

# Feshbach Resonances in Ultracold Quantum Gases

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# Outline

- Interactions in ultracold gases

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- Universal bound states

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- Diatomic Feshbach molecules

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- Universal bound states
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- Classification of Feshbach resonances
- Diatomic Feshbach molecules
- Outlook

# Interactions in ultracold gases

## Typical parameters

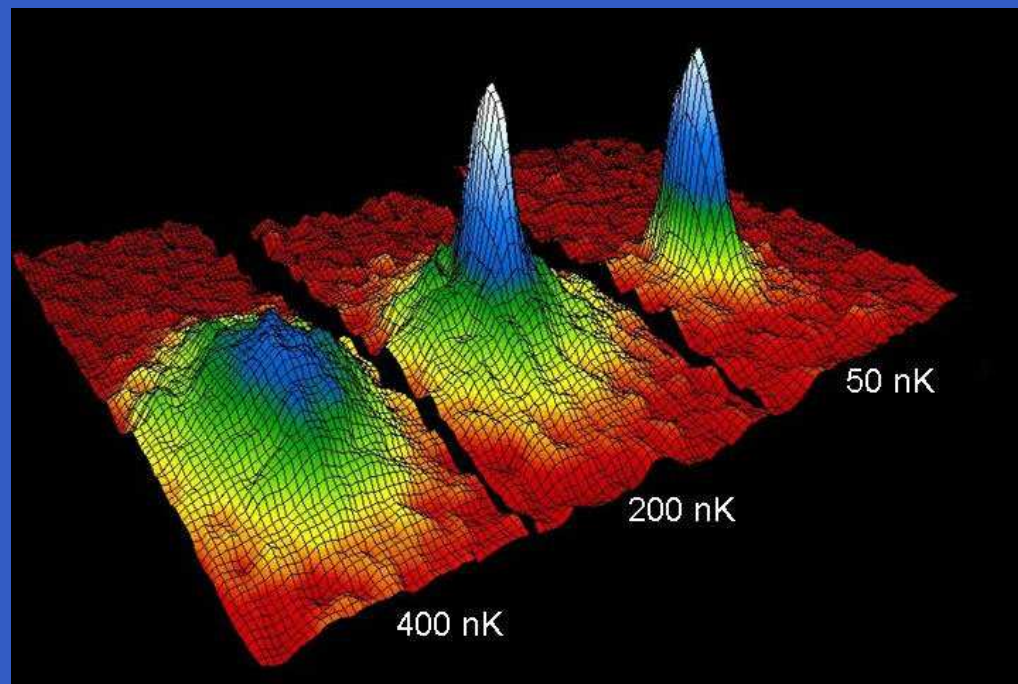
temperature 10 - 100 nK

density  $10^{13}$  -  $10^{14}$  cm $^{-3}$

number of atoms  $10^3$  -  $10^7$

size 10  $\mu$ m - 1 mm

lifetime 10 s



JILA & MIT 1995

## Condensed species

$^{87}\text{Rb}$  Na  $^7\text{Li}$  H  $^{85}\text{Rb}$   $^4\text{He}^*$   $^{41}\text{K}$  Cs Yb Cr  $^{39}\text{K}$

Quantum degenerate fermions  $^{40}\text{K}$   $^6\text{Li}$   $^3\text{He}^*$  etc.

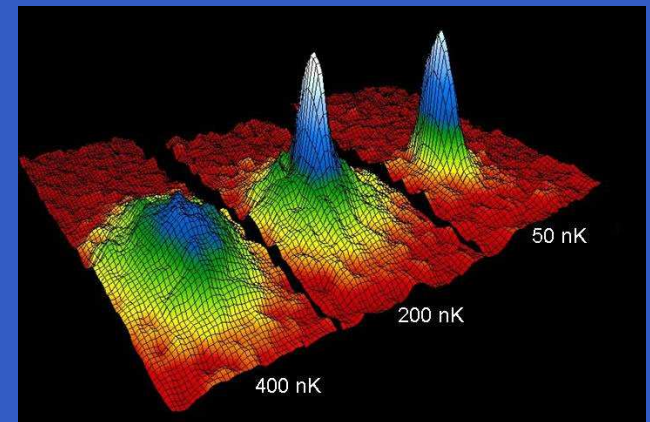


# Interactions in ultracold gases

How does one describe a Bose-Einstein condensate?

The Gross-Pitaevskii equation for the macroscopically occupied quantum state of each single atom, the **condensate wave function**:

$$i\hbar \frac{\partial}{\partial t} \Psi(\mathbf{x}, t) = \left[ -\frac{\hbar^2}{2m} \nabla^2 + V_{\text{trap}}(\mathbf{x}) \right] \Psi(\mathbf{x}, t) + \frac{4\pi\hbar^2}{m} a |\Psi(\mathbf{x}, t)|^2 \Psi(\mathbf{x}, t)$$



E.P. Gross, Nuovo Cimento **20**, 454 (1961)

L.P. Pitaevskii, Sov. Phys. JETP **13**, 451 (1961)

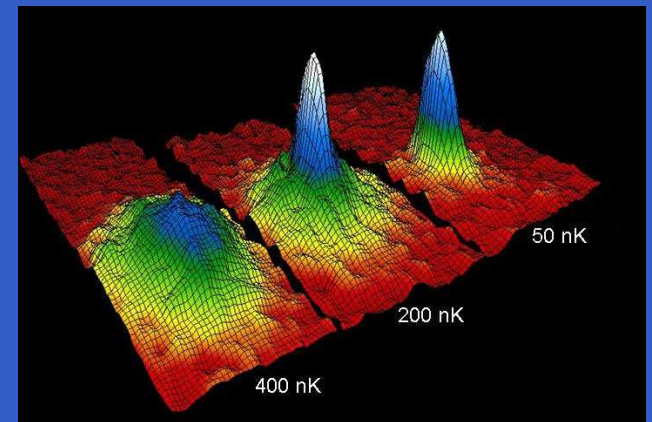
Review: F. Dalfovo, S. Giorgini, L.P. Pitaevskii, and S. Stringari, RMP **71**, 463 (1999)

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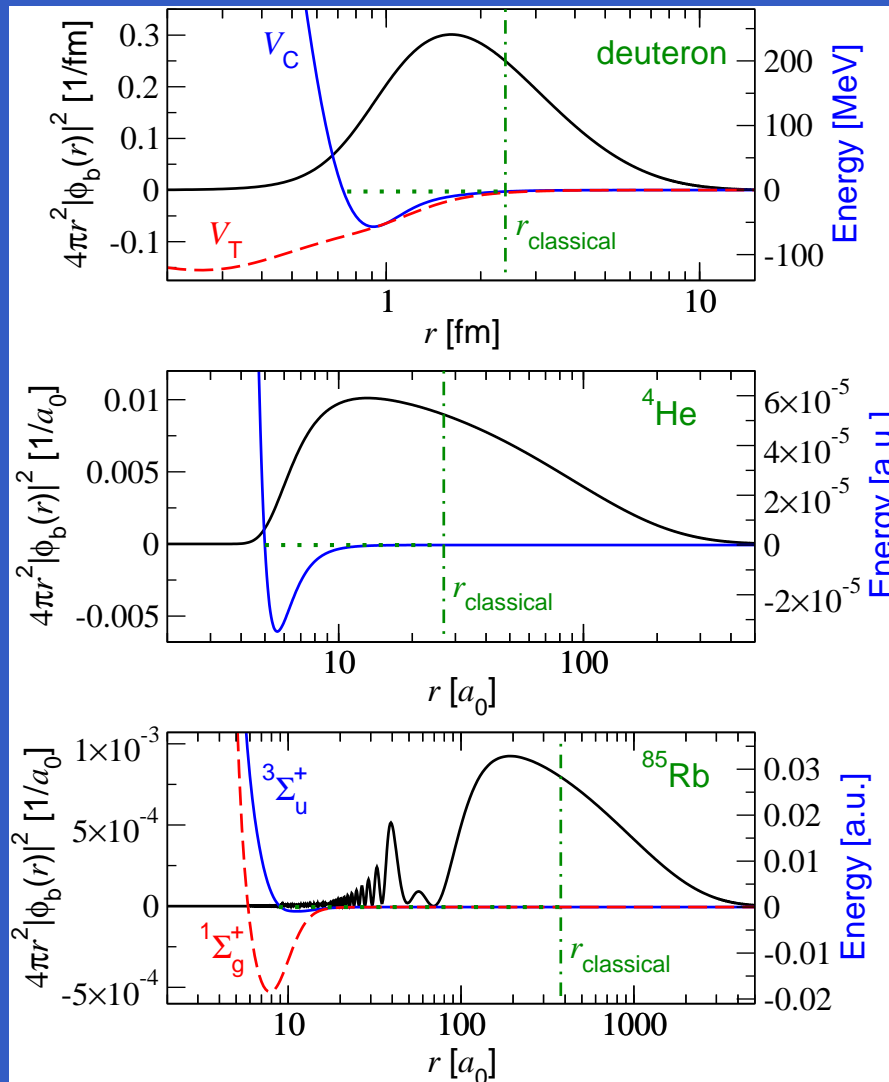
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The inter-atomic interactions are characterised solely by the ***s*-wave scattering length,  $a$ !**

# Universal bound states



The average separation between constituent particles exceeds the distance scale set by the classical motion in the potential well!

Classic examples: deuteron and  $^4\text{He}_2$

New example: Feshbach molecules

Review: A.S. Jensen, K. Riisager, D.V. Fedorov, and E. Garrido, RMP **76**, 215 (2004)

# Universal bound states

## Universal properties of diatomic halo molecules

● Hamiltonian:

$$H_{2B} = -\frac{\hbar^2}{m} \nabla^2 + V(r)$$

# Universal bound states

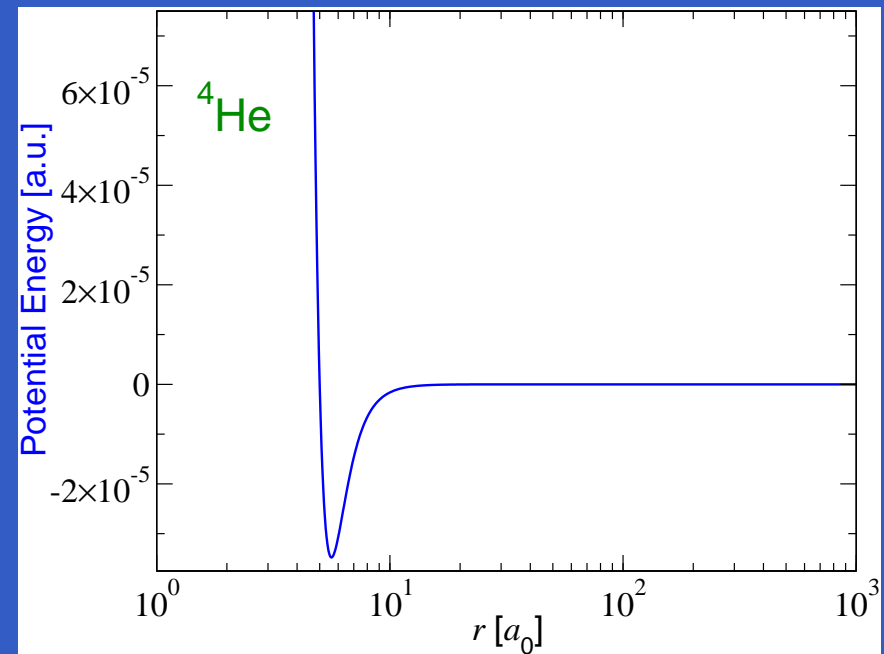
## Universal properties of diatomic halo molecules

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$$V(r) \underset{r \rightarrow \infty}{\sim} -C_6/r^6$$



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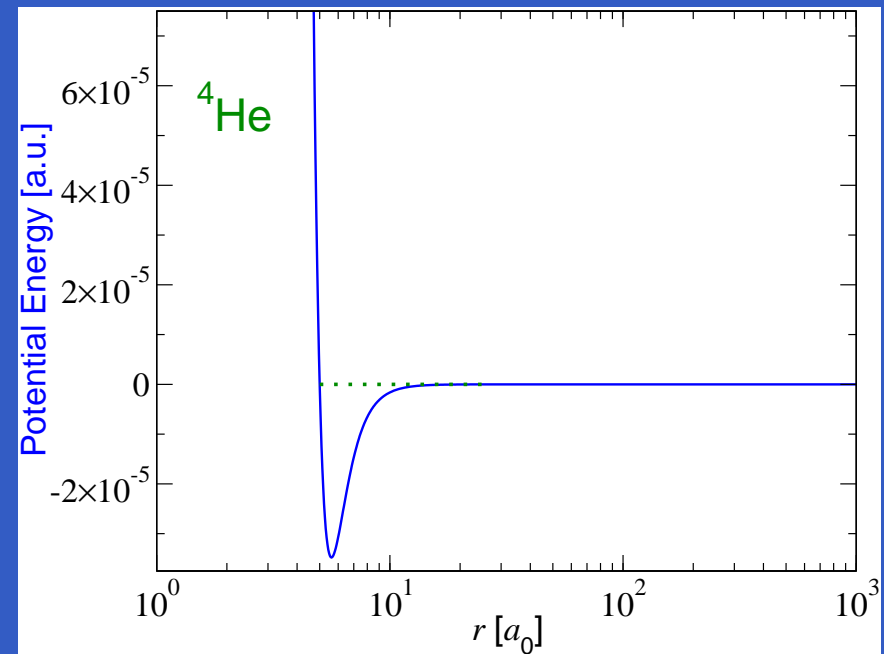
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- Bound-state energy:

$$E_b = -\hbar^2/(ma^2)$$



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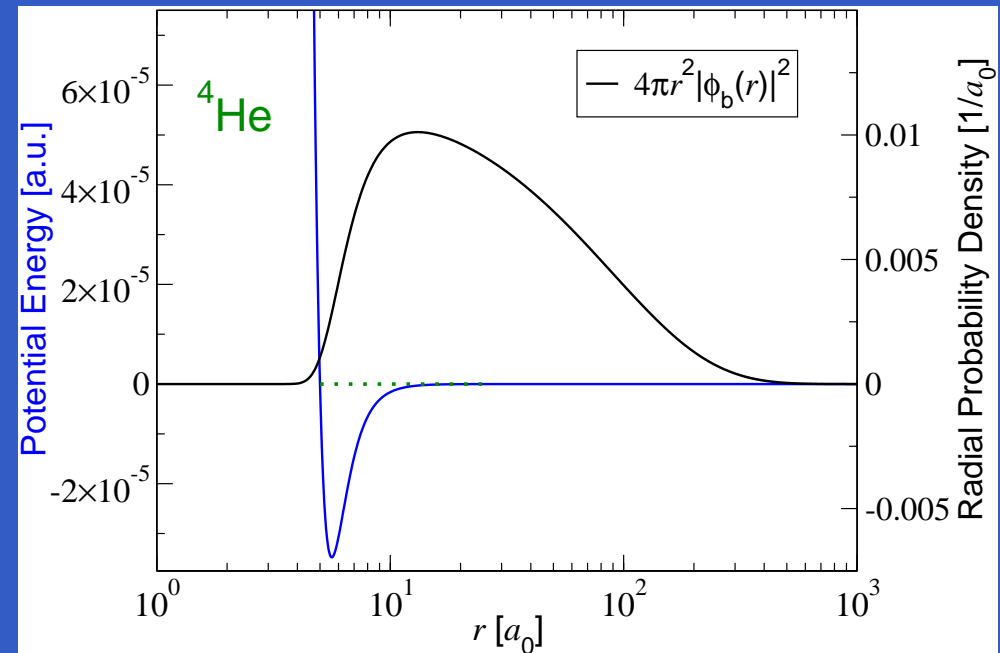
$$V(r) \underset{r \rightarrow \infty}{\sim} -C_6/r^6$$

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- Bound-state wave function:

$$\phi_b(r) = \frac{1}{\sqrt{2\pi a}} \frac{e^{-r/a}}{r}$$



# Universal bound states

**Universality: Physical properties are determined by a single length scale, the  $s$ -wave scattering length,  $a$ !**

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● van der Waals potential:

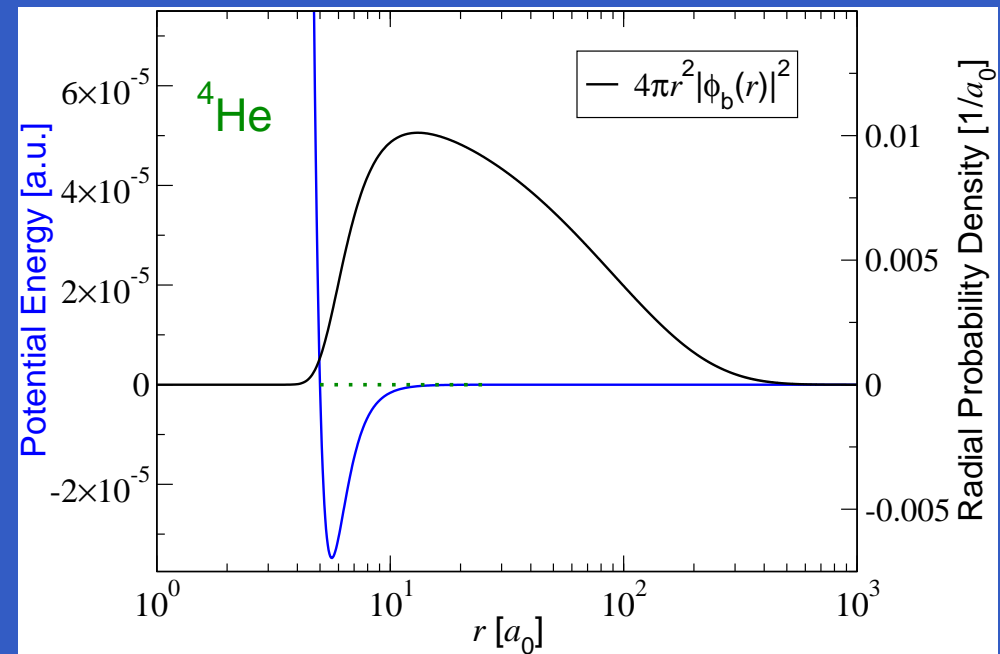
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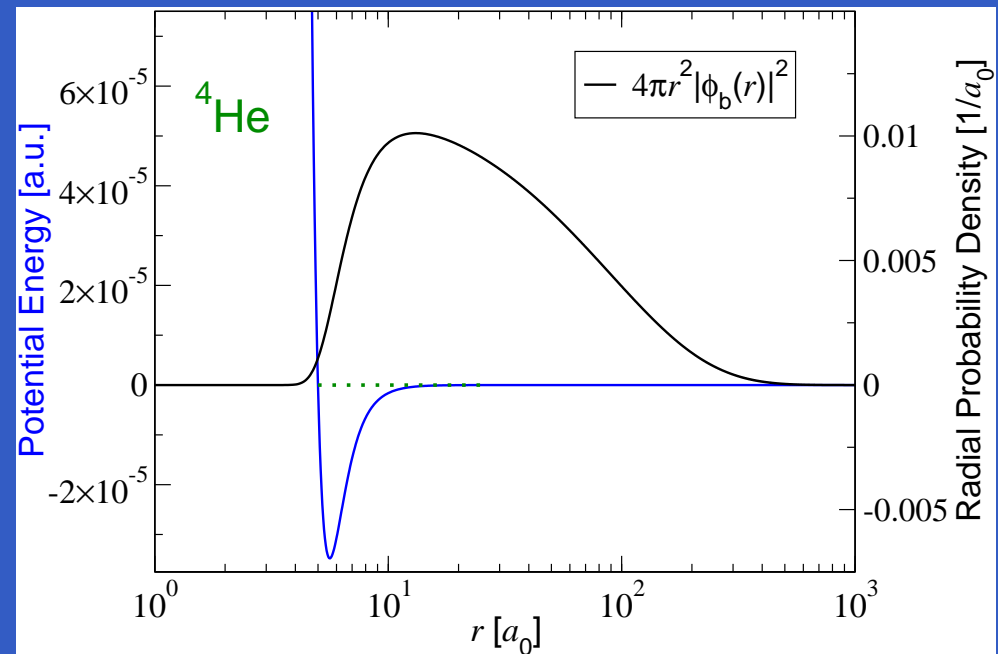


# Universal bound states

## Characteristic length scales

● Bond length:

$$\langle r \rangle = a/2$$



# Universal bound states

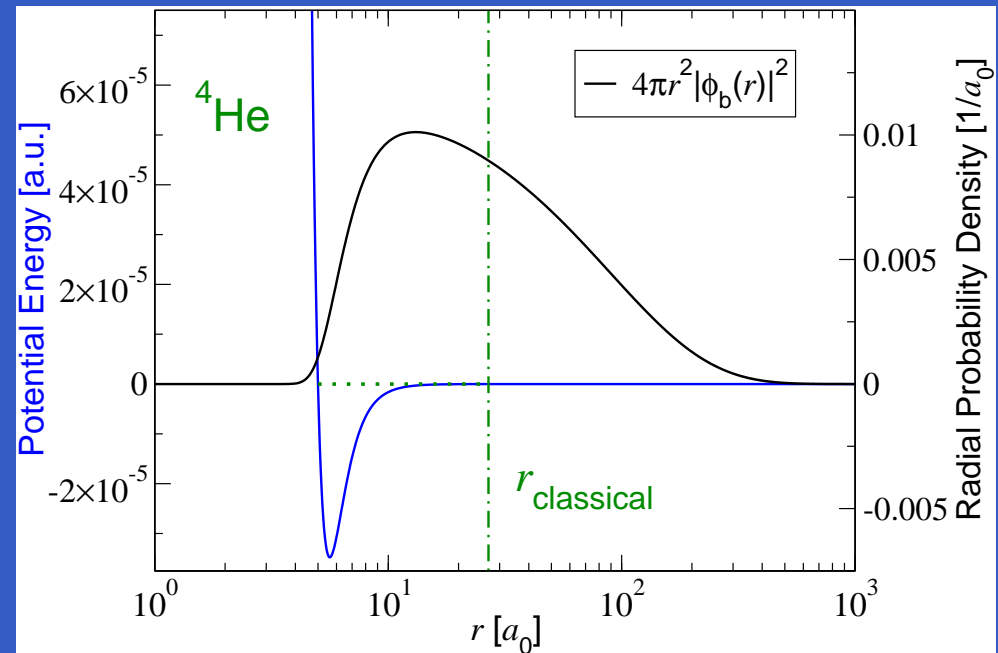
## Characteristic length scales

- Bond length:

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- Classical turning point:

$$r_{\text{classical}} = [a(2l_{\text{vdW}})^2]^{1/3}$$



# Universal bound states

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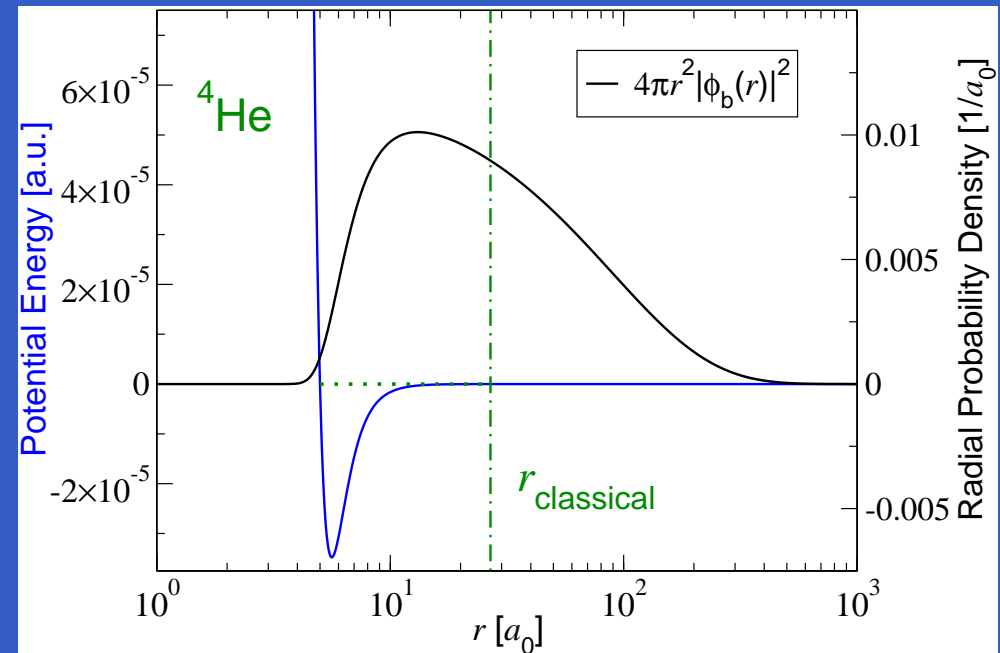
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- Classical turning point:

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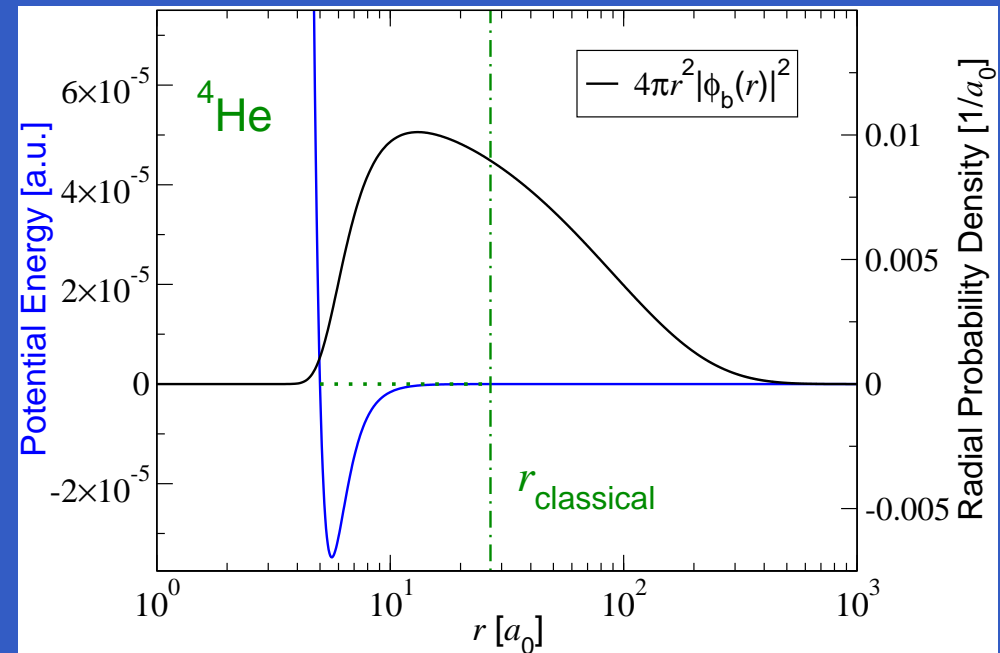
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- Characteristic property:

$$a \gg r_{\text{classical}}$$



# Universal bound states

## Properties of the helium dimer

- Bond length:

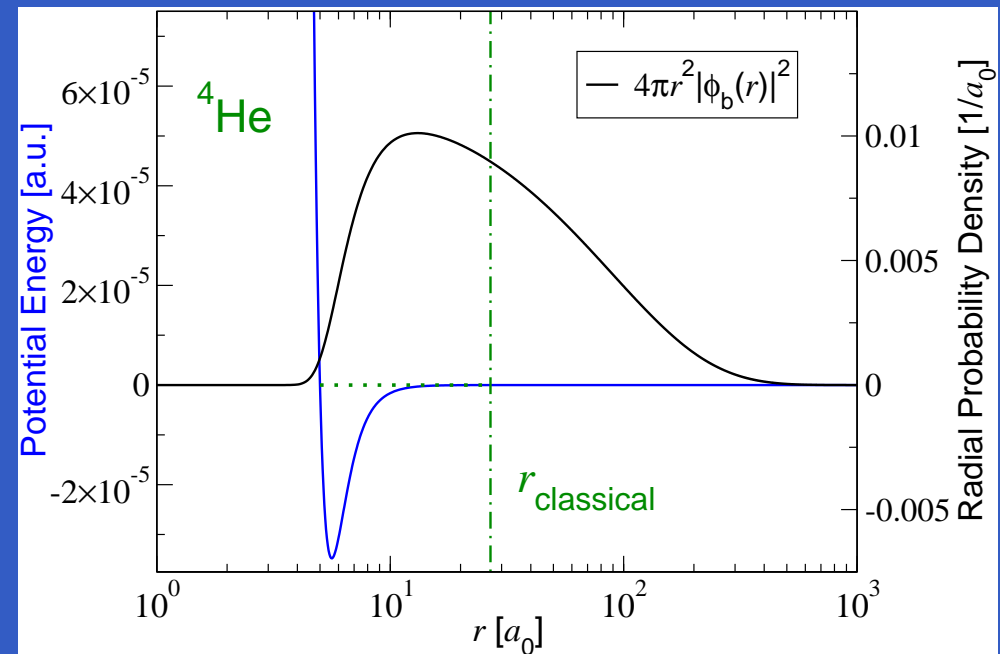
$$\langle r \rangle = 5.2 \pm 0.4 \text{ nm}$$

- Classical turning point:

$$r_{\text{classical}} = 1.4 \text{ nm}$$

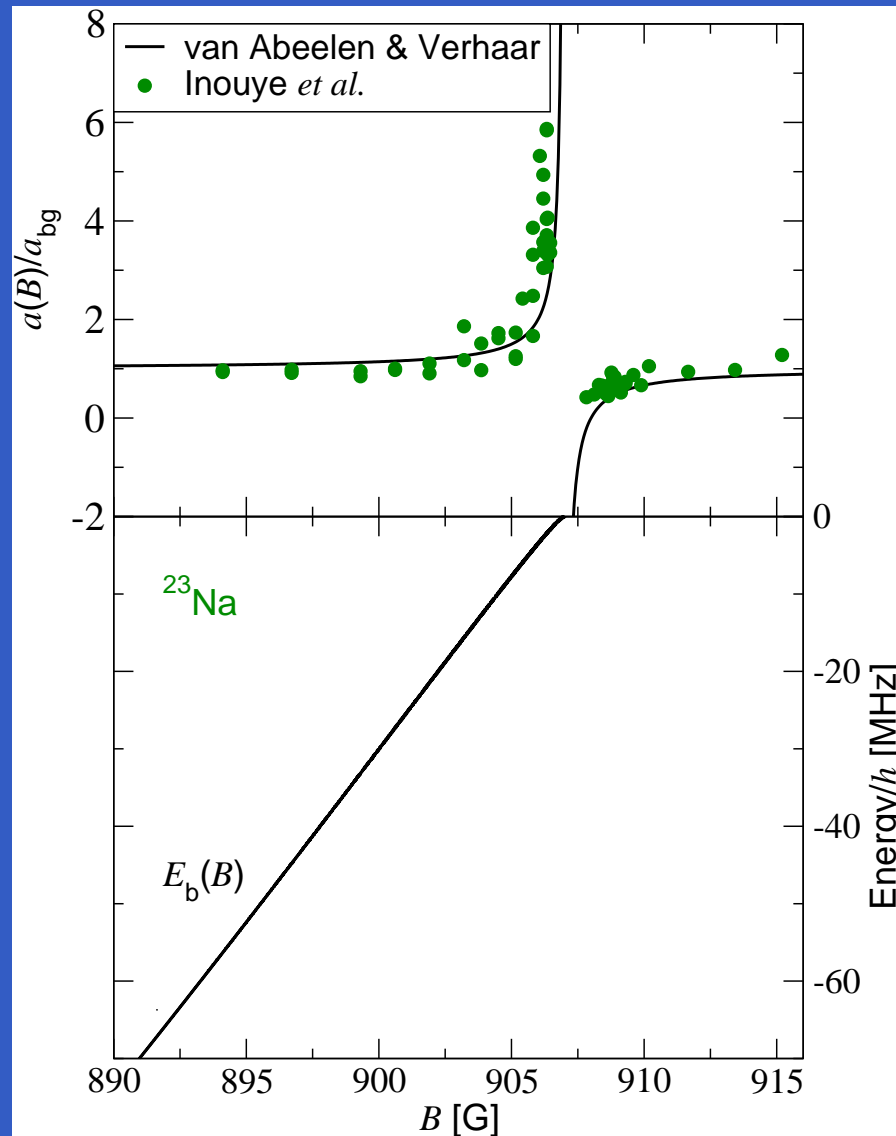
- Characteristic property:

$$a = 10.4 \text{ nm} \gg r_{\text{classical}}$$



R.E. Grisenti, W. Schöllkopf, J.P. Toennies, G.C. Hegerfeldt, TK, and M. Stoll, PRL **85**, 2284 (2000)

# Magnetically-tunable interactions



The scattering length diverges.

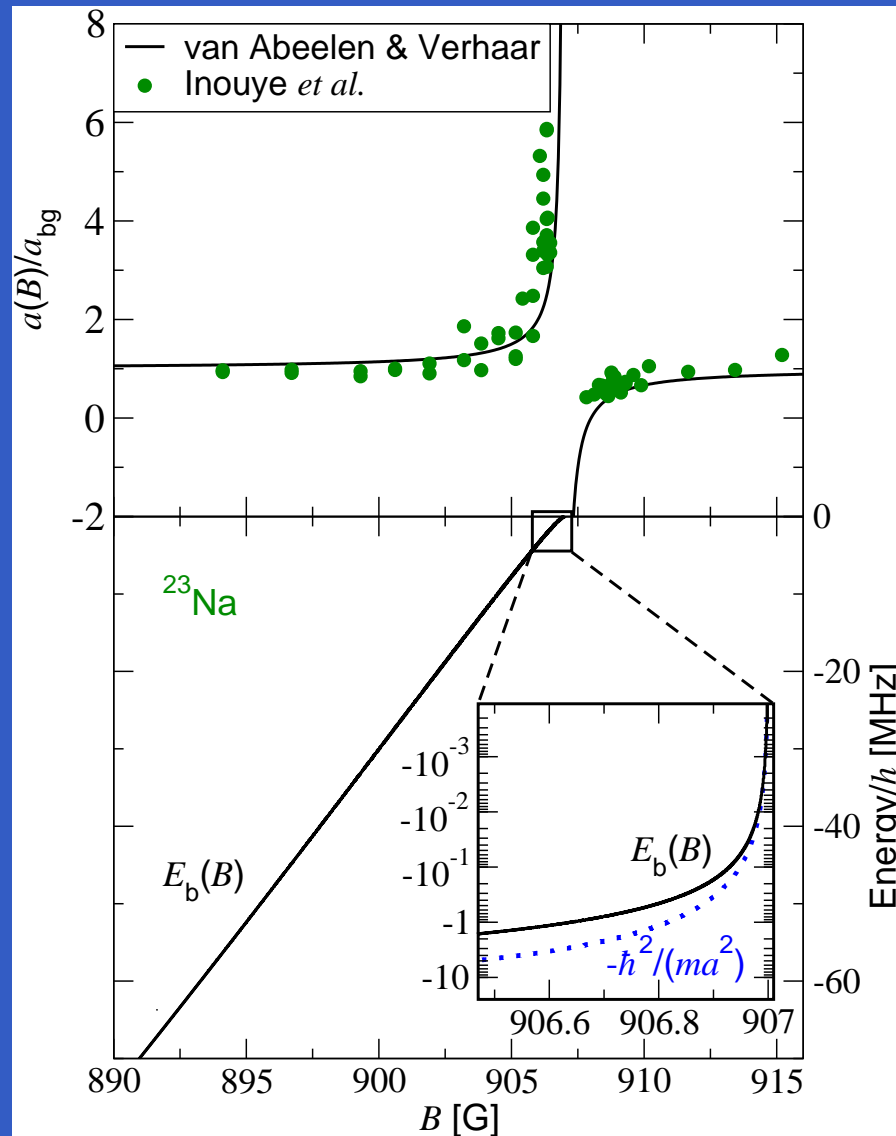


A new vibrational bound state with energy  $E_b(B)$  emerges at the collision threshold.

Theory: E. Tiesinga, B.J. Verhaar, and H.T.C. Stoof,  
PRA **47**, 4114 (1993)

Experiment: S. Inouyé *et al.*, Nature **392**, 151 (1998)

# Magnetically-tunable interactions



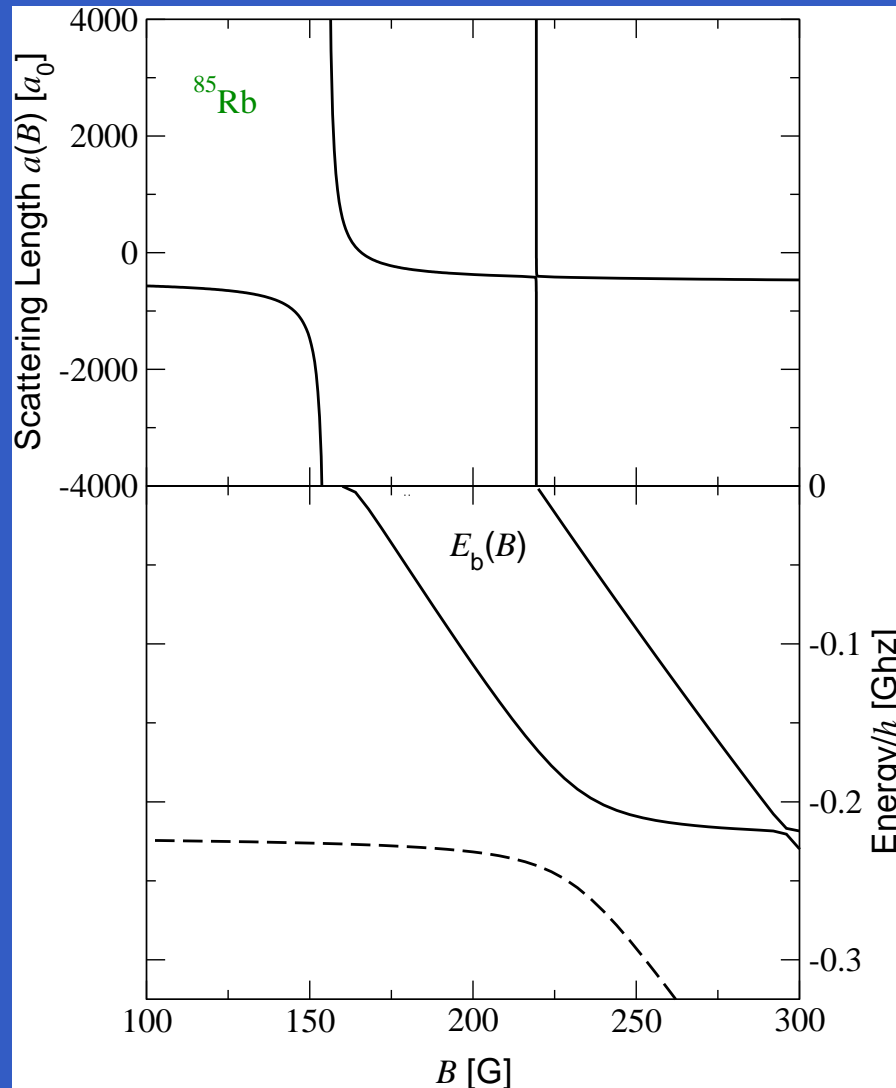
The scattering length diverges.



The energy of the Feshbach molecule,  $E_b(B)$ , becomes universal.

- Theory: E. Tiesinga, B.J. Verhaar, and H.T.C. Stoof,  
 PRA **47**, 4114 (1993)
- Experiment: S. Inouyé *et al.*, Nature **392**, 151 (1998)
- Review: TK, K. Góral, and P.S. Julienne,  
 RMP **78**, 1311 (2006)

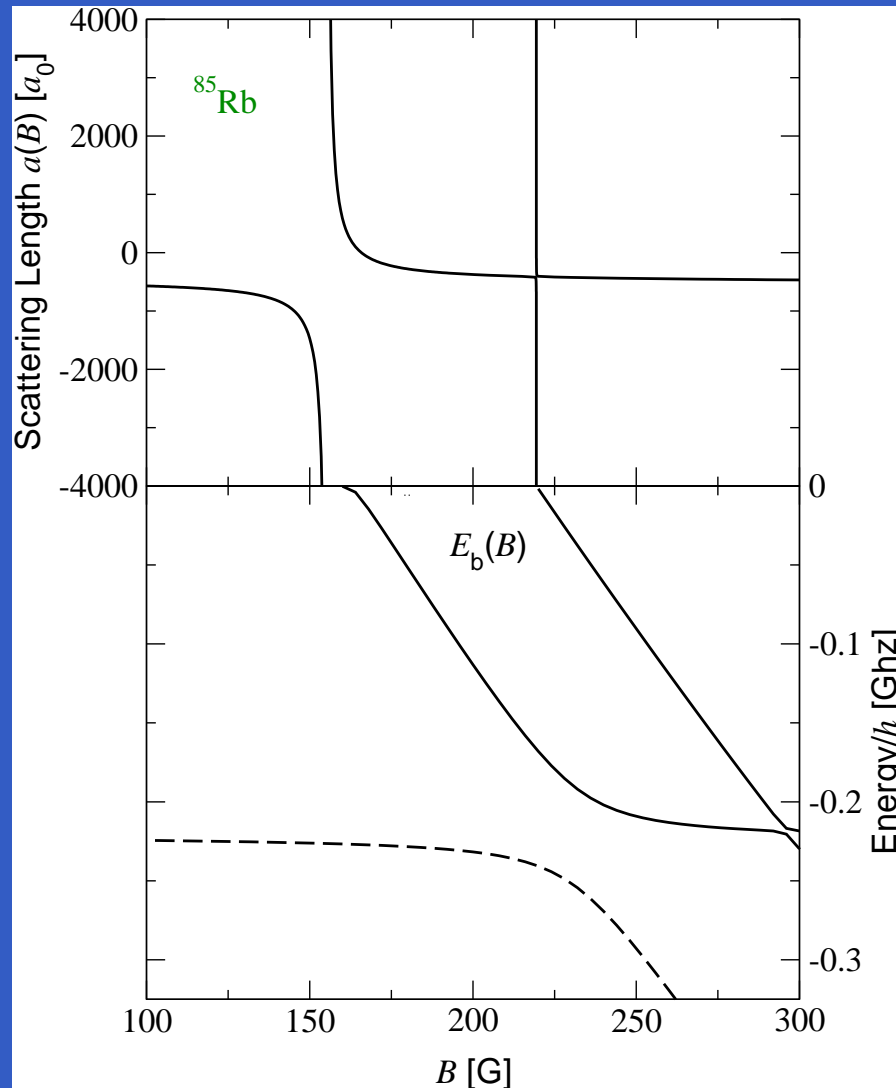
# Magnetically-tunable interactions



- The slope of  $E_b(B)$  with respect to the magnetic-field strength,  $B$ , varies between different atomic species.

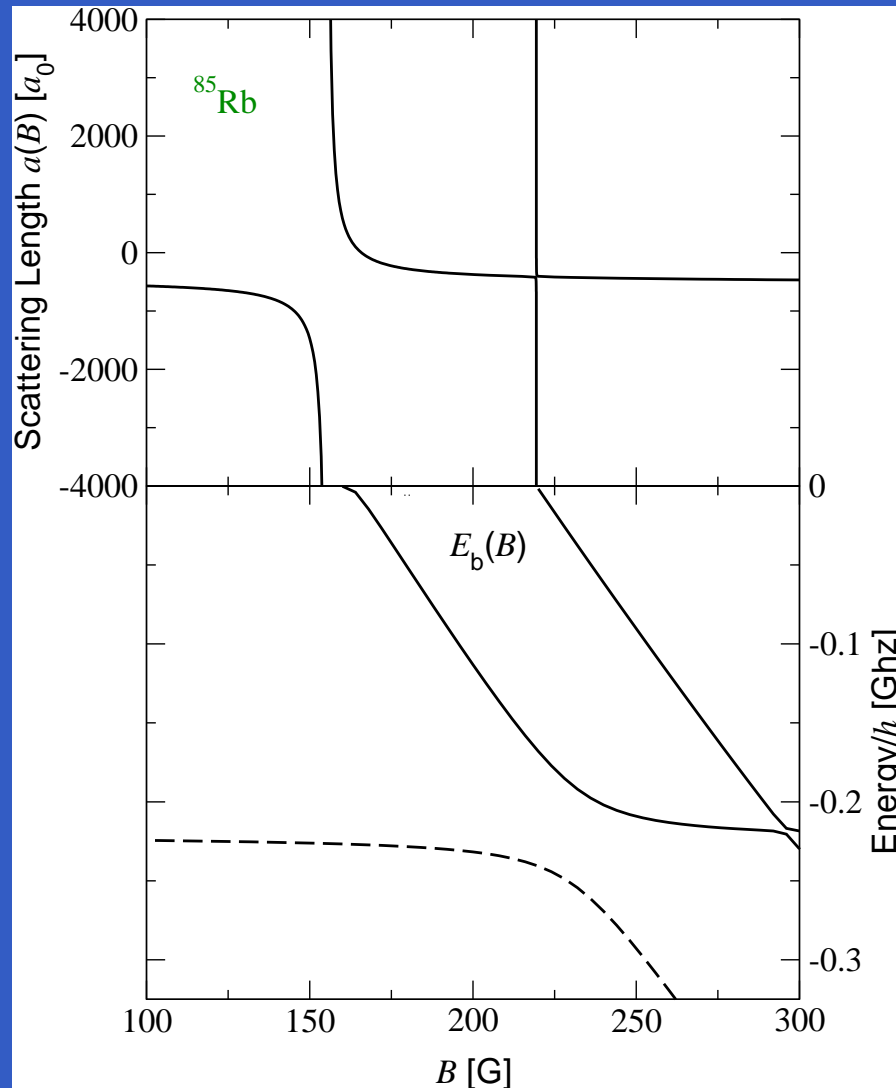


# Magnetically-tunable interactions



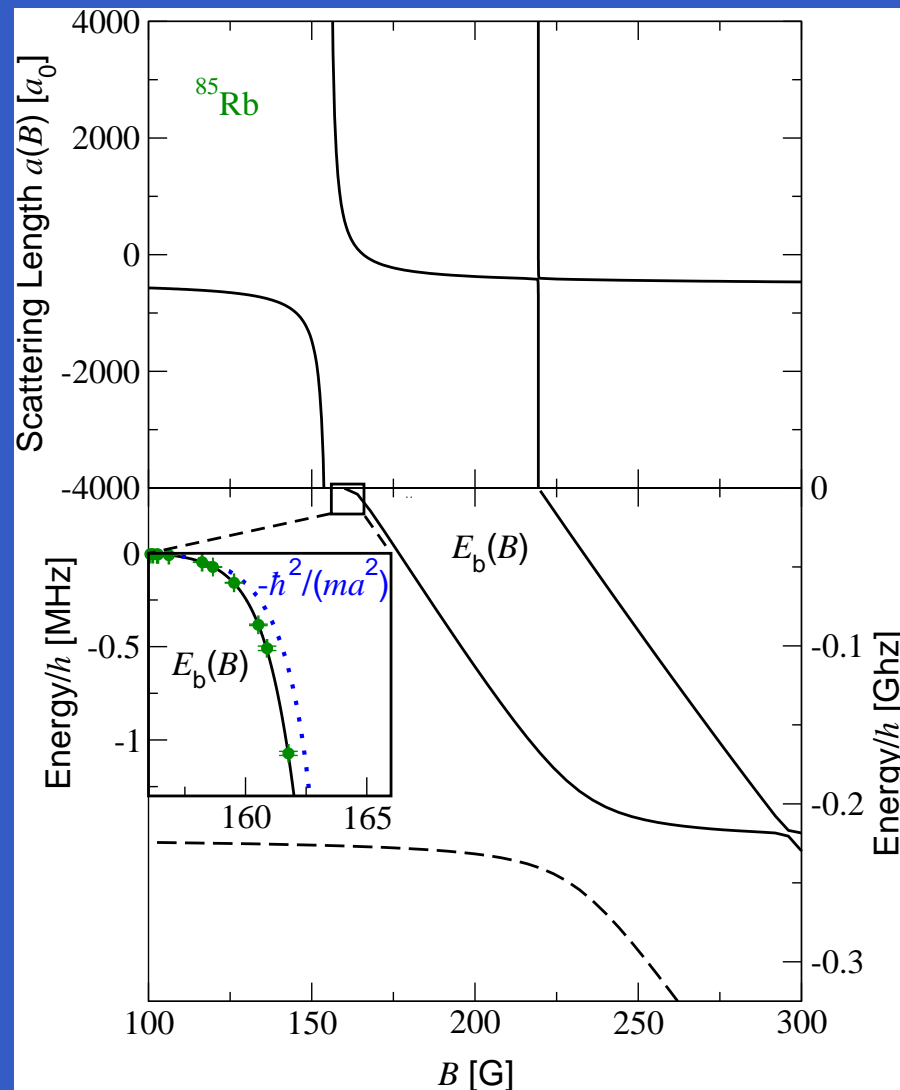
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- Singularities of the scattering length can have different widths.

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- Background-scattering lengths away from singularities can differ between atomic species.
- Singularities of the scattering length can have different widths.
- The region of universality can differ between Feshbach resonances.

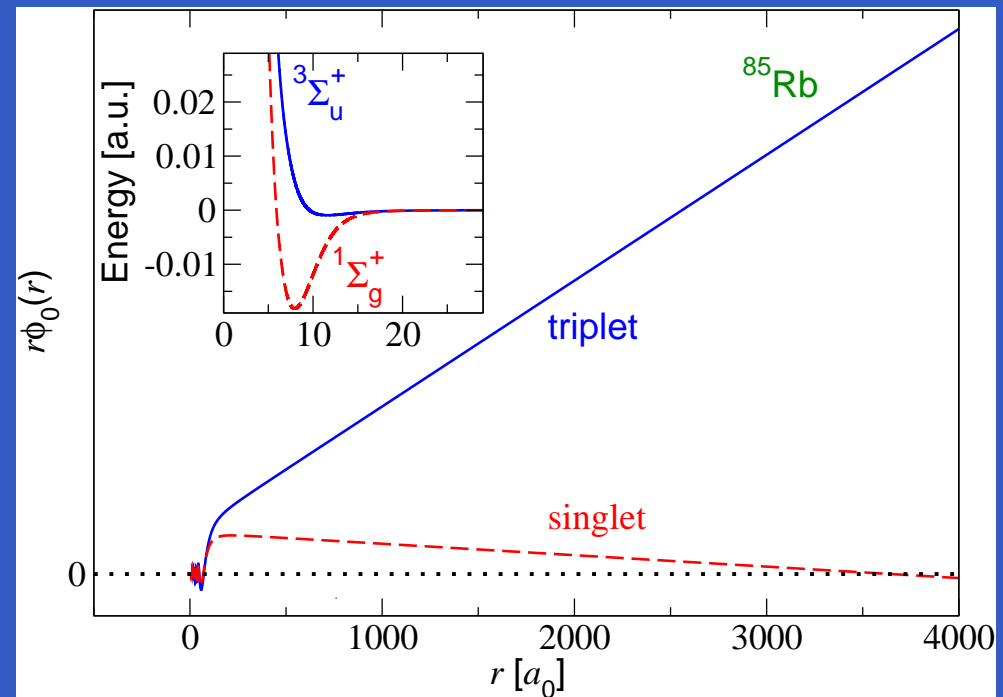
N.R. Claussen *et al.*, PRA **67**, 060701(R) (2003)

# Magnetically-tunable interactions

How does one determine a scattering length?

Radial Schrödinger equation:

$$\left[ -\frac{\hbar^2}{m} \frac{d^2}{dr^2} + V(r) \right] r\phi_0(r) = 0$$



J.R. Taylor, *Scattering Theory* (Wiley, New York, 1972)

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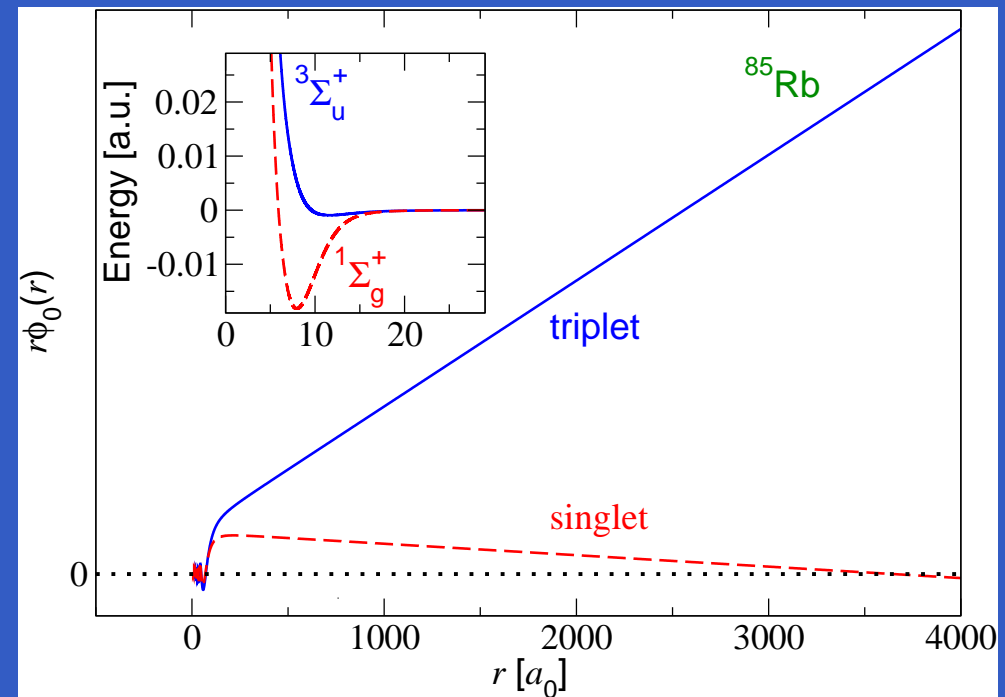
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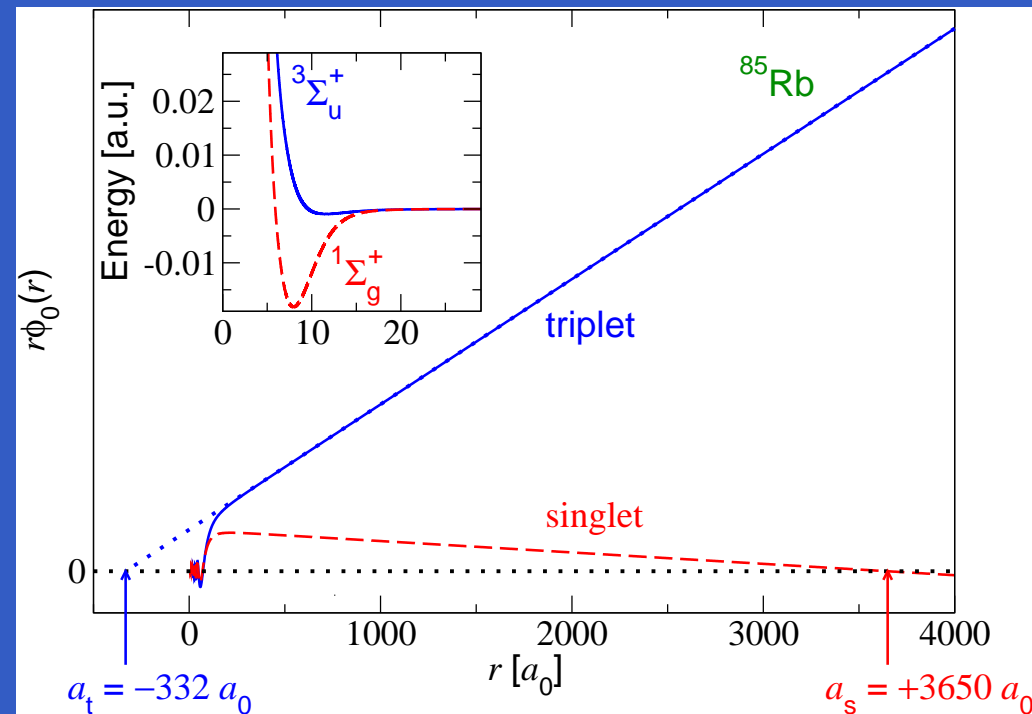
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- Asymptotic behaviour as  $r \rightarrow \infty$ :

$$r\phi_0(r) \propto r - a$$



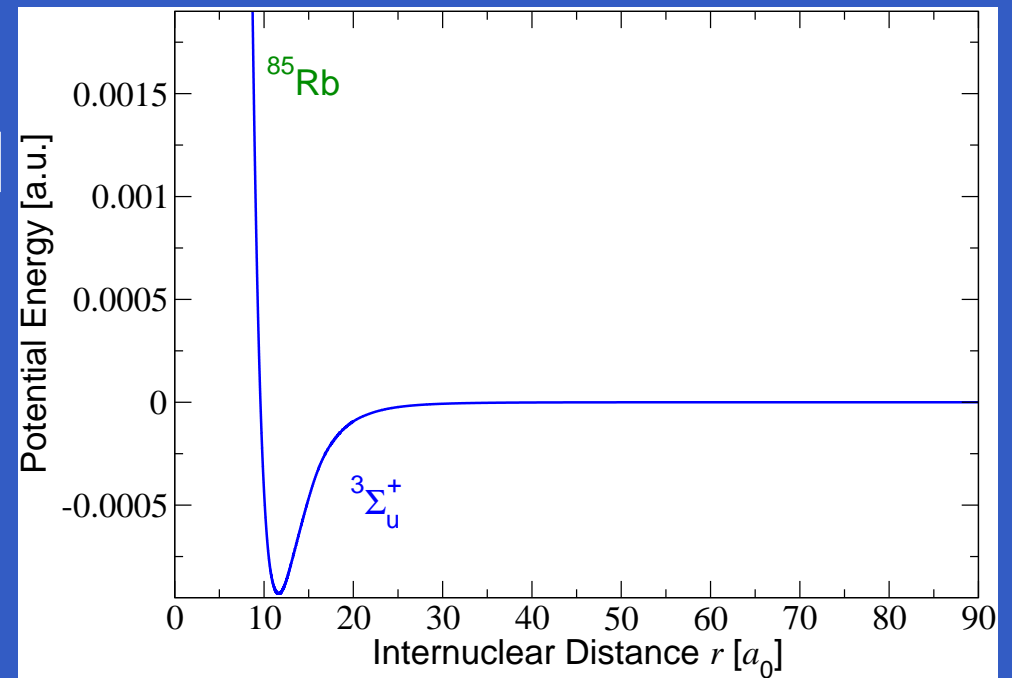
J.R. Taylor, *Scattering Theory* (Wiley, New York, 1972)

# Magnetically-tunable interactions

## Scattering length of an alkali potential

- Gibakin-Flambaum formula:

$$a = \bar{a} [1 - \tan(\varphi_{\text{WKB}} - \pi/8)]$$



G.F. Gribakin and V.V. Flambaum, PRA 48, 546 (1993)

# Magnetically-tunable interactions

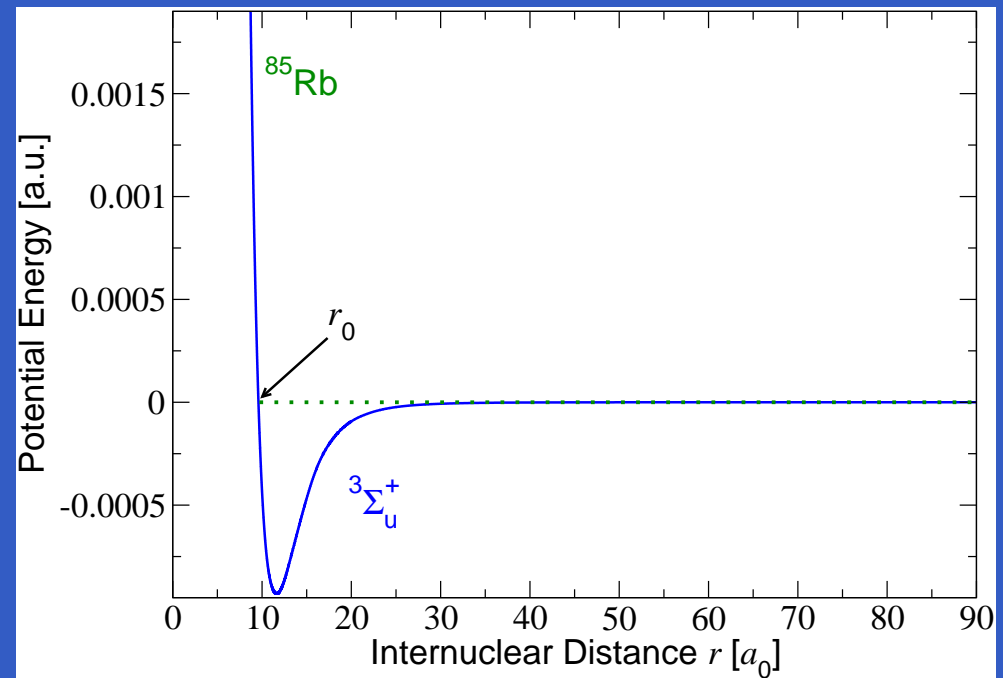
## Scattering length of an alkali potential

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- WKB phase shift:

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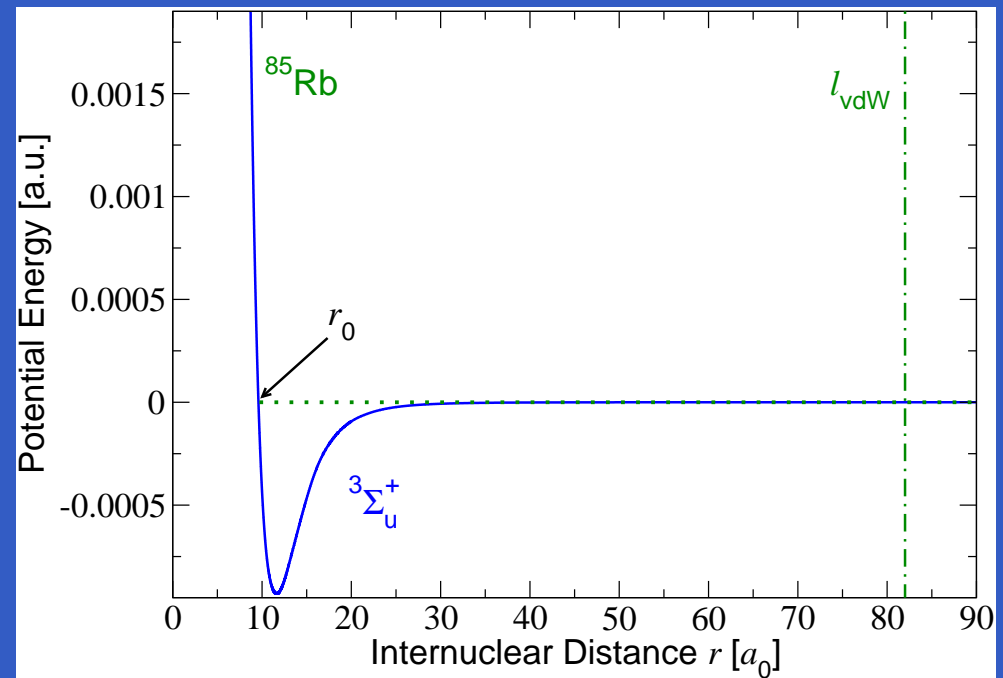
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- Mean scattering length:

$$\bar{a} = 0.95598 l_{\text{vdW}}$$

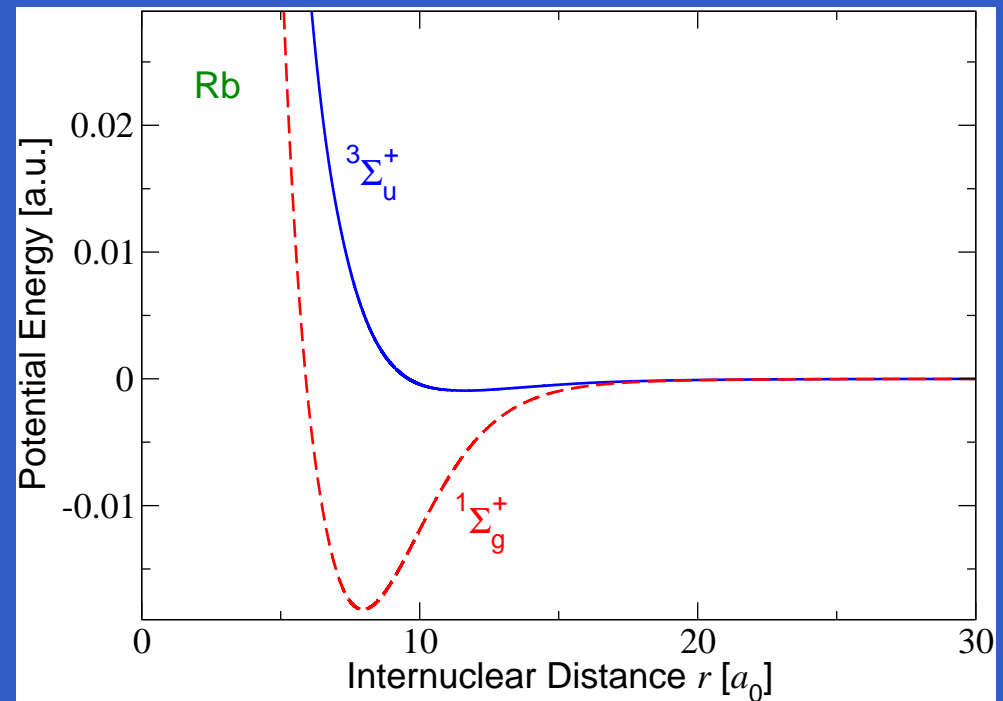


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# Magnetically-tunable interactions

## What is coupled-channels theory?

- Singlet and triplet potentials describe interactions for given total electronic spin quantum numbers of an alkali-atom pair,  $S = 0, 1$ , respectively.

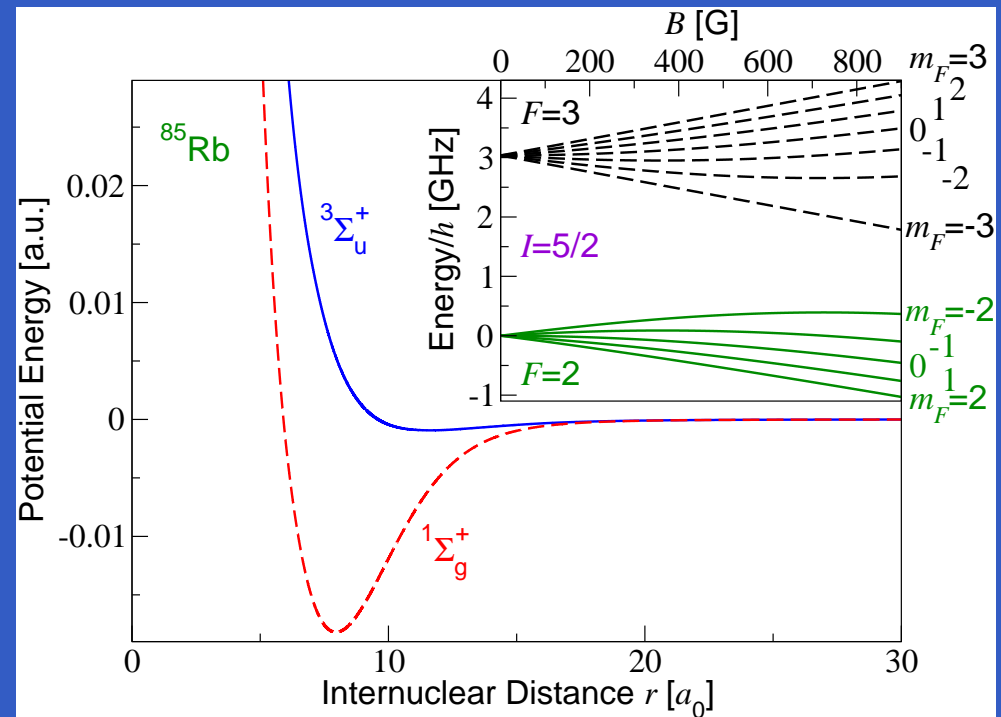


General theory: H.T.C. Stoof, J.M.V.A. Koelman, and B.J. Verhaar, PRB **38**, 4688 (1988)

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- Asymptotic scattering channels are described by pairs of Zeeman states in terms of angular momentum quantum numbers of each atom.



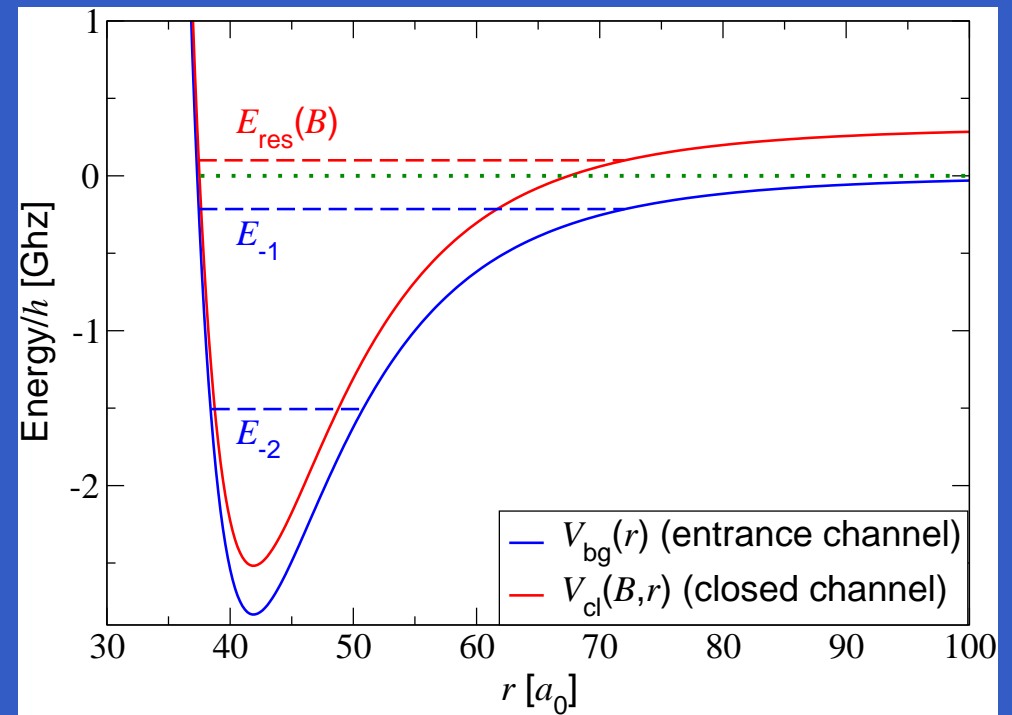
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Hyperfine structure in alkali atoms: E. Arimondo, M. Inguscio, and P. Violino, RMP **49**, 31 (1977)

# Magnetically-tunable interactions

## Feshbach resonances

- Two-channel Hamiltonian:

$$H_{2B} = \begin{pmatrix} H_{bg} & \\ & H_{cl}(B) \end{pmatrix}$$



A.J. Moerdijk, B.J. Verhaar, and A. Axelsson, PRA **51**, 4852 (1995)  
F.H. Mies, E. Tiesinga, and P.S. Julienne, PRA **61**, 022721 (2000)

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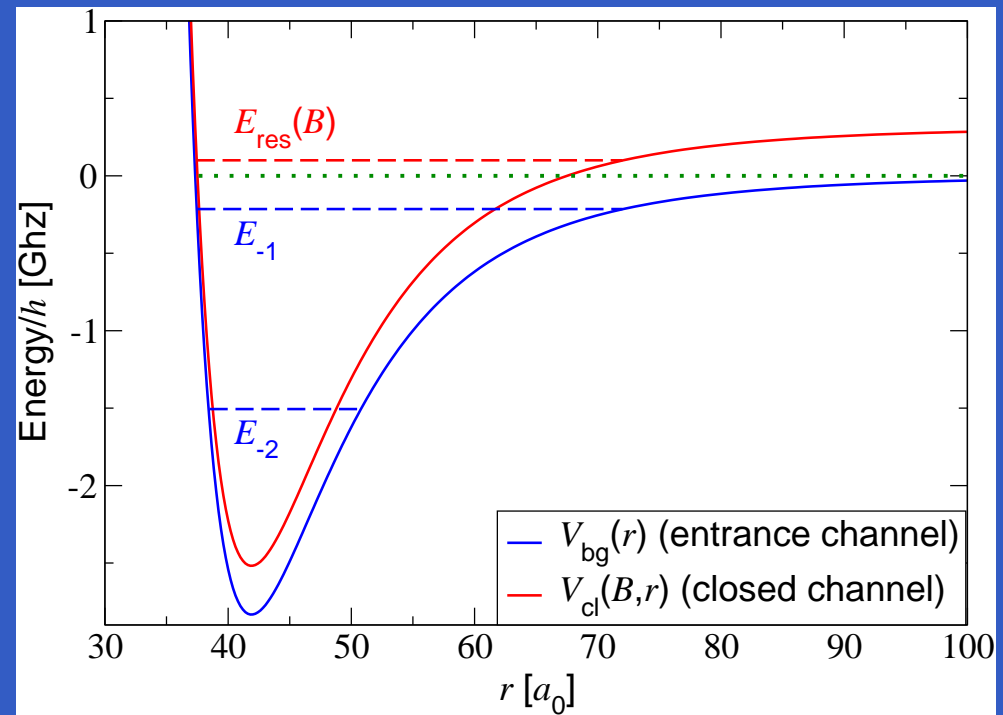
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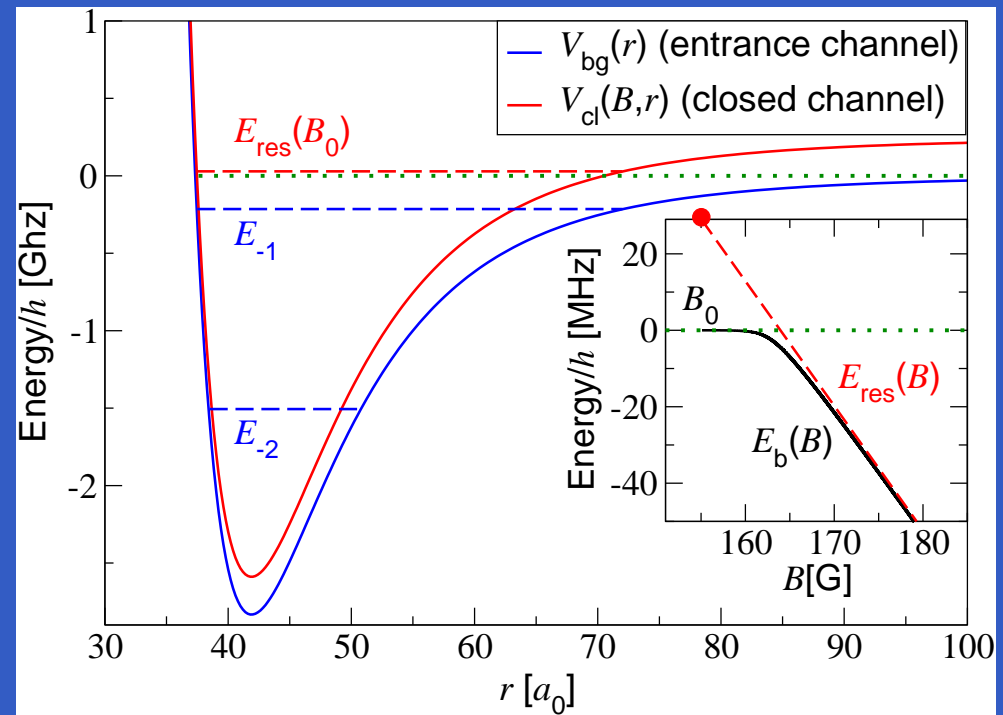
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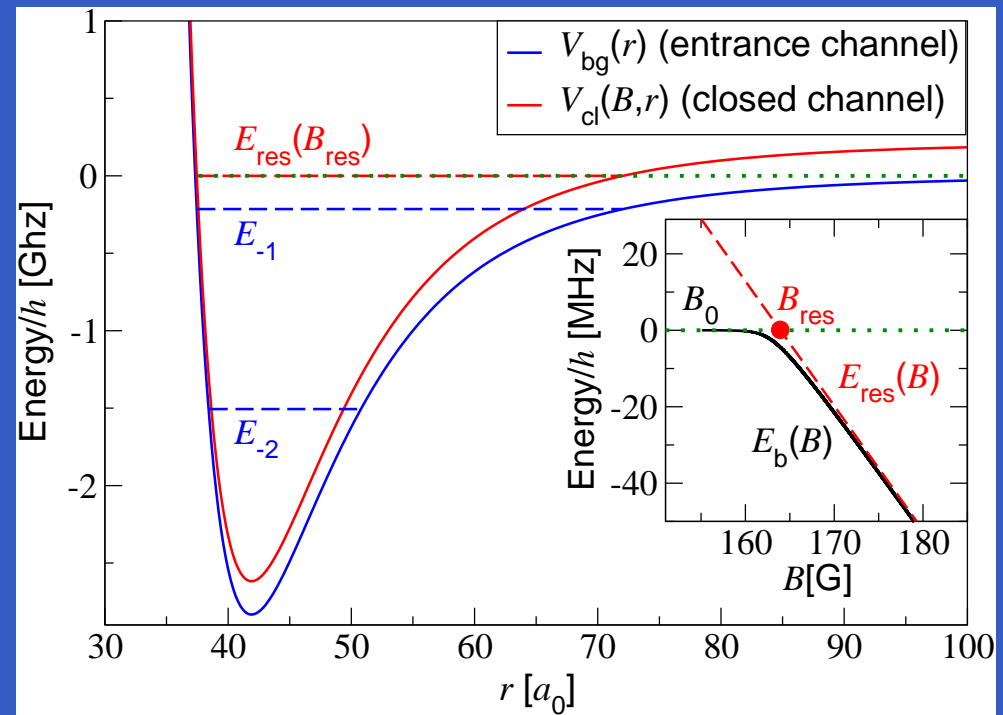
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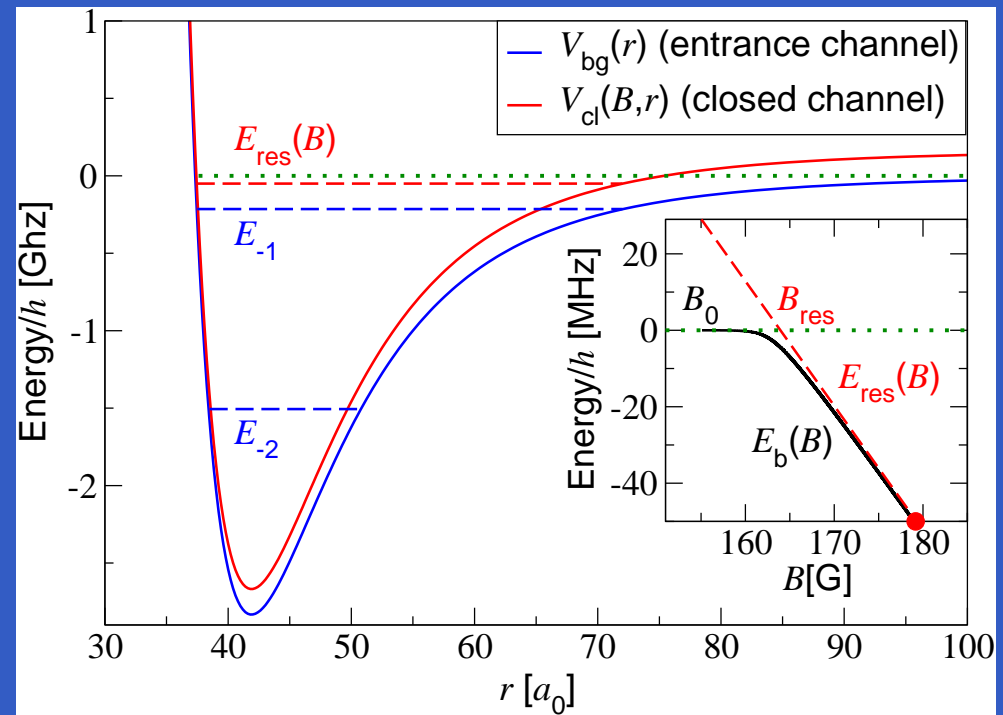
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# Magnetically-tunable interactions

## Two-channel versus coupled-channels theory

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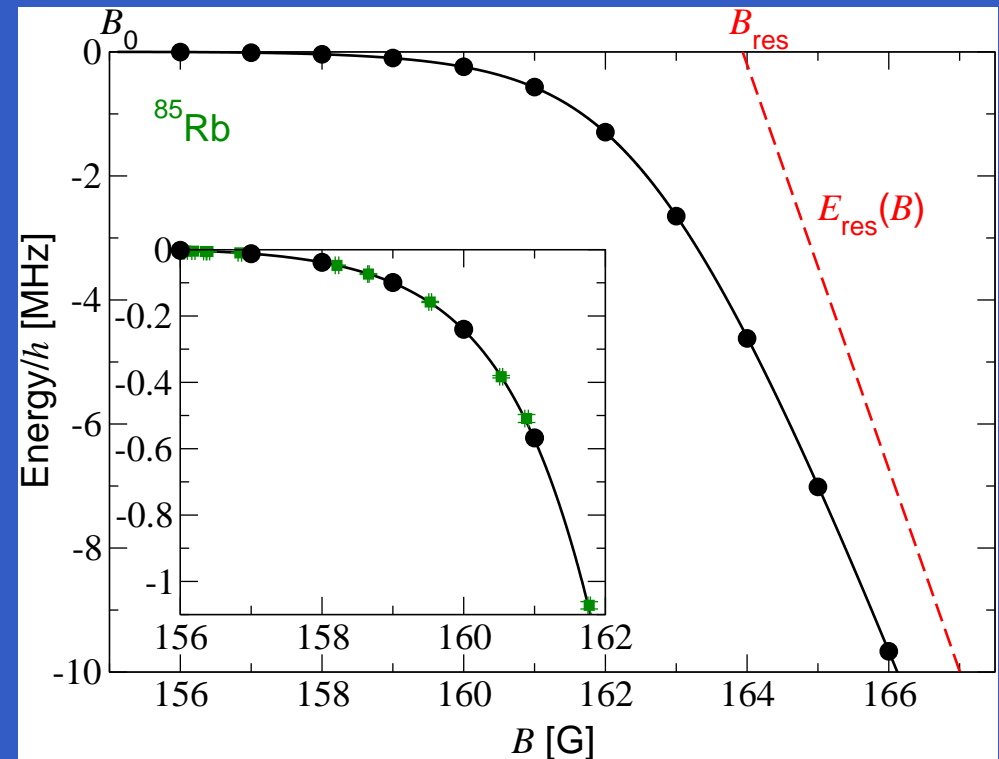
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Coupled-channels theory and experiment: N.R. Claussen, S.J.J.M.F. Kokkelmans, S.T. Thompson, E.A. Donley, E. Hodby, and C.E. Wieman, PRA **67**, 060701(R) (2003)

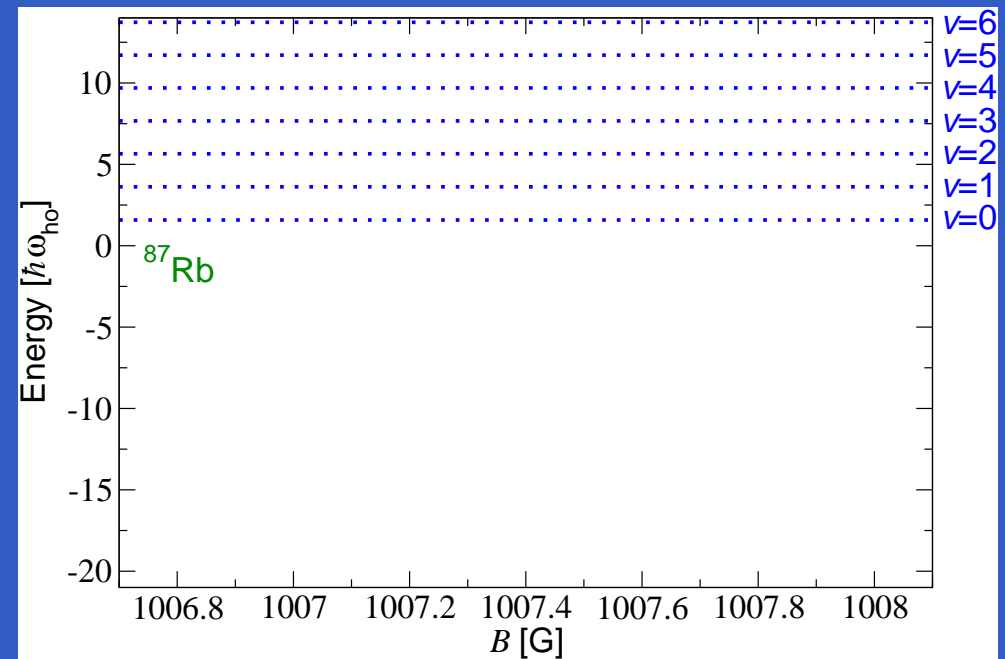
This two-channel approach: K. Góral, TK, and K. Burnett, PRA **71**, 023603 (2005)

# Magnetically-tunable interactions

## Feshbach-molecule association in a harmonic atom trap

Entrance-channel Hamiltonian:

$$H_{\text{bg}} = -\frac{\hbar^2}{m} \nabla^2 + V_{\text{bg}}(r) + V_{\text{ho}}(r)$$



Experiment: G. Thalhammer, K. Winkler, F. Lang, S. Schmid, R. Grimm, and J. Hecker-Denschlag,  
PRL **96**, 050402 (2006)

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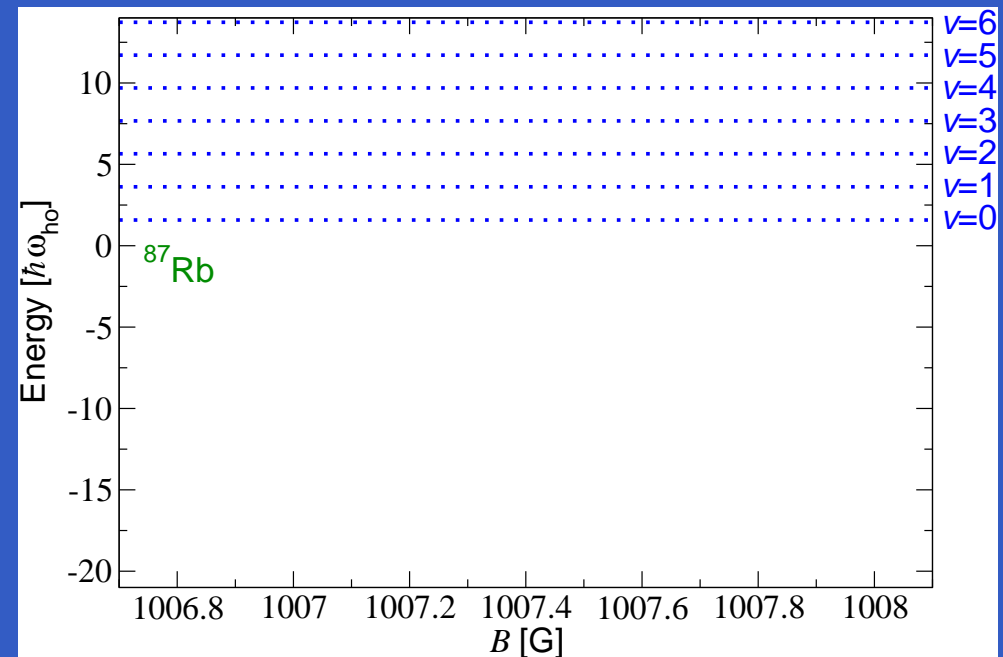
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- Single-resonance method:

$$H_{\text{cl}}(B) \rightarrow |\phi_{\text{res}}\rangle E_{\text{res}}(B) \langle \phi_{\text{res}}|$$



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PRL **96**, 050402 (2006)

# Magnetically-tunable interactions

## Feshbach-molecule association in a harmonic atom trap

- Entrance-channel Hamiltonian:

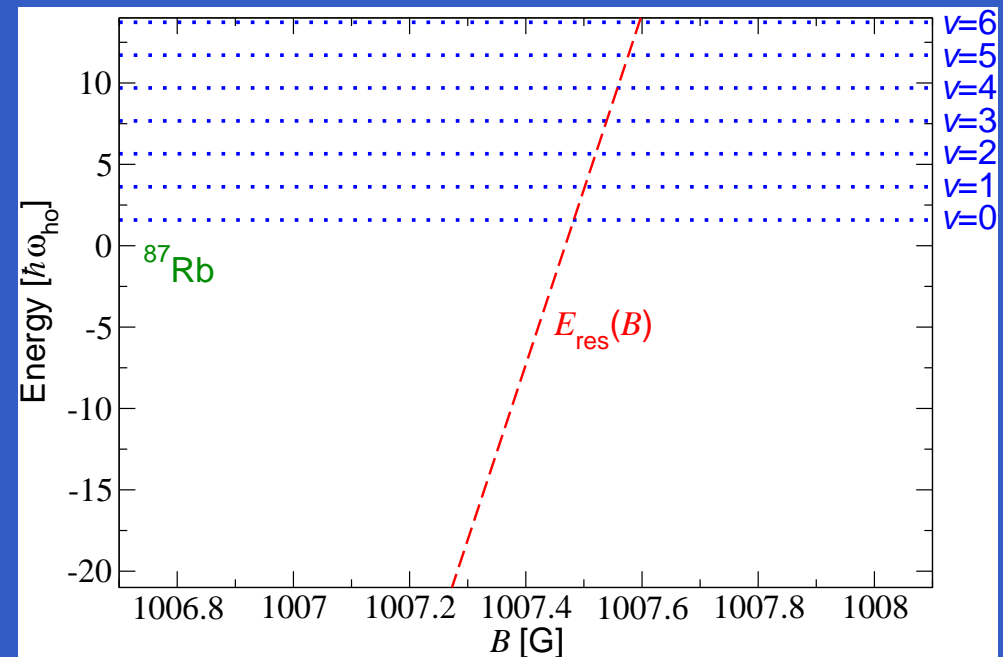
$$H_{\text{bg}} = -\frac{\hbar^2}{m} \nabla^2 + V_{\text{bg}}(r) + V_{\text{ho}}(r)$$

- Single-resonance method:

$$H_{\text{cl}}(B) \rightarrow |\phi_{\text{res}}\rangle E_{\text{res}}(B) \langle \phi_{\text{res}}|$$

- Resonance energy:

$$E_{\text{res}}(B) = \mu_{\text{res}}(B - B_{\text{res}})$$



Experiment: G. Thalhammer, K. Winkler, F. Lang, S. Schmid, R. Grimm, and J. Hecker-Denschlag,  
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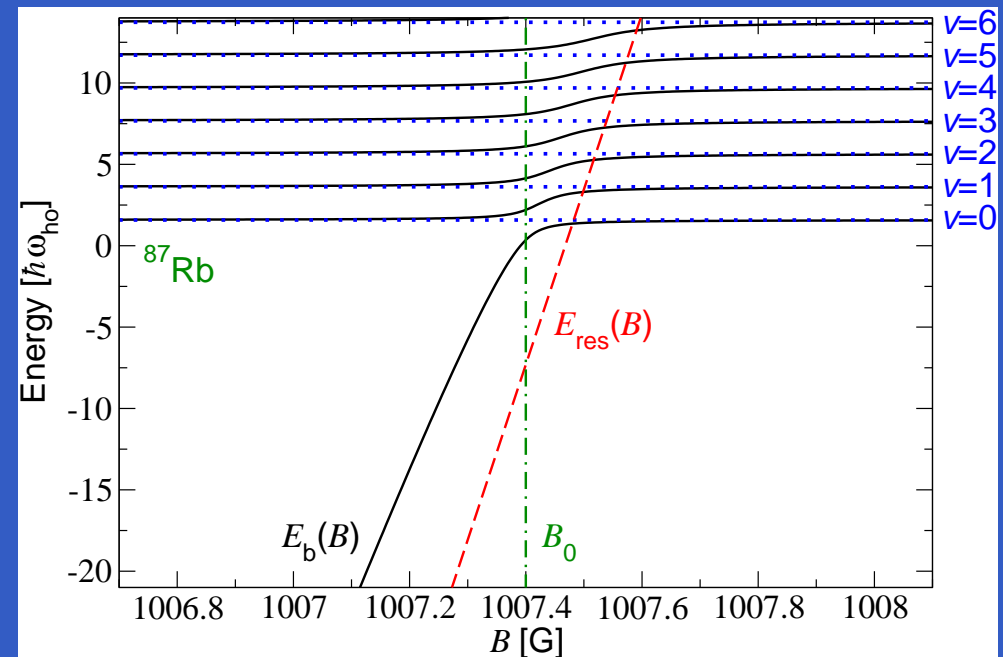
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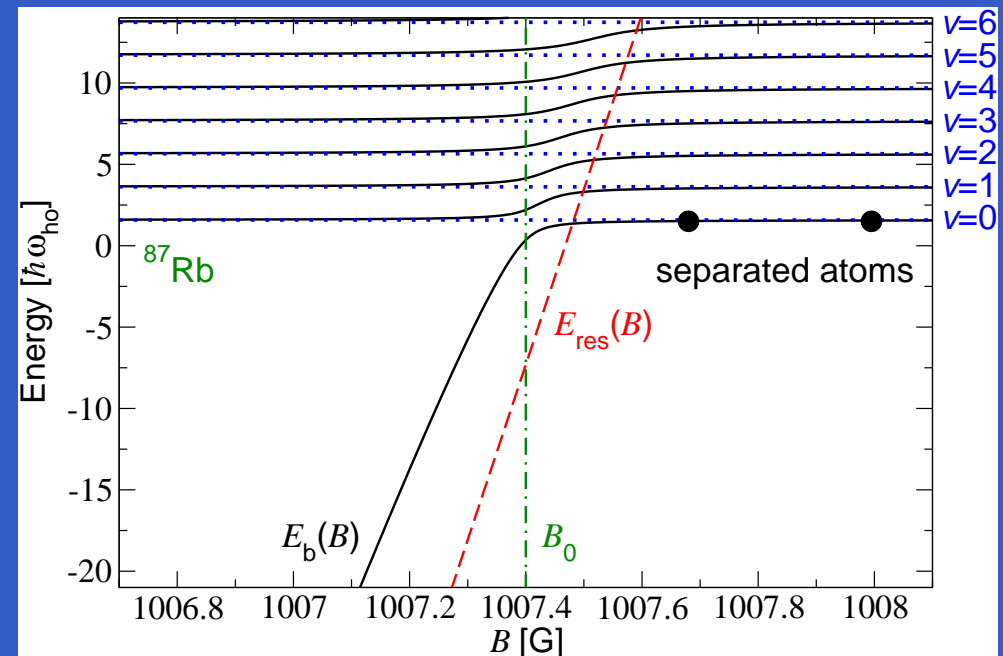
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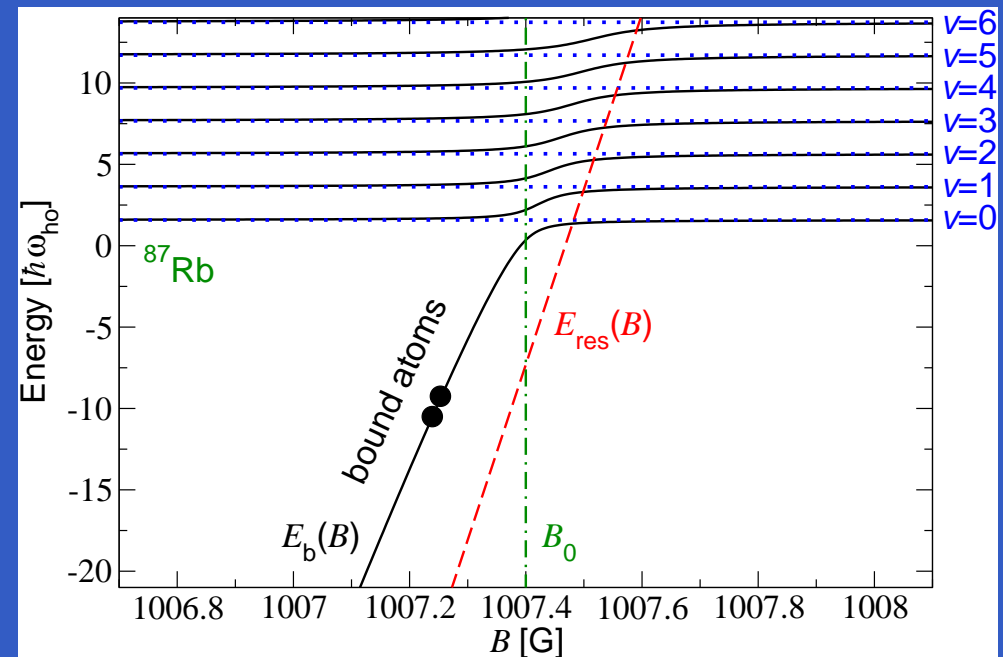
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# Magnetically-tunable interactions

## Resonance shape

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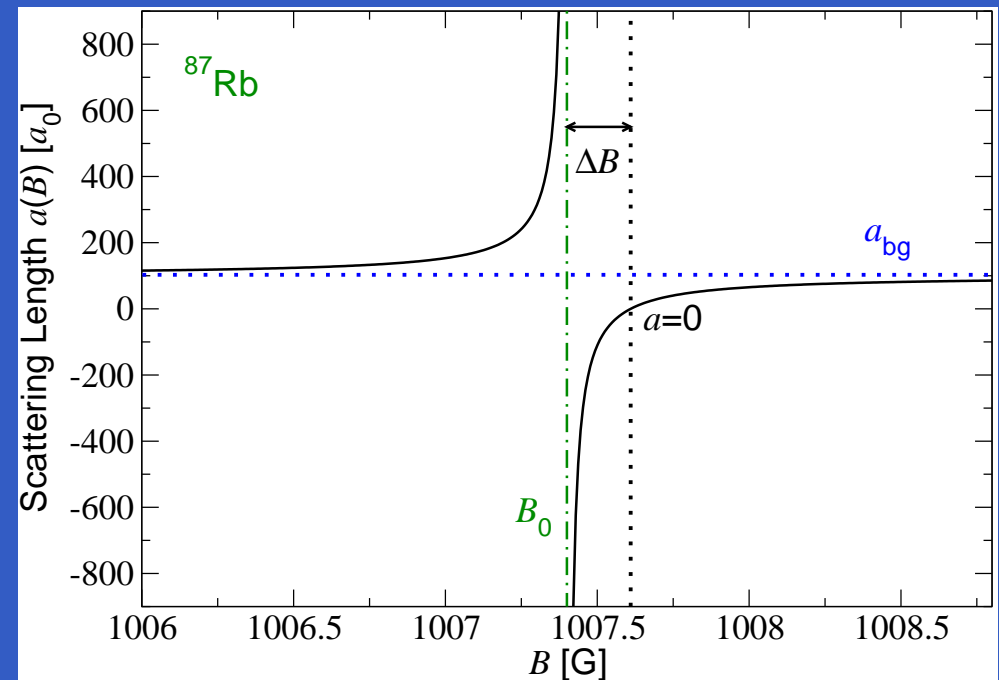
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- Scattering length:

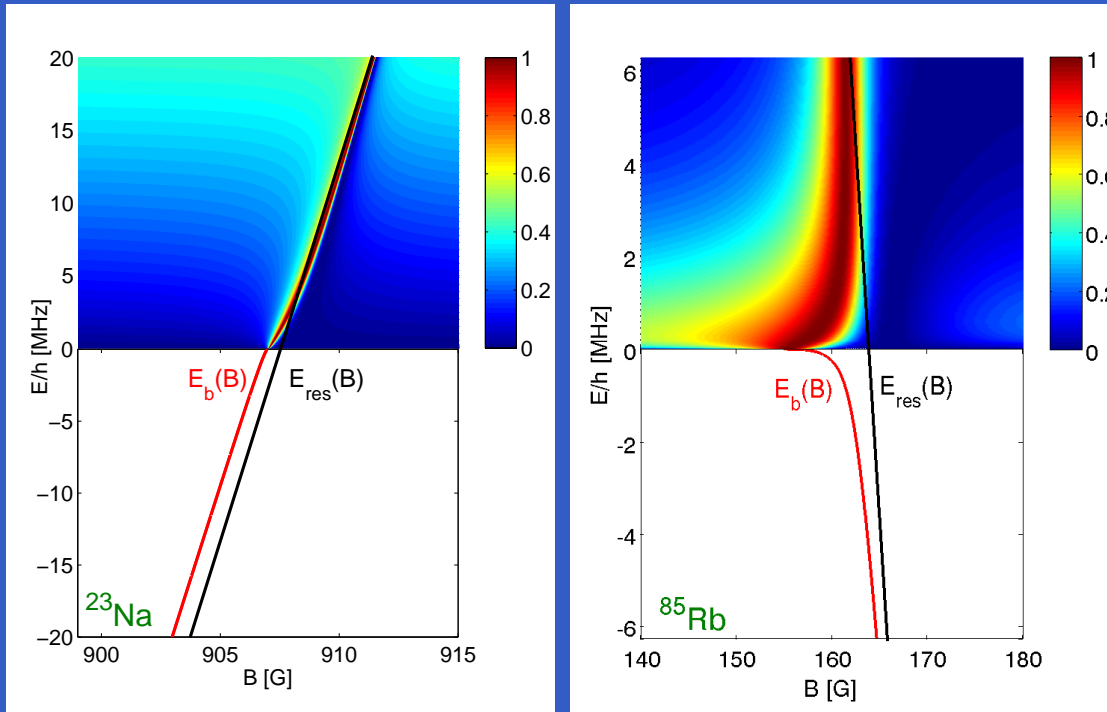
$$a(B) = a_{\text{bg}} \left( 1 - \frac{\Delta B}{B - B_0} \right)$$



A.J. Moerdijk, B.J. Verhaar, and A. Axelsson, PRA **51**, 4852 (1995)



# Classification of Feshbach resonances



$s$ -wave scattering cross section:

$$\sigma_0(k) = 8\pi k^{-2} \sin^2 \delta_0(k)$$

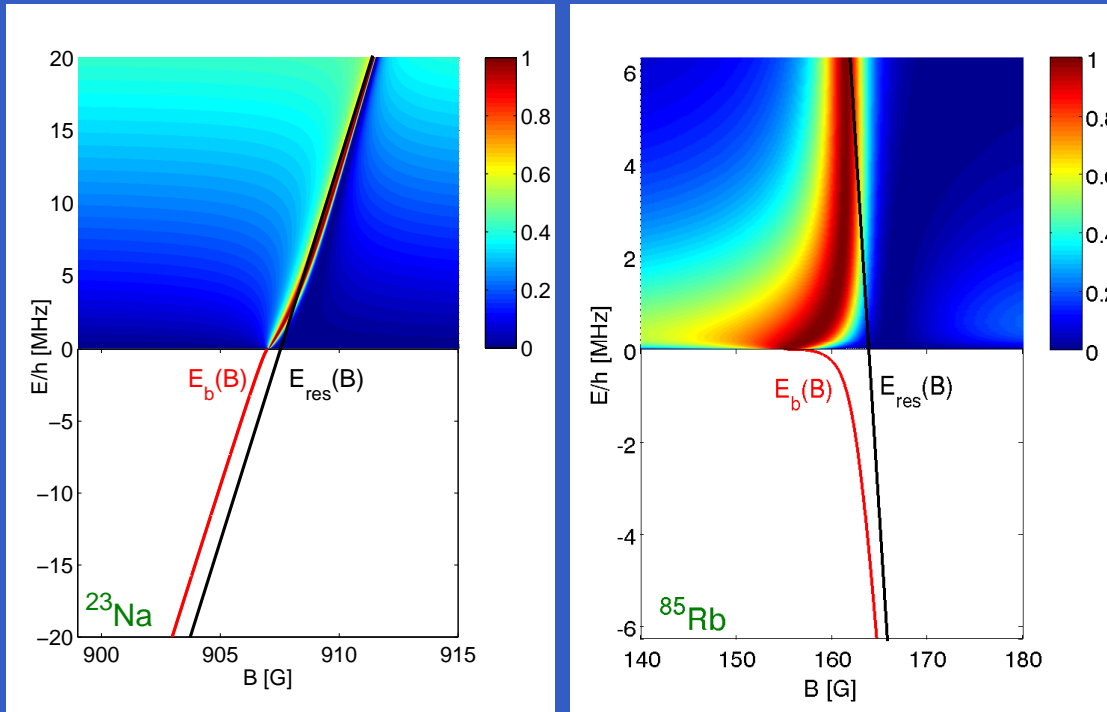
$$\underset{k \rightarrow 0}{\sim} 8\pi a^2$$

P.S. Julienne and B. Gao, *Simple theoretical models for resonant cold atom interactions*, in Atomic Physics **20**, ed. by C. Roos, H. Häffner, and R. Blatt (American Institute of Physics, Conference Proceedings **869**, 2006), pp. 261-268, e-print physics/0609013

B. Marcellis *et al.*, PRA **70**, 012701 (2004)

Figures: Tom Hanna

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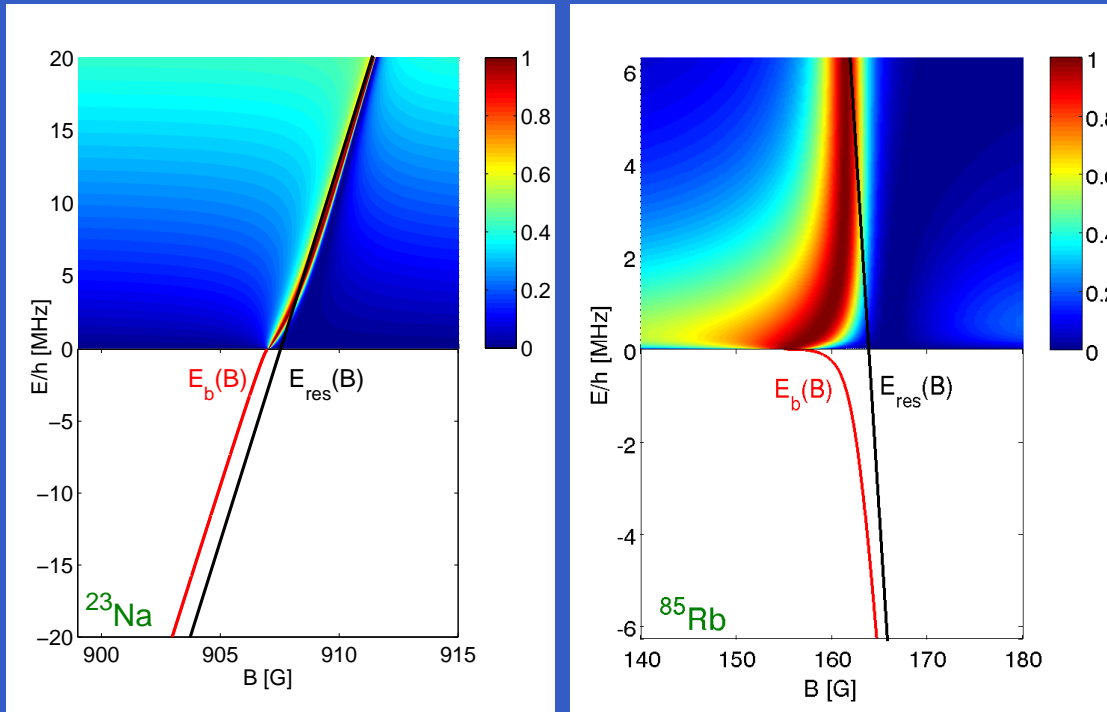
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P.S. Julienne and B. Gao, e-print physics/0609013

B. Marcelis *et al.*, PRA **70**, 012701 (2004)

Figures: Tom Hanna

# Classification of Feshbach resonances



P.S. Julienne and B. Gao, e-print physics/0609013

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The maximum  $\sigma_0(k)$  mirrors the behaviour of the bound-state energy,  $E_b(B)$ , at the collision threshold.

Width and behaviour of the resonance energy,  $E_{\text{res}}(B)$ , above the collision threshold differ between  $^{23}\text{Na}$  and  $^{85}\text{Rb}$ .

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When is a single-channel treatment appropriate?

- Single-channel Hamiltonian:

$$H_{2B}^{\text{eff}} = -\frac{\hbar^2}{m} \nabla^2 + V_{\text{eff}}(B, r)$$

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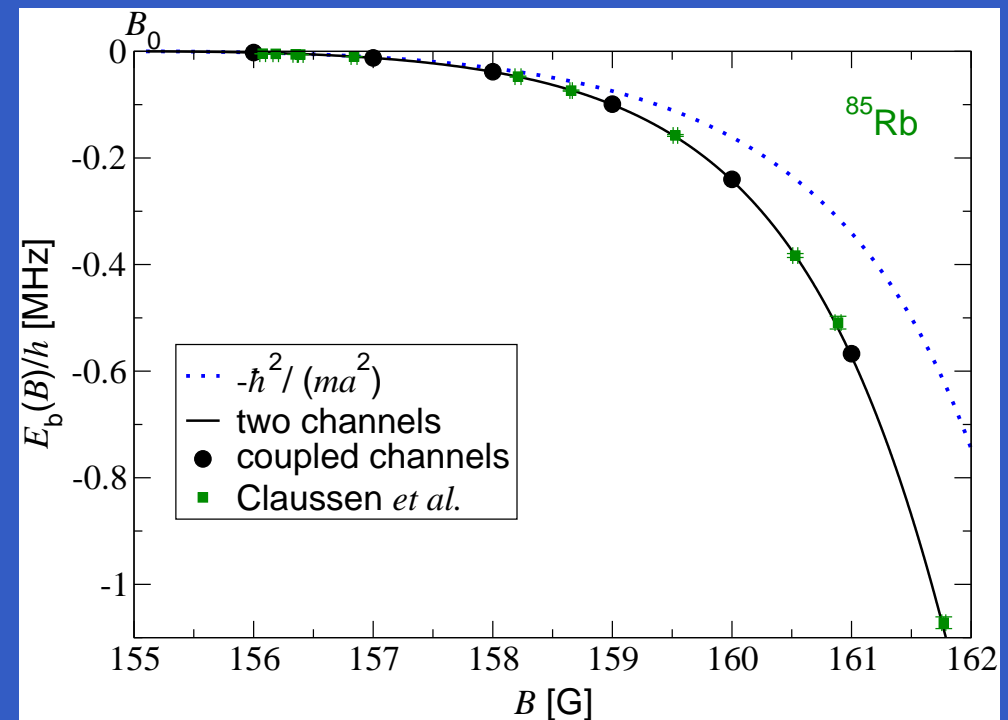
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$$E_b = -\hbar^2 / (ma^2)$$



Coupled-channels theory and experiment: N.R. Claussen, S.J.J.M.F. Kokkelmans, S.T. Thompson, E.A. Donley, E. Hodby, and C.E. Wieman, PRA **67**, 060701(R) (2003)

This two-channel approach: K. Góral, TK, and K. Burnett, PRA **71**, 023603 (2005)

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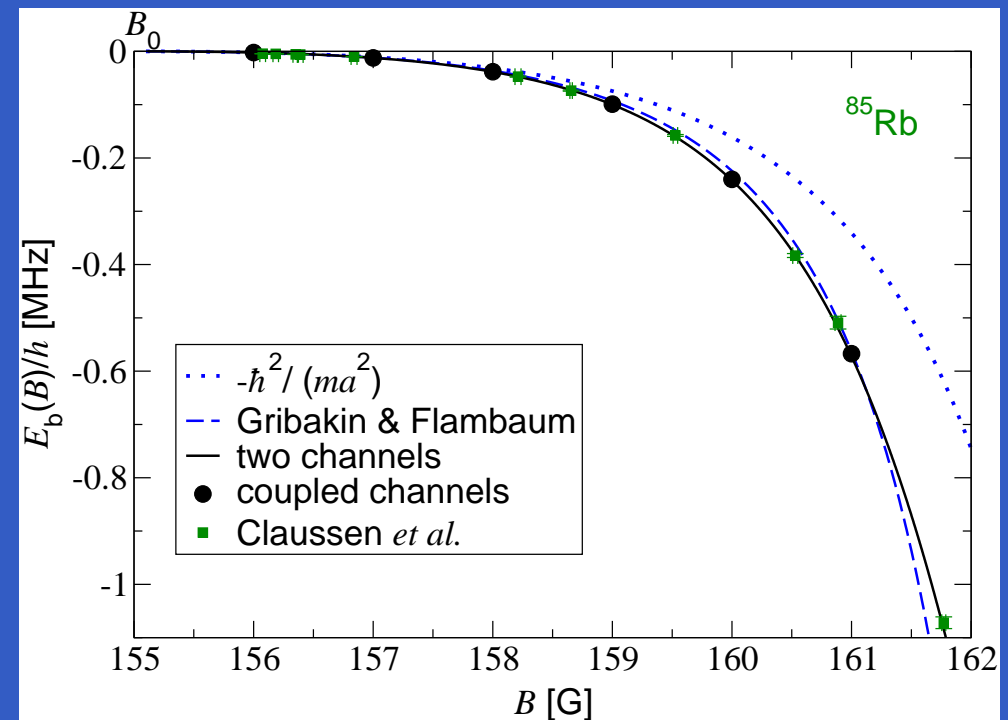
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$$E_b = -\hbar^2 / [m(a - \bar{a})^2]$$



G.F. Gribakin and V.V. Flambaum, PRA **48**, 546 (1993)

B. Gao, J. Phys. B **37**, 4273 (2004)

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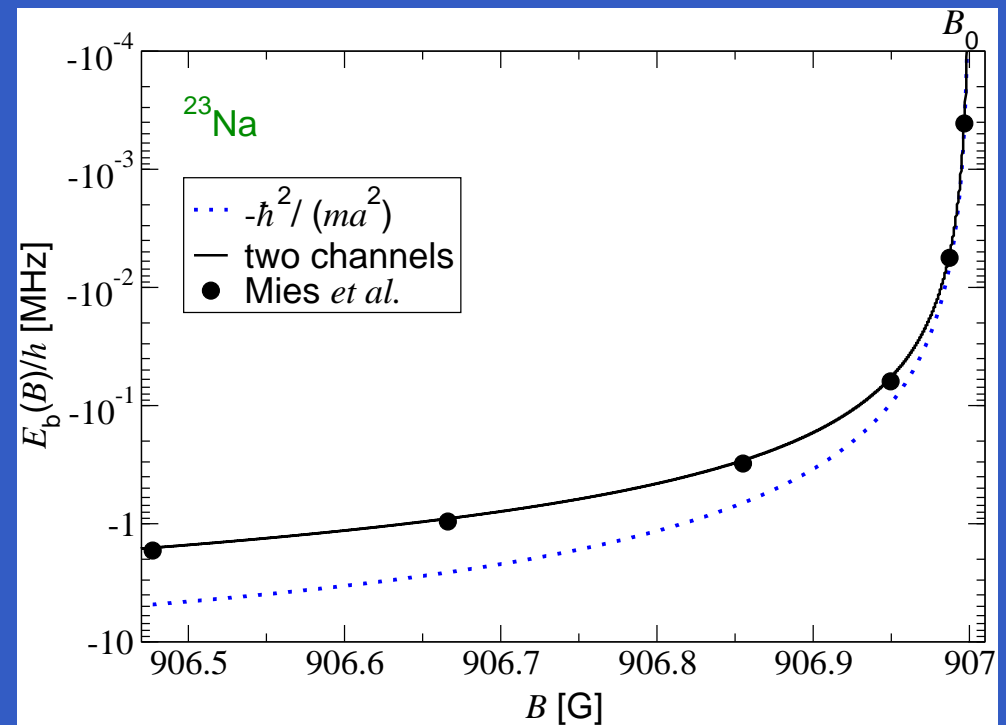
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This two-channel two-potential model: F.H. Mies, E. Tiesinga, and P.S. Julienne, PRA **61**, 022721 (2000)

This two-channel single-resonance approach: TK, K. Góral, and T. Gasenzer, PRA **70**, 023613 (2004)



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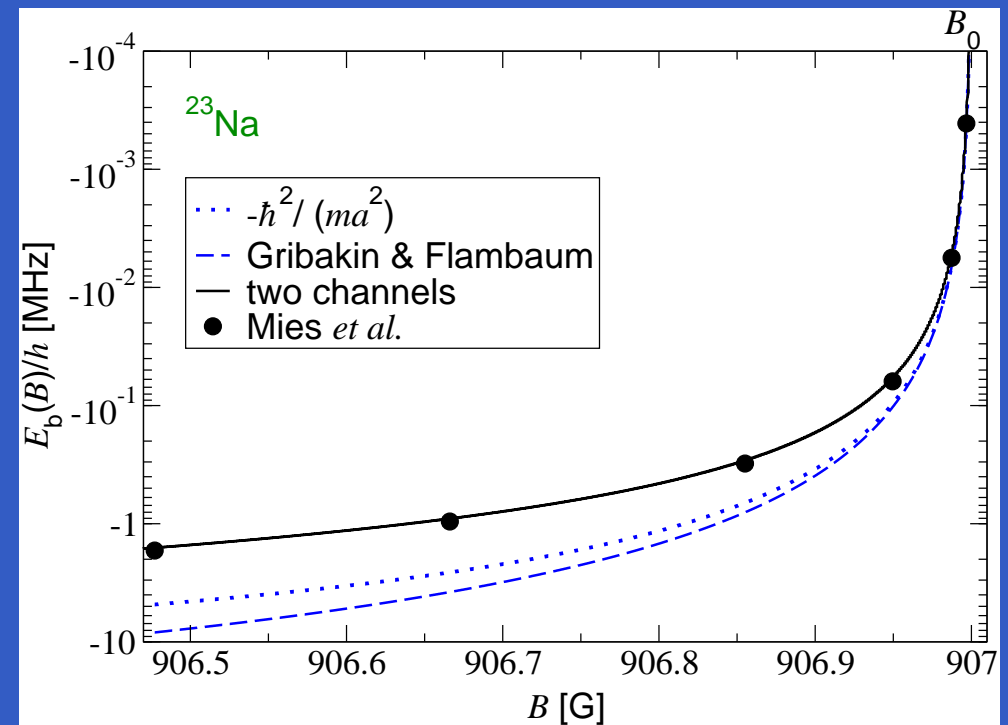
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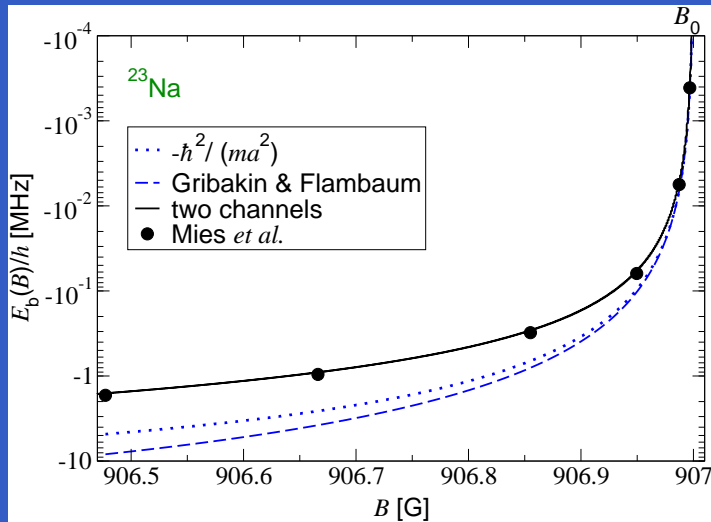


G.F. Gribakin and V.V. Flambaum, PRA **48**, 546 (1993)

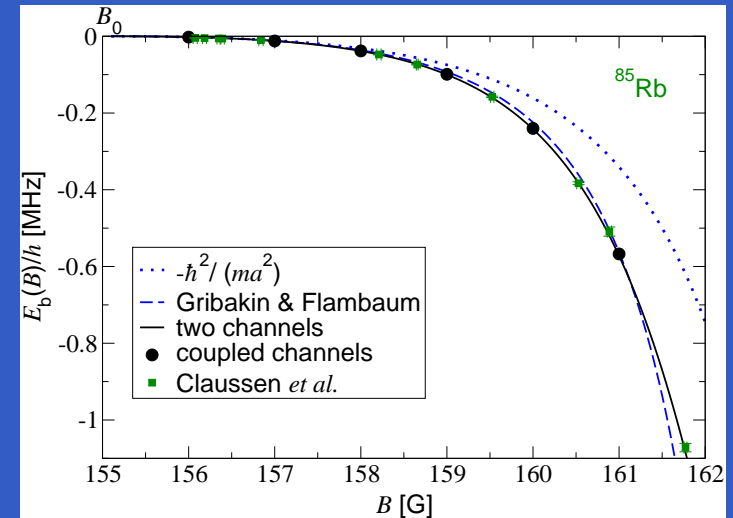
B. Gao, J. Phys. B **37**, 4273 (2004)

# Classification of Feshbach resonances

When is a single-channel treatment appropriate?



closed-channel dominated



entrance-channel dominated

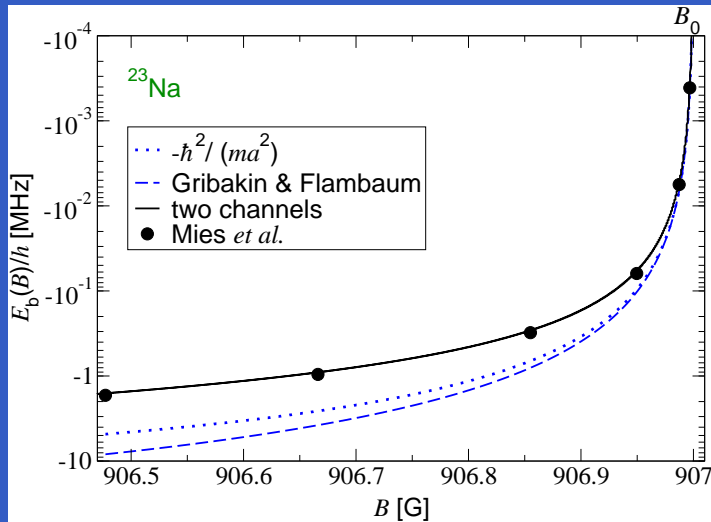
$$\left| E_b(B) + \frac{\hbar^2}{m[a(B) - \bar{a}]^2} \right| < \left| E_b(B) + \frac{\hbar^2}{ma^2(B)} \right|$$

TK, K. Góral, and T. Gasenzer, PRA **70**, 023613 (2004)

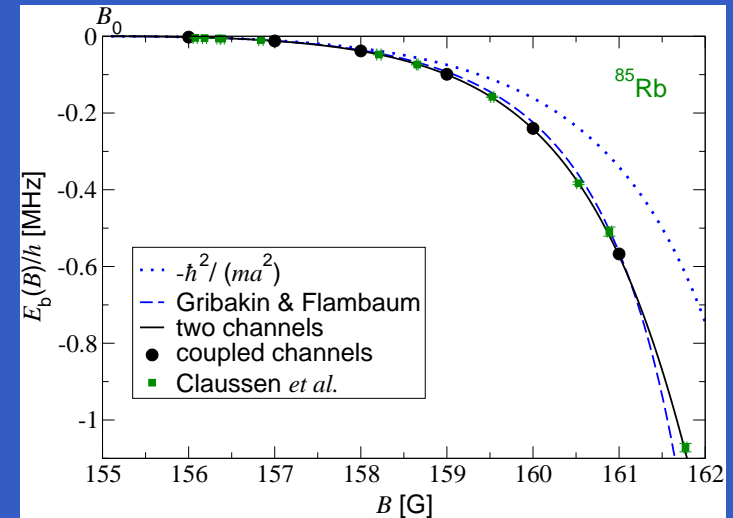
See also: D.S. Petrov, PRL **93**, 143201 (2004)

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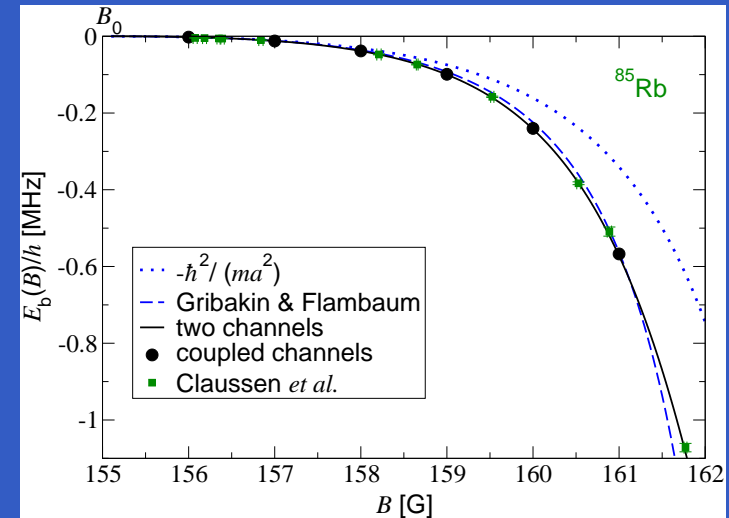
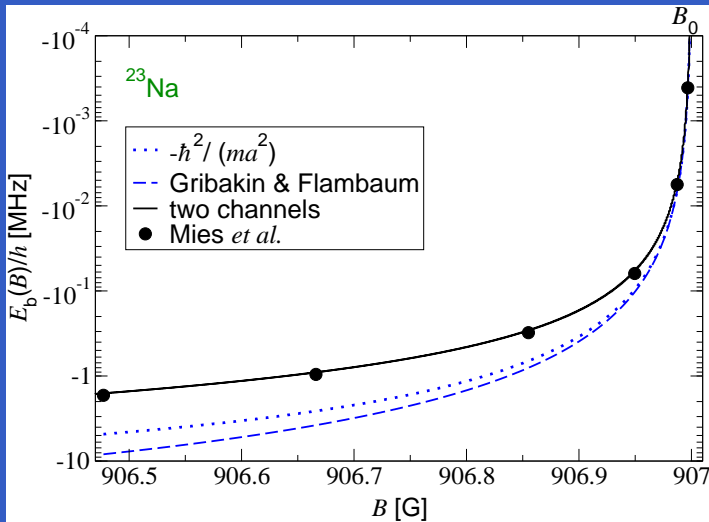
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$\Updownarrow$

$$\eta = \frac{\hbar^2/(m\bar{a}^2)}{a_{\text{bg}}\mu_{\text{res}}\Delta B/\bar{a}} = \frac{E_{\text{vdW}}}{\Gamma_{\text{res}}(E_{\text{vdW}})} < 1$$

# Classification of Feshbach resonances

When is a single-channel treatment appropriate?



closed-channel dominated

$$\eta > 1$$

${}^6\text{Li}$  543 G      1215

${}^{23}\text{Na}$  907 G      11

${}^{87}\text{Rb}$  1007 G    5.9

entrance-channel dominated

$$\eta < 1$$

${}^{40}\text{K}$  202 G      0.46

${}^{85}\text{Rb}$  155 G      0.04

${}^6\text{Li}$  834 G      0.02

Review: TK, K. Góral, and P.S. Julienne, RMP **78**, 1311 (2006)

# Diatomic Feshbach molecules

## Two-component Feshbach-molecular bound states

- Schrödinger equation:

$$H_{2B} \begin{pmatrix} \phi_b^{\text{bg}} \\ \phi_b^{\text{cl}} \end{pmatrix} = E_b \begin{pmatrix} \phi_b^{\text{bg}} \\ \phi_b^{\text{cl}} \end{pmatrix}$$

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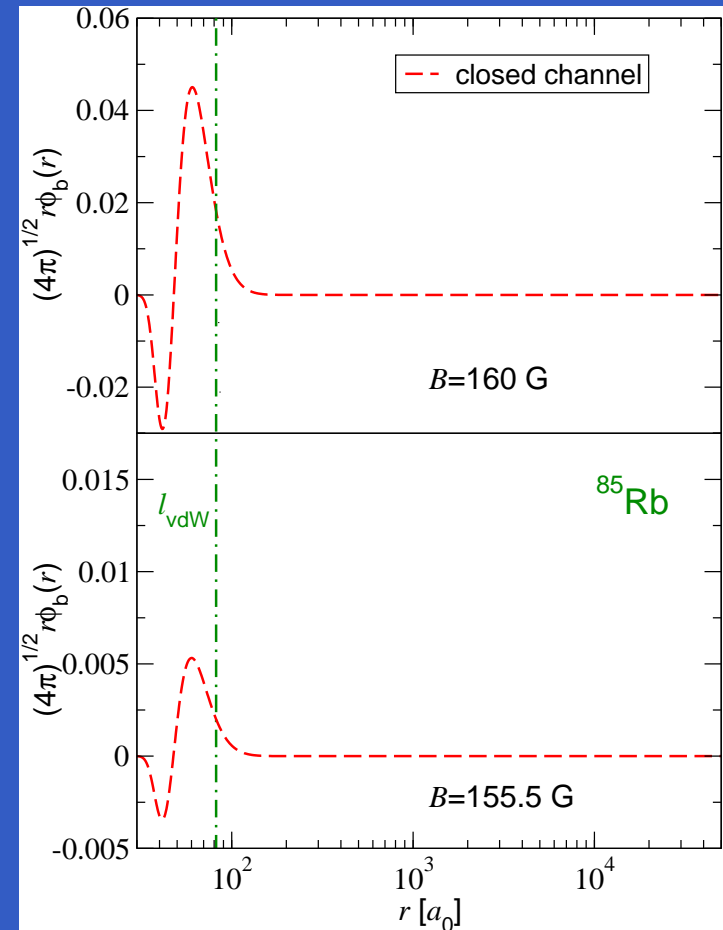
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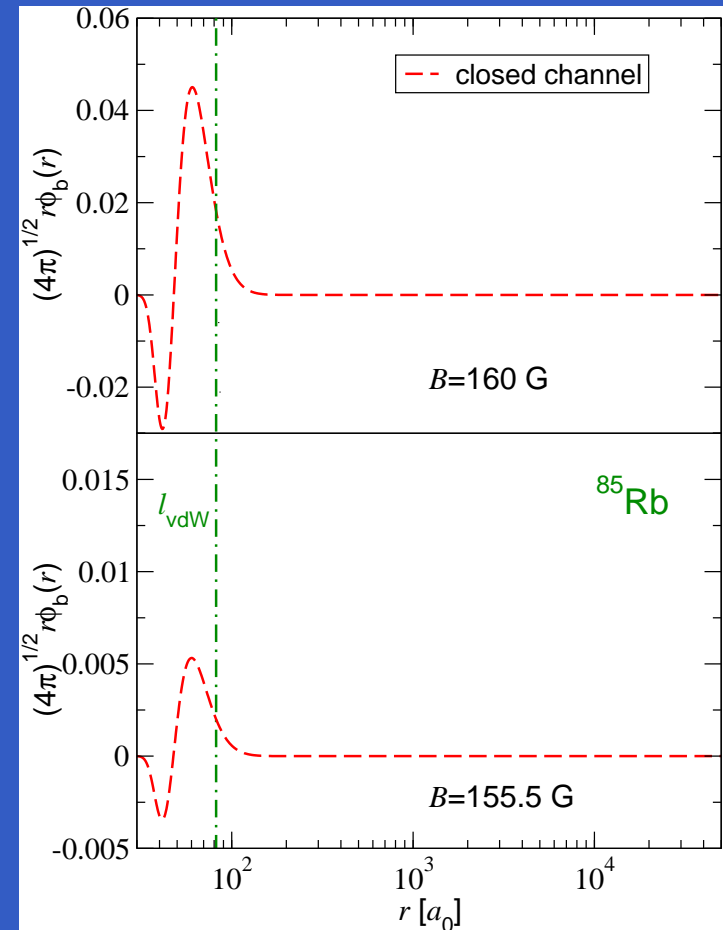
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# Diatomic Feshbach molecules

Near-resonant Feshbach-molecules become universal!

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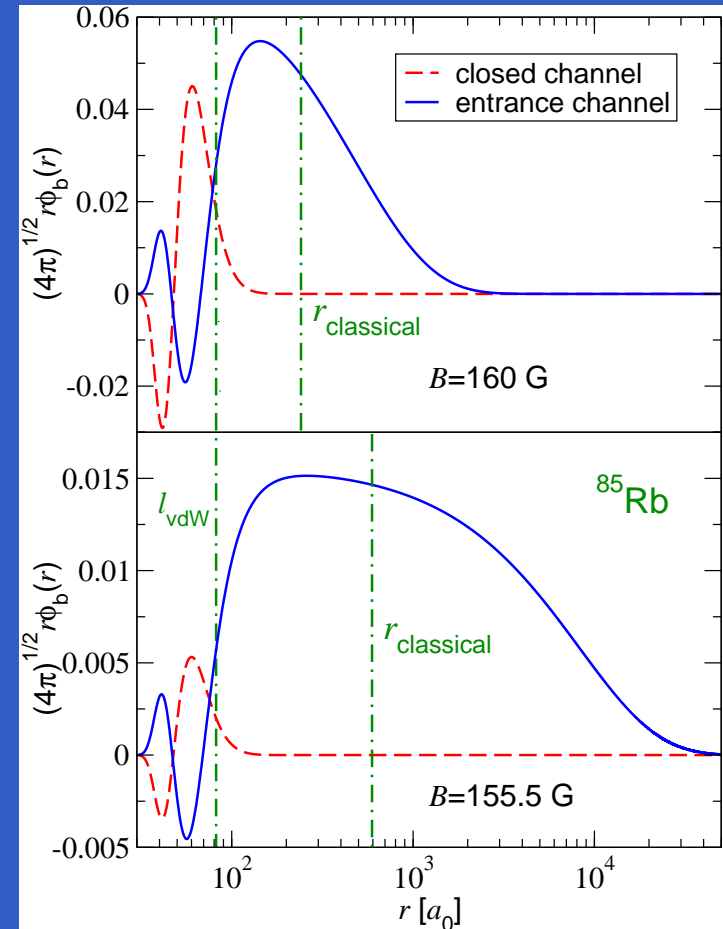
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# Diatomic Feshbach molecules

## Stern-Gerlach separation of $^{87}\text{Rb}$ atoms and molecules

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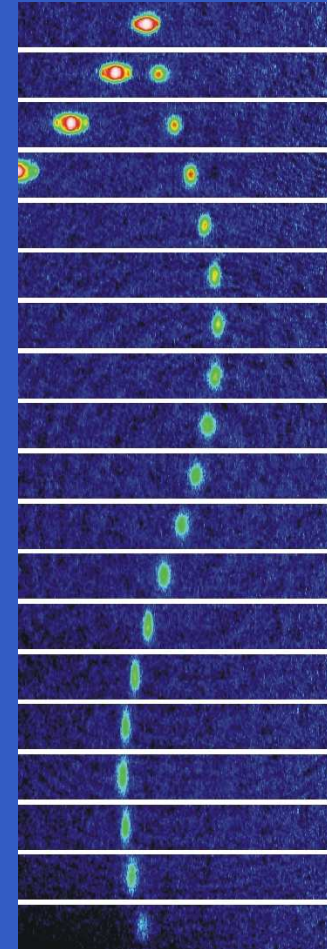
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Experiment: S. Dürr, T. Volz, A. Marte, and G. Rempe, PRL **92**, 020406 (2004)

See also: J. Herbig *et al.*, Science **301**, 1510 (2003)

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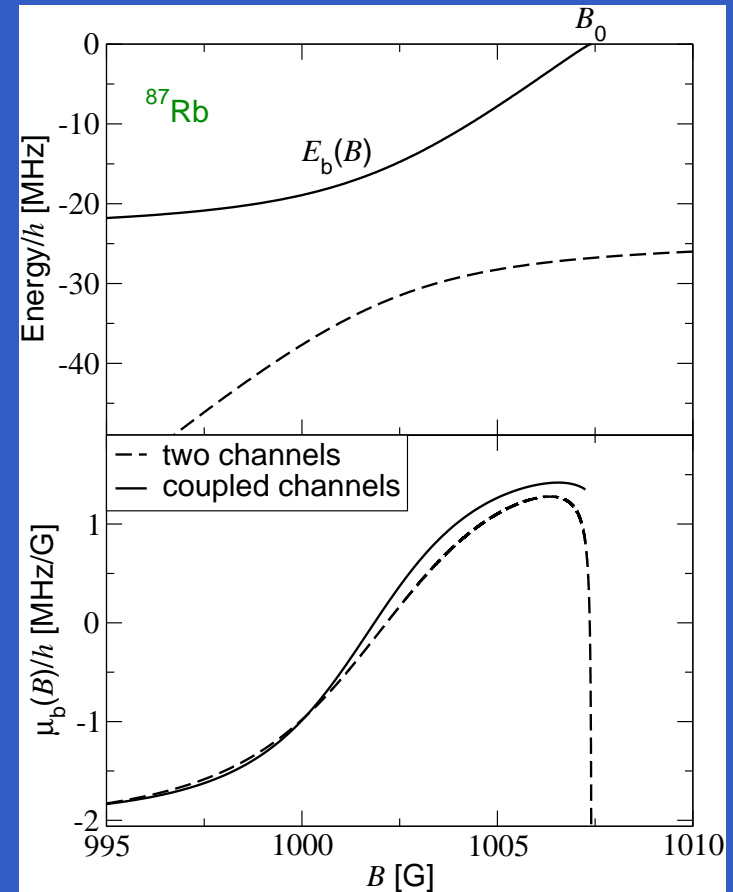
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Coupled-channels theory: E.G.M van Kempen and B.J. Verhaar  
 This two-channel approach: K. Góral *et al.*, J. Phys. B **37**, 3457 (2004)

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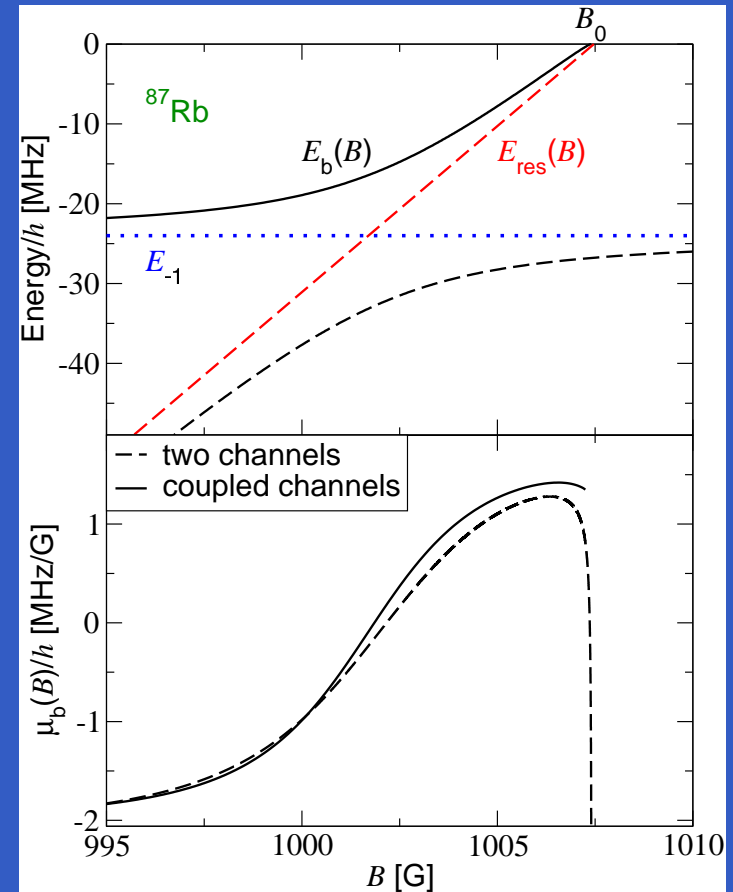
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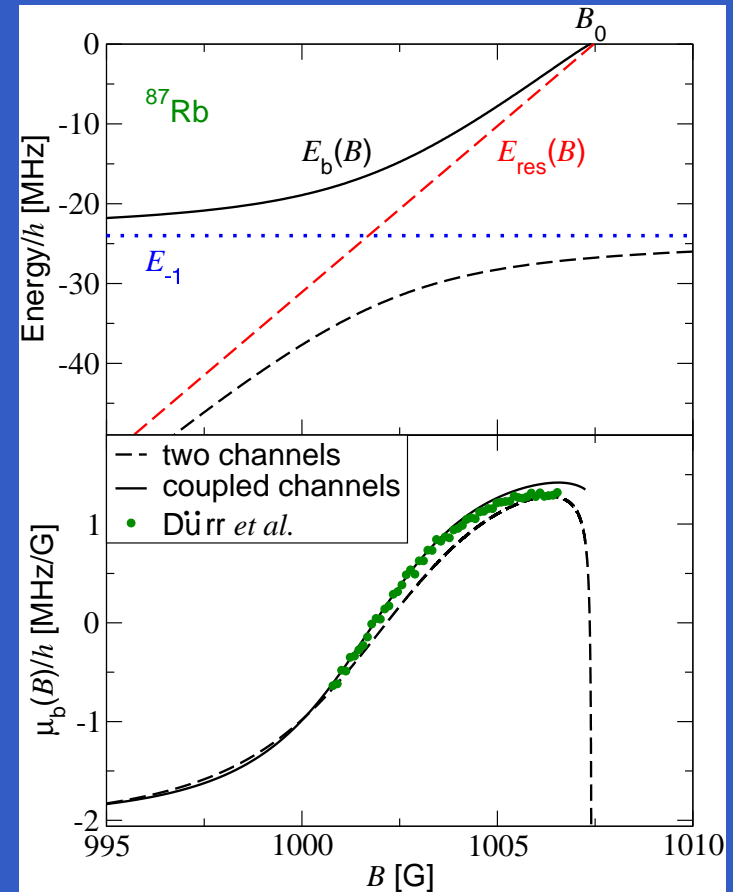
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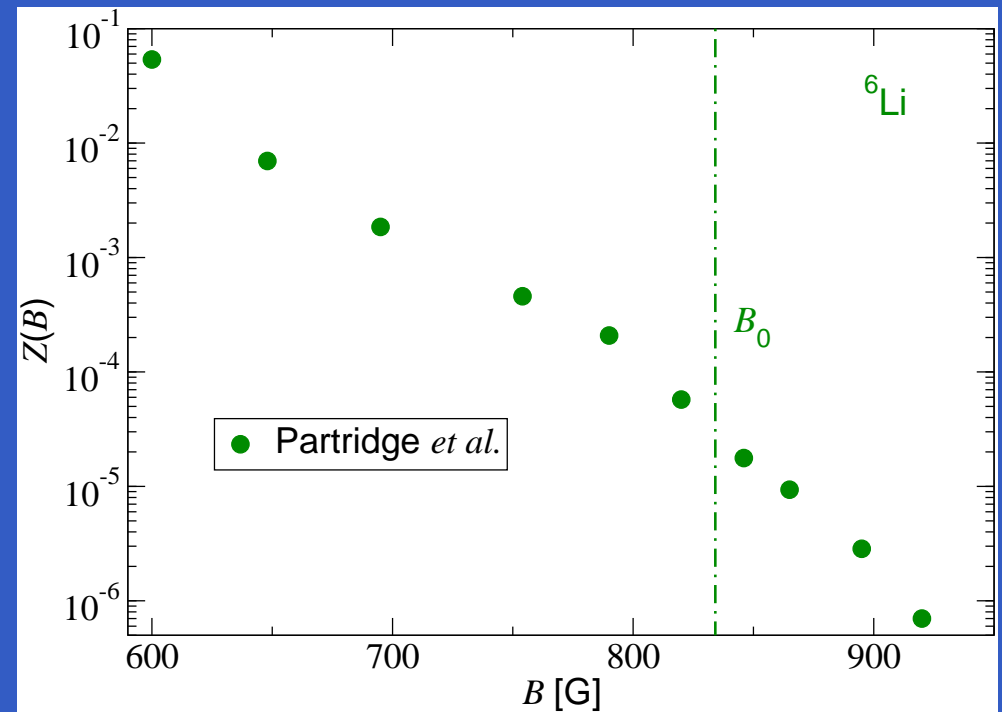
## Direct measurement of closed-channel admixtures

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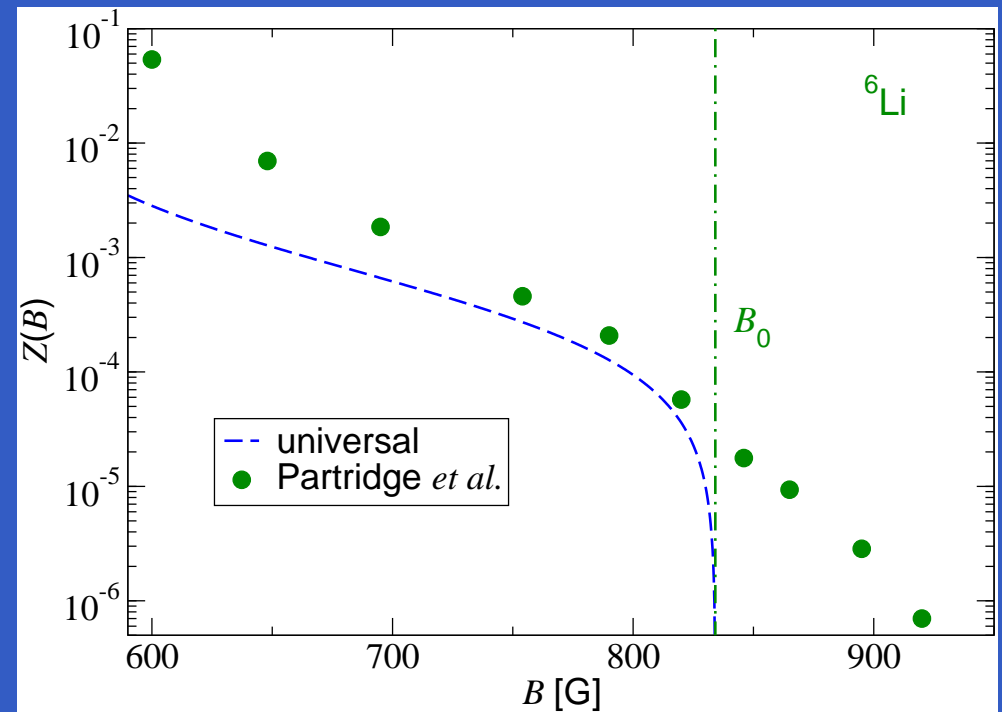
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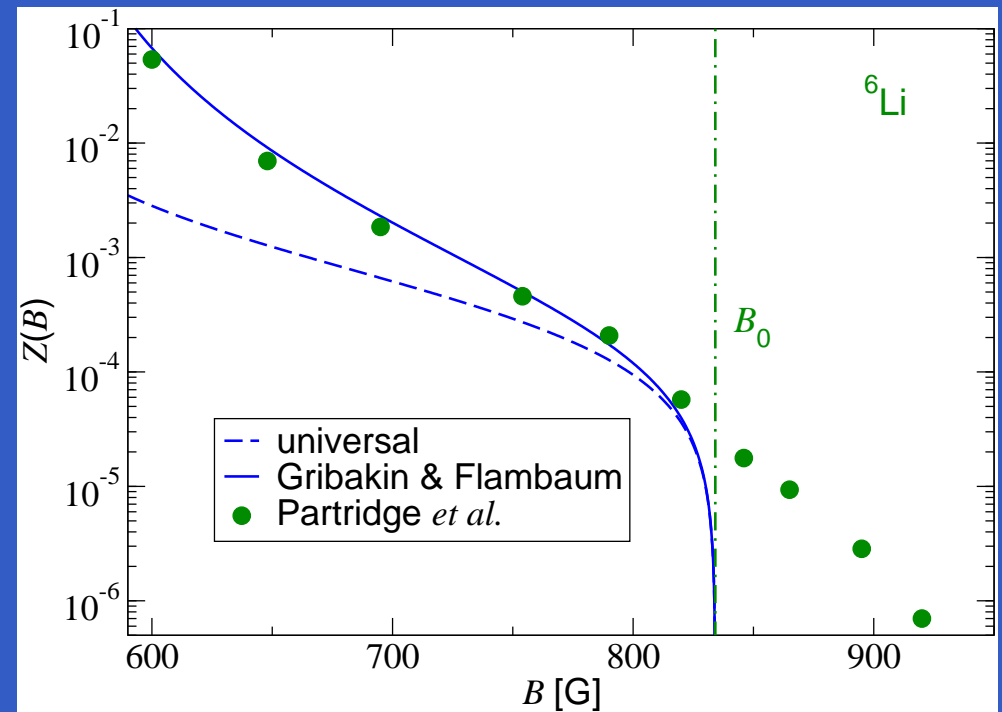
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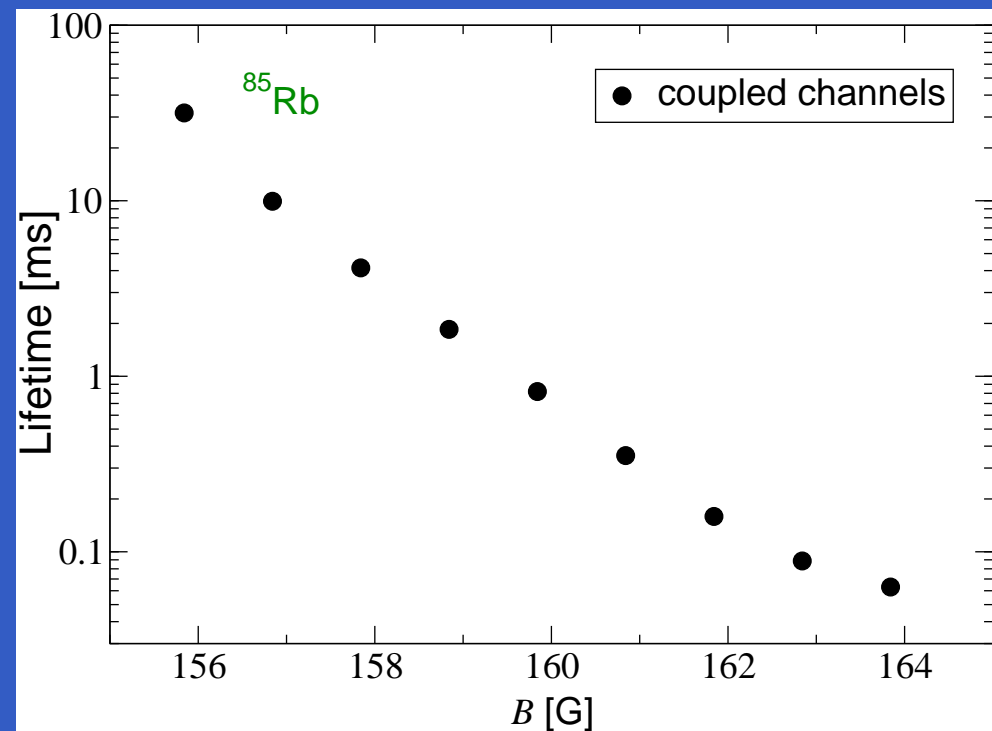
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$^{85}\text{Rb}_2$  Feshbach molecules decay due to spin relaxation!

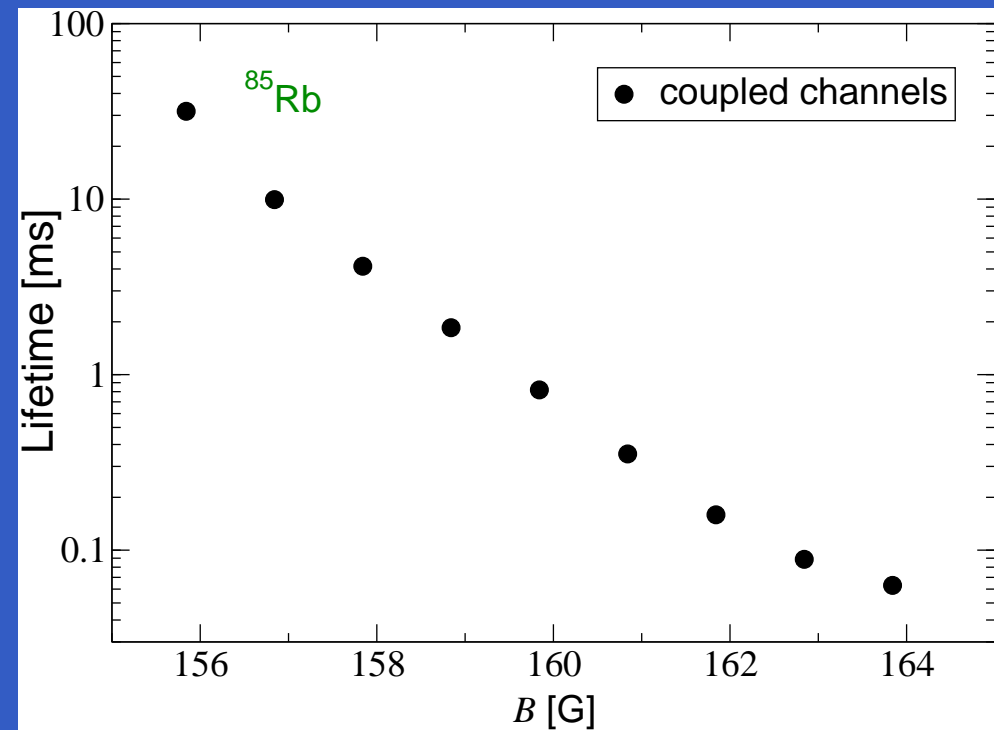


# Diatomic Feshbach molecules

The lifetime depends on the size of the molecule!

● Average volume of a halo state:

$$\mathcal{V} = 4\pi\langle r^3 \rangle / 3 = \pi a^3$$



# Diatomic Feshbach molecules

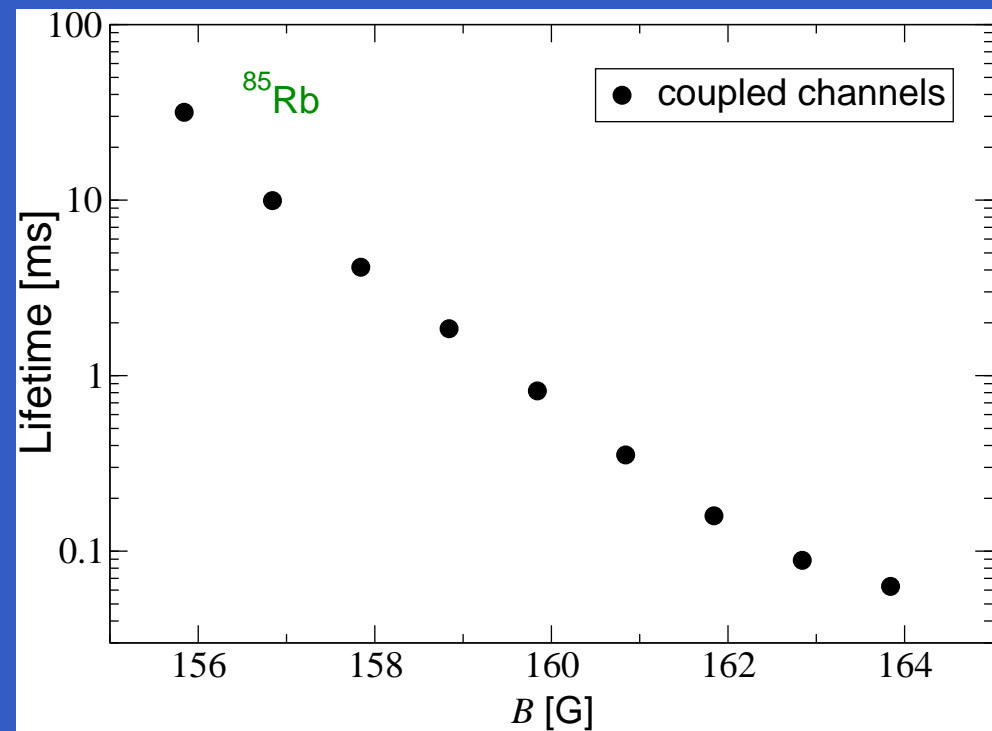
The lifetime depends on the size of the molecule!

- Average volume of a halo state:

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- Rate equation for spin relaxation:

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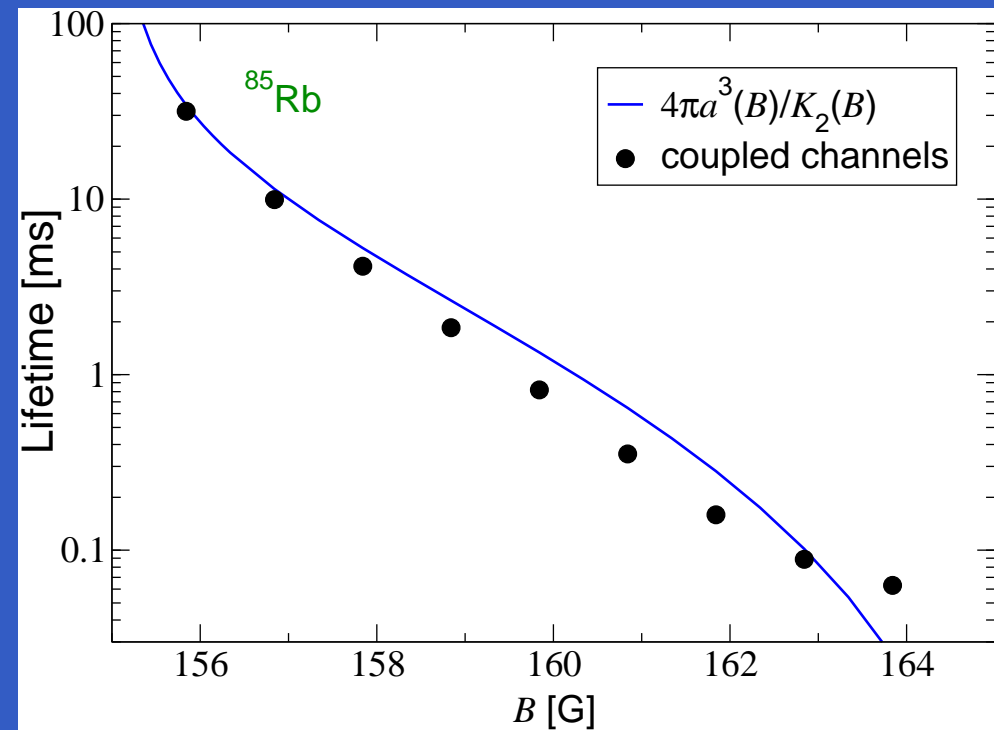
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# Diatomic Feshbach molecules

Experiment and theory agree!

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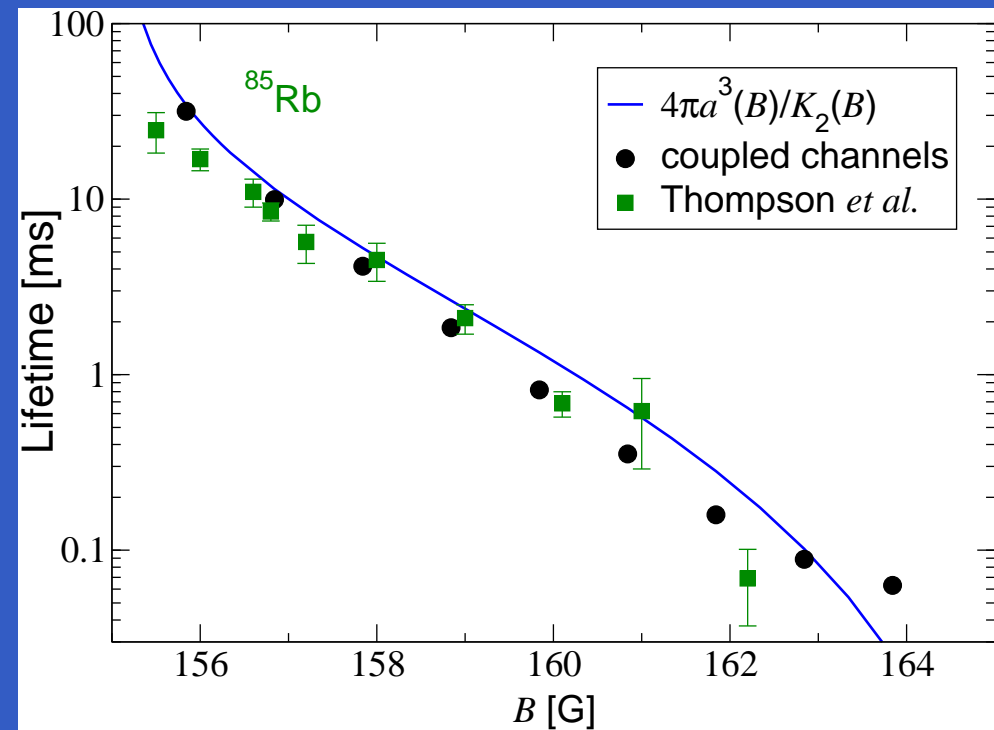
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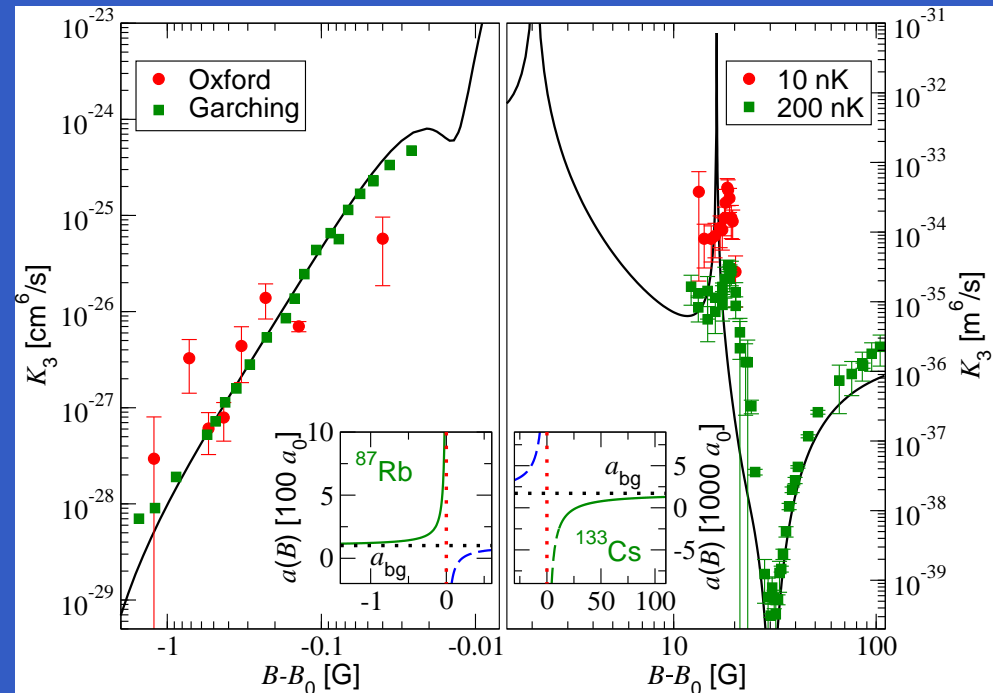


Experiment: S.T. Thompson, E. Hodby, and C.E. Wieman, PRL **94**, 020401 (2005)

Theory: TK, E. Tiesinga, and P.S. Julienne, PRL **94**, 020402 (2005)

# Outlook

## ● Universal trimer molecules



General concept: E. Nielsen and J.H. Macek, PRL **83**, 1566 (1999)

B.D. Esry, C.H. Greene, and J.P. Burke, PRL **83**, 1751 (1999)

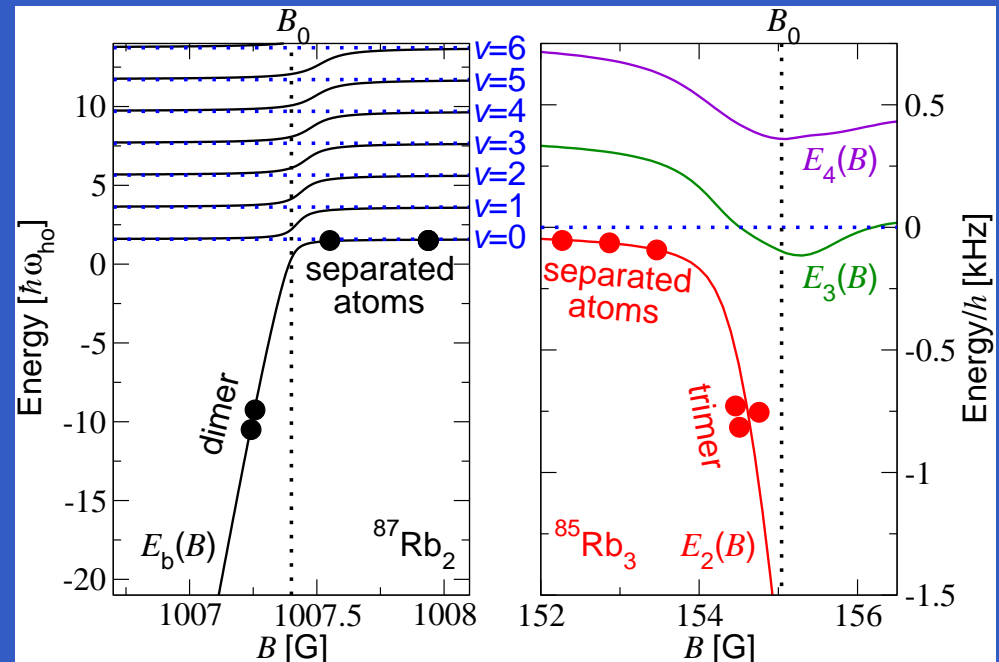
Experiment: T. Kraemer *et al.*, Nature **440**, 315 (2006)

See also: G. Smirne *et al.*, PRA **75**, 020702(R) (2007)

J. Stenger *et al.*, PRL **82**, 2422 (1999)

# Outlook

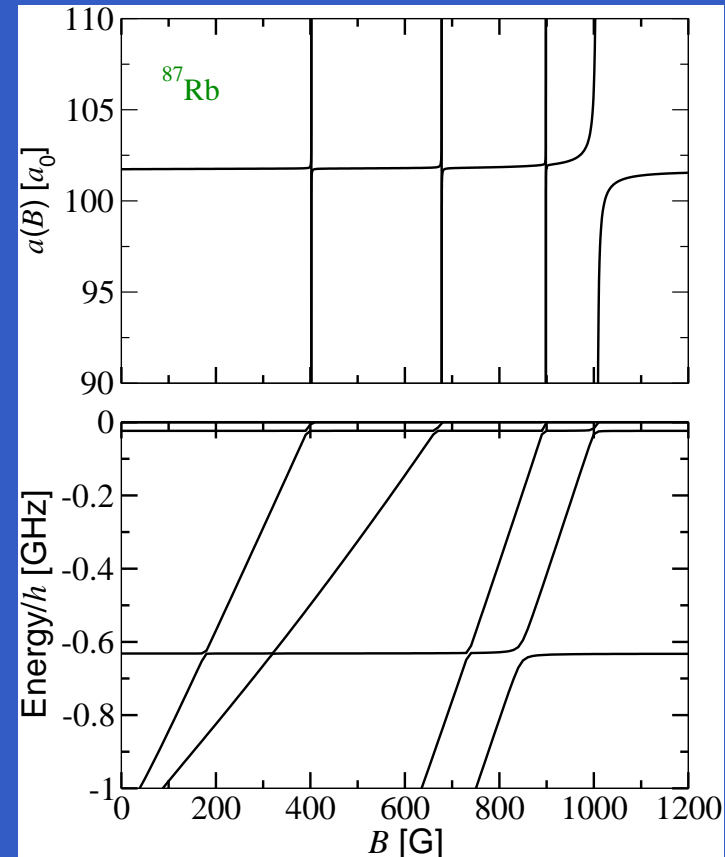
## Universal trimer molecules



General idea: V. Efimov, Phys. Lett. **33B**, 563 (1970)  
This concept: M. Stoll and TK, PRA **72**, 022714 (2005)  
See also: V. Efimov, Nucl. Phys. A **362**, 45 (1981)  
W. Schöllkopf and J.P. Toennies, Science **266**, 1345 (1994)

# Outlook

- Universal trimer molecules
- Ground-state and polar molecules

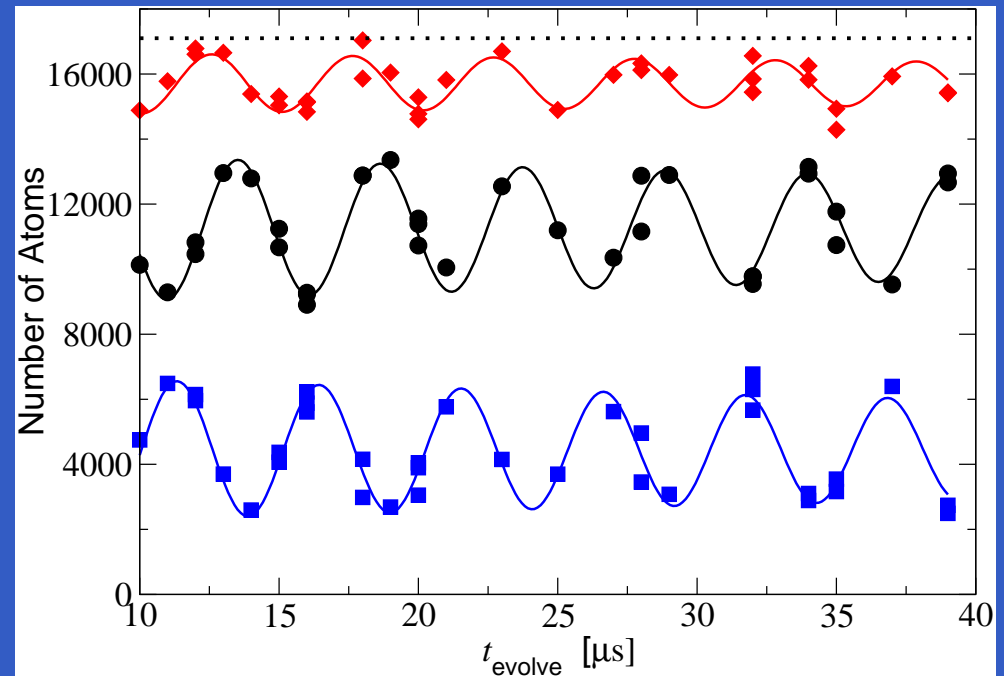


This concept: D. Jaksch *et al.*, PRL **89**, 040402 (2002)  
This experiment: K. Winkler *et al.*, PRL **98**, 043201 (2007)  
Figure: E. Tiesinga and P.S. Julienne



# Outlook

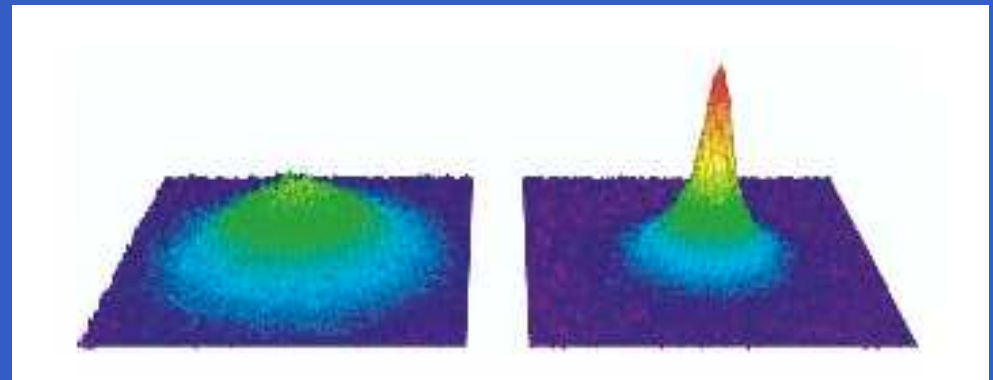
- Universal trimer molecules
- Ground-state and polar molecules
- Dynamics of pairing and molecule formation



Adapted from: E.A. Donley *et al.*, Nature 412, 295 (2002)

# Outlook

- Universal trimer molecules
- Ground-state and polar molecules
- Dynamics of pairing and molecule formation
- **Superfluid Fermi gases**



Adapted from: M. Greiner *et al.*, Nature **426**, 537 (2003)

# Thanks to

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- Experimental groups at: Oxford, JILA, Innsbruck, MIT, Rice, and Munich

