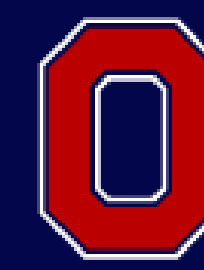


Electron dynamics in strong field trajectory models

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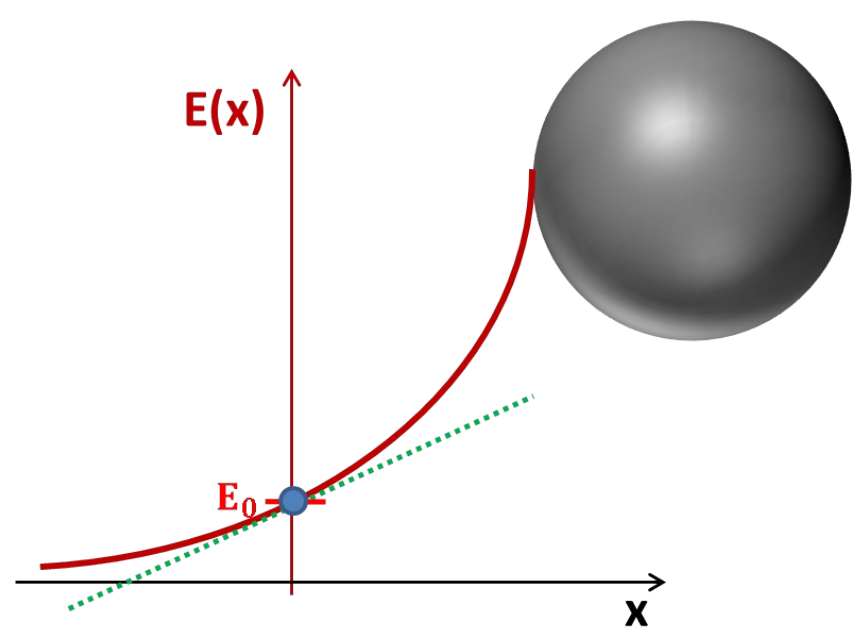
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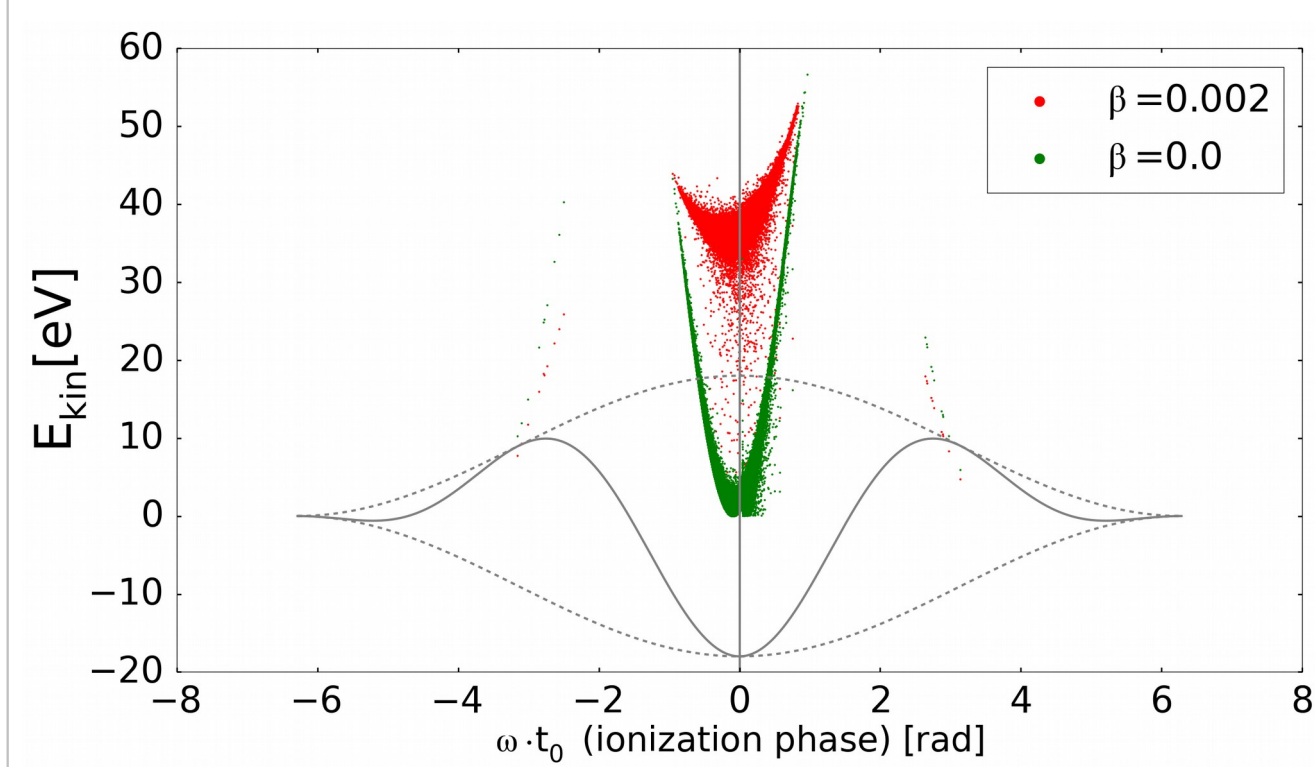
Emergence of a higher energy structure (HES) In inhomogeneous strong fields [H1,H2]

Setting:



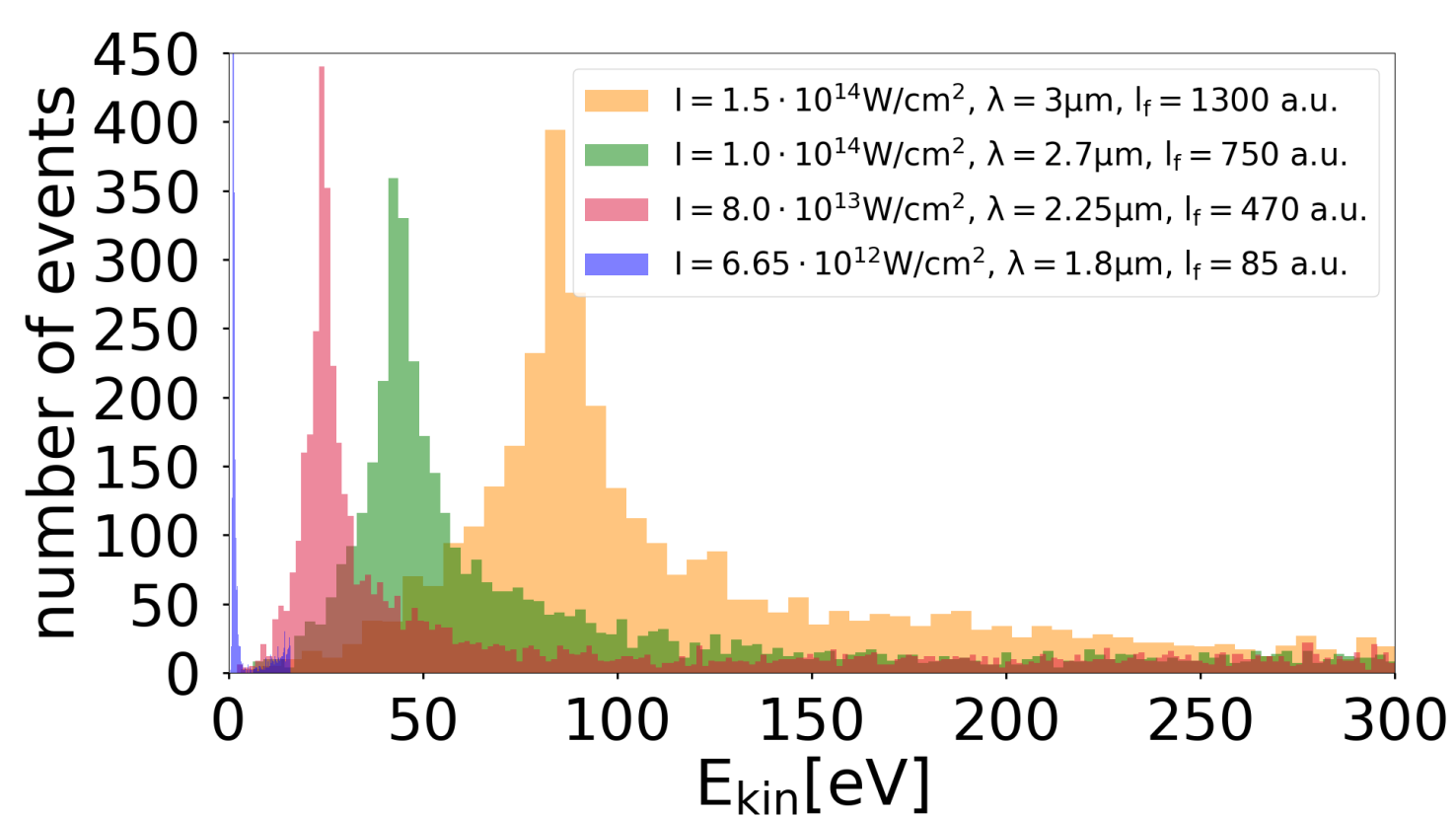
Ionization of gas in the inhomogeneous field in the vicinity of a nanostructure (linear approximation of field decay).

Trajectory analysis:

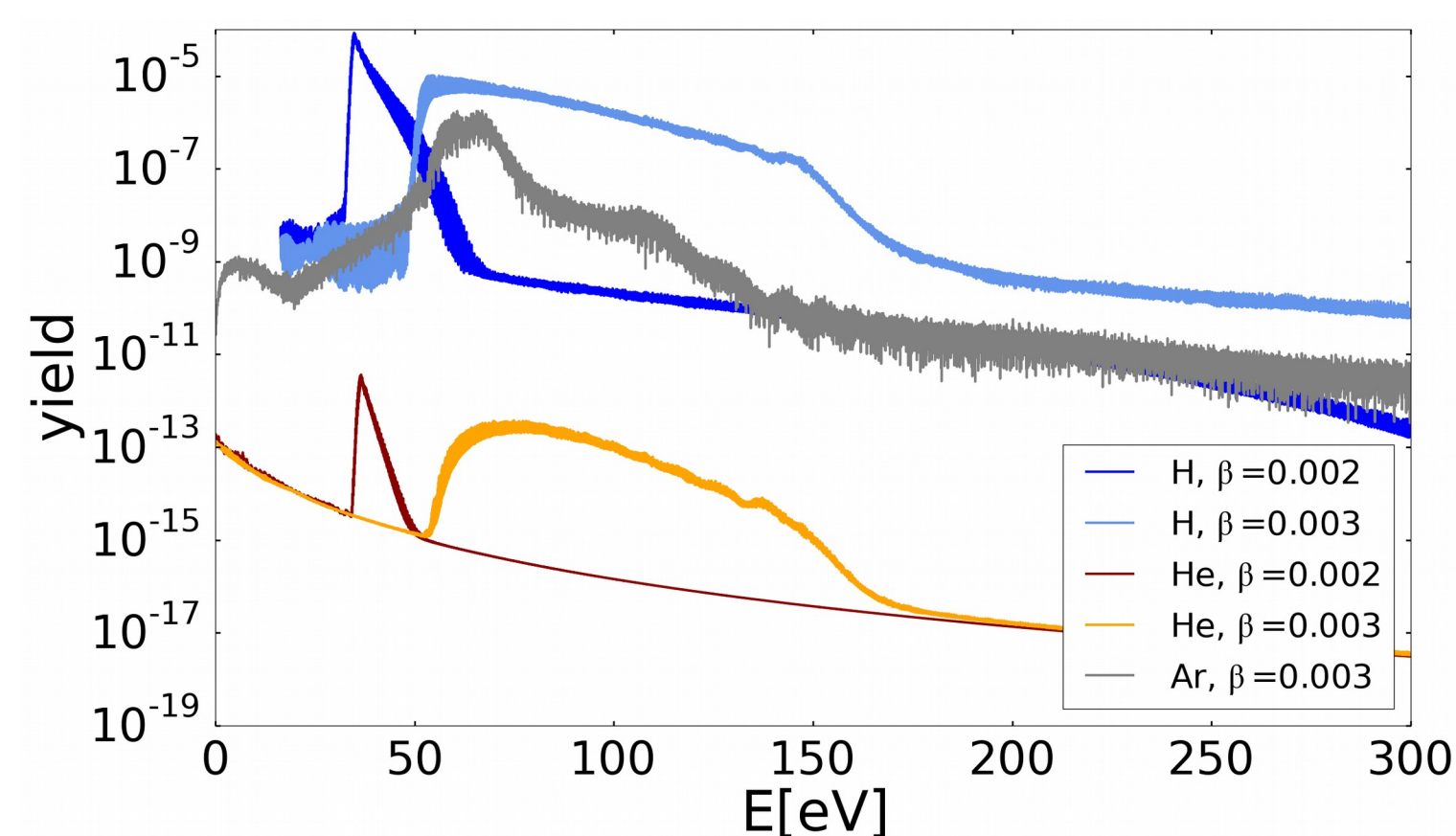


The electrons that form the HES are born within a narrow time window → creation of almost monoenergetic electron beams with femtosecond duration.

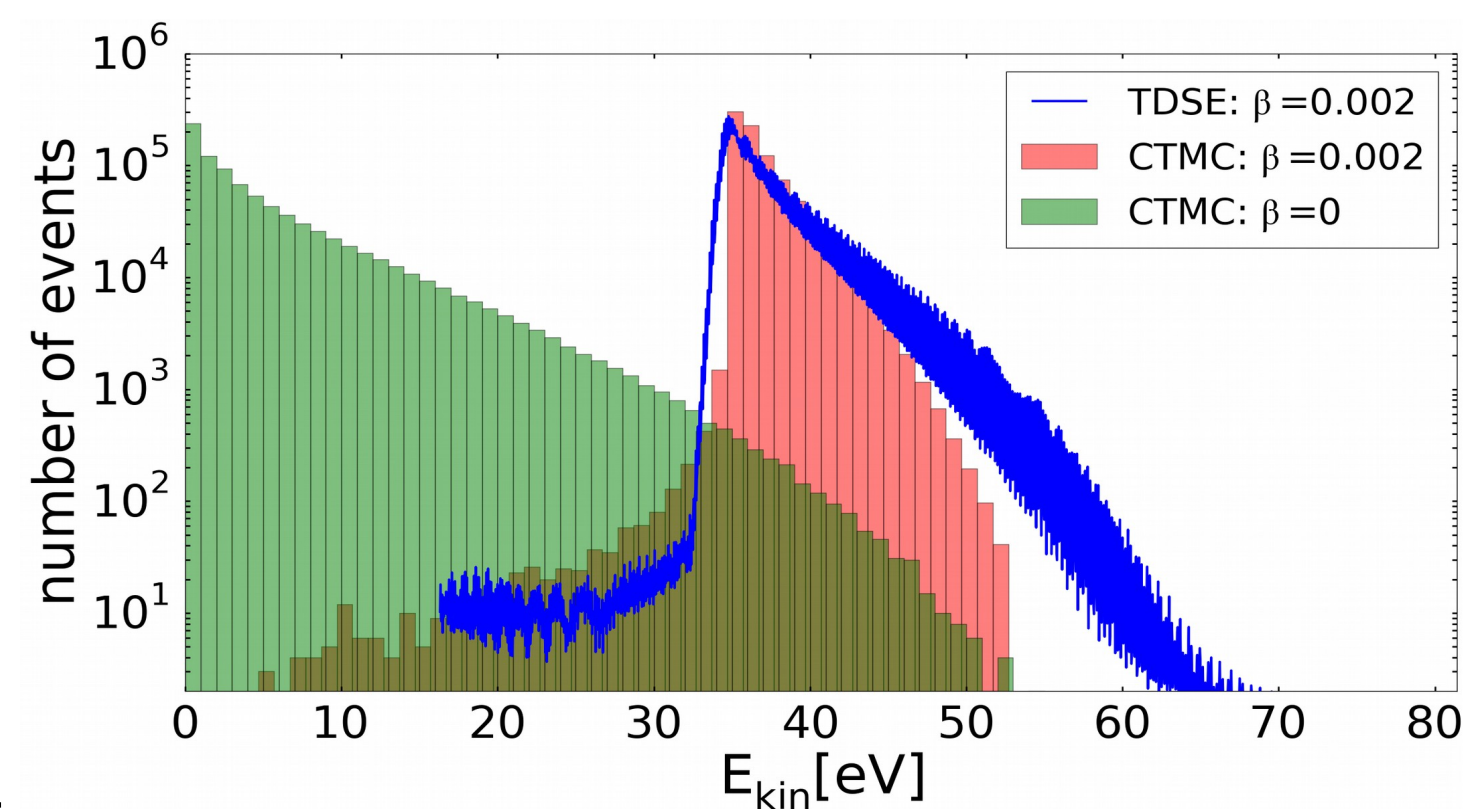
Generalization:



Photoelectron spectrum:

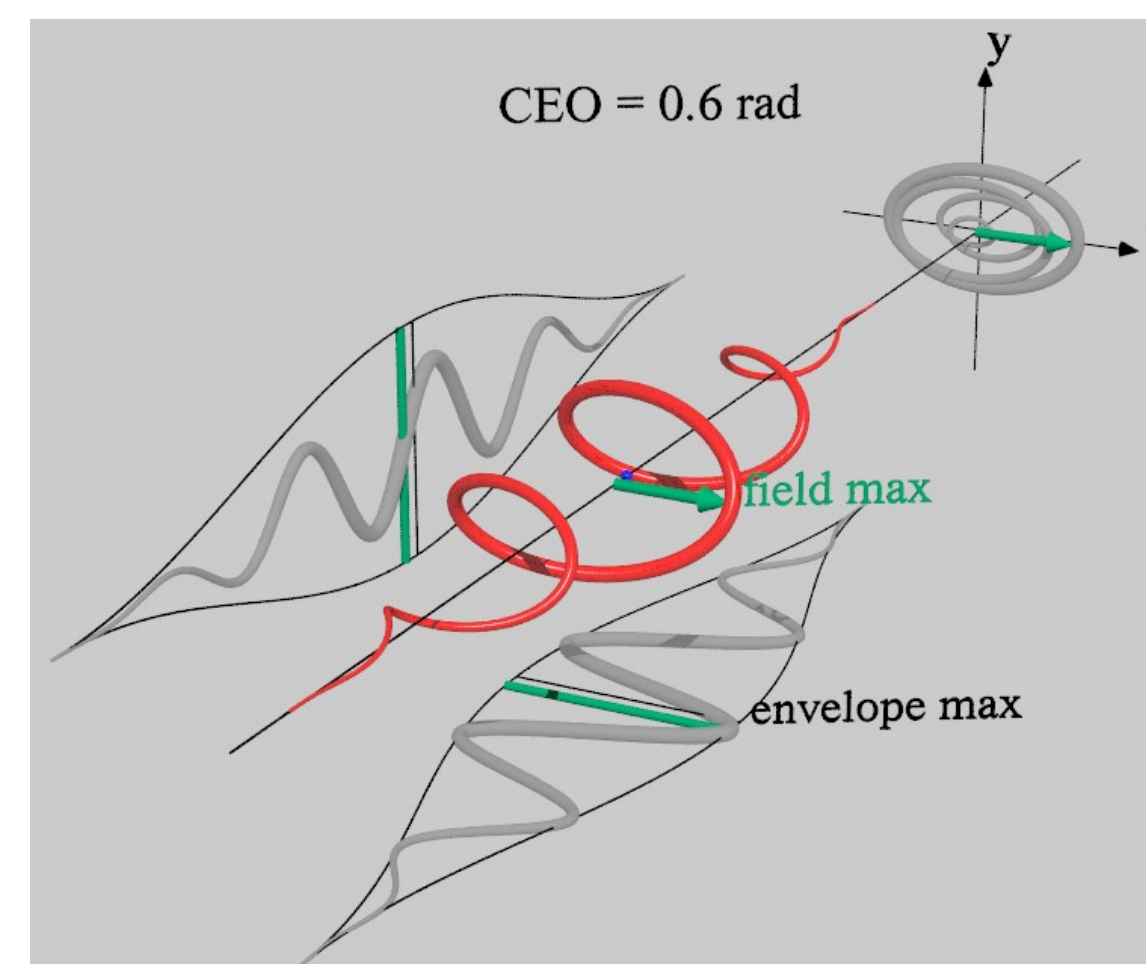


TDSE spectrum: Peaks emerge at a higher energy. The peak's width and position depend on the inhomogeneity parameter β [H1,H2].



The effect is captured by CTMC simulations.

Attoclock revisited on electron tunnelling time [A1]



Time Reconstruction:

- Single Classical Trajectory (SCT) simulation assuming instantaneous tunnelling at field maximum
- offset angle = ellipticity correction + delay

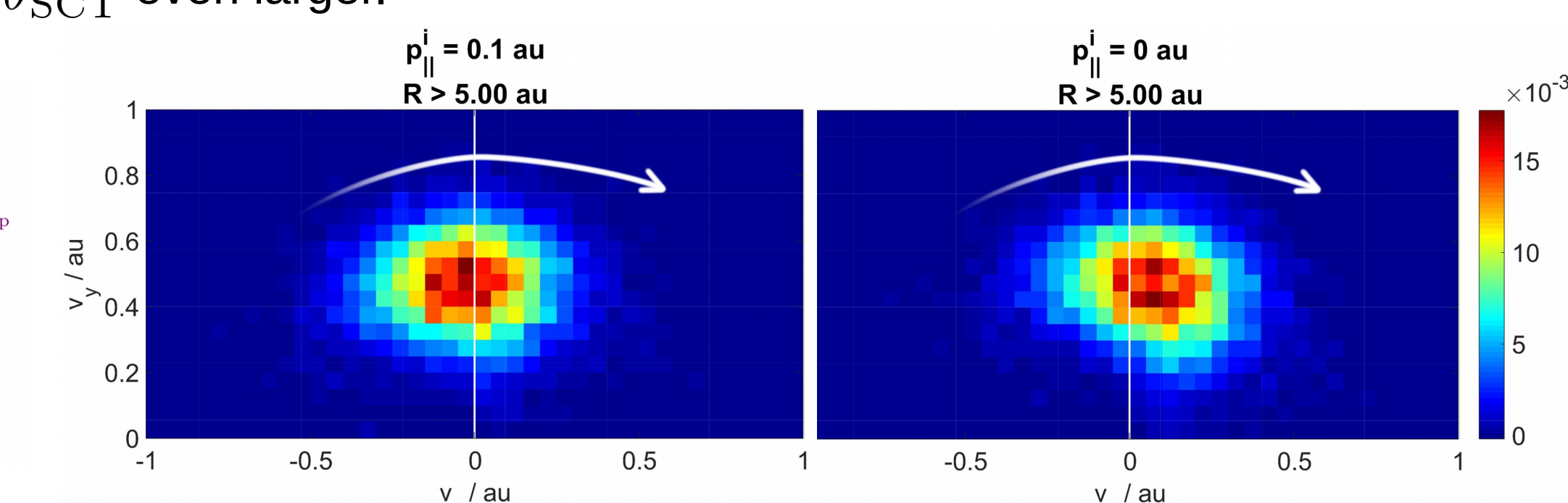
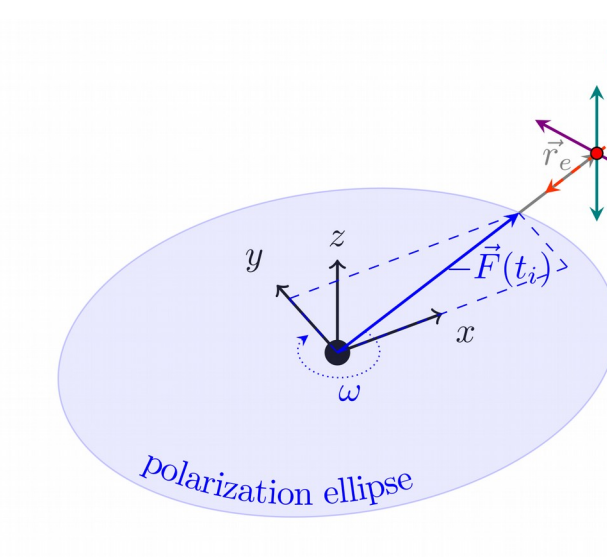
$$\theta - \theta_{SCT} = \frac{(1 - \epsilon^2)\omega t_i}{\epsilon} + \epsilon\omega t_i$$

Nonadiabatic effects [A4]:

- initial transverse momentum → lower field strength calibration for same data [A5]
- energy gain → shorter exit radius

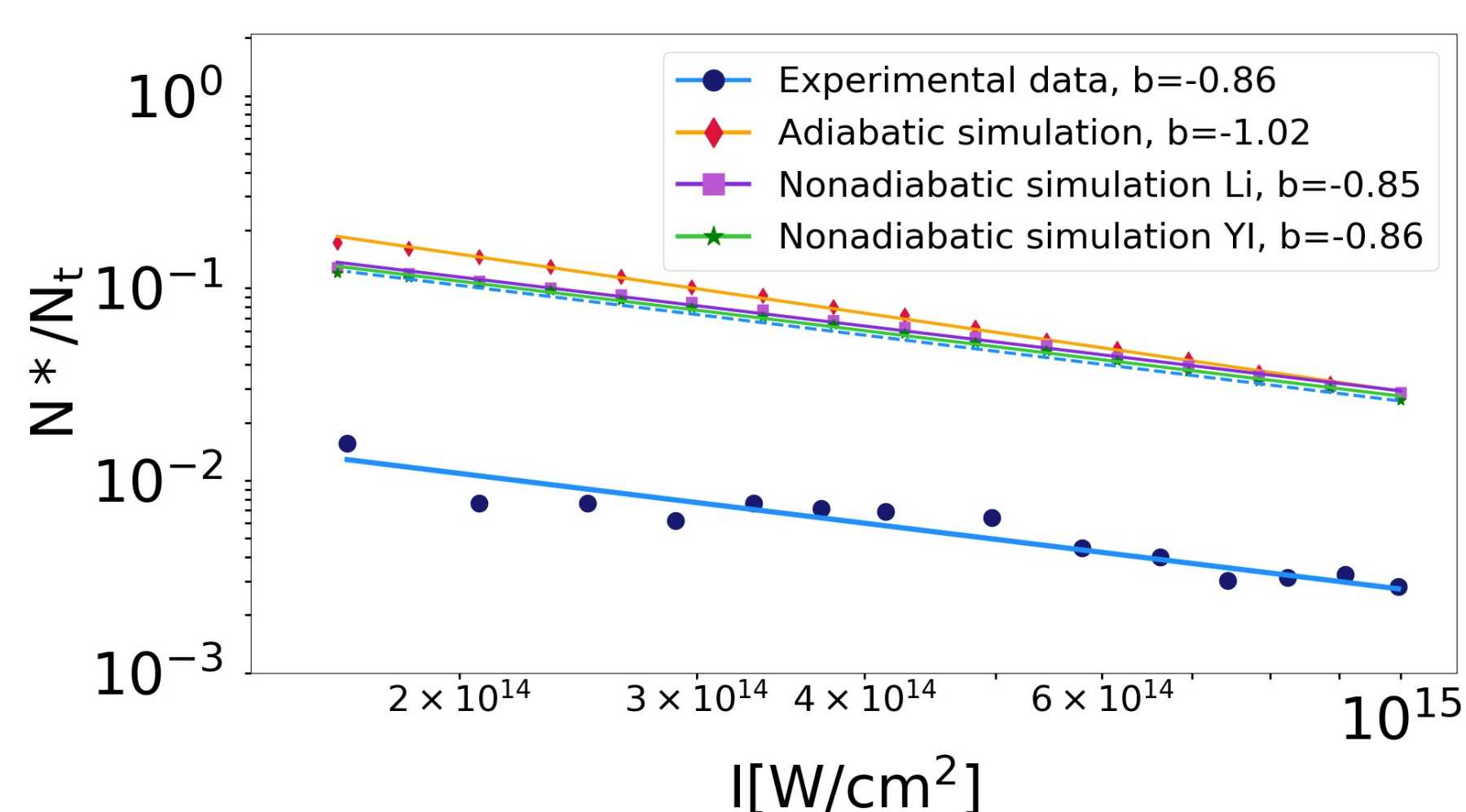
Initial condition dependence in classical trajectories

An initial momentum distribution centered around finite longitudinal momentum $p_{||}^i$ [A6,A7,R5] reduces $\theta_{SCT} \rightarrow \theta - \theta_{SCT}$ even larger.

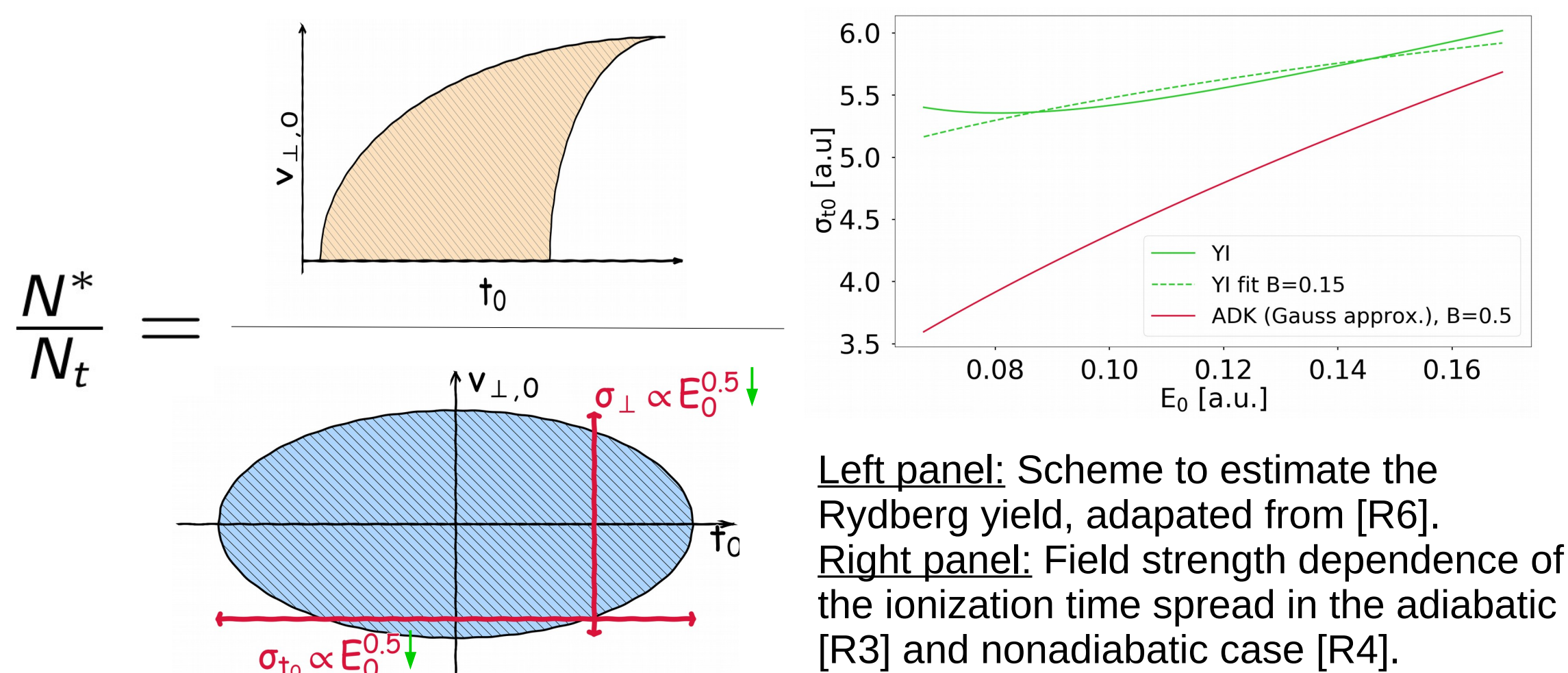


Rydberg states

Intensity dependence [R2]:



Experimental data extracted from [R1], the adiabatic and nonadiabatic data were obtained in CTMC simulations [R2] using [R3], [R4] and [R5] for the theories labelled ADK, YI, and Li respectively.

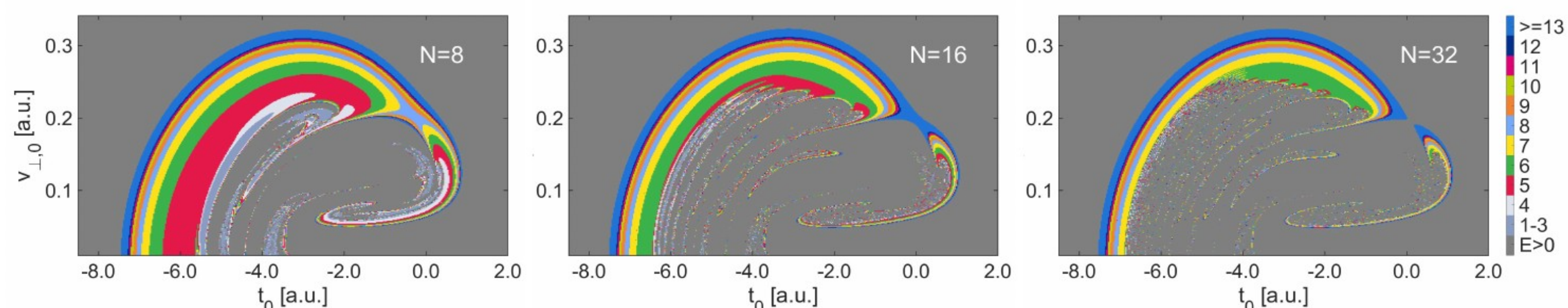


Left panel: Scheme to estimate the Rydberg yield, adapted from [R6]. Right panel: Field strength dependence of the ionization time spread in the adiabatic [R3] and nonadiabatic case [R4].

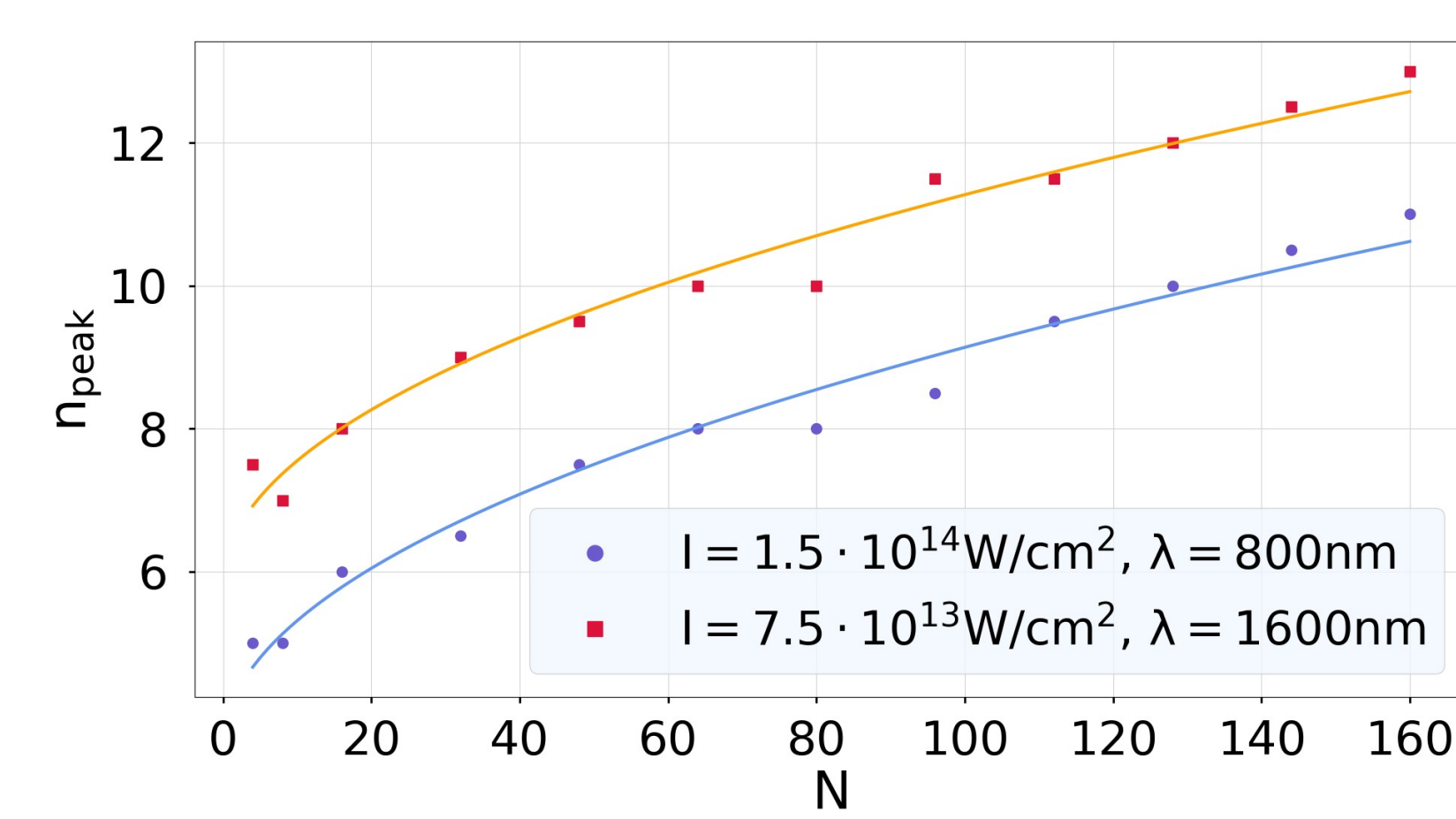
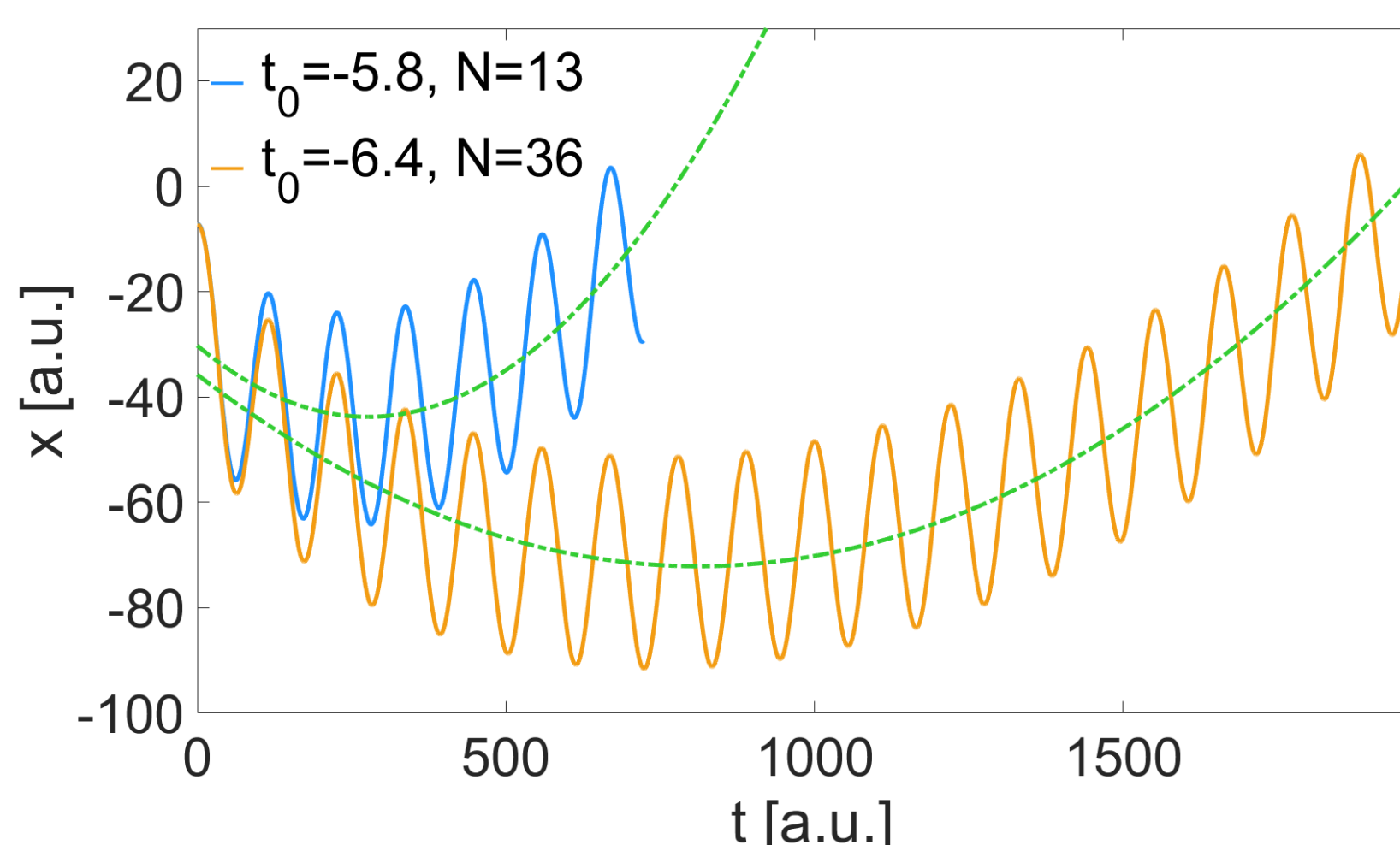
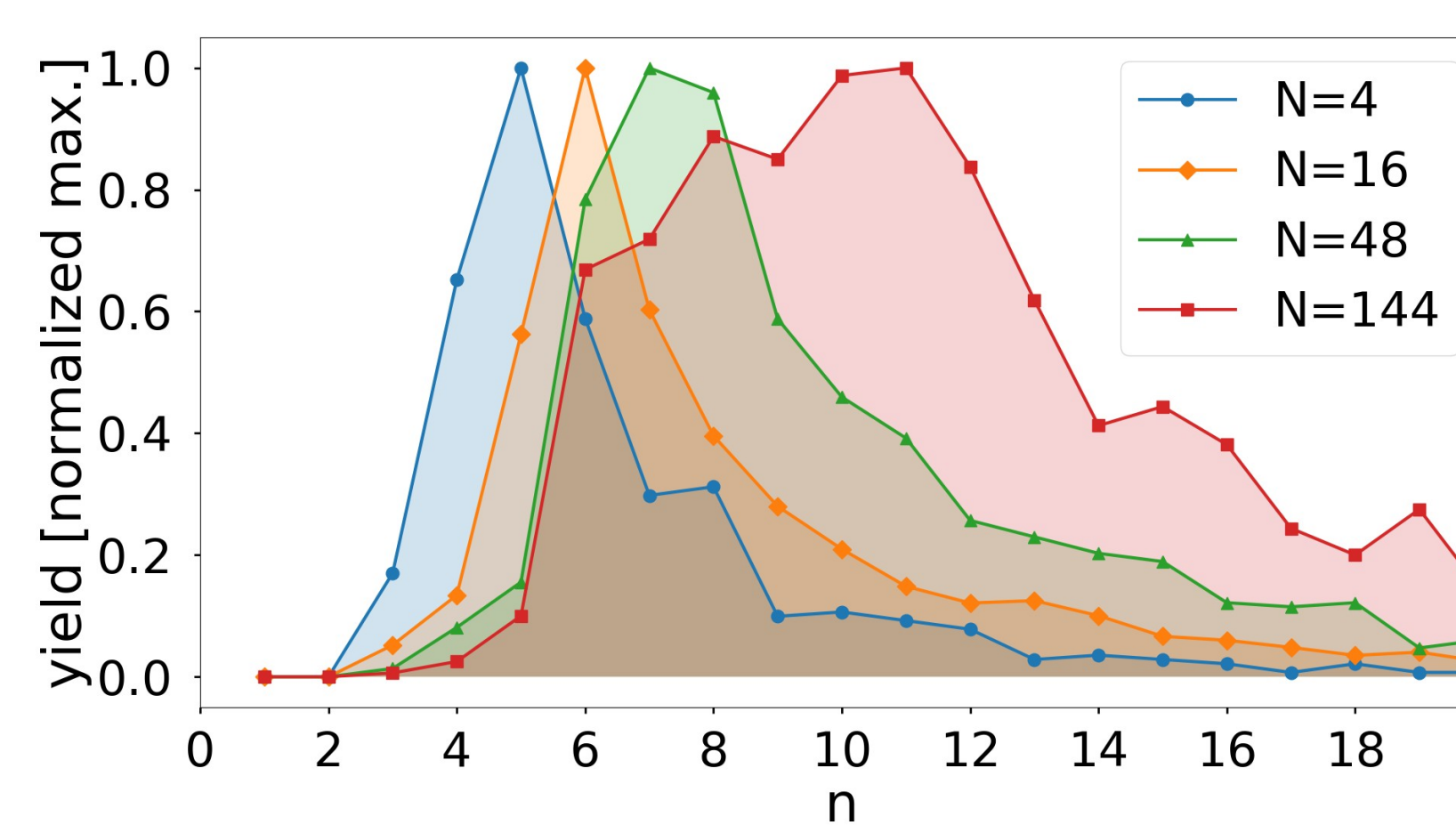
Nonadiabatic Rydberg yield decreases more slowly with increasing intensity, mainly because the time spread grows more slowly with increasing intensity in the nonadiabatic theory (due to nonadiabatic broadening at smaller intensities).

→ New nonadiabaticity test without calibration of absolute intensity

Pulse duration dependence [R7]:



Initial conditions for classical trajectories ending up in Rydberg states, colour coded by principal quantum number n , become fewer with increasing pulse duration.



Principal quantum number distribution shifting to larger values for longer pulses because:

- born closer to the field maximum ($t_0=0$)
- driven back to the parent ion in fewer optical cycles
- only 'survives' short pulse as Rydberg state

The same mechanism also explains Rydberg yield decreasing with increasing pulse duration.

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 [R7] Ortmann, Hofmann and Landsman, *arxiv:1810.07164* (2018)