

Laser-driven excitation schemes for the ^{229}Th isomer

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Abstract

The nucleus of ^{229}Th has attracted great interest in recent years as a candidate for a nuclear clock with outstanding properties [1, 2]. This is related to its low excitation energy of 8.36 eV and its exceptionally long lifetime of a few 10^3 s. In this talk, we investigate theoretically driving and quenching schemes that support the development of a nuclear clock.

Motivated by recent advances in VUV laser technology [3], the first part of this talk is devoted to the photoexcitation of the nuclear isomer using structured light [kir1]. We focus on spatial and temporal excitation patterns induced by Bessel and higher-order Gaussian modes for the magnetic dipole channel of the nuclear transition. Nuclear excitation in both solid-state targets and single ions is investigated, and the differences to plane wave driving are highlighted.

In the second part, we investigate laser-assisted internal conversion (IC) and electronic bridge (EB) schemes in the large band gap crystal $^{229}\text{Th}:\text{LiSrAlF}_6$. Doping-induced symmetry breaking within the pristine crystal leads to the emergence of electronic defect states around the ^{229}Th nucleus that can facilitate processes like EB and IC [5, 6]. We find that targeted laser-assisted EB and IC schemes for nuclear deexcitation prove to be significantly more efficient than spontaneous nuclear decay. This offers promising prospects for the operation of a solid-state clock.

References

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