field at the position of the critical electron density [18]. With our parameter set, we obtain on the order of $\Theta_{sf} \sim 10^{\circ}$, which alters already the interaction significantly. Because of the bent critical surface, quite different electron trajectories and acting forces are possible such that the simple assumption of parallel electron velocity and B-field vectors does not hold anymore. In comparison with our experimental results, also in theoretical studies modulations are visible with CP pulses (cf. PIC simulation visualized in Figs. 2–4 of a very recent paper [19]). From our experimental results alone, we cannot conclude on the modulation period. Our analysis in Ref. [12] suggests a period of half a laser cycle due to the oscillatory part of the Lorenz force in the case of LP pulses. The PIC simulations for CP pulses in Ref. [19] suggest a modulation close to a laser cycle in a certain parameter range. We did not observe a clear difference between LP and CP pulses in our experiments. We have to keep this question open.

These findings point to other effects and also suggest that more parametric experimental studies and respective highresolution simulations are necessary to resolve the detailed plasma dynamics at the temporal scale of the driving laser cycle.

The density of the electron layer at the target surface and the creation of the hot-electron population is a function of the temporal contrast. Therefore, it is interesting to study how the modulation effect changes in respect to the temporal contrast of the laser pulse. Applying UHC laser pulses to 5- μ m-thick Ti foils leads to visible though much weaker modulations. If now an ultrathin (here a 30 nm plastic foil) target is exposed with UHC pulses, the modulation effect vanishes. This comparison is illustrated in Fig. 3. The visibility or occurrence of modulations is not an effect of the signal strength, as it is visible in the

equal-measure color-coded photos and profile scans in Fig. 3.

In the case of ultrathin foils, which have low mass, the whole target starts to move during the irradiation, and, probably more importantly, the illuminated target area is bent, which changes the sheath field geometry at the target rear. This is different in the case of thick targets where bending (or hole boring) influences the illuminated target front side only. Bending of the whole target foil might hinder small variations of the sheath field to become effective in an ion signal that is always obtained by integration over a finite emission area and volume.

In addition, we studied how a target surface structure can influence fast ion generation within the TNSA regime. Laser-induced periodic surface structures (LIPSS) were created with the strongly attenuated driving laser in situ, and a different parametric study was performed [20]. Therefore, always the structured surface is exposed with the high-intensity laser pulse. Here, the laser-irradiated surface is modified to enhance the absorption of the highintensity laser pulse which drives the ion beam from the plane rear surface of the target. This was clearly observed at UHC conditions [20]. In the case of HC pulses, a plasma gradient appears such that the absorption is enhanced too, and the difference to the structured surface is less apparent in maximum proton energies but is still clearly visible in x-ray emission [20]. Two characteristic proton spectra obtained with HC pulses are depicted in Fig. 4. An in-depth discussion about the underlying hot-electron dynamics providing an explanation for this finding is worked out [21]. The use of the slitspectrograph configuration revealed that the surface structure introduces observable smearing and distortions to the regular modulation structure of the proton beam (cf. inserted pictures in Fig. 4). The insets in Fig. 4 show that a slight regular bending of the line features in both traces appeared. In this case, the used slit was longer, and the spatial dependence of the magnetic field inside the U-type magnet becomes visible. The change of the modulation appearance when structured targets were used is another proof of the effect itself. All this demonstrates that the observed phenomenon is not an artifact but shows dependencies on target and laser parameters.

This is quite understandable, because the spatial scale of the created surface structure is on the range of several hundred nanometers, which translates to an illumination variation at the order of one femtosecond. This is close to the field modulation cycle of about 1.35 fs (half of the laser field period). Figure 4 shows the comparison between proton spectra obtained with a structured and a nonstructured target surface of 5-µm-thick Ti foils. Here, the proton numbers are evaluated with respect to the detected energy intervals. The signal shows a variation over 3 orders of magnitude, and thus the density modulations, which are at a few percent level of the total signal only, are hardly visible in the graphs without zooming in. Comparative studies between structured and nonstructured targets were performed [20]. Also these results gave motivation to investigate the time dependence of the hot-electron distribution further, which determines ion acceleration and x-ray emission.

IV. ACCOMPANYING PARTICLE-IN-CELL SIMULATION

PIC simulations were performed at a laser intensity of 5×10^{19} W/cm² using the two-dimensional modified PSC code [22]. The simulation box $(15000 \times 5000 \text{ cells})$ is $30 \times 25 \ \mu\text{m}^2$ large with step sizes of 2 nm in the longitudinal and 5 nm in the transversal direction, and 30 particles per cell are included. The time step is 2 nm/c. The interaction is collisionless, and the simulation uses periodic boundary conditions. A high-contrast laser pulse [35 fs, 4 μ m spot size, super-Gaussian (Gaussian) in space (time)] interacts with a Ti target (density 6×10^{22} cm⁻³). An average ionization state $\langle Z \rangle = 10$ is evaluated from the commonly called Ammossov-Delone-Krainov (ADK) model formula [23,24] well grounded on theoretical work [25–27]—the PPT model. The target thickness is restricted to 1 μ m due to numerical reasons. The simulation yielded a maximum sheath field of $0.55E_L$ (E_L , electrical field strength of the laser pulse). This corresponds to $20 \times$ $10^{12} \text{ V/m using } I_L = 5 \times 10^{19} \text{ W/cm}^2 \text{ and } I_L = \varepsilon_0 c E_L^2.$

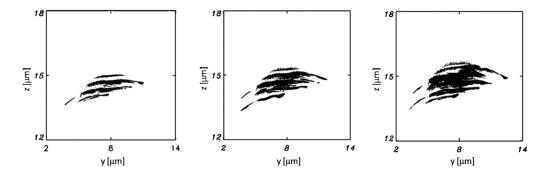


FIG. 5. 2D-PIC simulation of electron density evolution at time instants 21.3, 22.65, and 23.9 fs (from left to right), initial target position at $z = 13.5 \ \mu m$, and laser propagation in the z direction.

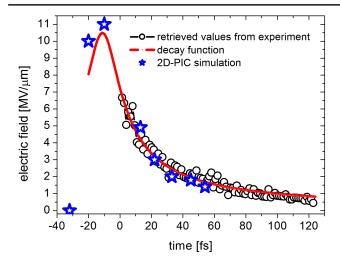


FIG. 6. Evaluated decrease of the acceleration field strength (black circles) from the experimentally obtained density modulation in the energy distribution of protons (cf. Fig. 1—90 signal maxima of the proton EDF); the red curve represents a nonlinear fit with a decay function (cf. the text) and blue stars results from 2D-PIC simulations (cf. the text).

The field strengths obtained from the theory and deduced from the experiment agree quite well (cf. Fig. 6).

We complement the results shown in Ref. [12] with Fig. 5, which shows the generation and evolution of electron bunches at every laser half cycle in the simulation. These bunches modulate the accelerating sheath field [28], and modulation of the density in the emitted proton beam becomes visible [12].

V. SIMPLE APPROACH TO RETRIEVE THE DECAY CHARACTERISTIC OF THE ACCELERATION FIELD

As briefly mentioned in our previous work [12], one can use the recorded modulation maxima $(E_{p_{\max,i}})$ or minima in the proton EDF in order to estimate the acceleration (a) of protons (mass m_p) with a distinct kinetic energy. In the following, in the data interpretation we restrict to a simplified description of the acceleration scenario. It neglects spatial dependencies of the acceleration field, and it assumes that different ion energies can be described with a different effective acceleration time in a temporally varying field. We will show that this data interpretation leads to a prediction of the temporal decay of the field which is in accordance to PIC simulation and suggestions of analytical models.

Every laser half-cycle $\tau_p/2 \sim 1.35$ fs, which is the modulation period of the oscillatory part of the Lorenz force at twice the laser frequency, a maximum and minimum density variation in the proton EDF is formed. The modulation gives a time stamp for the elapsing acceleration time of the ions in an electrical field with decreasing strength. We assume that the acceleration time is $i \tau_p/2$.

Thus, considering i acceleration cycles, we can approximate the ion (proton) velocities $v_{n,i}$ with

$$v_{p,I} - v_{p,i-1} = \sum_{n=1}^{i} a_n \tau_p / 2 - \sum_{n=1}^{i-1} a_n \tau_p / 2 = a_i \tau_p / 2.$$
 (1)

Doing so, we can take the experimentally determined energies of neighbor peaks i and i+1 $(W_{p_{\max,i}})$ for estimating the acceleration a_i for each i:

$$(W_{p_{\max i}})^{1/2} - (W_{p_{\max i-1}})^{1/2} = (m_p/2)^{1/2} (a_i \tau_p/2).$$
 (2)

The simple formula uses the approximation that the final EDF of the ions is a result of different acceleration times. This gives a value of the acceleration field strength E_i :

$$E_i = F_i / e_0 = m_p a_i / e_0 \tag{3}$$

(e_0 , elementary charge) which is plotted in Fig. 6. Because we do not know the absolute point in time when the acceleration starts, we chose t=0 for the first visible modulation peak at high kinetic energy. The proton cutoff energy is higher than the energy of the first peak. Between the cutoff and the first visible peak, no modulation is visible. Therefore, extrapolated field values concerning the cutoff, as well as the maximum field value in the PIC simulation (cf. Fig. 6), appear artificially with negative time values. The time, which is assigned to the field decay, is then $i \tau_p/2$ (i, number of recorded density maxima in the proton trace, $i_{\rm max}=90$).

The observables are the signal maxima in the proton trace which can be correlated with Eq. (1), and field strength calculation follows with Eqs. (2) and (3). These retrieved values of the electrical field strength are compared with an analytical function derived in Mora's work [29], which describes the temporal decay of the field. This function reads as $E = P_1 \{1 + P_2[(t - t_0) + P_3]^2\}^{-1/2}$ (E, electrical field strength; t, time; parameters $P_{1,2,3}$ and t_0 , which are used in the fit procedure). We determine P_1 with the following simple linear extrapolation: For the first observed maximum in the proton trace at 6.3 MeV, a field value of 6.7 MV/ μ m is retrieved. This ratio gives for the observed cutoff at 10 MeV a field value of 10.6 MV/ μ m, which we use for P_1 . The free parameters $P_{2,3}$ and t_0 are determined using a nonlinear fitting routine, and the obtained red curve is plotted in Fig. 5. Here, the used free parameters are not brought into further context with Mora's model. Apparently, this analytical function describes well the temporal field decay derived from the experiment.

But Mora's model [29] uses an isothermal approach which holds for the duration of the laser pulse. In our case, the laser pulse is much shorter than the derived timescale for the acting acceleration field. Therefore, a hybrid model approach [30], incorporating also a second adiabatic expansion phase when the laser pulse is off, is more appropriate. Such a situation is simulated in 2D-PIC

calculations covering a time window of 90 fs. The result is also plotted in Fig. 4 (blue stars): The data represent the maximum field strength which occurs in the acceleration sheath in the target normal direction at different times. Because the values derived from the experiment (black circles) do not allow timing of the laser peak (or pulse switch on), the maximum position of the PIC data and the decay function curve approximating the experiment were shifted to overlap. The comparison shows reasonable agreement.

Using the same timescale as in Fig. 6, we plot the energy of the protons at the appearing density peaks in Fig. 7.

In order to also compare the retrieved temporal development of the proton energies E_p with model predictions incorporating adiabatic cooling, the well-known interconnection between the laser-produced hot-electron distribution T_h and the TNSA protons is used. The following set of equations (cf. the analytical model in Ref. [21])—

$$E_p \approx 2k_B T_h,$$
 (4)

$$\eta(I) \approx 0.2 + \left(0.1 + 0.06 \frac{L}{\lambda}\right) I_{18}^{0.2},$$
(5)

$$T_h(I,t) \approx \frac{T_{0h} [\eta(I)I\lambda^2/1.37 \times 10^{18} \text{ (W} \mu\text{m}^2/\text{cm}^2)]^{0.5}}{1 + (2t^2T_{0h}/d^2m_i)[\eta(I)I\lambda^2/1.37 \times 10^{18} \text{ (W} \mu\text{m}^2/\text{cm}^2)]^{0.5}}$$
(6)

with k_B the Boltzmann constant, η the calculated laser absorption as a function of laser intensity I, laser wavelength λ , and scale length L of the gradient of the electron density at the laser-irradiated target surface, I_{18} the multiple of 10^{18} W/cm², d the target thickness, m_i the ion mass, and $T_{\rm oh}$ the model parameter—is used to calculate the proton energy as a function of time.

In order to reproduce the cutoff at $E_p = 10$ MeV for an intensity of 5×10^{19} W/cm² (FWHM), $\lambda = 800$ nm, and L = 80 nm [at a time $(t + t_0) = 0$, where t_0 is the fit parameter], we set $T_{0h} = 1.58$ MeV. This compares well with the ponderomotive energy of 1.4 MeV; cf., e.g., Ref. [31]. With $t_0 = 25$ fs, $m_i = m_p$, and $d^2 = 1.1 \ \mu\text{m}^2$,

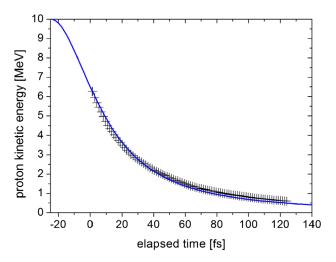


FIG. 7. Proton kinetic energy (crosses, experimental data) at the observed density maximum plotted at a temporal position concerning the maximum number i and a following elapsed time $i\tau_p/2 \sim i \times 1.35$ fs. The blue line represents a model (cf. the text) for how the proton energy decreases as a function of the hotelectron temperature.

the model curve in Fig. 6 is calculated. Here, d^2 is used as a fit parameter.

We can estimate the energy balance between the stored energy in the sheath field and the total amount of energy released with fast ions via acceleration in the sheath field. This estimation is made on the basis of a simple capacitor model for the sheath field. Calculating the energy stored in a capacitor and using an initialized field $E_{\rm sheathfield}$ of $10~{\rm MV}/{\mu m}$, an area $A_{\rm sheath}$ with a diameter of $10~{\mu m}$ at the target rear (set with a divergent hot-electron current from the focal region of the target front), and a Debye length λ_d of $1~{\mu m}$ [6] for the charge separation distance in the field (ε_0 , the vacuum dielectric constant) yields

$$W_{\rm sheath} = \frac{1}{2} \varepsilon_0 A_{\rm sheath} \lambda_d (E_{\rm sheathfield})^2 \approx 35 \text{ mJ}.$$
 (7)

This is at the order of 2% of the incident laser energy, also a consistent value when compared with experimentally determined laser-to-ion energy conversion efficiencies [32] for our laser and target parameters.

VI. CONCLUSION AND SUMMARY

One can conclude that the ion acceleration itself is a main channel of energy dissipation for the sheath acceleration structure. This "adiabatic expansion and cooling" is coupled to a fast decline of the acceleration field strength which we could trace with our experimental observation. This behavior sets limits for optimization in laser-driven ion acceleration, e.g., if the laser light absorption is enhanced via surface modification or other means. The initially very high values of the acceleration sheath field drop down very fast, nearly 50% in about 30 fs in our case. The fading of the acceleration field lasts much longer. The field decay is connected to the dynamics of the electron EDF. Because different branches of the hot-electron distribution develop differently, the effect on different types of secondary fast ion and x-ray emission is also different.

In summary, we can state that we can trace the temporal development of the electrical acceleration field at a femto-second timescale. The experimentally obtained density modulation in proton spectra and its interpretation give a key to study the temporal field decay which determines the distribution of ion energies. Our results support a fast decay of the acceleration field in the femtosecond timescale in accordance with plasma expansion and ion acceleration models as well as with simulation results. Especially the peak values of the sheath field are much higher as averaged values which are often used for estimations.

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