

# Ultracold doubly-polar molecules: on the way to create them via a Laser-Assisted Self-Induced Feshbach Resonance

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Controlling the scattering properties between ultracold alkali-metal atoms and alkaline-earth or ytterbium atoms (with a closed-shell ( $^1S$ ) ground state) offers fascinating perspectives like the realization of topological phases of matter [1], novel Efimov states with systems of large mass imbalance, or the formation of ultracold polar paramagnetic molecules which are promising for instance for quantum simulation of lattice-spin systems [2]. It has long been thought that if one of the atoms has no magnetic moment, magnetic Feshbach resonances (MFR) could not be observed. A subtle coupling mechanism has been recently invoked to predict MFRs between paramagnetic  $Rb(^2S)$  and non-magnetic  $Sr(^1S)$  atoms [3, 4]. The observation of the resulting very narrow MFR remains however challenging [4].

We propose a new type of Feshbach resonance (FR): the Light-Assisted Self-Induced Feshbach Resonance (LASIFR), where the closed and open coupled channels characterizing the FR are both associated with the electronic ground state of a heteronuclear diatomic molecule, which possesses an intrinsic permanent electric dipole moment (PEDM) varying with the internuclear distance. A sub-THz photon couples the pair of ultracold atoms, occupying for instance the lowest motional state of an optical lattice site, to a bound rovibrational state close to the dissociation limit. The resonant pattern becomes obvious in the field-dressed-state picture, which formally maps the scheme of a MFR.

The interspecies scattering length can then be controlled by both the intensity and the detuning of the photon frequency with respect to the transition one. A frequency chirp across the resonance can be engineered to fulfill the adiabaticity criterion for the complete population transfer toward the molecular bound level. Like with a "regular" MFR, a STIRAP can be followed for transferring the population to the absolute ground state. We exemplify this proposal with the case of  $RbSr$ , based on our investigations on the structure of this molecule [5, 6].

The similarities of the proposed approach with previous works (see for instance [7, 8]) will also be discussed.

## References

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