

Spatially Modulated Interaction and Dipolar Interaction Induced Resonances

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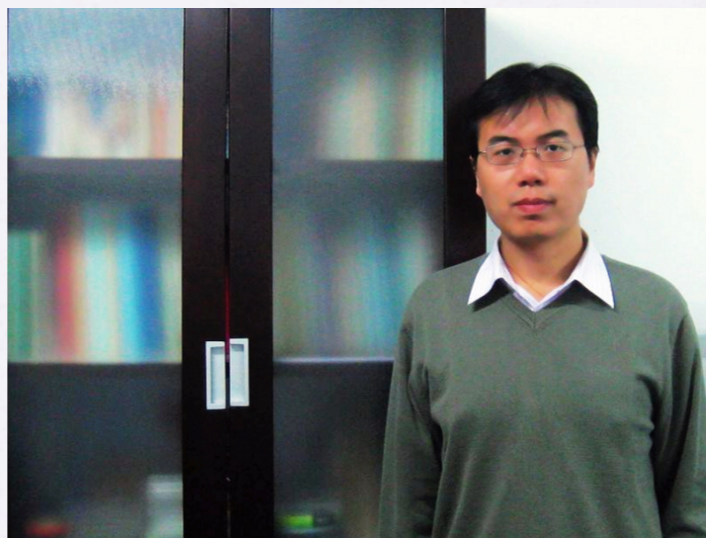
Three major recent topics:

1. Quantum gases in synthetic gauge field (with spin-orbit coupling)
2. New strongly interacting systems in quantum gases
3. Magnetism in quantum gases

Main Group Members



Ran Qi



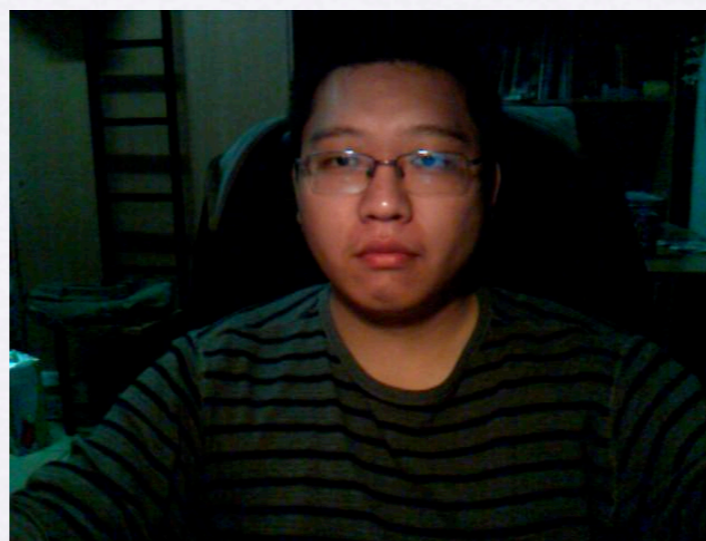
Zeng-Qiang Yu



Xiaoling Cui



Chao Gao

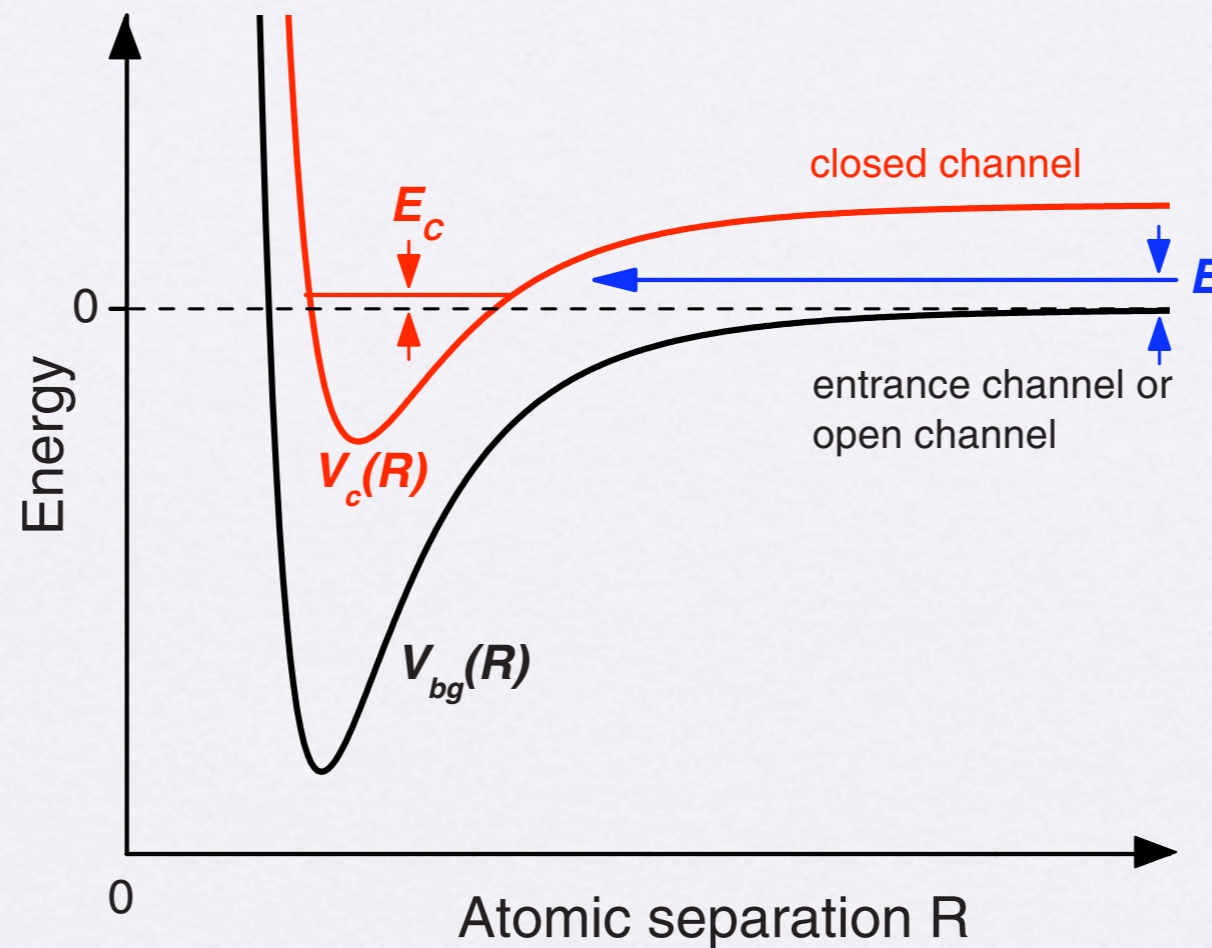


Zheyu Shi



Chao-Ming Jian

Feshbach resonance is an important tool to achieve strong interactions in ultracold Fermi gases



magnetic Feshbach resonance; optical Feshbach resonance; confinement induced resonance

**New Way to achieve scattering resonance
in New cold atom systems
with Novel features**

A: Spatially modulated interaction induced resonance

Alkali-earth-(like) atomic gases

B: Strong dipolar interaction induced resonance

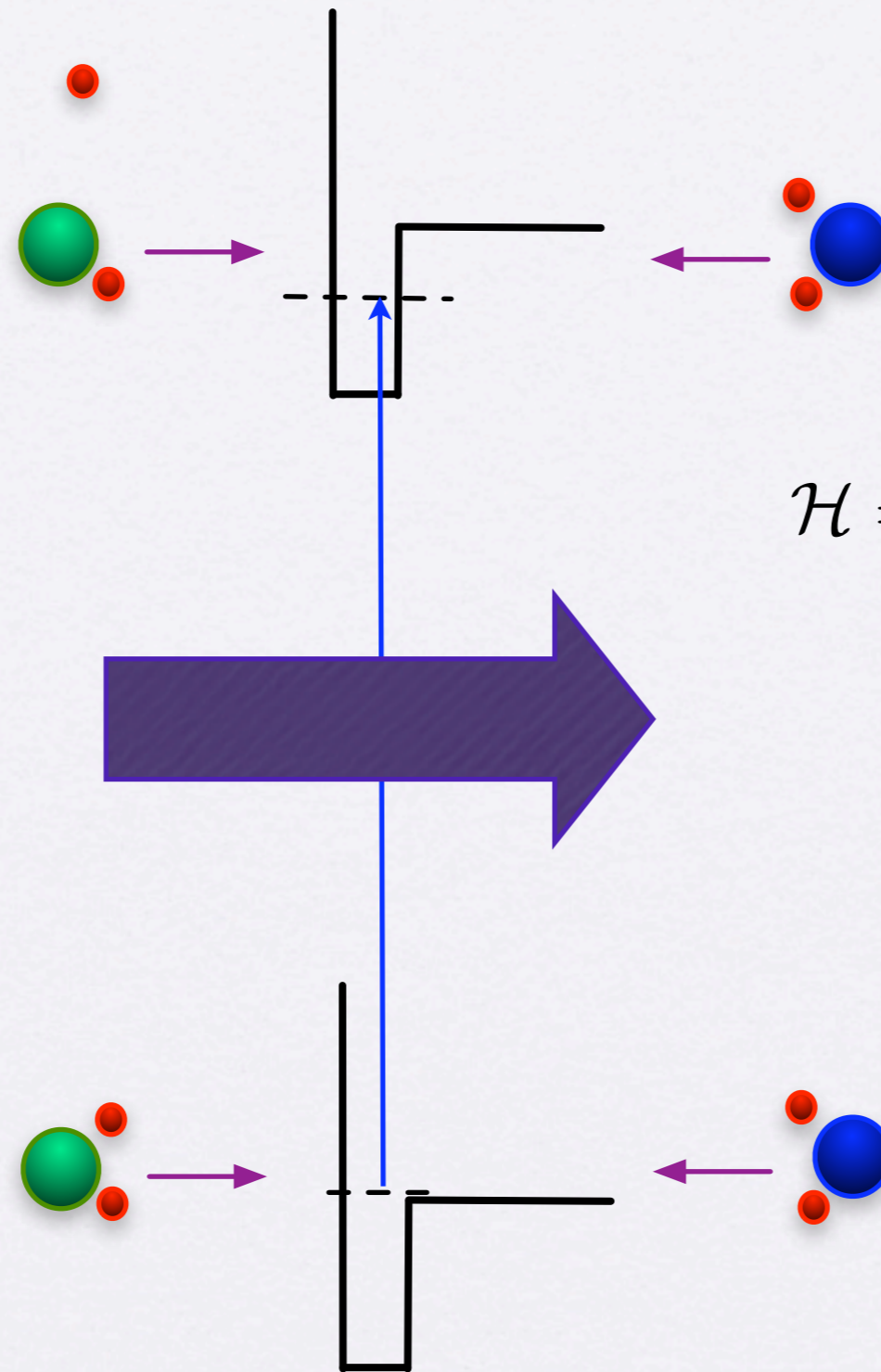
Polar molecular gases

**New Way to achieve scattering resonance
in New systems
with New features**

A: Spatially modulated interaction induced resonance

Alkali-earth-(like) atomic gases

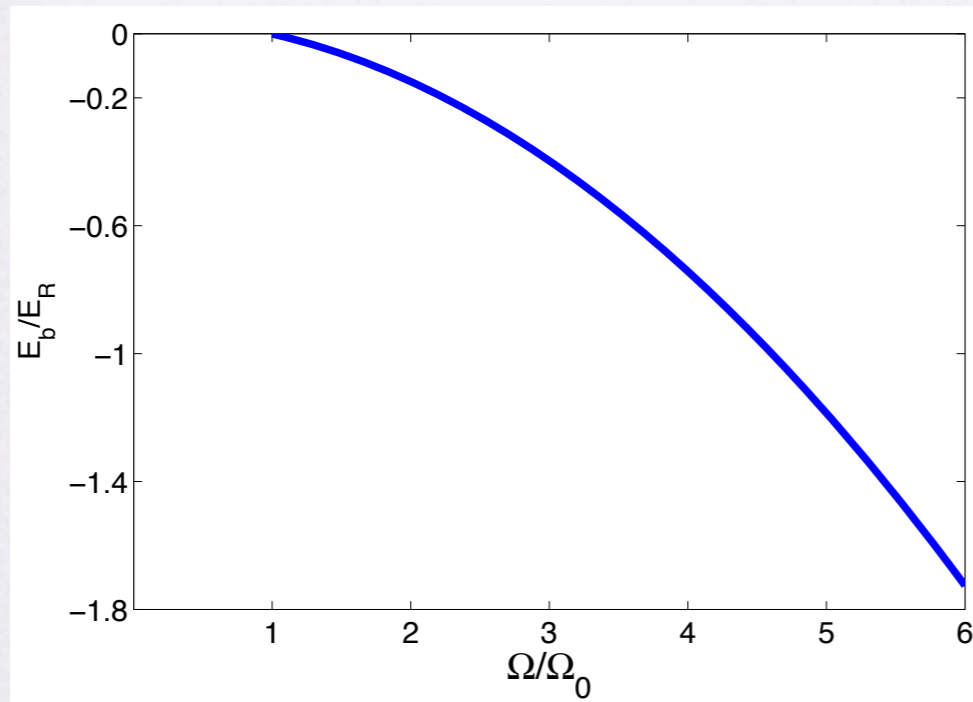
Idea of Optical Feshbach resonance



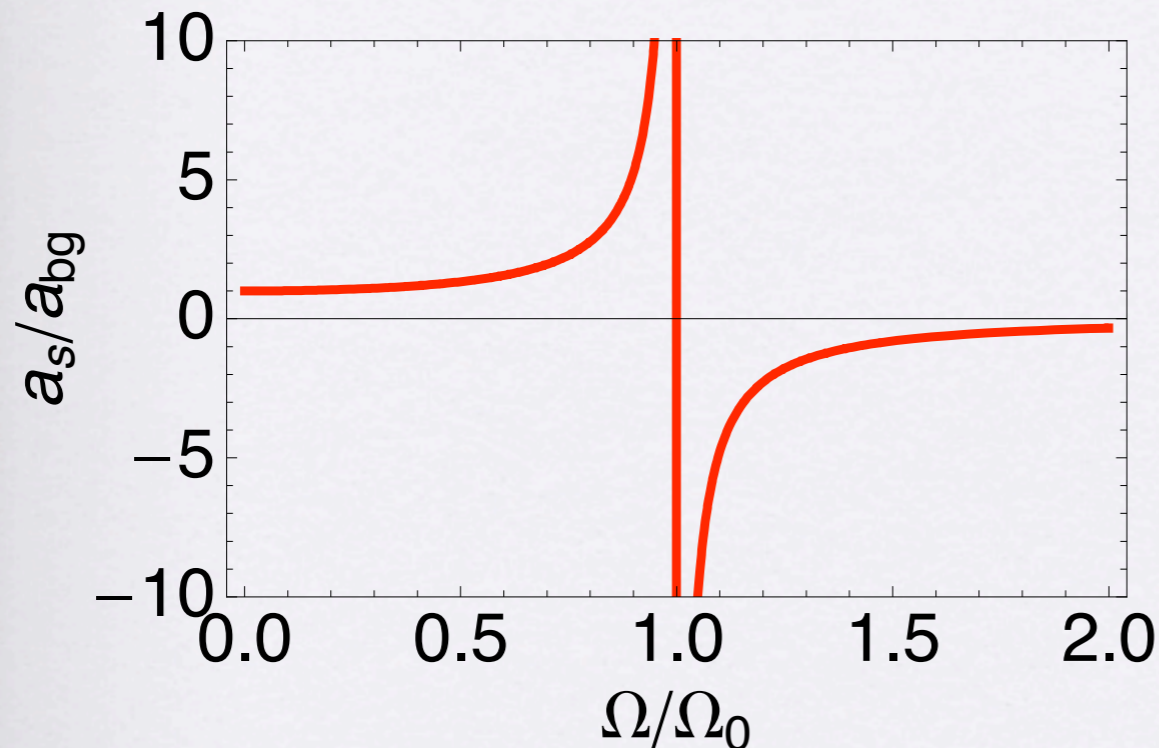
$$\mathcal{H} = -\frac{\hbar^2}{4m} \nabla_{\mathbf{R}}^2 - \frac{\hbar^2}{m} \nabla_{\mathbf{r}}^2 + v(\mathbf{r})$$

$$v(\mathbf{r}) = \begin{pmatrix} -V_0 & \hbar\Omega \\ \hbar\Omega & -V_c \end{pmatrix}$$

Optical Feshbach resonance



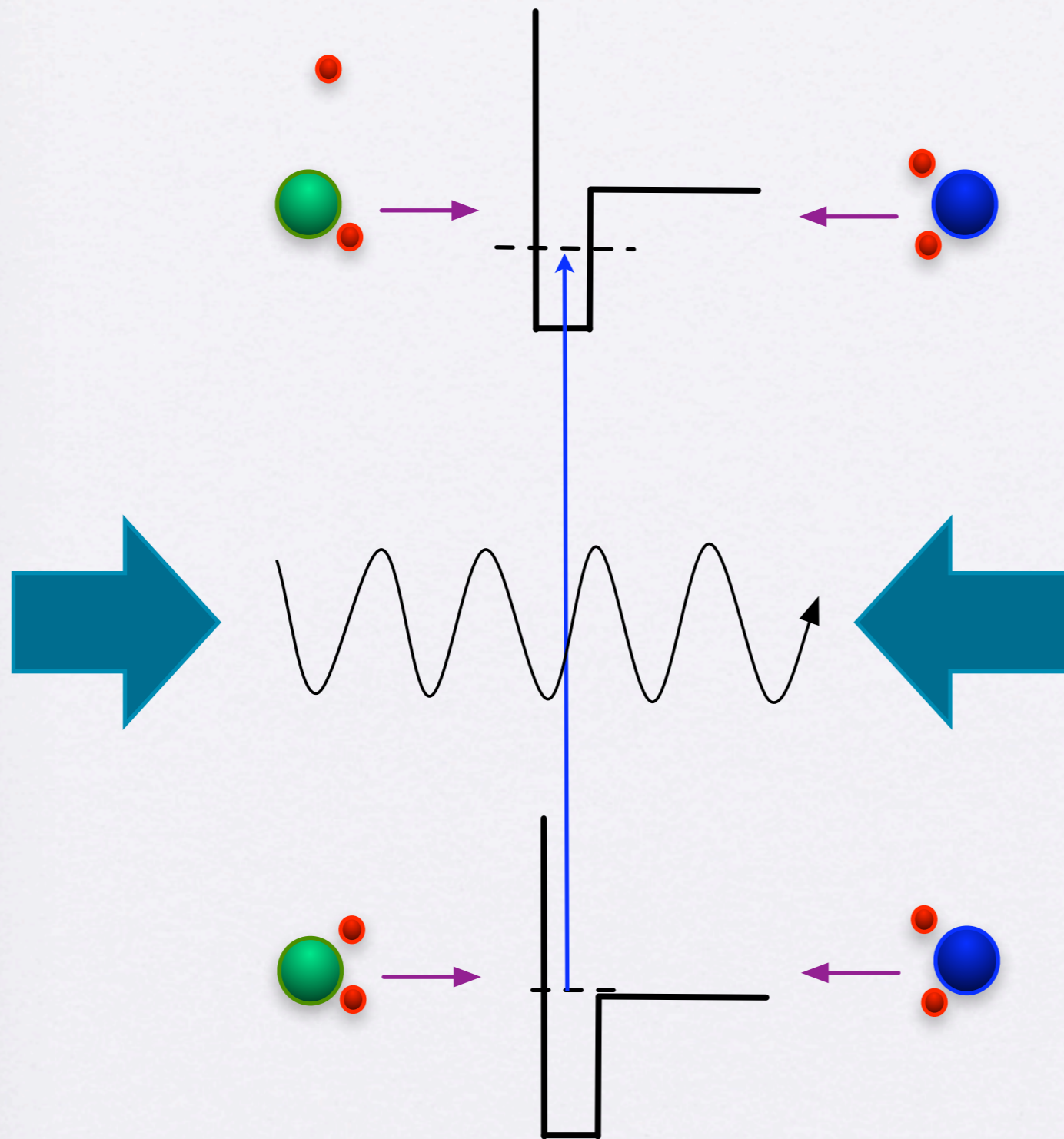
$$\mathcal{H} = -\frac{\hbar^2}{4m} \nabla_{\mathbf{R}}^2 - \frac{\hbar^2}{m} \nabla_{\mathbf{r}}^2 + v(\mathbf{r})$$



$$v(\mathbf{r}) = \begin{pmatrix} -V_0 & \hbar\Omega \\ \hbar\Omega & -V_c \end{pmatrix}$$

$$a_s = a_{bg} \left(1 - \frac{\Omega^2}{\Omega^2 - \Omega_0^2} \right)$$

Optical Feshbach resonance with Standing wave



Spatial dependent interaction

Two-body interaction potential:

$$V(\mathbf{r}_1, \mathbf{r}_2) = V(\mathbf{r}_1 - \mathbf{r}_2)$$

Spatial independent



Spatial dependent interaction

Two-body interaction potential:

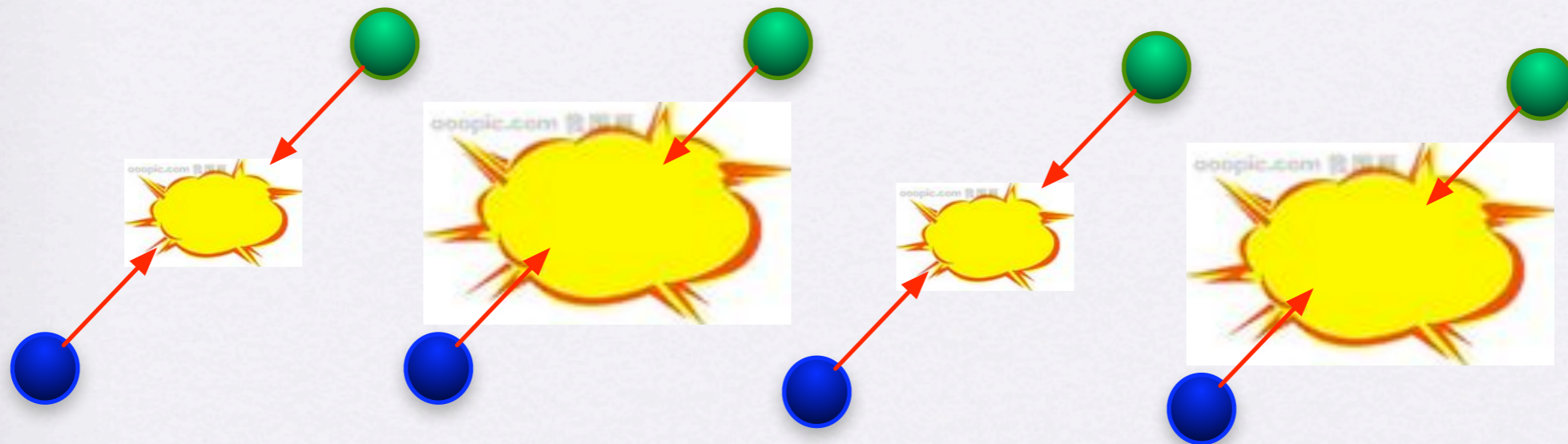
$$V(\mathbf{r}_1, \mathbf{r}_2) = V\left(\mathbf{r}_1 - \mathbf{r}_2, \frac{\mathbf{r}_1 + \mathbf{r}_2}{2}\right)$$

$$\mathcal{H} = -\frac{\hbar^2}{4m} \nabla_{\mathbf{R}}^2 - \frac{\hbar^2}{m} \nabla_{\mathbf{r}}^2 + v(\mathbf{r}, \mathbf{R})$$

$$v(\mathbf{r}) = \begin{pmatrix} -V_0 & \hbar\Omega(\mathbf{R}) \\ \hbar\Omega(\mathbf{R}) & -V_c \end{pmatrix}$$

Spatially periodically modulated

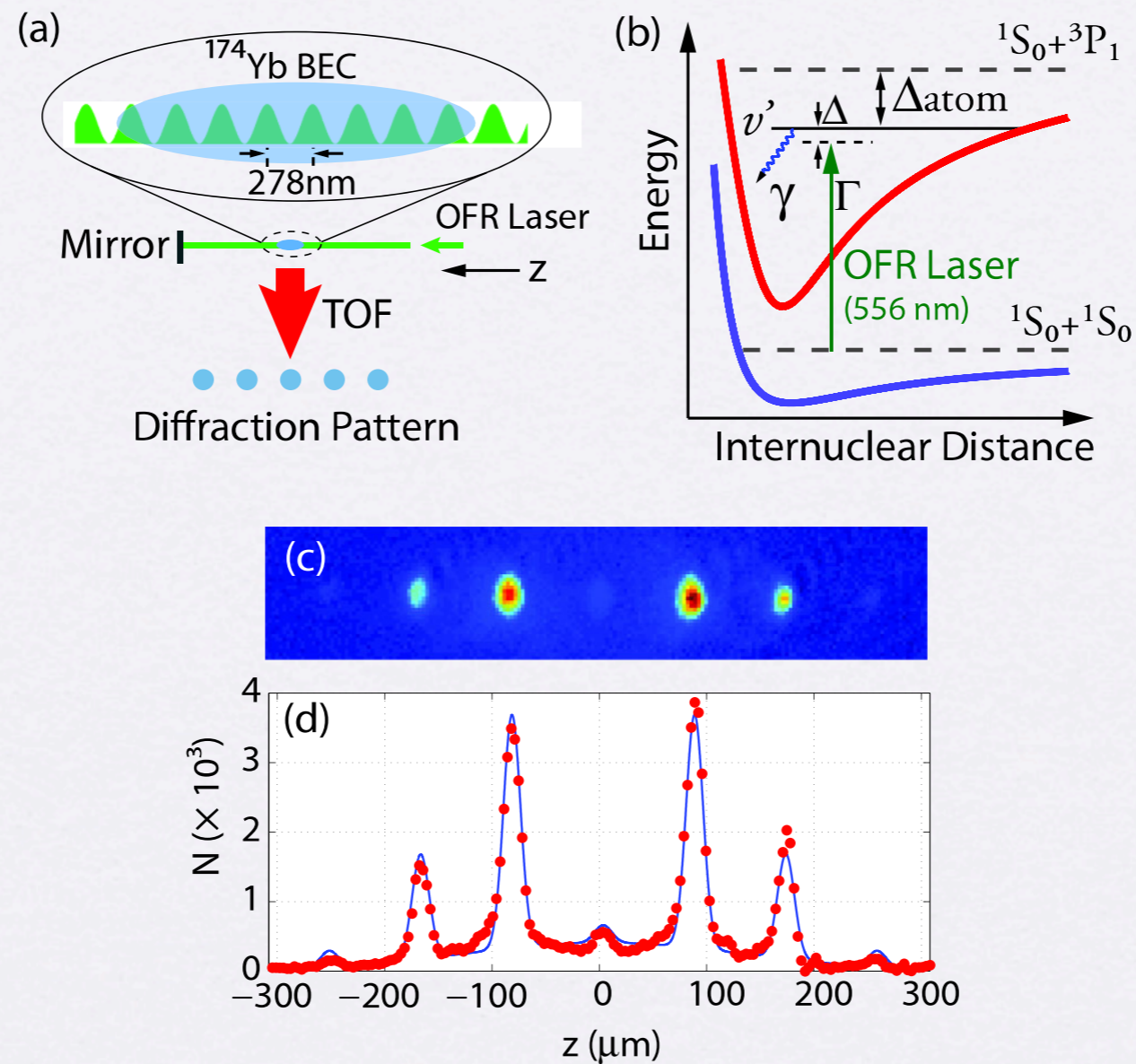
$$\Omega(\mathbf{R}) = \Omega \cos(Kx)$$



$a_s(x)$ is spatially dependent and modulates periodically in space

Experimental Realization

Submicron spatial modulation of an interatomic interaction in a Bose-Einstein condensate, PRL, 105, 050405 (2010) Kyoto group



How $a_s(x)$ **modulates in space?**

$$a_s = a_{\text{bg}} \left(1 - \frac{\Omega^2}{\Omega^2 - \Omega_0^2} \right) \quad \Omega(\mathbf{R}) = \Omega \cos(Kx)$$



$$a_s(x) = a_{\text{bg}} \left(1 - \frac{\Omega^2 \cos^2(Kx)}{\Omega^2 \cos^2(Kx) - \Omega_0^2} \right)$$



Any other physics effects?

What we have done:
Solve two-body problem of this Hamiltonian

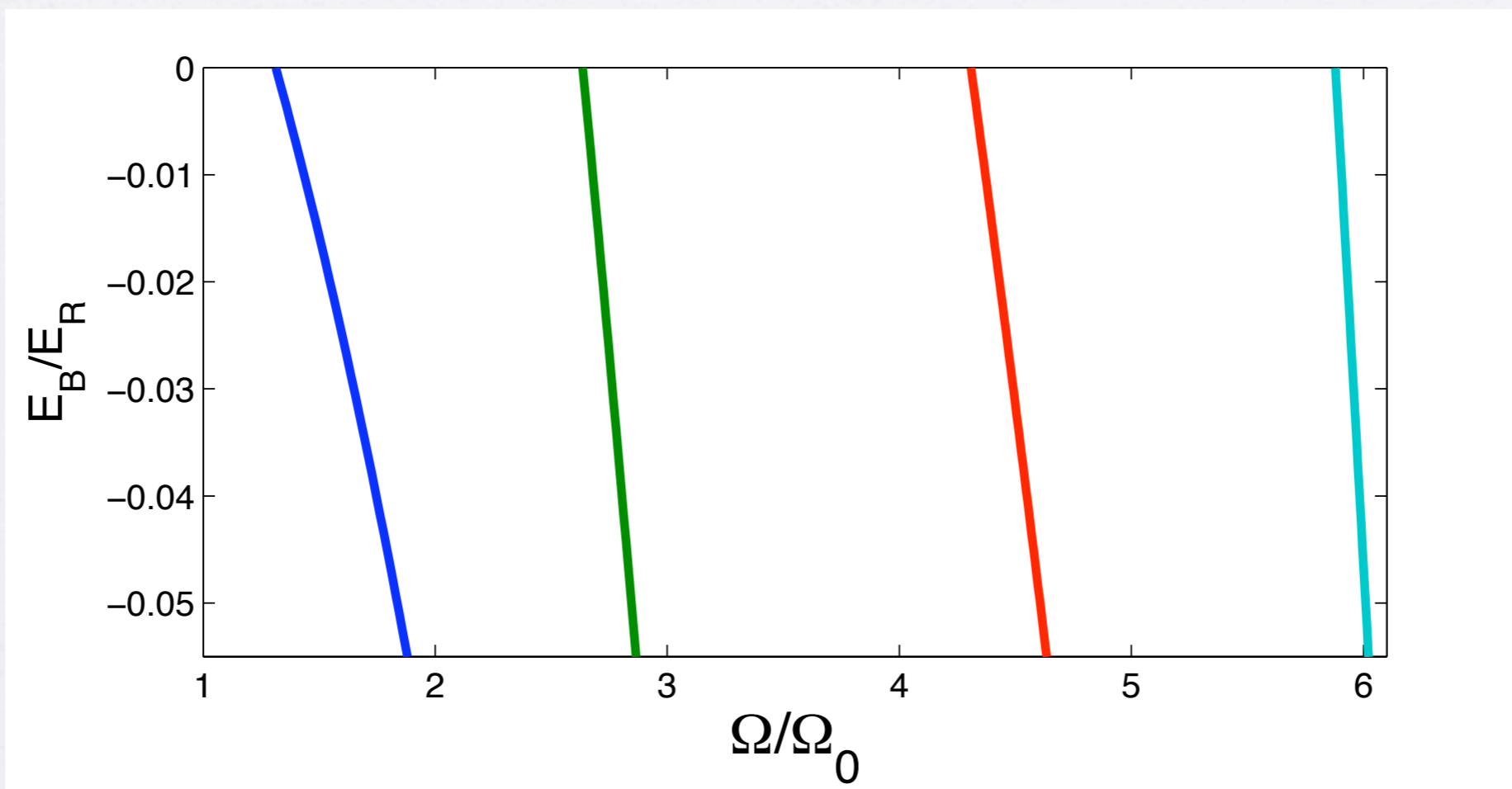
$$\mathcal{H} = -\frac{\hbar^2}{4m} \nabla_{\mathbf{R}}^2 - \frac{\hbar^2}{m} \nabla_{\mathbf{r}}^2 + v(\mathbf{r})$$

$$v(\mathbf{r}) = \begin{pmatrix} -V_0 & \hbar\Omega(\mathbf{R}) \\ \hbar\Omega(\mathbf{R}) & -V_c \end{pmatrix}$$

$$\Omega(\mathbf{R}) = \Omega \cos(Kx)$$

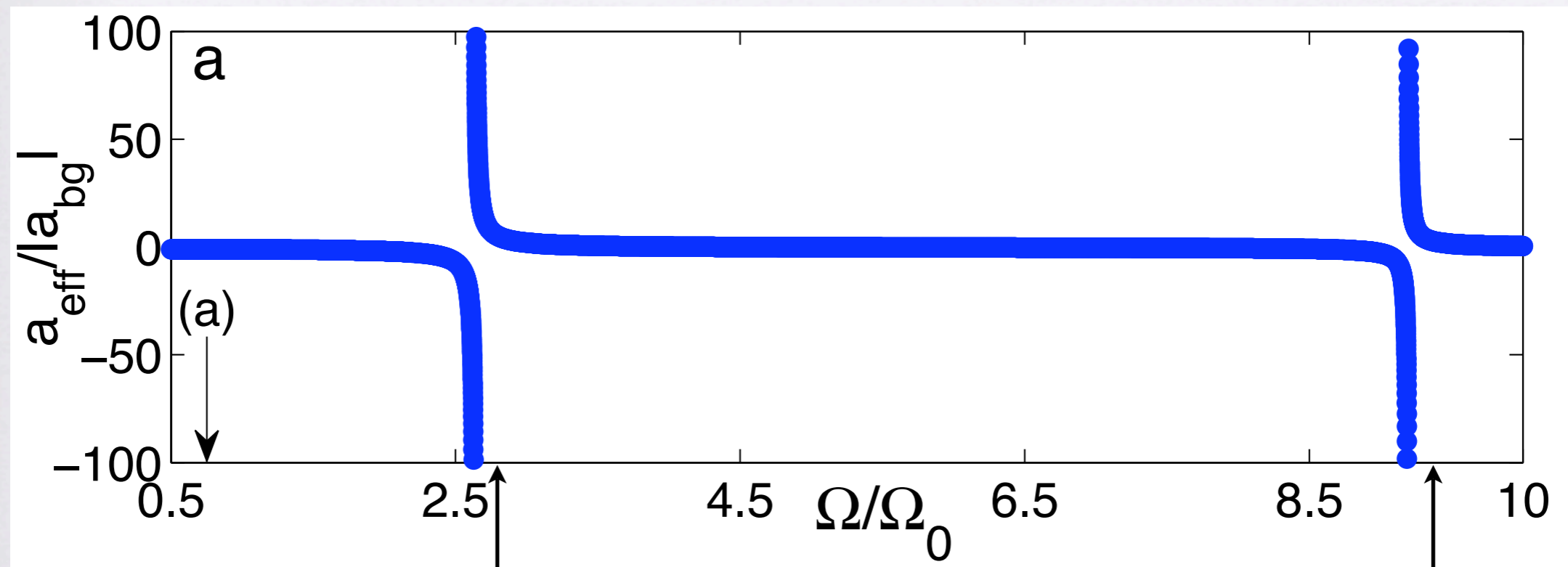
Qi Ran and HZ, arXiv: 1101.4464

Results I: Bound States



Results II: Scattering Resonances

$$a_{\text{eff}} = \lim_{k \rightarrow 0} \frac{\tan \delta(k)}{k}$$



Resonances

Resonances

**Strongly interacting many-body system !!
Universal Behavior ?**

Results III: Local Scattering Length --- related to local interaction energy

Bethe-Peierls condition:

$$\lim_{r \rightarrow 0} \psi(r, x) = \frac{1}{r} - \frac{1}{a_{loc}(x)}$$

Local scattering length

$$a_{loc}(x) = - \lim_{r \rightarrow r_0} \frac{r\psi_0(x, r)}{\partial_r (r\psi_0(x, r))}$$

The mean-field energy for a BEC:

$$\mathcal{E} = \int dx \left[-\frac{\hbar^2}{2m} \varphi^* \nabla^2 \varphi + \frac{4\pi\hbar^2}{m} a_{loc}(x) n^2(x) \right]$$

Results III: Local Scattering Length

**Exact
formula:**

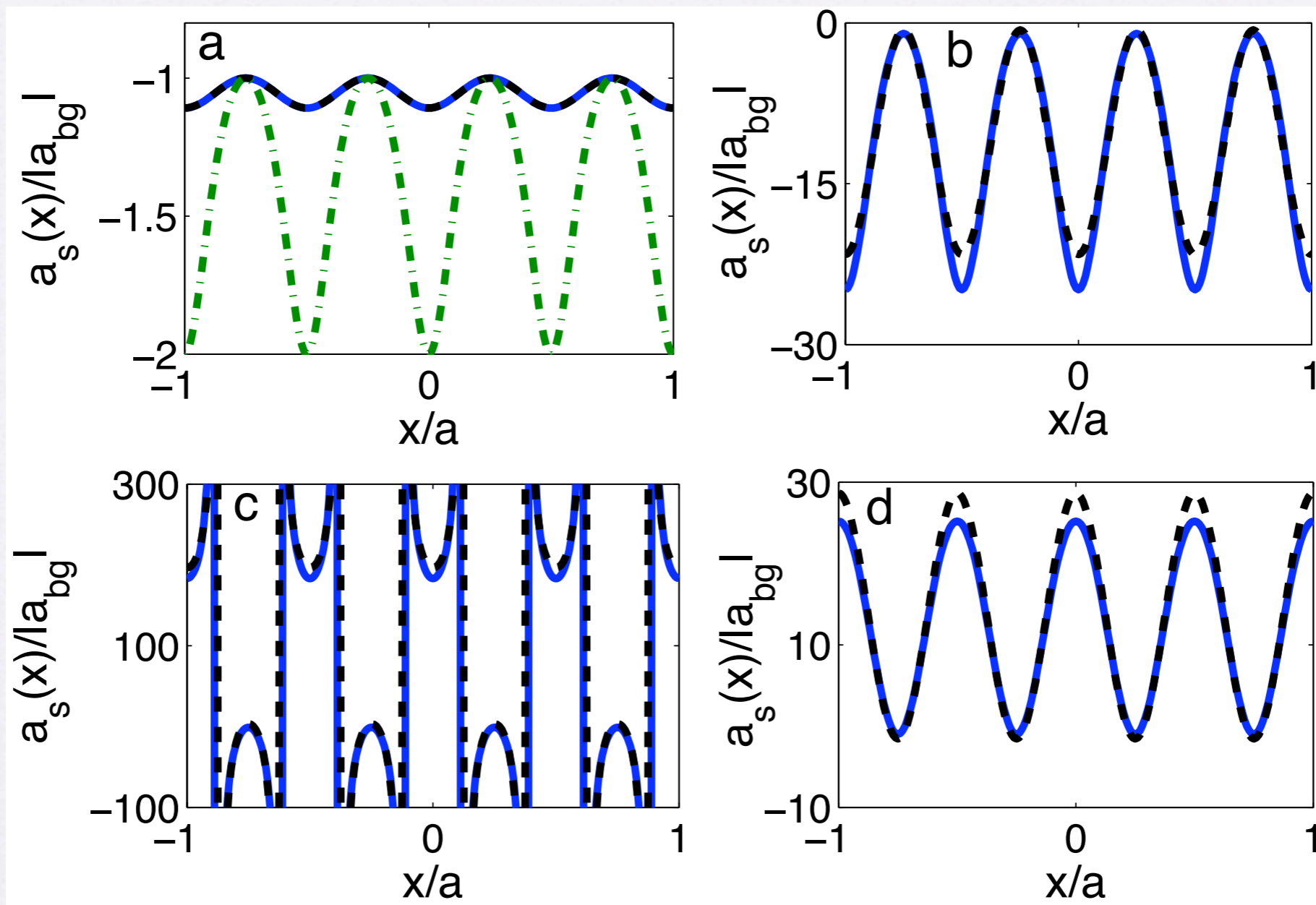
$$a_{\text{loc}}(x) = \frac{1 - \sum_{m \neq 0} U_m \cos(mKx)/U_0}{a_{\text{eff}}^{-1} - \sum_{m \neq 0} U_m |m|K \cos(mKx)/(2U_0)}$$

**Simplified
formula**

$$Ka_{\text{eff}} \ll 1 \quad a_{\text{loc}}(x) = a_{\text{eff}} \left[1 - \frac{2U_2}{U_0} \cos(2Kx) \right]$$

$$Ka_{\text{eff}} \gg 1 \quad a_{\text{loc}}(x) = \frac{1}{K} \left[1 - \frac{U_0}{2U_2 \cos(2Kx)} \right]$$

Results III: Local Scattering Length



**Summary:
Take Home Message**

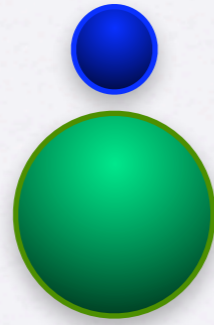
New Mechanism	New System	New Features
Two-body interaction potential has center-of-mass dependence	Alkali-earth-(like) atomic gases: Sr, Ca, Yb	Spatially dependent local scattering length

**New Way to achieve scattering resonance
in New systems
with New features**

B: Strong dipolar interaction induced resonance

Polar molecular gases

Polar molecular gases



large dipole moment: d

Hard to cool it directly !!

Polar molecular gases

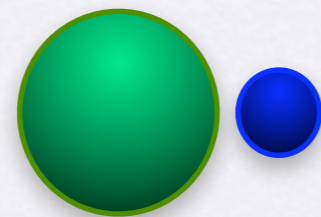


Feshbach molecule



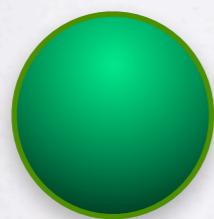
Difficulties:

- 1. Large energy detuning: 10-100 THz**
- 2. Small transition matrix element**

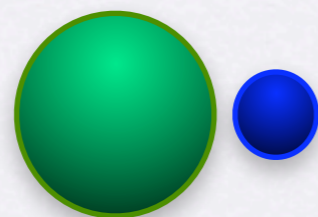


Ground state molecule

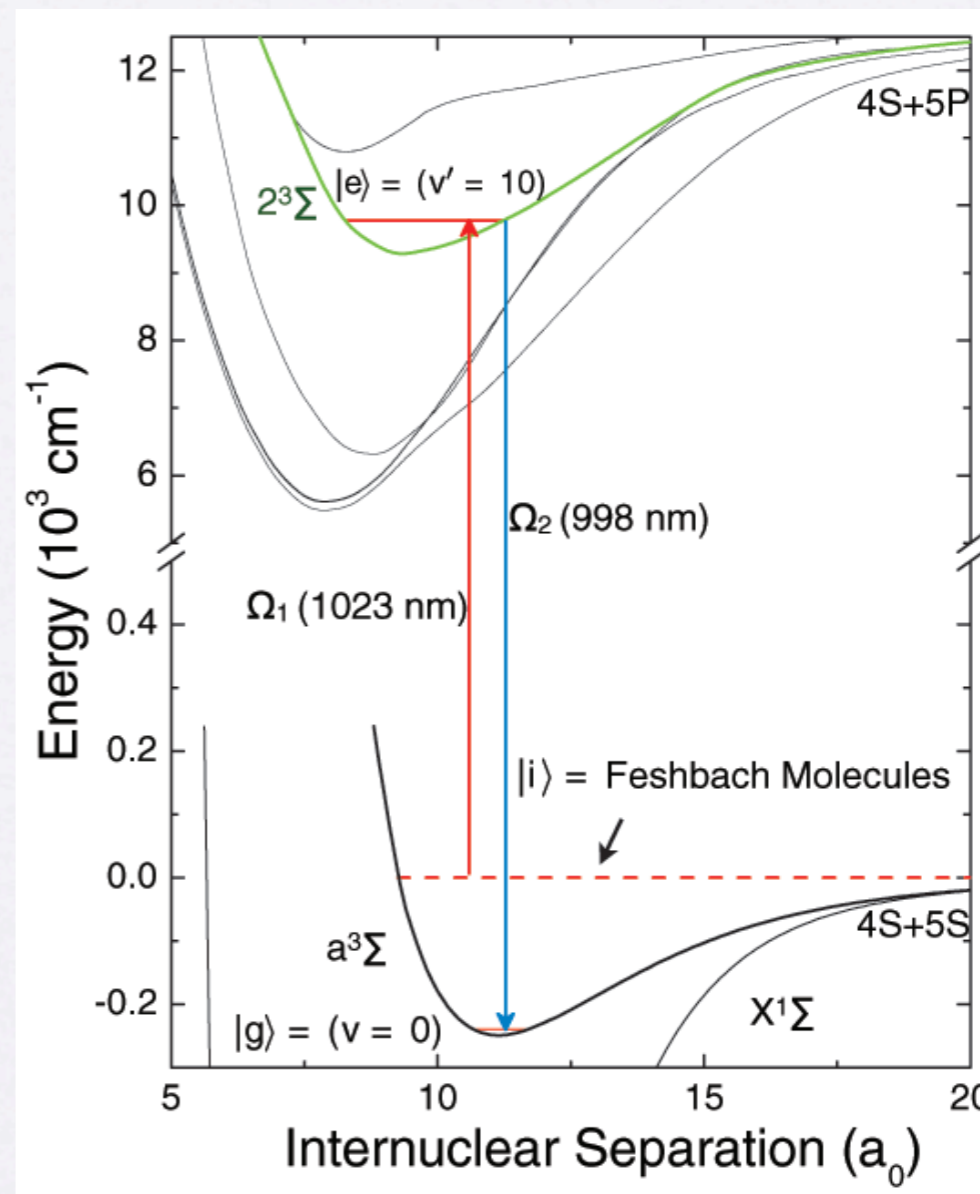
Polar molecular gases



Feshbach molecule



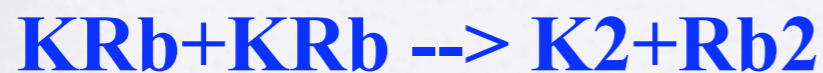
Ground state molecule



STimulated **R**aman **A**diabatic **P**assage

D. S. Jin and Jun Ye's group

Polar molecular gases



Chemically unstable !!

	Na	K	Rb	Cs
Li	-328(2)	-533.9(3)	-618(200)	-415.38(2)
Na		74.3(3)	45.5(5)	236.75(20)
K			-8.7(9)	37.81(13)
Rb				29.1(1.5)

Zuchowski and Hutson, PRA (2010)

$$E_D \sim \frac{D}{\langle r \rangle^3} \sim k_F^3 D \quad E_F \sim k_F^2$$

$$k_F D \sim 1$$

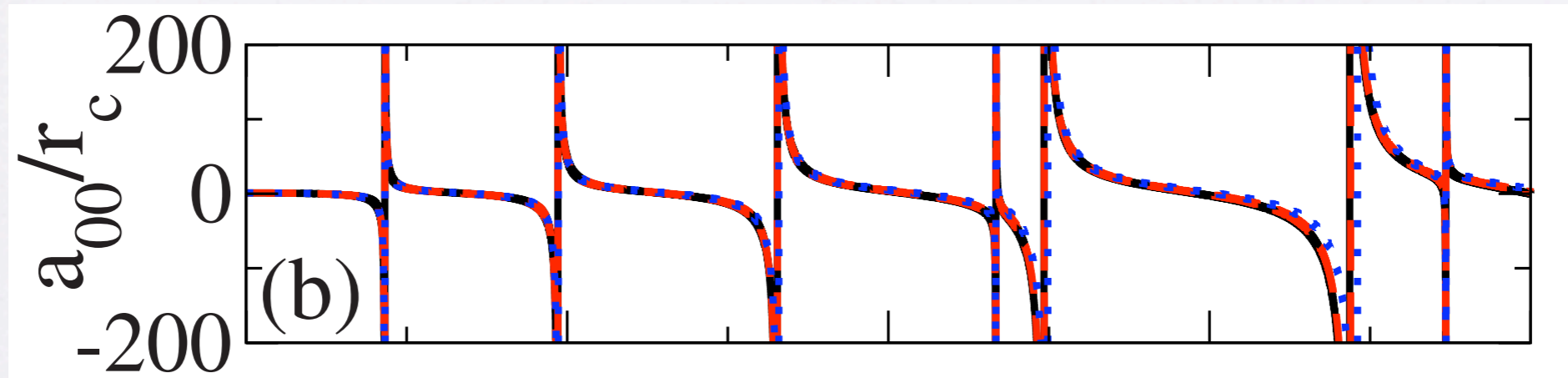
$$E_D \sim E_F$$

Dipole moments of vibrational ground state: $\nu=0$

Mixture	[Debye]
Li-Na	0.56
Li-K	3.6
Li-Rb	4.2
Li-Cs	5.5
Na-K	2.8 ←
Na-Rb	3.3 ←
Na-Cs	4.6 ←
K-Rb	0.6
K-Cs	1.9 ←
Rb-Cs	1.2 ←

M. Aymar and O. Dulieu, J. Chem. Phys., 122, 204302 (2005)

s-wave resonances with strong dipolar interactions

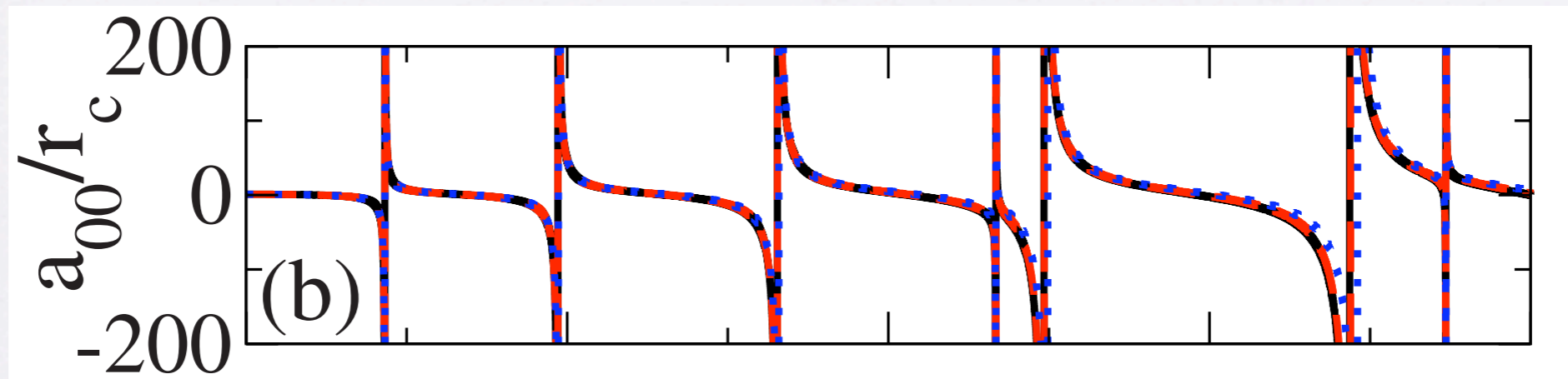


Kanjilal and Blume PRA (2008)

$$V_D = \frac{D(1 - 3 \cos^2 \theta)}{r^3}$$

$$\langle Y_{00} | V_D | Y_{00} \rangle = 0$$

s-wave resonances with strong dipolar interactions

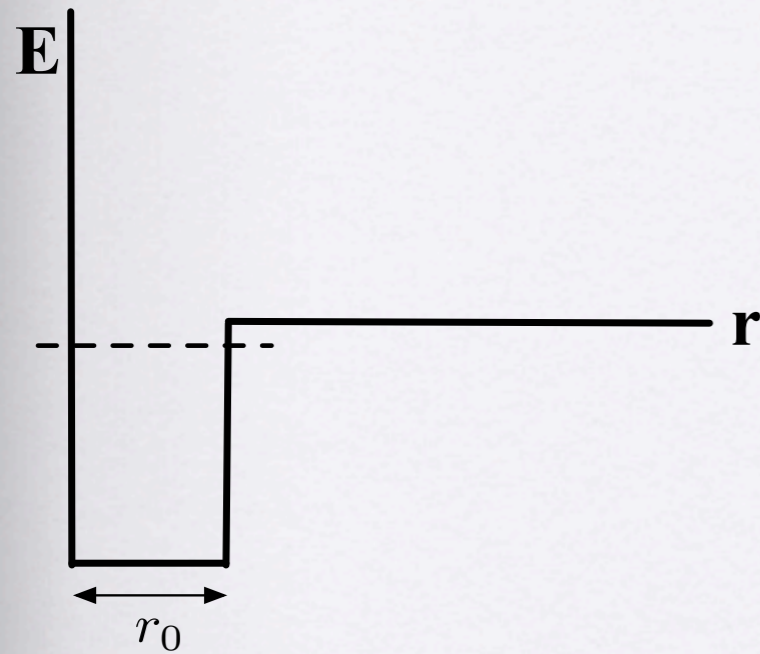


Kanjilal and Blume PRA (2008)

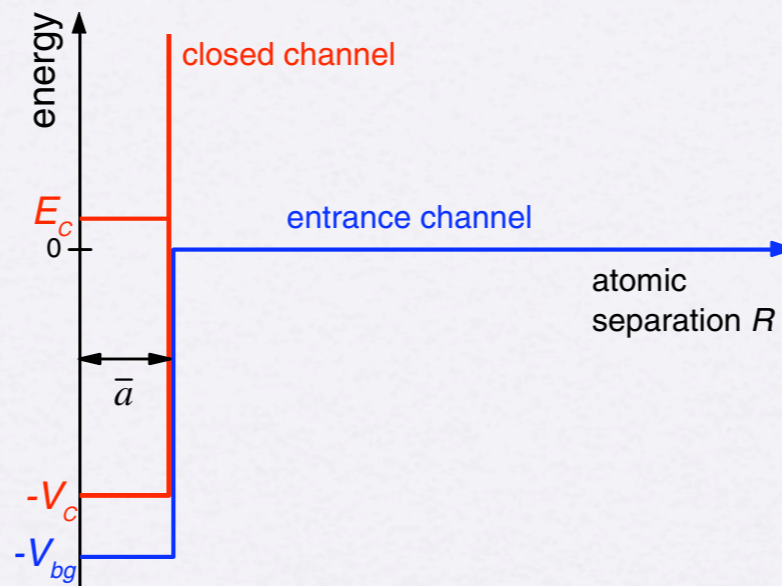
$$\langle Y_{20} | V_D | Y_{00} \rangle \neq 0$$

$$V_{\text{eff}} = - \frac{\langle Y_{00} | V_D | Y_{20} \rangle \langle Y_{20} | V_D | Y_{00} \rangle}{l(l+1)/r^2} \propto - \frac{D^2}{r^4}$$

How well can we tune a (**positive**) effective range in cold atoms ?
 -- from the wiki

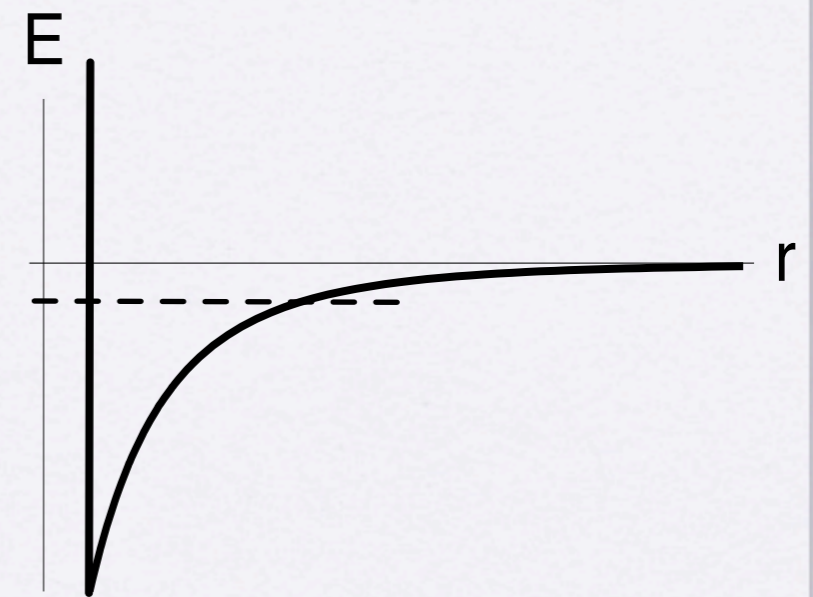


$$k_F r_{\text{eff}} \ll 1$$



$$k_F r_{\text{eff}} \gtrsim 1$$

$$r_{\text{eff}} < 0$$



$$k_F r_{\text{eff}} \sim 1$$

$$r_{\text{eff}} > 0$$

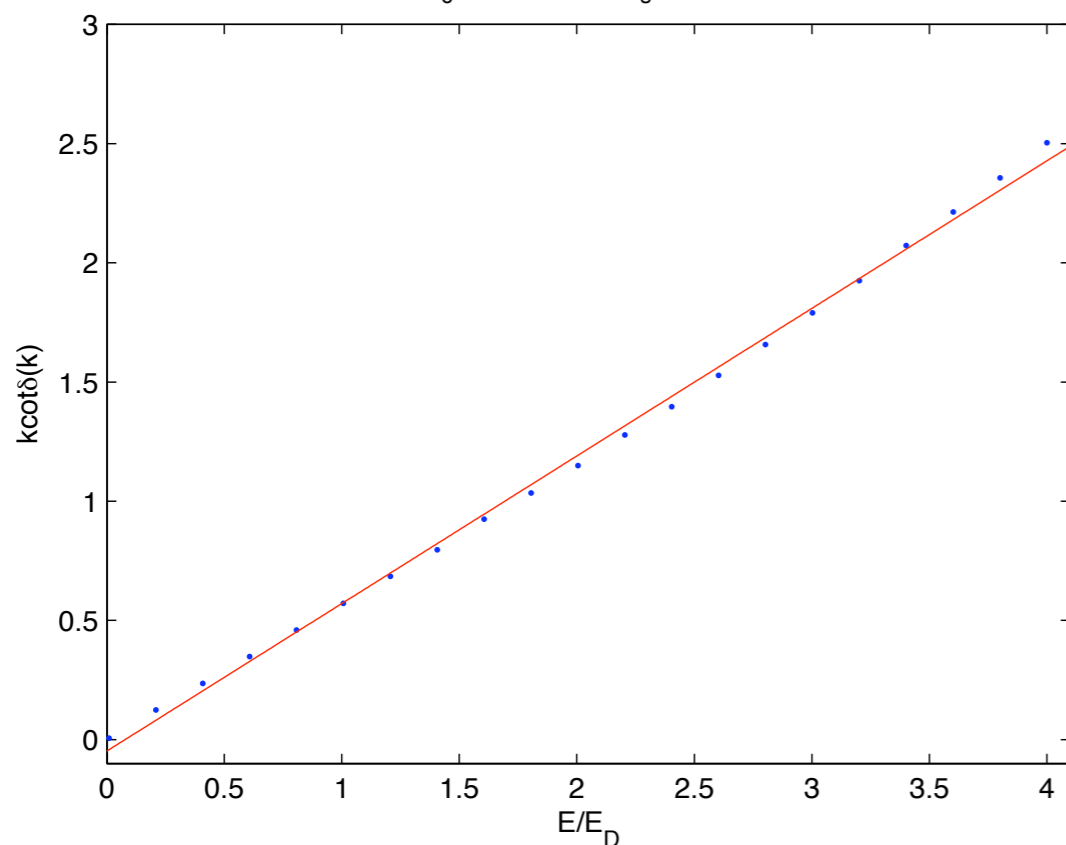
Narrow resonance !

Controlled by D, D can be tuned by electronic field

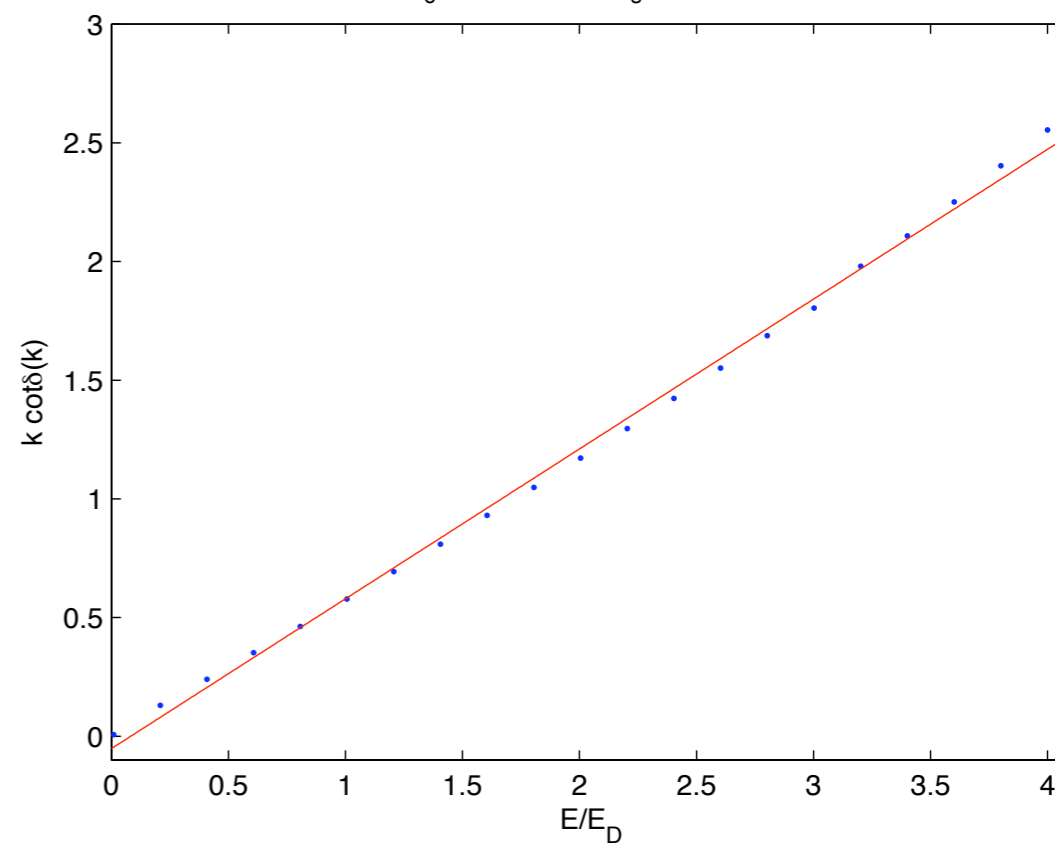
Positive effective range from a dipole induced resonance

$$\frac{k}{\tan \delta(k)} = -\frac{1}{a_s} + \frac{1}{2}r_{\text{eff}}k^2$$

$r_c=0.04803 \text{ D}, a_s=869 \text{ D}$



$r_c=0.2093 \text{ D}, a_s=-5340 \text{ D}$

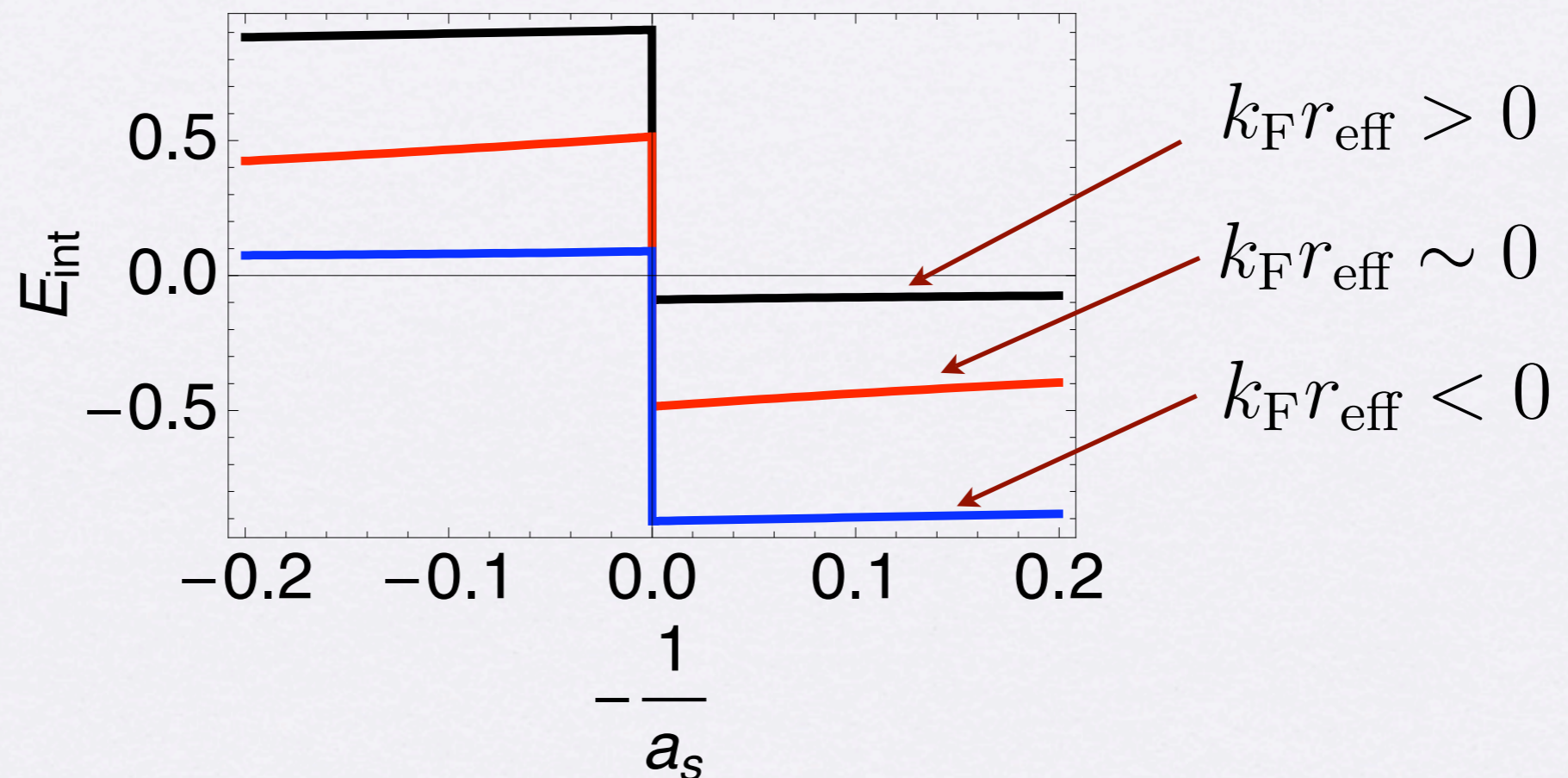


What is the physical effect of effective range being positive or negative

High temperature regime:

$$b_2 = \int_0^{+\infty} \frac{dk}{\pi} \frac{d\delta(k)}{dk} e^{-\lambda^2 k^2 / (2\pi)}$$

$$E_{\text{int}} = \frac{3k_B T n}{2} (n\lambda^3) \left[-\frac{b_2}{\sqrt{2}} + \frac{\sqrt{2}}{3} T \frac{\partial b_2}{\partial T} \right]$$



What is the physical effect of effective range being positive or negative

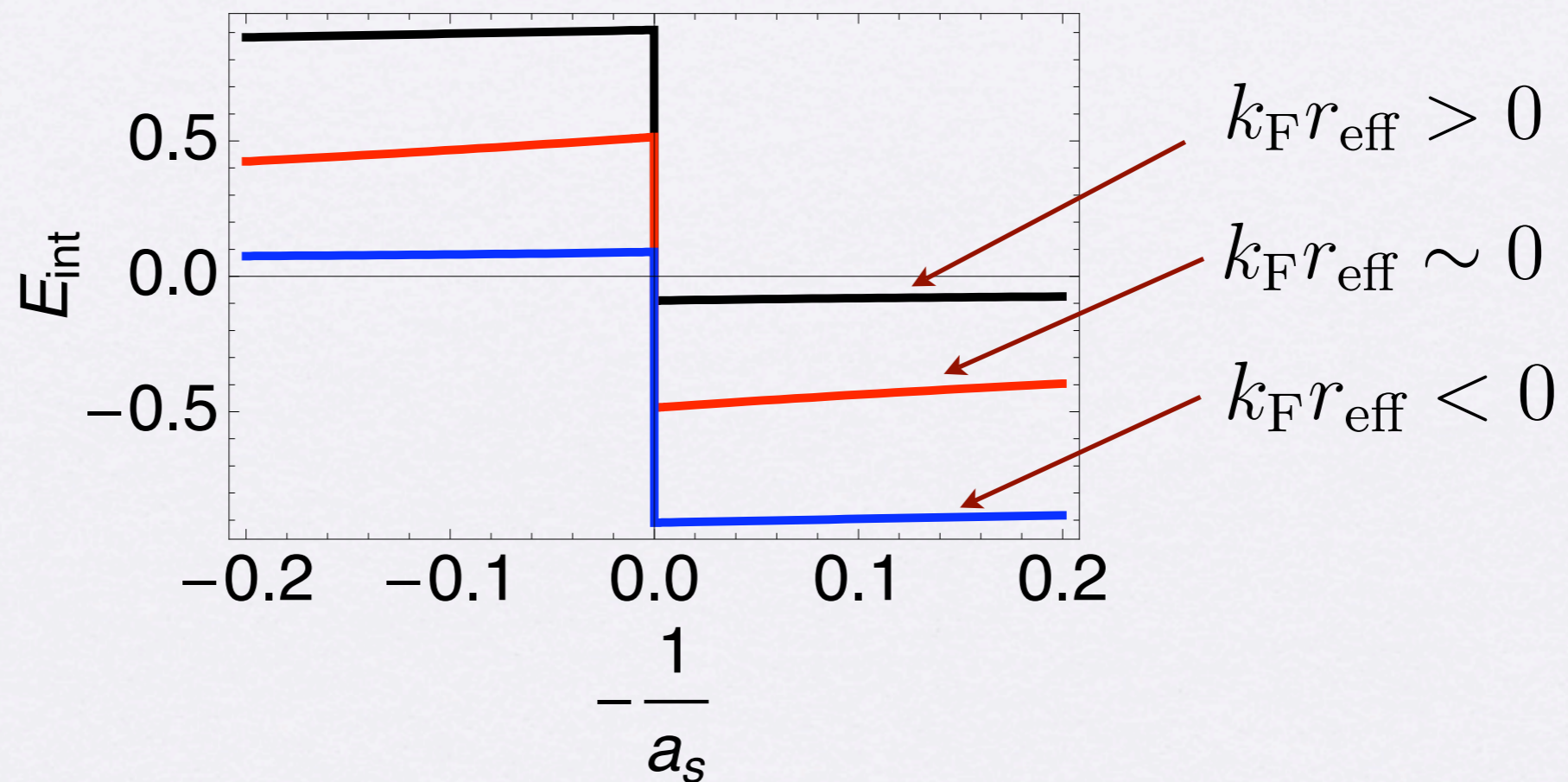
$$\frac{1}{a_{\text{avg}}} = -\frac{1}{a_s} + \frac{1}{2} r_{\text{eff}} \langle k^2 \rangle$$

$$a_s < 0 \quad r_{\text{eff}} < 0 \quad \rightarrow \quad a_{\text{avg}} < a_s$$

$$a_s < 0 \quad r_{\text{eff}} > 0 \quad \rightarrow \quad a_s < a_{\text{avg}}$$

$$a_s > 0 \quad r_{\text{eff}} < 0 \quad \rightarrow \quad a_{\text{avg}} < a_s$$

$$a_s > 0 \quad r_{\text{eff}} > 0 \quad \rightarrow \quad a_s < a_{\text{avg}}$$



**Summary:
Take Home Message**

New Mechanism	New System	New Features
Two-body interaction potential has center-of-mass dependence	Alkali-earth-(like) atomic gases: Sr, Ca, Yb	Spatially dependent local scattering length
Strong long range dipolar interactions	Polar molecules	positive and sizable effective range

Thank you very much for your attention !