

強く相互作用するフェルミ粒子系の熱力学

向山 敬 (Takashi Mukaiyama)



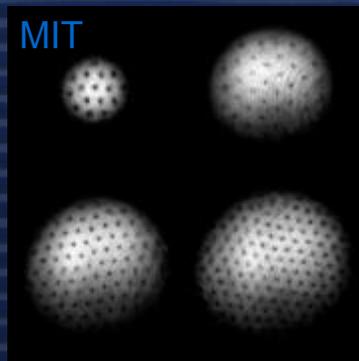
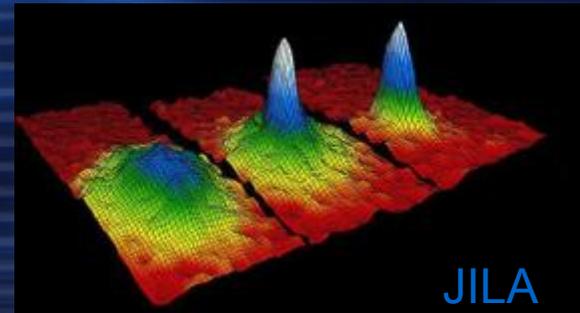
電気通信大学



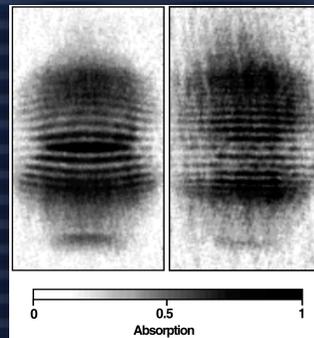
JST-ERATOマクロ量子制御プロジェクト

BEC in a cold atom system

1995 Realization of atomic gas Bose-Einstein condensation

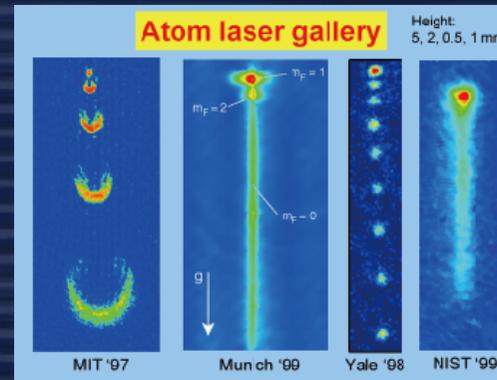


vortices

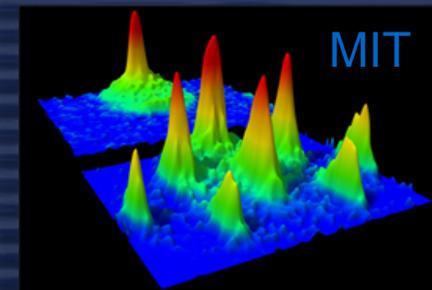


Science 275, 637 (1997).

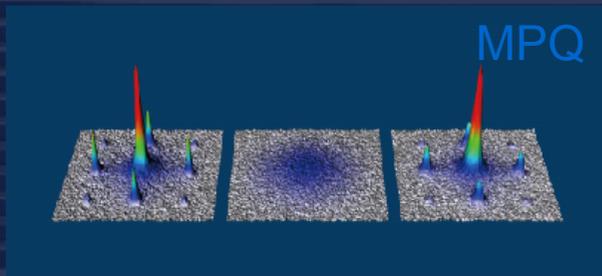
interference



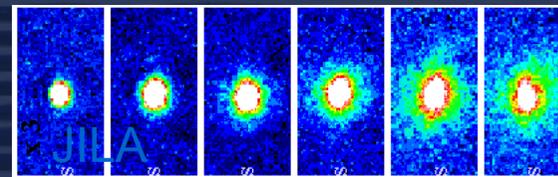
Atom laser



Super-radiance



Mott-insulator phase



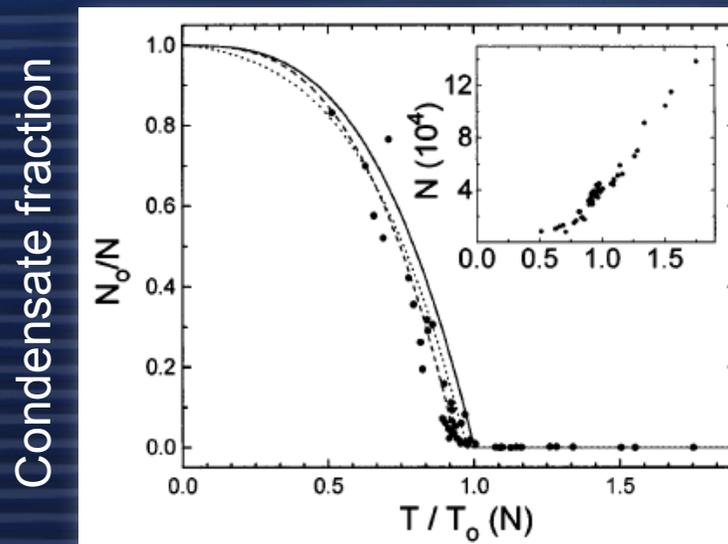
Bose nova

BEC in a cold atom system

Cold atoms are

- very dilute ($10^{11} \sim 10^{14} \text{ cm}^{-3}$),
- with no impurities, no defects.

➔ Amenable to simple theoretical description



J. R. Ensher, et al.,
Phys. Rev. Lett. **77**, 4984
(1996).

5% deviation of critical temperature
from theoretical predictions



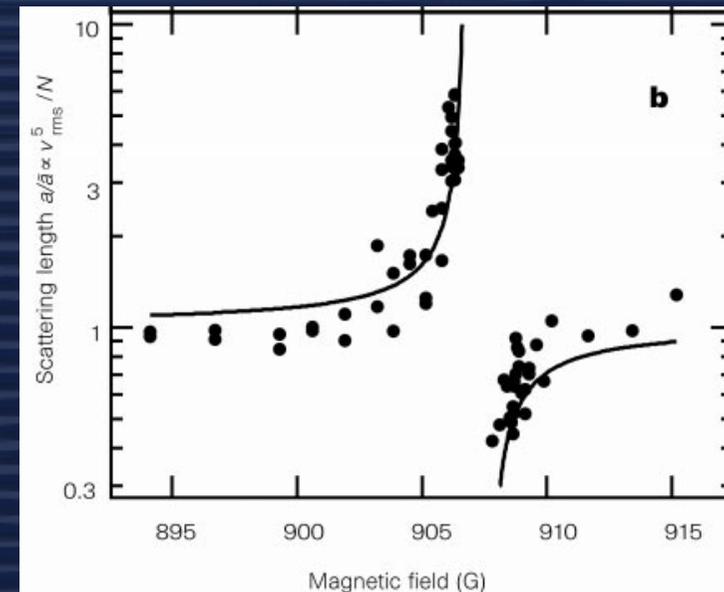
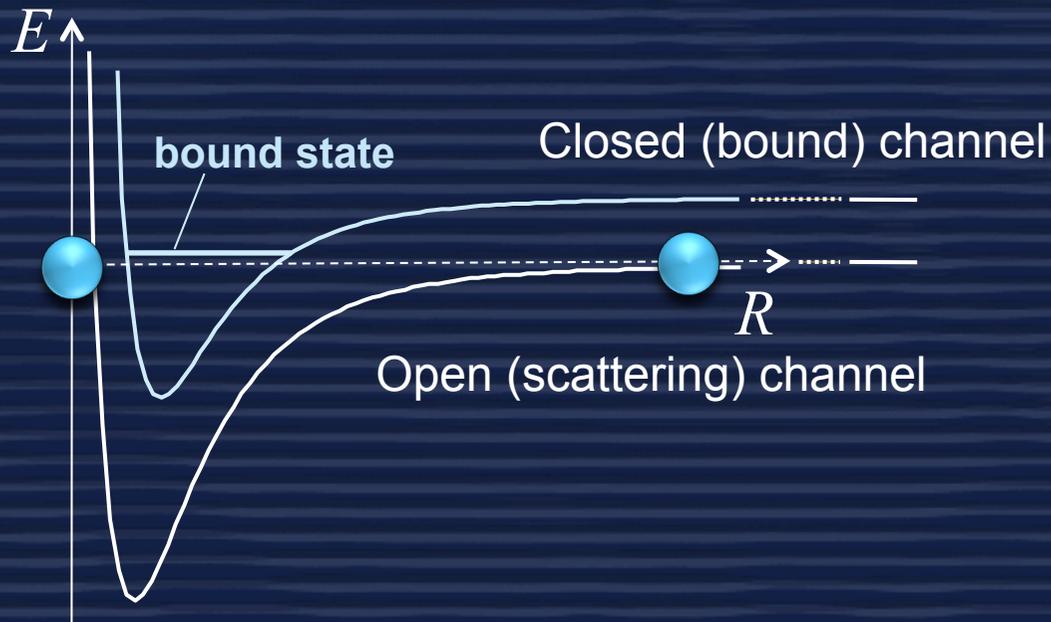
- 3% shift due to finite number correction
- 2% shift due to interaction

Inter-atomic interaction is tunable !!

Feshbach resonance

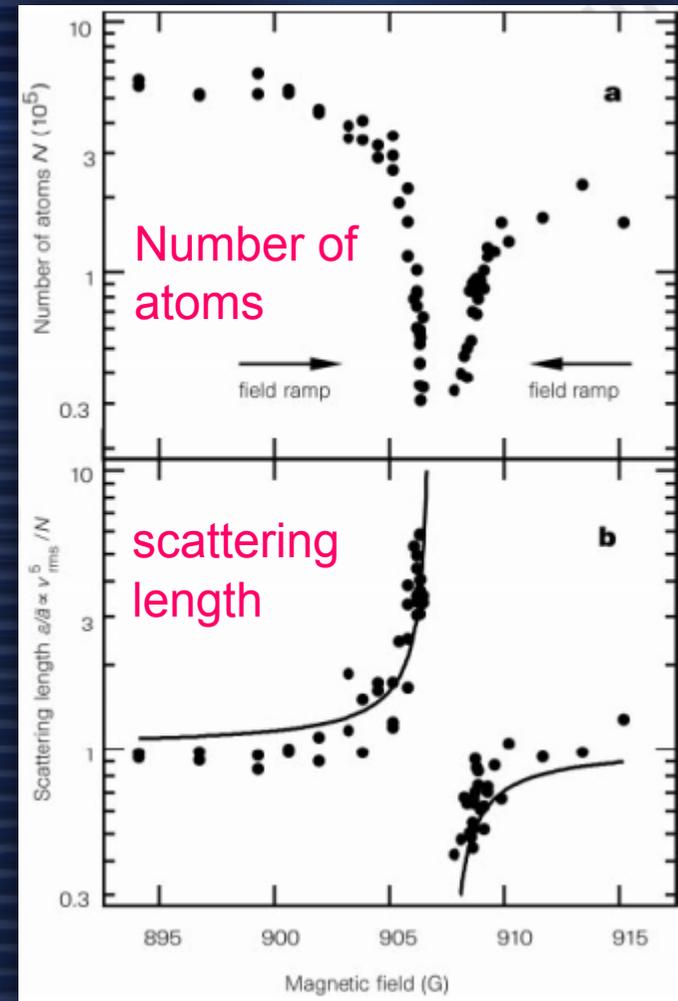
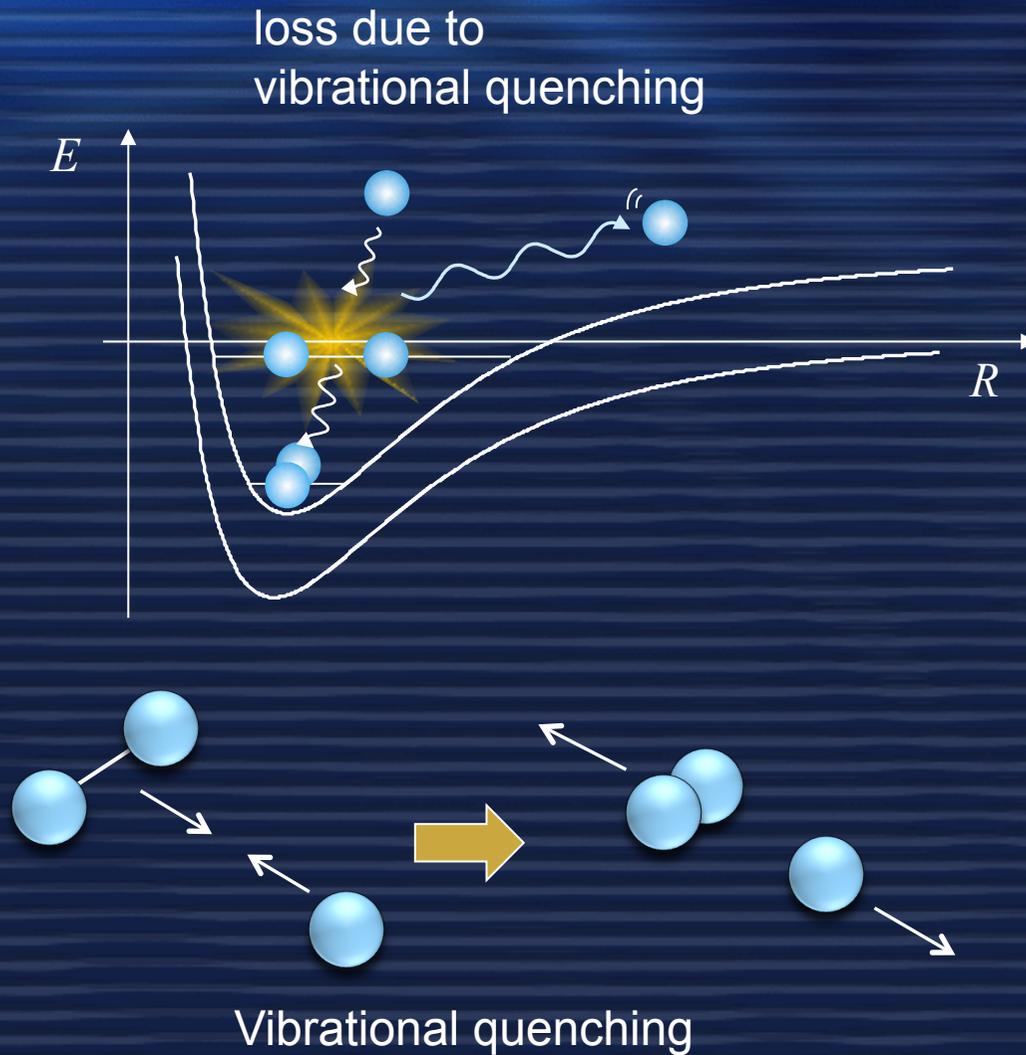
There are two channels corresponding to different spin states.

Resonance occurs when open and closed channel are energetically degenerate.



S. Inouye, et al.,
Nature **392**, 151 (1998).

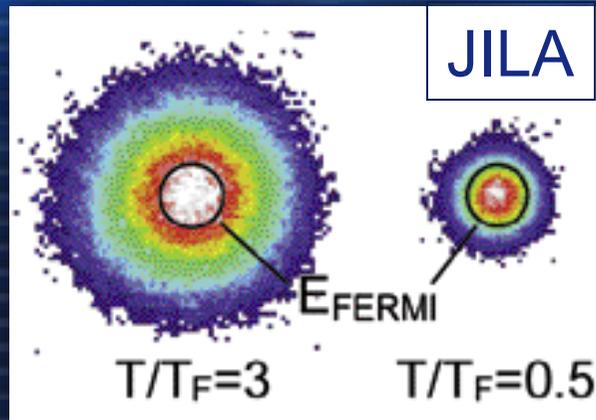
Loss near Feshbach resonance



S. Inouye et al.,
Nature, **392**, 151 (1998).

ultracold fermionic atoms

1999 Fermi degenerate gas



Collision channel

Identical bosons : $l=0$ (*s-wave*), $l=2$ (*d-wave*), ...

Identical fermions: $l=1$ (*p-wave*), $l=3$ (*f-wave*), ...

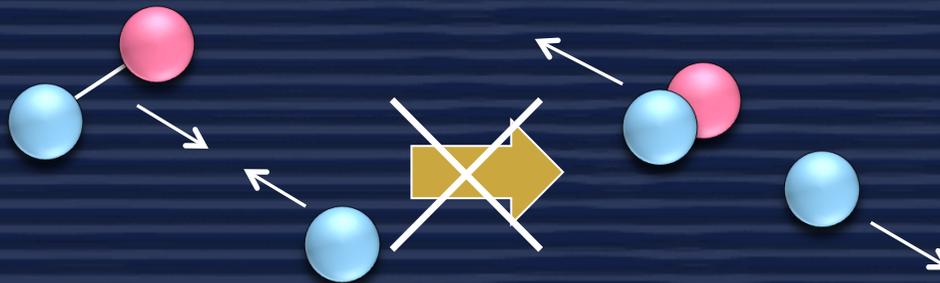
ultracold : *s-wave* is the dominant collision channel.

→ Identical fermions do not collide.

Think about two-component fermions



At the Feshbach resonance for  and , no loss occurs due to **Pauli exclusion principle**.

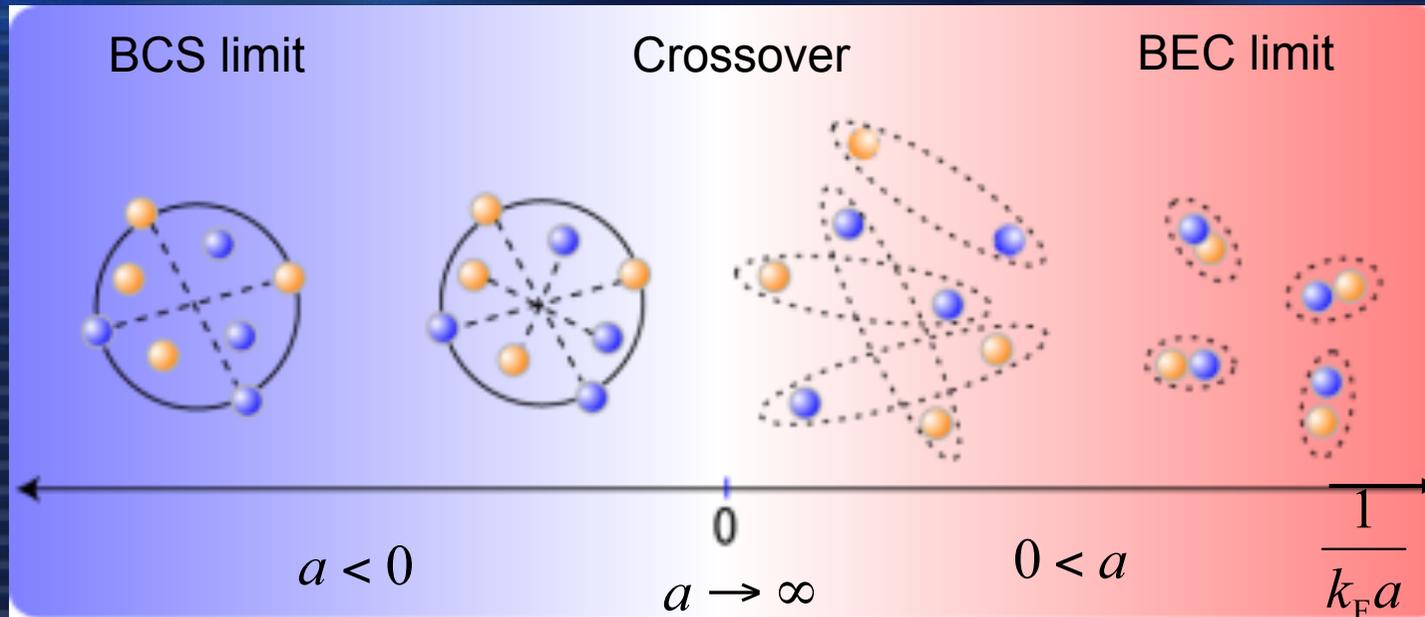


Therefore two-component fermions are stable even at a Feshbach resonance.

We are able to prepare an interacting
(reasonably stable) **two-component** Fermi gas of
atoms with an **arbitrary interaction strength !!**

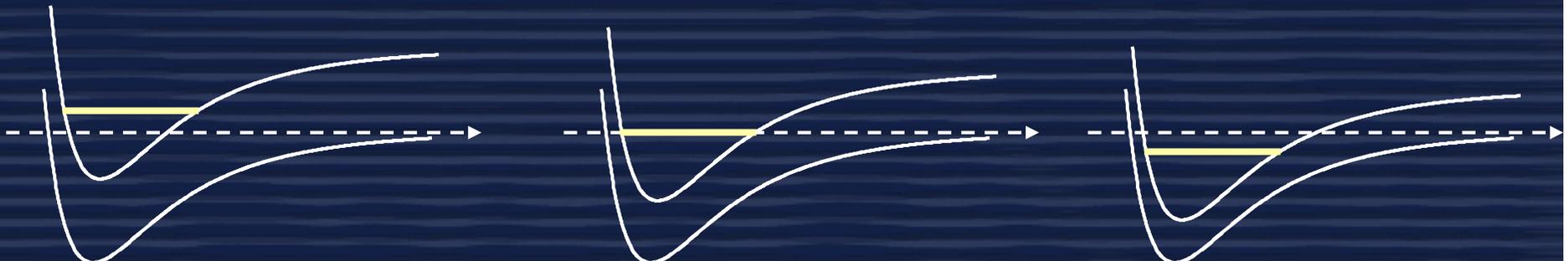
BCS-BEC crossover

Idea : BEC and BCS type superfluid are the opposite extreme of the same phase



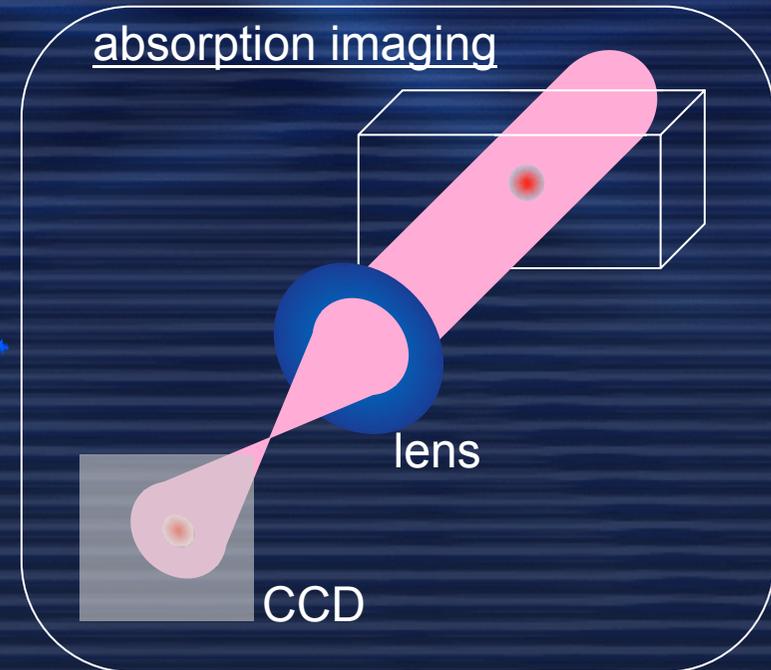
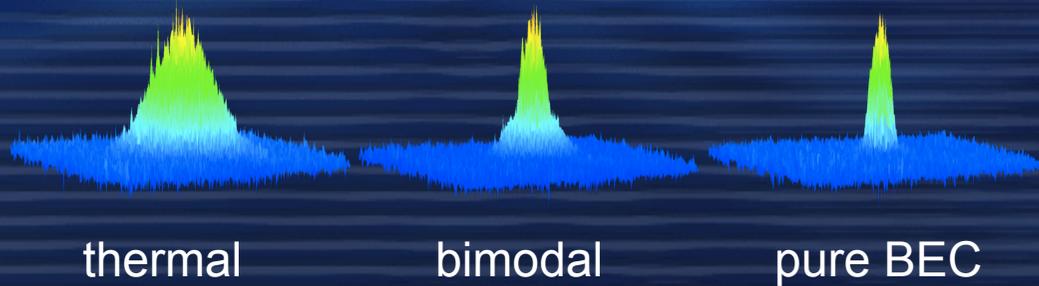
Pairs of fermions - Cooper pairs
Momentum space pairing

Tightly bound bosons - molecules
Real space pairing



momentum distribution measurement

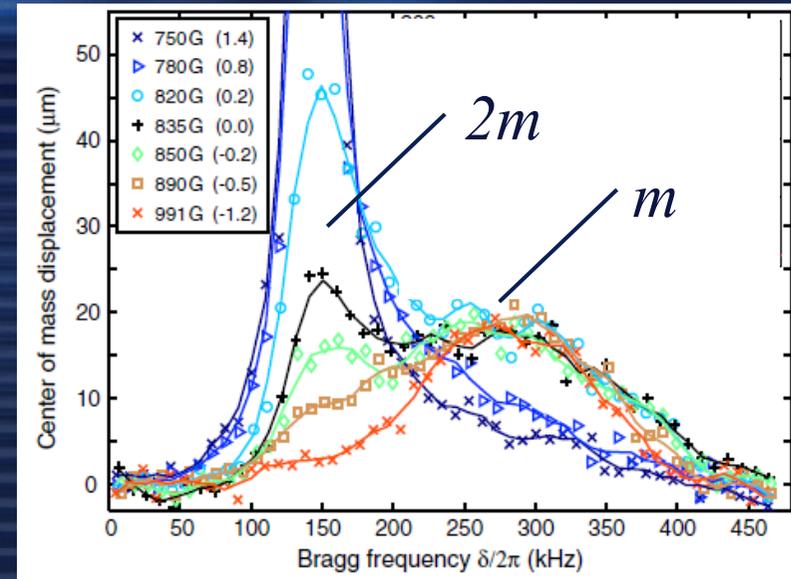
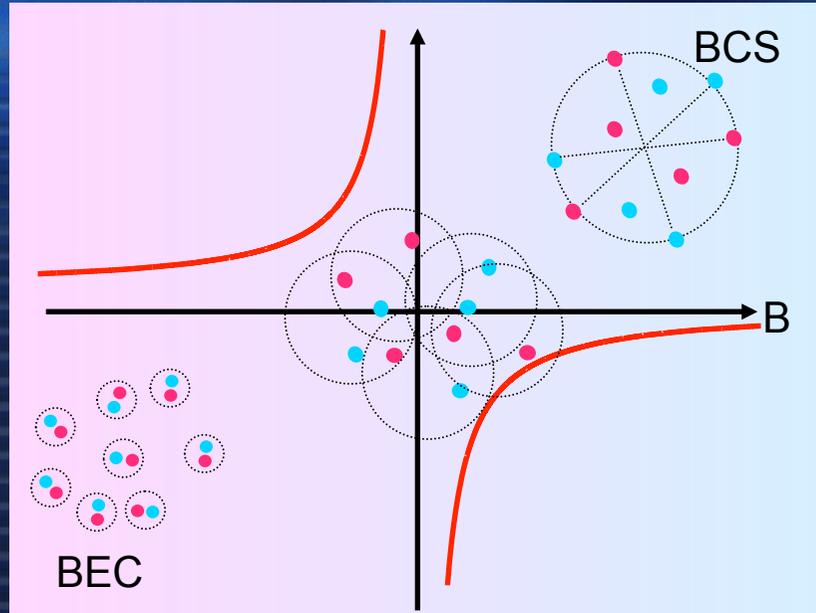
Bosonic case



Condensed or not?
See the bimodal profile !!

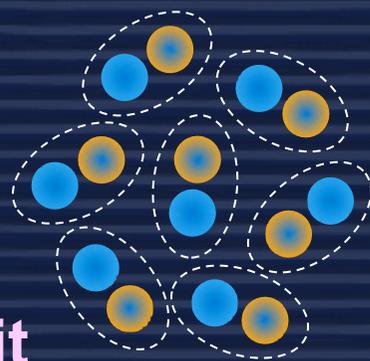
... unfortunately this scheme does not work.

Fermion pair condensate

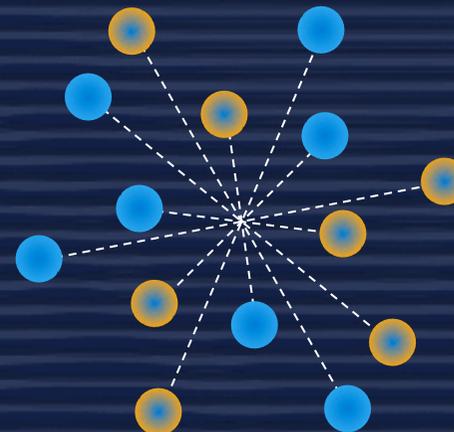


G. Veeravalli et al.
 Phys. Rev. Lett.
 101, 250403 (2008)

BEC limit



spatially
 correlated pair



BCS limit

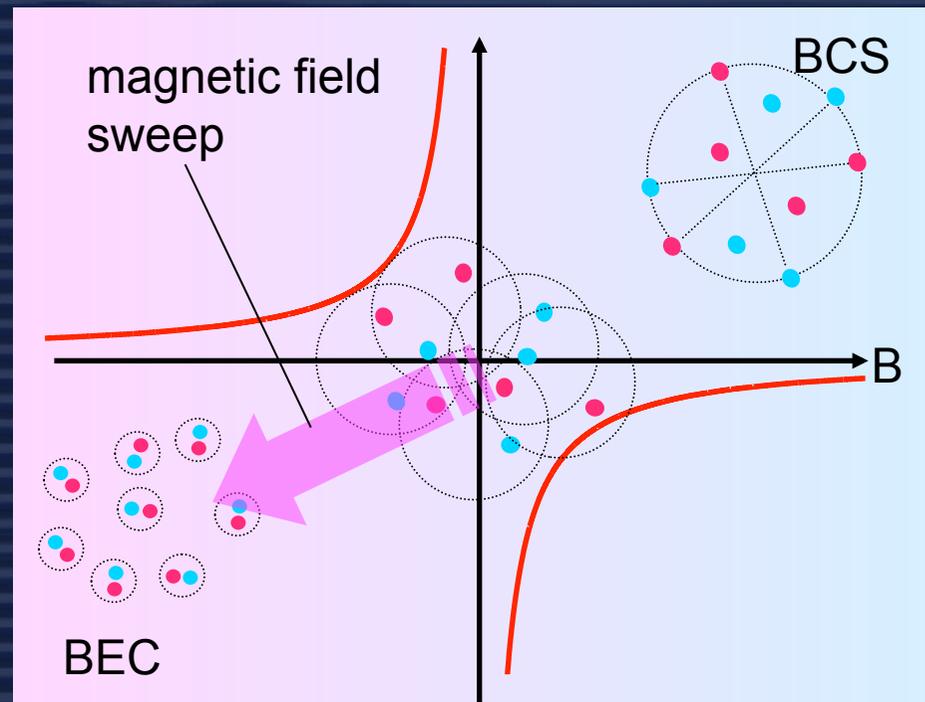
momentum
 correlated pair

“projection”

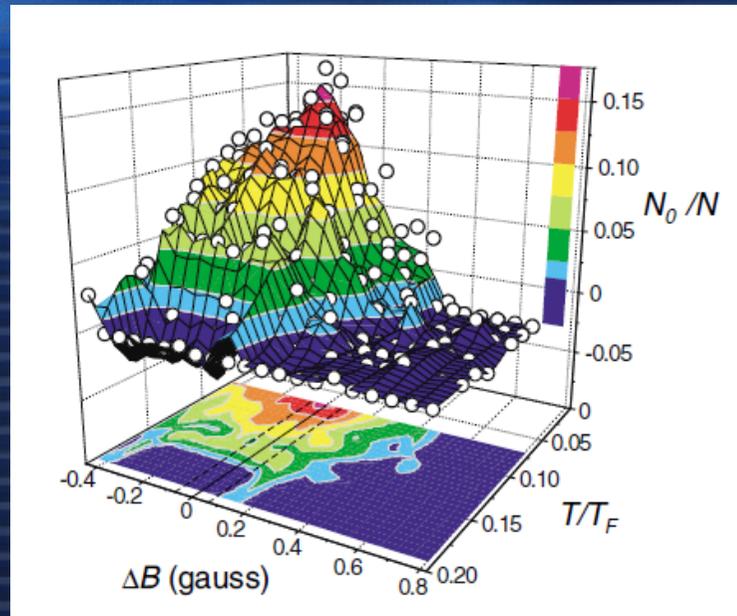
C. A. Regal et al.
Phys. Rev. Lett., **92**, 040403 (2004)

If we sweep the magnetic field

- **slow** enough to convert atom pairs into molecules
- **fast** enough such that the momentum distribution of the projected molecules reflects that of pairs prior to the sweep



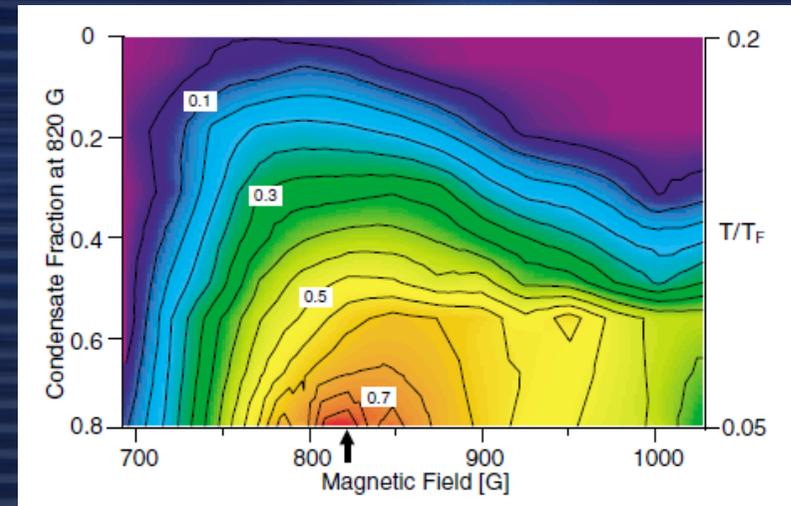
BCS-BEC crossover (experiment)



C. A. Regal *et al.*, PRL 92, 040403 (2004)

T/T_F is measured under ideal gas condition (magnetic field where $a=0$).

Adiabatic does keep entropy constant, but **not** T/T_F .

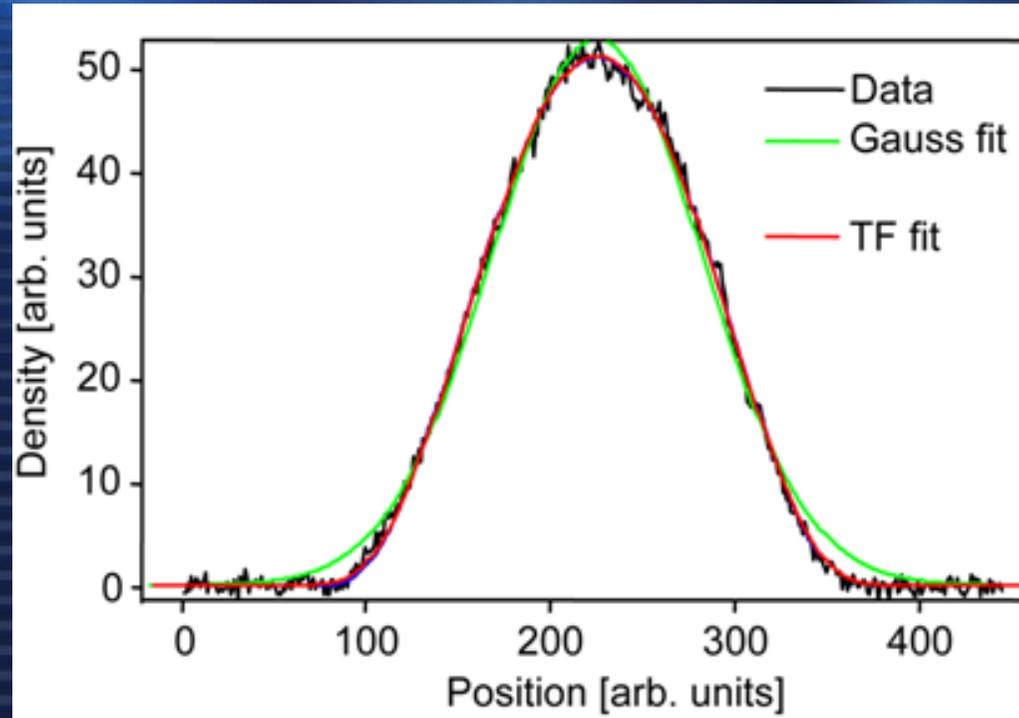


M. W. Zwierlein *et al.*, PRL 92, 120403 (2004)

T/T_F was measured at 1025 G (magnetic field where $a=-4000a_0$).

Again, T/T_F is not the one measured at unitarity, but somewhere different.

Conventional thermometry



Let the cloud expand.



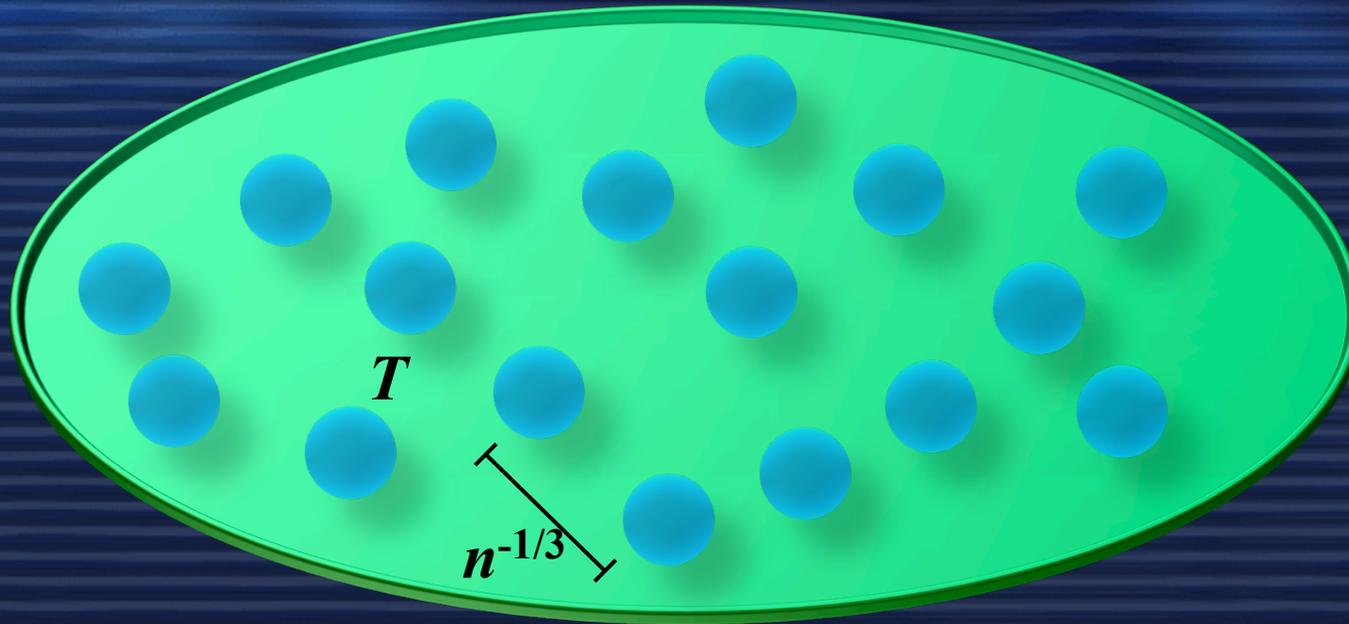
Fit the profile with a reasonable function.

$$n(z) = -\frac{3N}{\sqrt{\pi}\sigma_{TF,z}} \left(\frac{T}{T_F}\right)^{5/2} \cdot Li_{5/2} \left[-\xi \cdot \exp\left(-\frac{1}{2} \frac{z^2}{\sigma_z^2}\right) \right]$$

This scheme is good only when interaction energy \ll kinetic energy.

*Thermodynamic
of an ideal Fermi gas*

Thermodynamics of an ideal Fermi gas



Thermodynamic behavior of an ideal Fermi gas is described by its **temperature T** and **density n** .

Thermodynamic of an ideal Fermi gas

Fermi-Dirac distribution

$$n(\varepsilon) = \frac{1}{z^{-1} e^{\beta\varepsilon} + 1} \quad \left(z \equiv e^{\beta\mu}, \beta = (k_B T)^{-1} \right)$$

$$N = \int_0^\infty \frac{D(\varepsilon)}{z^{-1} e^{\beta\varepsilon} + 1} d\varepsilon \quad \longrightarrow \quad \left(\frac{k_B T}{E_F} \right)^{-3/2} = -\frac{3\sqrt{\pi}}{4} Li_{3/2} \left[-\exp \left(\frac{\mu/E_F}{k_B T/E_F} \right) \right]$$

$\left(-Li_s(-z) = \frac{1}{\Gamma(s)} \int_0^\infty \frac{t^{s-1}}{z^{-1} e^t + 1} dt \right)$

$$\frac{\mu}{E_F} = f_\mu \left(\frac{k_B T}{E_F} \right)$$

Thermodynamic of an ideal Fermi gas

$$E = \int_0^{\infty} \frac{\varepsilon D(\varepsilon)}{z^{-1} e^{\beta\varepsilon} + 1} d\varepsilon$$

$$\frac{E}{NE_F} = -\frac{3\sqrt{\pi}}{4} \left(\frac{k_B T}{E_F} \right)^{5/2} Li_{5/2}(-z)$$


$$\frac{E}{NE_F} = f_E \left(\frac{k_B T}{E_F} \right)$$

Other thermodynamic functions also have this similarity.

$$\frac{S}{k_B} = f_S \left(\frac{k_B T}{E_F} \right)$$

$$\frac{F}{NE_F} = f_F \left(\frac{k_B T}{E_F} \right)$$

Thermodynamic of an ideal Fermi gas

Internal energy : $\frac{E}{NE_F} = f_{E,ideal} \left(\frac{k_B T}{E_F} \right)$

Helmholtz free energy : $\frac{F}{NE_F} = f_{F,ideal} \left(\frac{k_B T}{E_F} \right)$

Chemical potential : $\frac{\mu}{E_F} = f_{\mu,ideal} \left(\frac{k_B T}{E_F} \right)$

Entropy : $\frac{S}{Nk_B} = f_{S,ideal} \left(\frac{k_B T}{E_F} \right)$

Dimensionless
functions

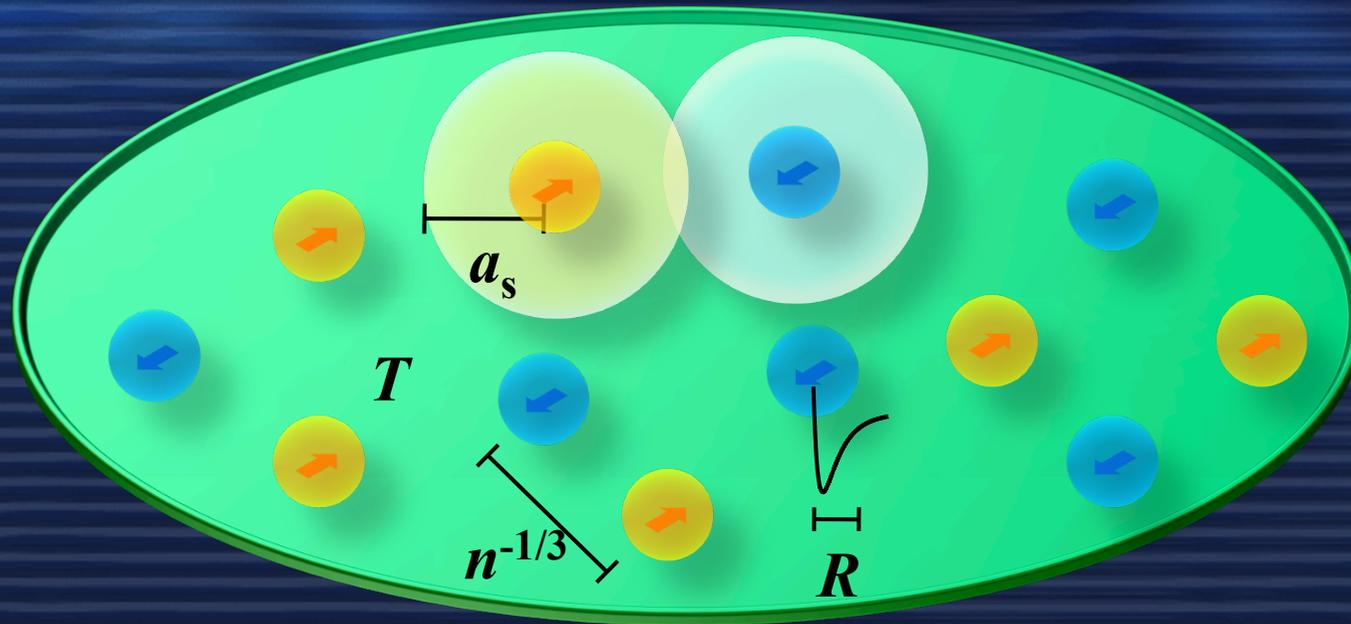
Material specific parameter, such as m , is taken up by E_F (T_F).
(Shape of the functions do not depend on the particle's nature.)



Universal thermodynamics

*Thermodynamic
of an interacting Fermi gas*

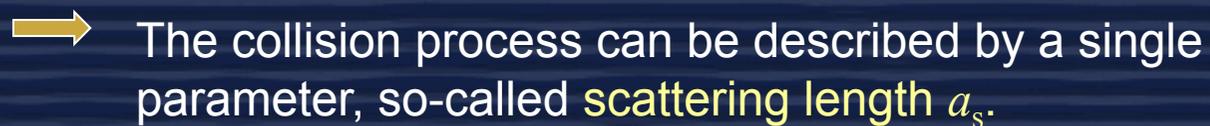
Ultracold, dilute, interacting Fermi gases



- ultracold : s-wave is the dominant channel.



- dilute : details of the potential is much smaller than $n^{-1/3}$



Thermodynamic of an interacting Fermions

Ideal Fermi gas

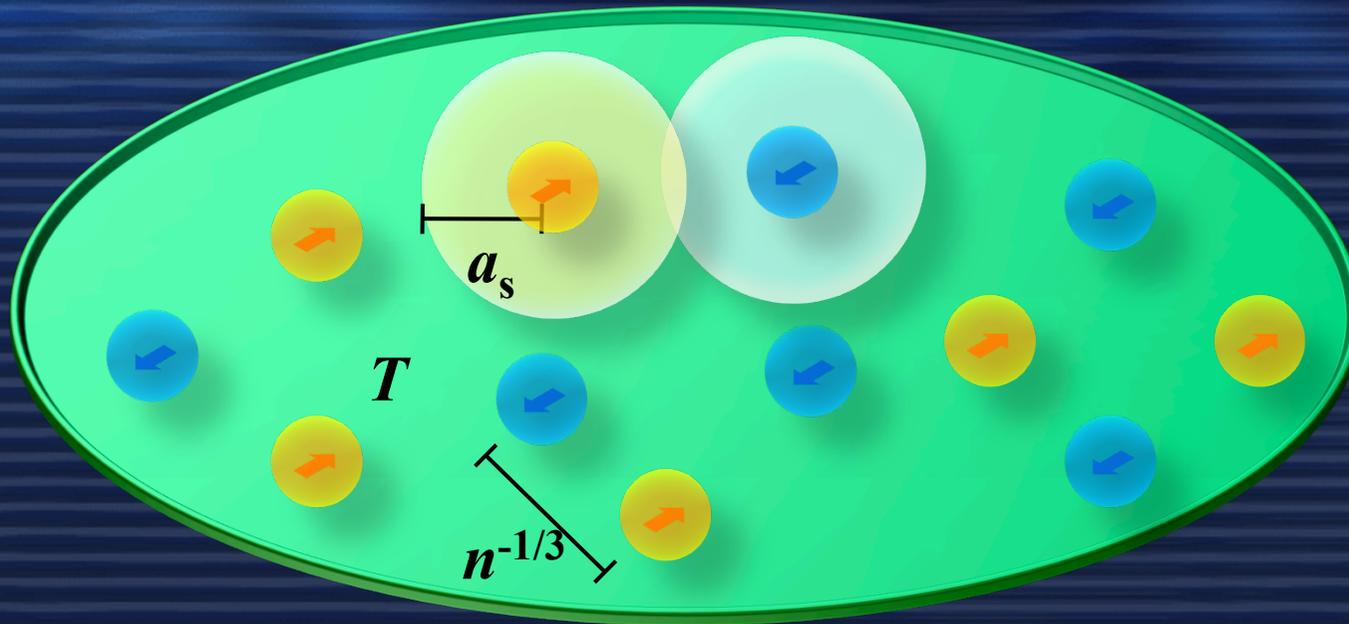
$$\frac{E}{NE_F} = f_{E,ideal} \left(\frac{k_B T}{E_F} \right)$$



Fermi gas with interaction

$$\frac{E}{NE_F} = f_E \left(k_B T, E_F, E_{\text{int}}(a_s) \right)$$

Ultracold dilute Fermi gas



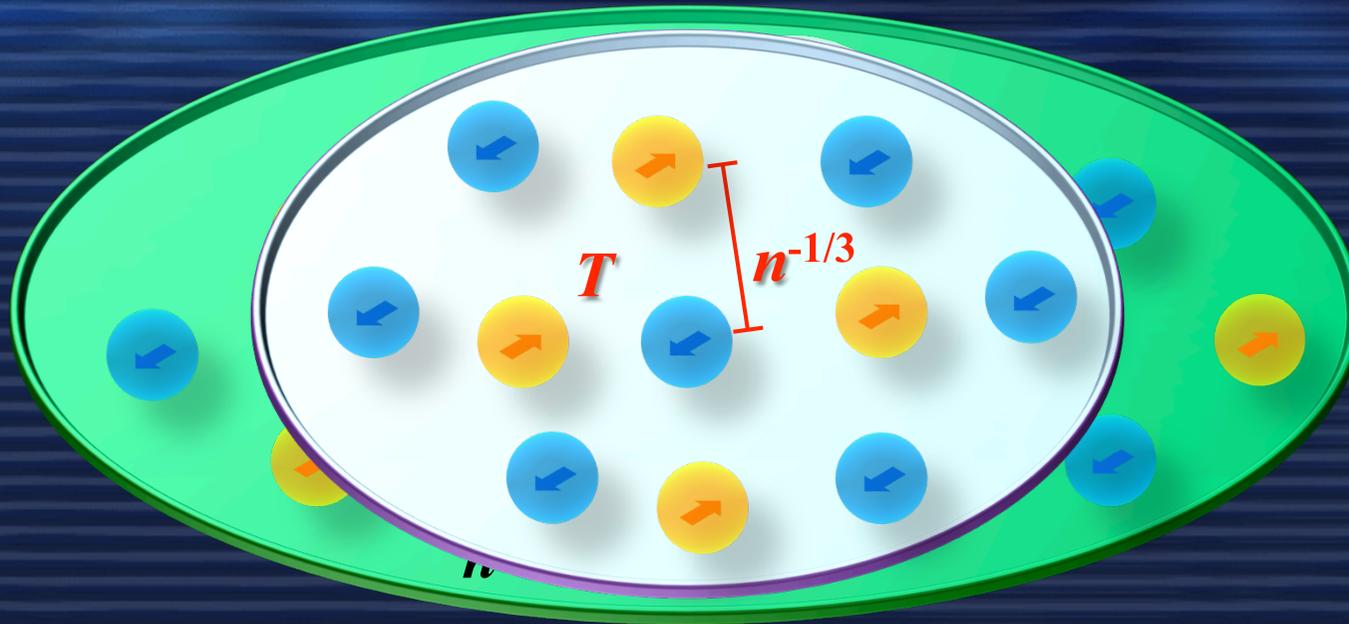
Remember the fact that a_s *is tunable!!*

Then, what happens when...

$$|a_s| \longrightarrow \infty$$

This situation is called unitarity limit.

Unitarity limit and Universality



a_s drops out of the description of the thermodynamics.

Thermodynamics depends only on the **density n** and **temperature T** .

Universal thermodynamics holds again...?

Thermodynamic of an interacting Fermions

Ideal Fermi gas

$$\frac{E}{NE_F} = f_{E,ideal} \left(\frac{k_B T}{E_F} \right)$$

Fermi gas with interaction

$$\frac{E}{NE_F} = f_E (k_B T, E_F, U(a))$$

When the scattering length
diverges...

$$\frac{E}{NE_F} = f_E (k_B T, E_F, \cancel{U(a)}) \Rightarrow f_{E,|a|=\infty} (k_B T, E_F) = f_{E,|a|=\infty} \left(\frac{k_B T}{E_F} \right)$$

There is a hypothesis that the thermodynamic functions
again have **the universal form**.

➔ **Universal hypothesis**

Universal thermodynamics

According to **universal hypothesis**, all thermodynamics should obey the universal functions:

$$\text{Internal energy : } \frac{E}{NE_F} = f_E \left(\frac{k_B T}{E_F} \right)$$

$$\text{Helmholtz free energy : } \frac{F}{NE_F} = f_F \left(\frac{k_B T}{E_F} \right)$$

$$\text{Chemical potential : } \frac{\mu}{E_F} = f_\mu \left(\frac{k_B T}{E_F} \right)$$

$$\text{Entropy : } \frac{S}{Nk_B} = f_S \left(\frac{k_B T}{E_F} \right)$$

Dimensionless universal functions, (shape of the function is different from those for an ideal gas)

System looks like a non-interacting Fermi gas.
(no other dimensional parameters involved in the problem)

Universal thermodynamics

Bertsch's Many-Body X challenge, Seattle, 1999

What are the ground state properties of the many-body system composed of spin $\frac{1}{2}$ fermions interacting via a zero-range, **infinite scattering-length** contact interaction.

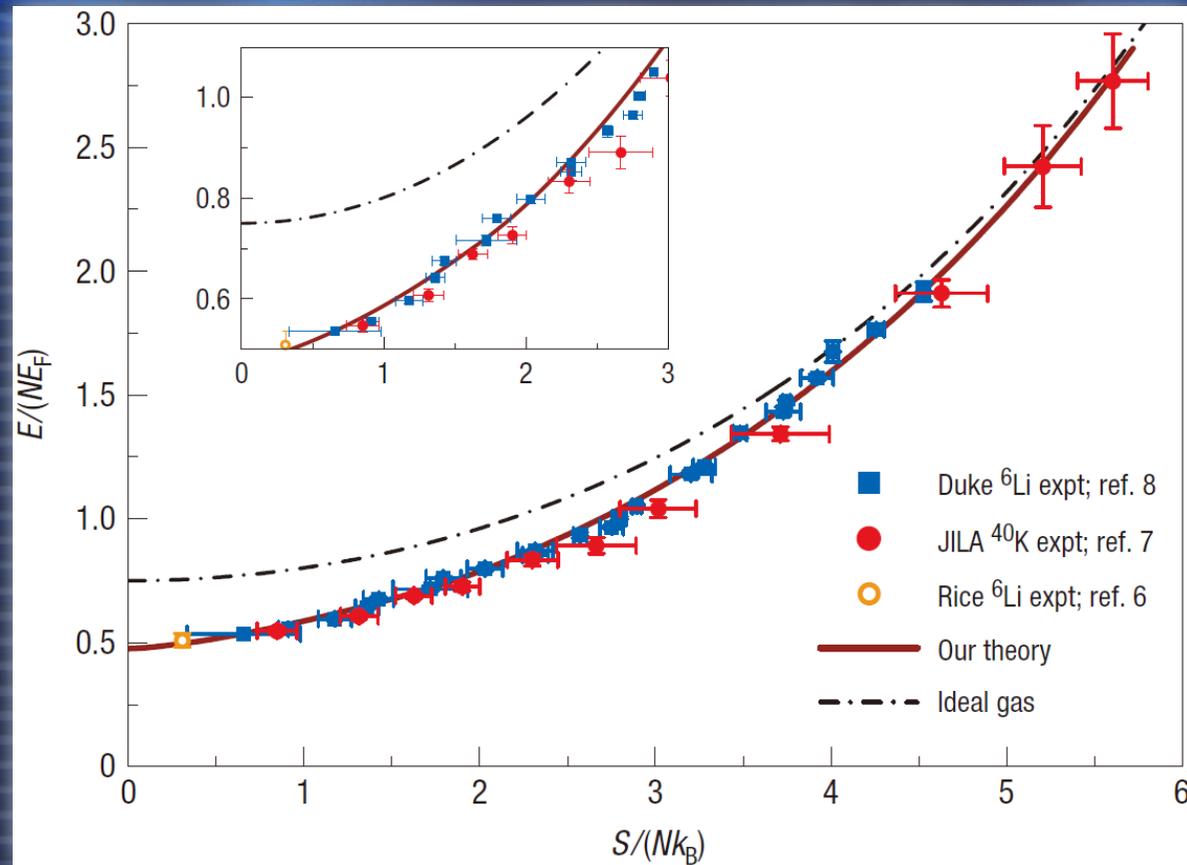
$$E_{gs} = f(N, V, m) = N \cdot E_F \times \xi \leftarrow \text{pure number}$$

Besides pure theoretical curiosity, this problem is relevant to neutron stars!



Universal thermodynamics

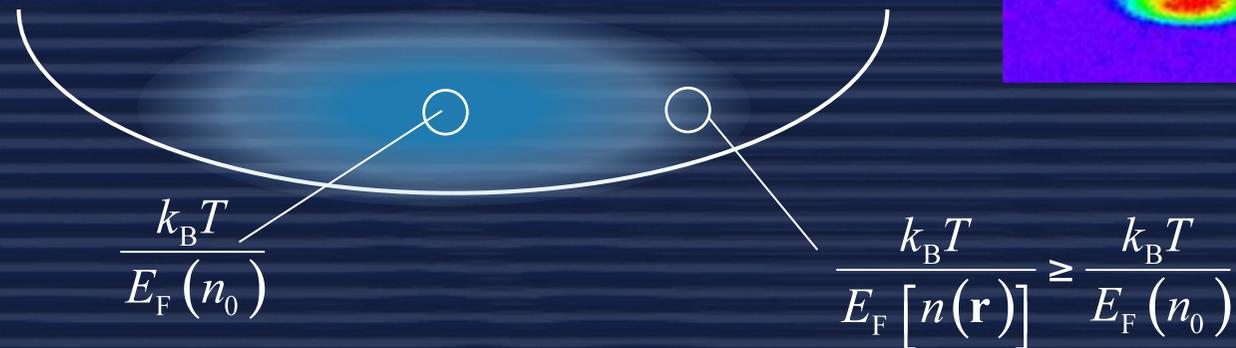
H. Hu, P. D. Drummond & X.-J. Liu,
Nature Physics 3, 469 - 472 (2007)



$f_E \left(\frac{k_B T}{E_F} \right)$ is still not known...

T is constant over the cloud (thermal equilibrium).
 E_F depends on the position (local density).

→ $\frac{k_B T}{E_F}$ is position-dependent.



Global measurement only gives the integration of all the different phases.

Goal of this experiment

Measurement of **local** thermodynamic quantities

and

the determination of the universal thermodynamic function.

$$\frac{E}{NE_F} = f_E \left(\frac{k_B T}{E_F} \right) \quad \Rightarrow \quad \frac{\varepsilon(\mathbf{r})}{n(\mathbf{r})E_F[n(\mathbf{r})]} = f_E \left(\frac{T}{T_F[n(\mathbf{r})]} \right)$$

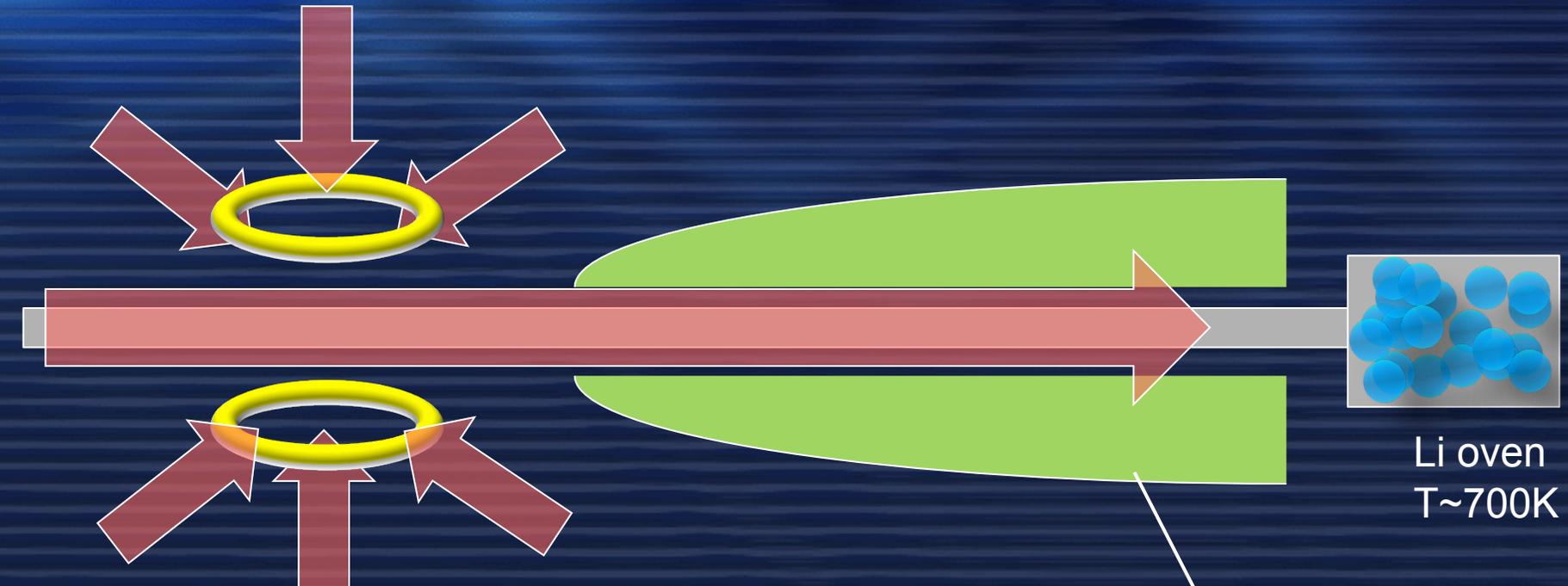
ε : local energy density

$$E_F = k_B T_F$$

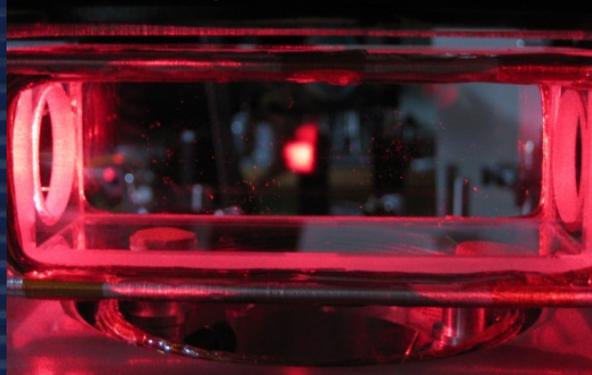
Experiment setup



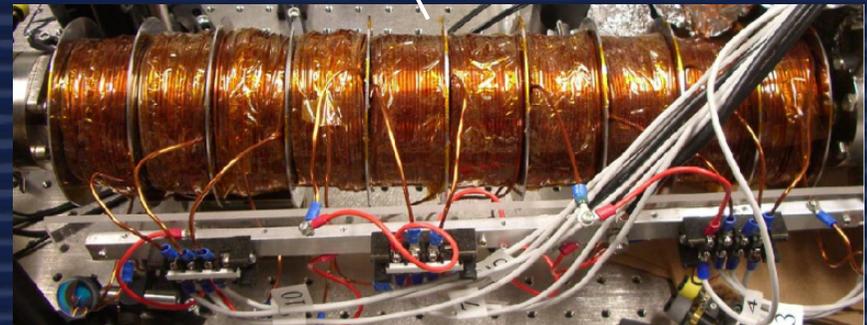
Deceleration and trapping



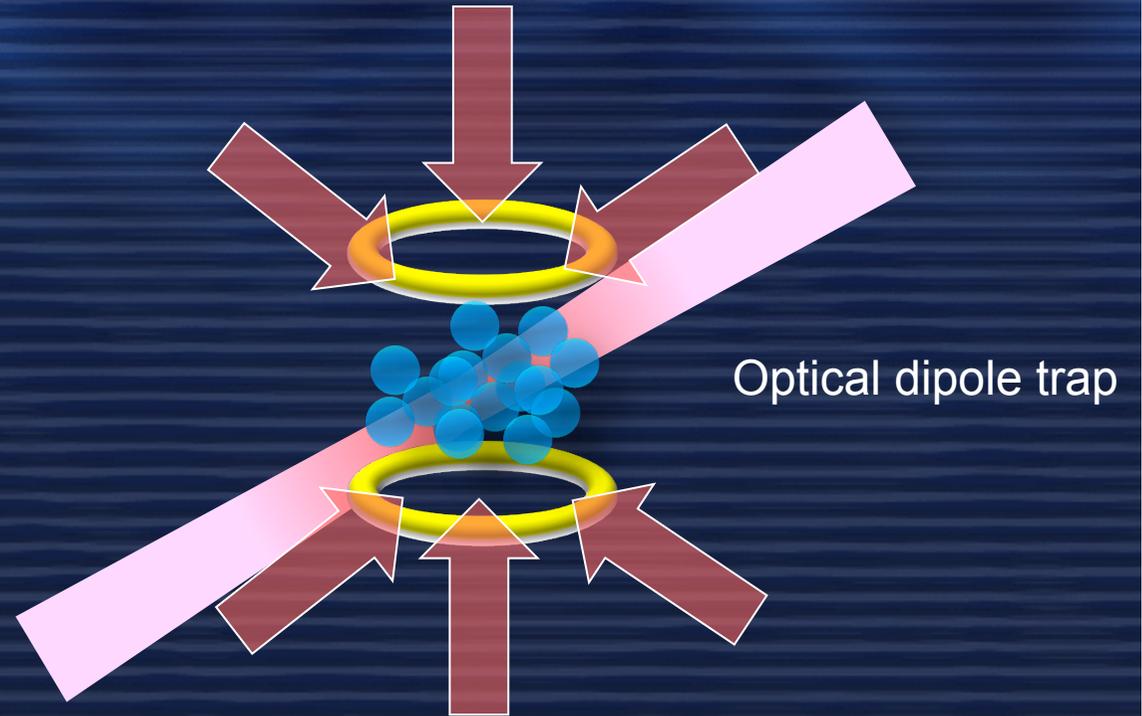
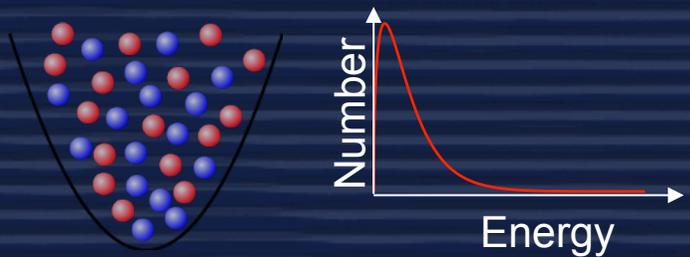
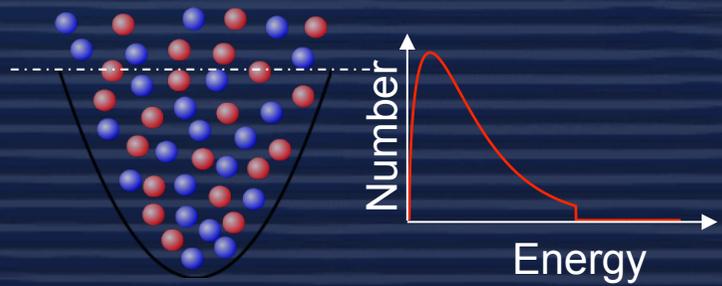
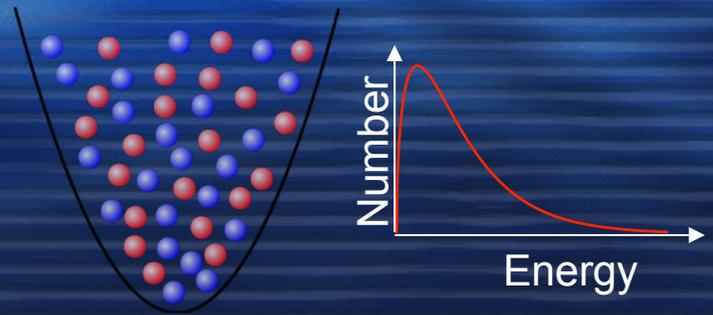
MOT (magneto-optical trap)



$T = 200 \mu\text{K}$
 $N > 10^8$



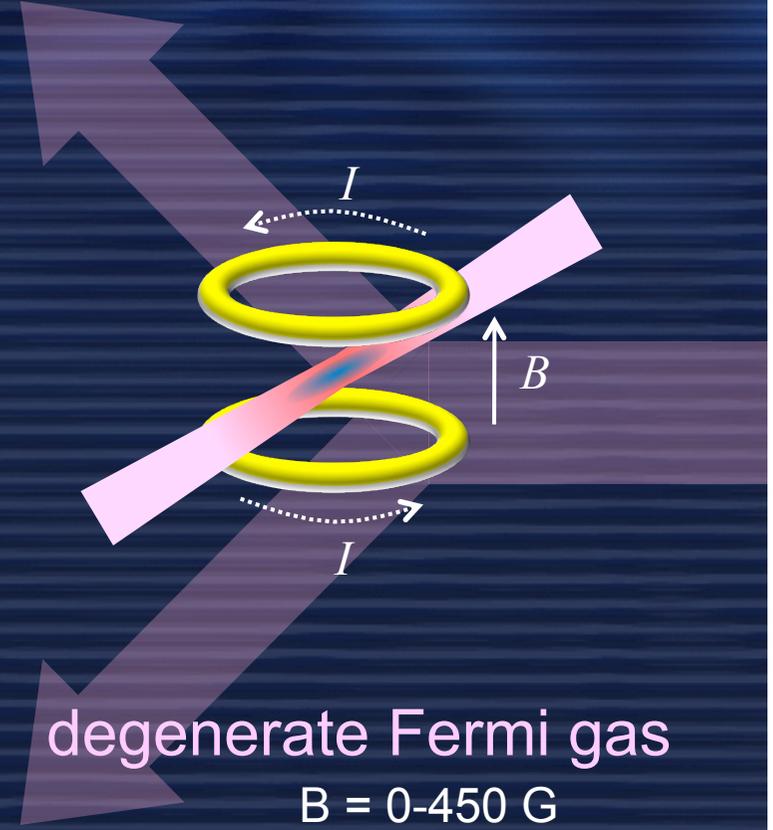
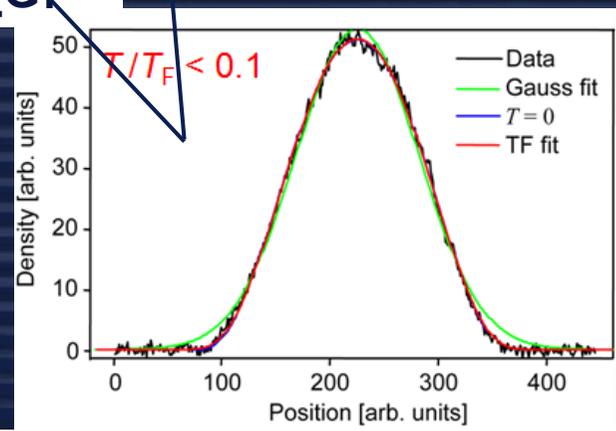
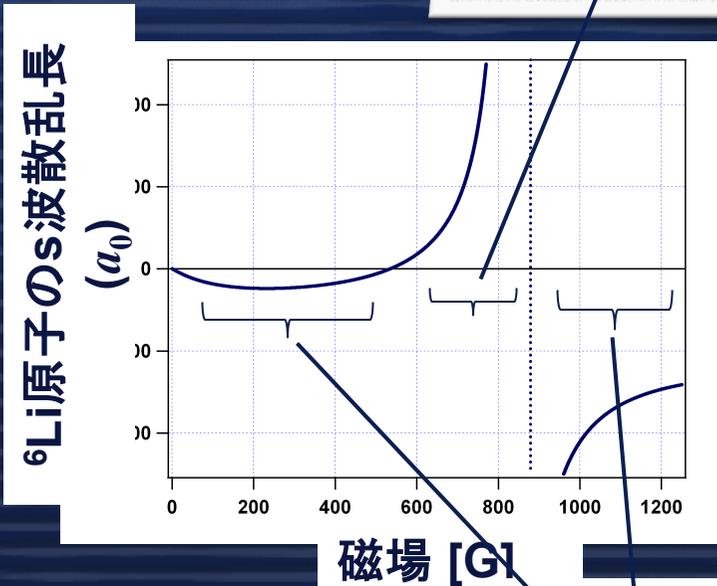
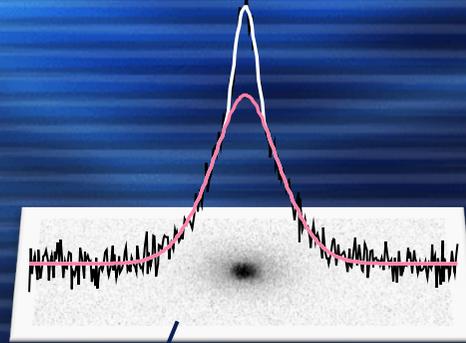
Optical dipole trap

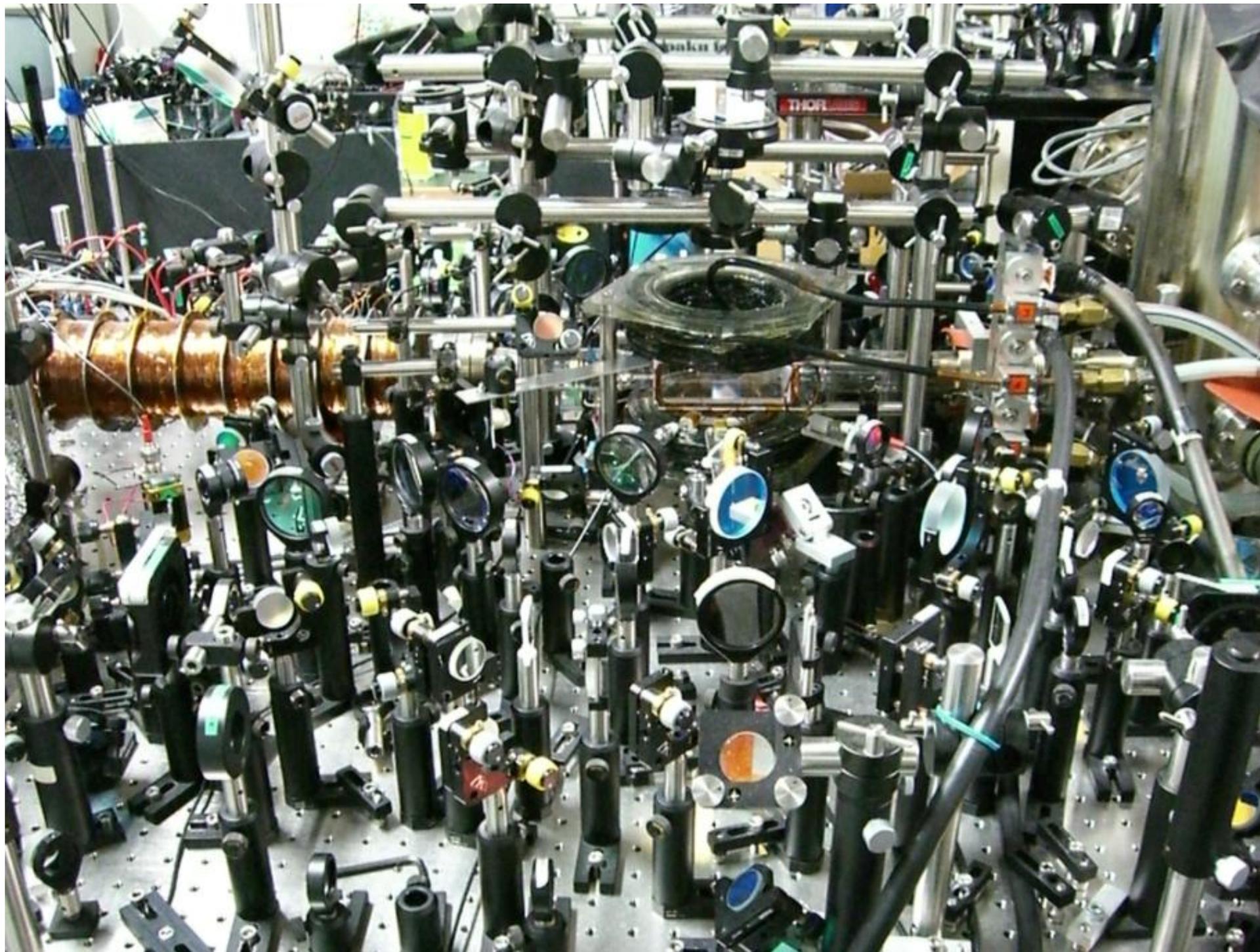


Evaporative cooling

molecular BEC

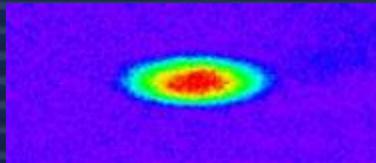
$B = 650 - 800 \text{ G}$





Determination of local energy $\varepsilon(\mathbf{r})$

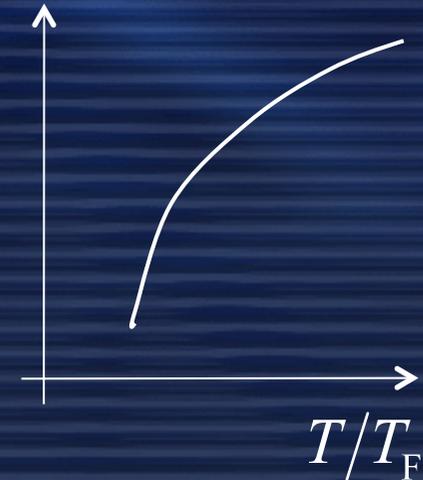
$$\frac{\varepsilon(\mathbf{r})}{n(\mathbf{r}) E_F[n(\mathbf{r})]} = f_E[T/T_F]$$



density profile

$n(\mathbf{r})$

$f_E[T/T_F]$



Useful equations :

- Equation of state of unitary gas : $p(\mathbf{r}) = \frac{2}{3} \varepsilon(\mathbf{r})$
- mechanical equilibrium (eq. of force balance) :



$$\nabla p(\mathbf{r}) + n(\mathbf{r}) \nabla V_{\text{Trap}}(\mathbf{r}) = 0$$

$$n(\mathbf{r}) \Rightarrow p(\mathbf{r}) \Rightarrow \varepsilon(\mathbf{r})$$

Determination of temperature T

$$\frac{\varepsilon(\mathbf{r})}{n(\mathbf{r}) E_F [n(\mathbf{r})]} = f_E [T/T_F]$$

$$p(\mathbf{r}) = \frac{2}{3} \varepsilon(\mathbf{r}) \text{ and } \nabla p(\mathbf{r}) + n(\mathbf{r}) \nabla V_{\text{Trap}}(\mathbf{r}) = 0 \implies E_{\text{total}} = 2 \times E_{\text{potential}}$$

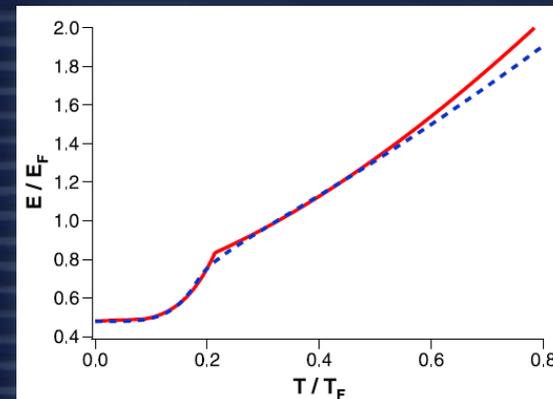
Adiabatic B-field sweep to turn off the interaction

\implies entropy S

$E_{\text{total}} \text{ vs } S$

$\implies E_{\text{total}} \text{ vs } T$

$$\uparrow \frac{1}{T} = \partial S / \partial E$$



Le Luo and J.E. Thomas,
J Low Temp Phys **154**, 1 (2009).

Our scheme

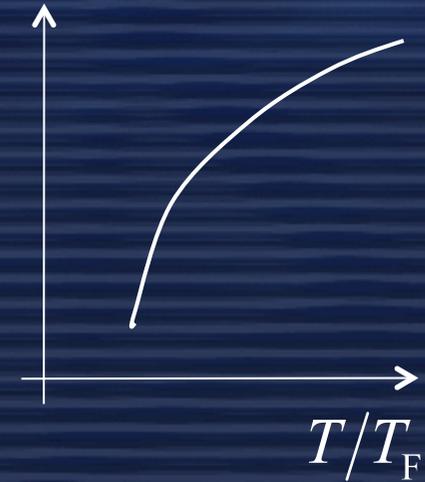
$$\frac{\varepsilon(\mathbf{r})}{n(\mathbf{r}) E_F [n(\mathbf{r})]} = f_E [T/T_F]$$

$$p = \frac{2}{3} \varepsilon$$

$$\nabla p(\mathbf{r}) + n(\mathbf{r}) \nabla V_{\text{Trap}}(\mathbf{r}) = 0$$

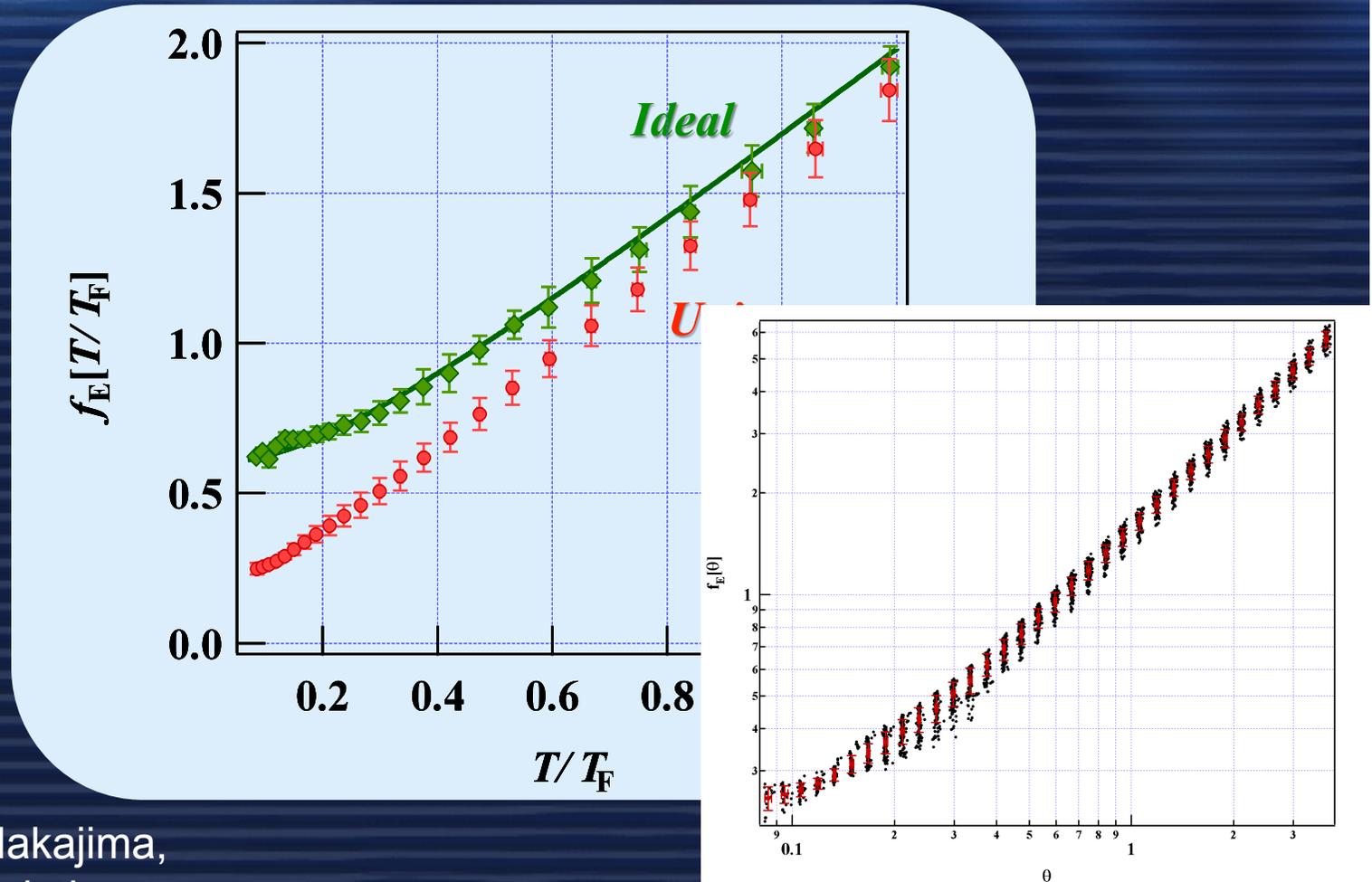


$$f_E [T/T_F]$$



Experimental determination of $f_E [T/T_F]$

$$\frac{\varepsilon(\mathbf{r})}{n(\mathbf{r}) E_F [n(\mathbf{r})]} = f_E [T/T_F]$$



M. Horikoshi, S. Nakajima,
M. Ueda and T. Mukaiyama,
Science, **327**, 442 (2010).

About 800 images are analyzed.

Verification of the determined $f_E [T/T_F]$

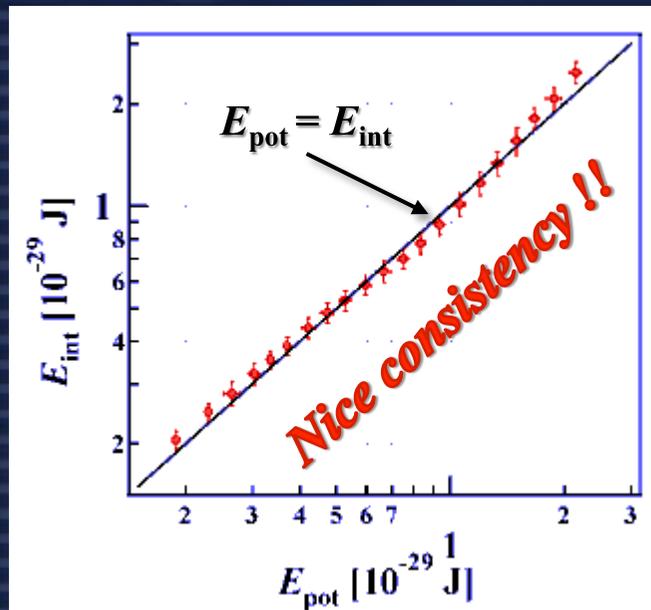
1. Energy comparison

$$E_{\text{total}} = 2 \times E_{\text{potential}} \longrightarrow E_{\text{pot}} = E_{\text{internal}}$$

Comparison

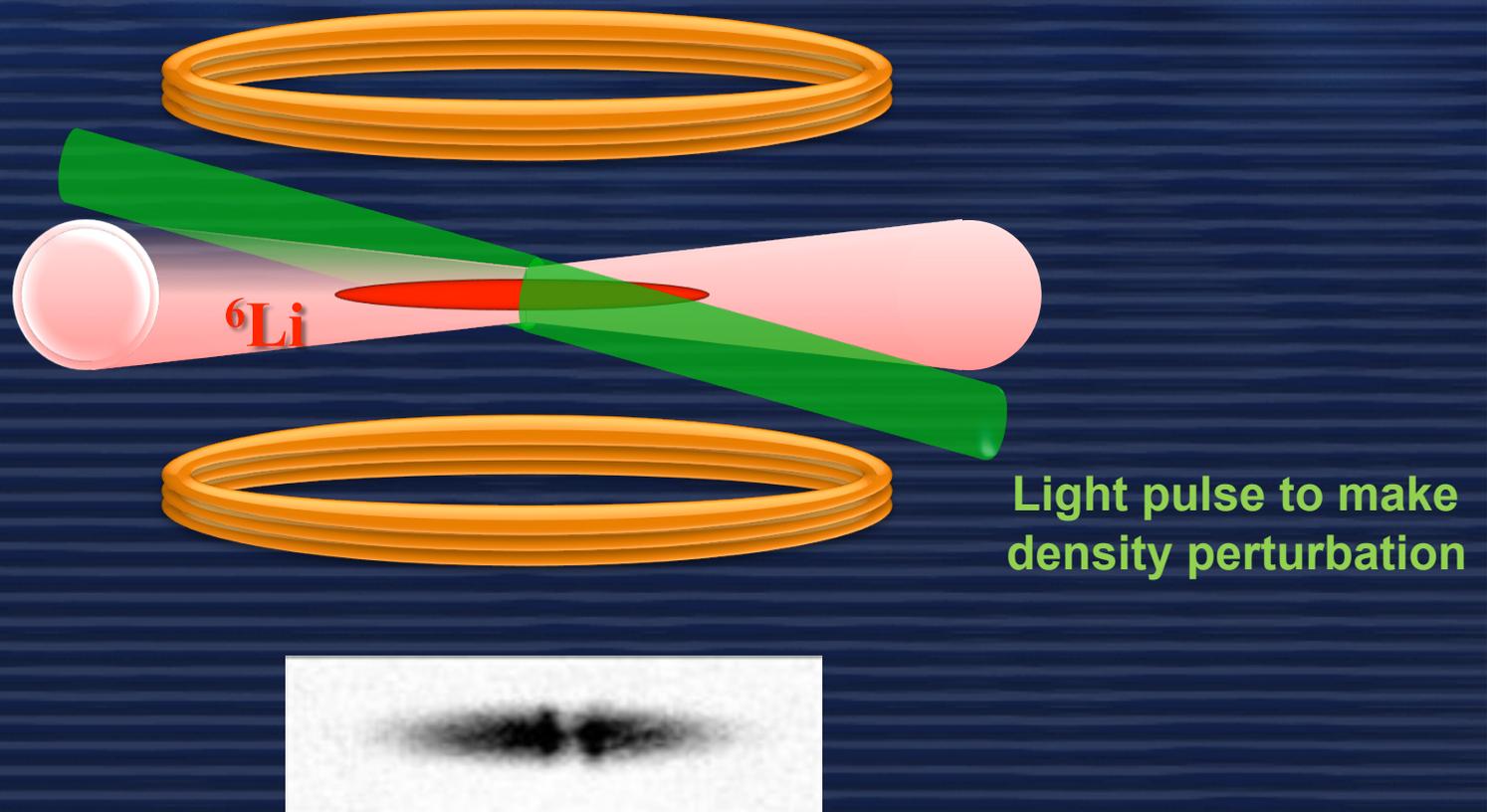
✓ Potential energy par particle : $E_{\text{pot}} = \frac{3}{2} m \omega_z^2 \langle z^2 \rangle$

✓ Internal energy par particle : $E_{\text{internal}} = \int n \varepsilon_F(n) \underline{f_E[\theta]} dV / N$



Verification of the determined $f_E [T/T_F]$

2. Effective speed of the first sound



Verification of the determined $f_E [T/T_F]$

2. Effective speed of the first sound

Propagation time

0.1ms

1.1ms

2.1ms

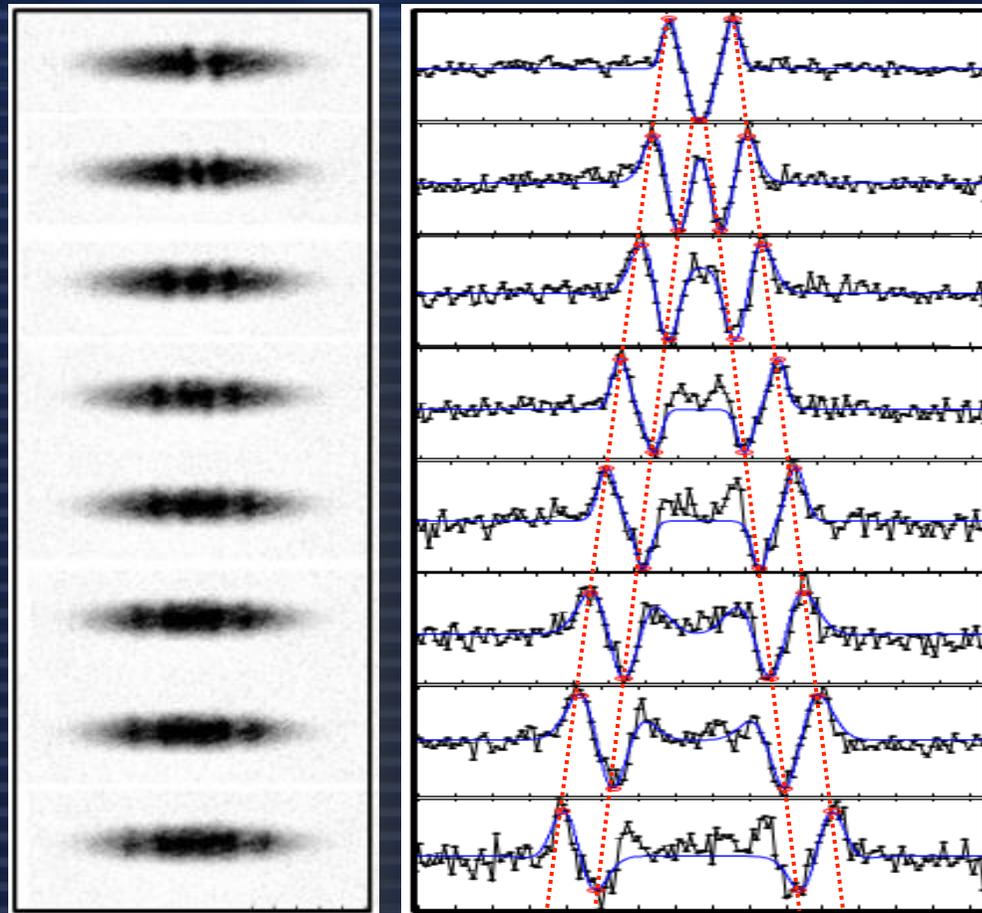
3.1ms

4.1ms

5.1ms

6.1ms

7.1ms



Verification of the determined $f_E [T/T_F]$

2. Effective speed of the first sound

Unitary gas shows hydrodynamic behavior due to the large collision rate

Effective speed of the first sound : $\bar{u}_1^2 [n, \theta] = \frac{\int \int \dot{n} \, dx dy}{m \int \int \dot{n} \left(\frac{\partial p}{\partial n} \right)^{-1} dx dy} \Bigg|_{z=0}$

$$p = \frac{2}{3} \varepsilon \propto f_E [T/T_F]$$

[P. Capuzzi, PRA 73, 021603(R) (2006)]

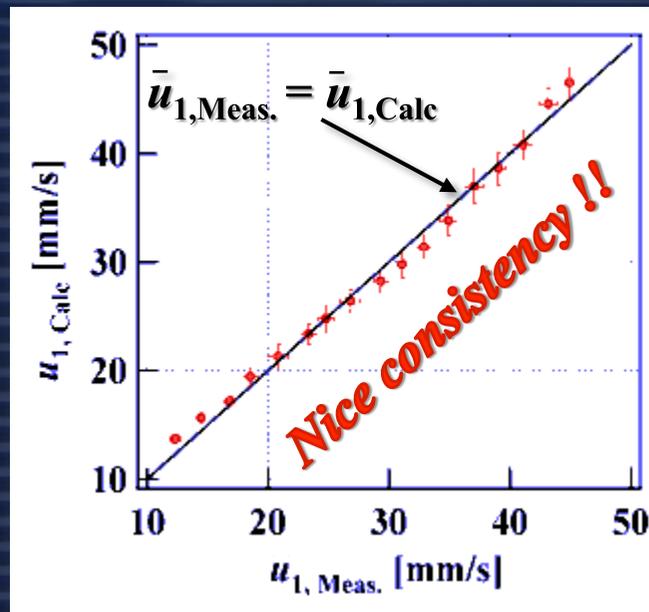
Comparison

Experiment

Verification of the determined $f_E [T/T_F]$

2. Effective speed of the first sound

Experimental values vs. calculated values from $f_E [\theta]$

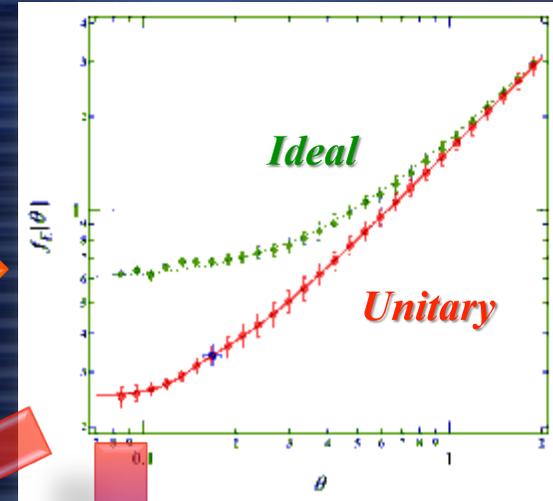


The universal function of the internal energy $f_E [T/T_F]$

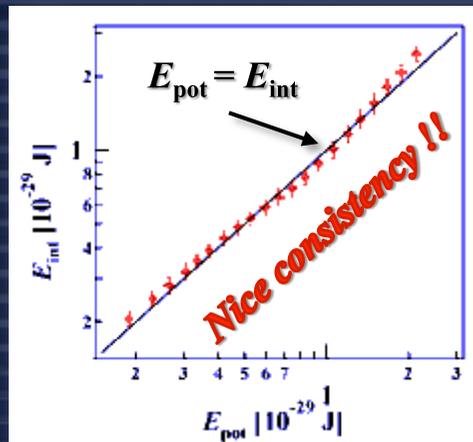
Universal hypothesis : $\frac{\varepsilon}{nE_F} = f_E [T/T_F]$

Equation of state : $p = \frac{2}{3} \varepsilon$

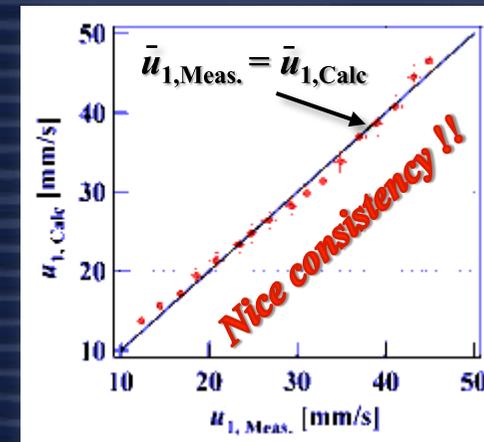
Mechanical equilibrium : $\nabla p(\mathbf{r}) + n(\mathbf{r}) \nabla V_{\text{Trap}}(\mathbf{r}) = 0$



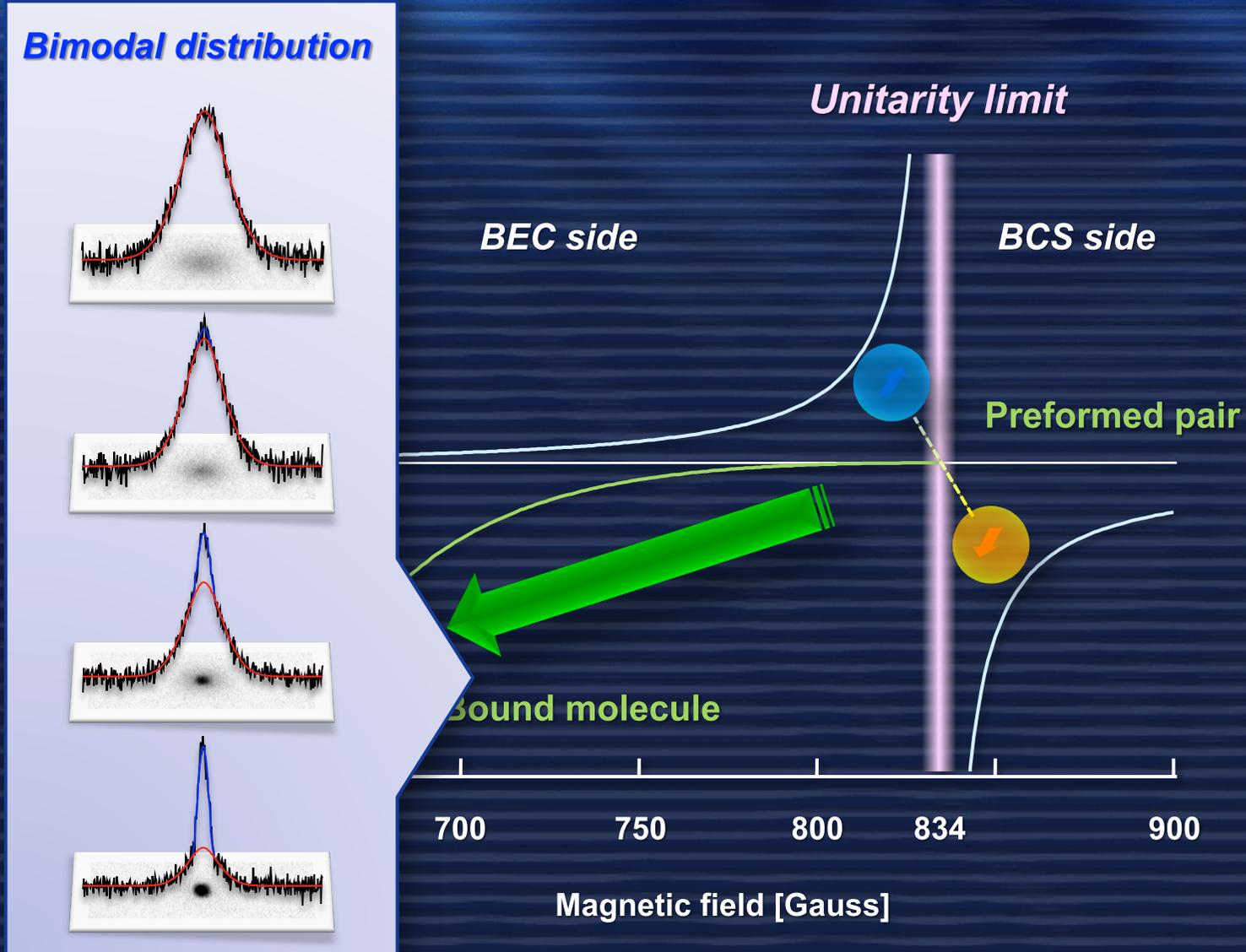
Energy comparison



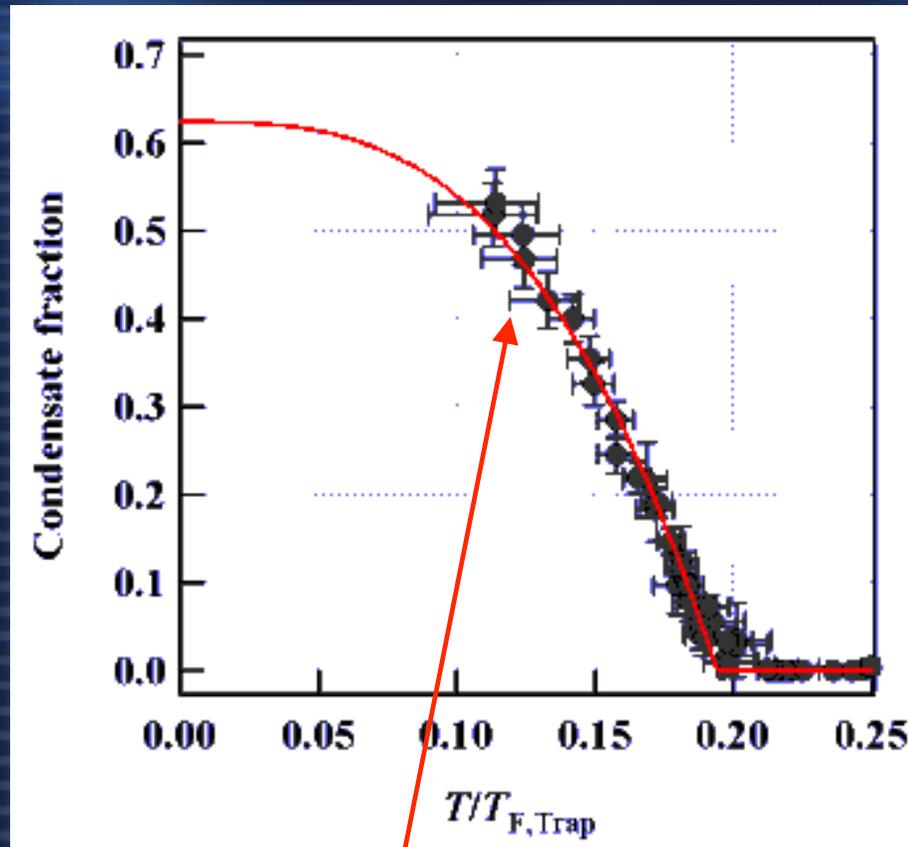
Speed of the first sound



Bimodal distribution of a fermion pair condensate



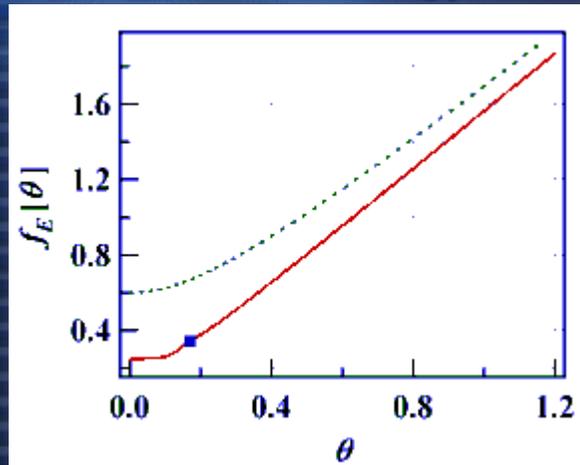
Condensate fraction vs Temperature



$$f(x) = CF_{\text{Max}} \cdot \left\{ 1 - \left(\frac{T}{T_C} \right)^{\underline{\underline{3.0(1)}}} \right\}$$

Universal thermodynamic functions

Internal energy

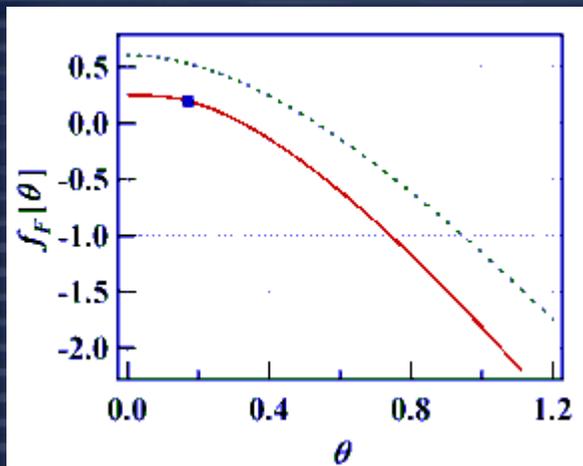


$$f_E = f_F - \theta f'_F$$

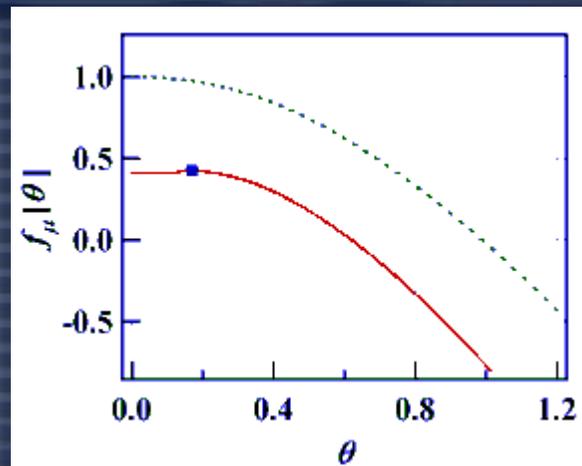
$$f_\mu = (5f_E - 2\theta f'_F)/3$$

$$f_S = -f'_F$$

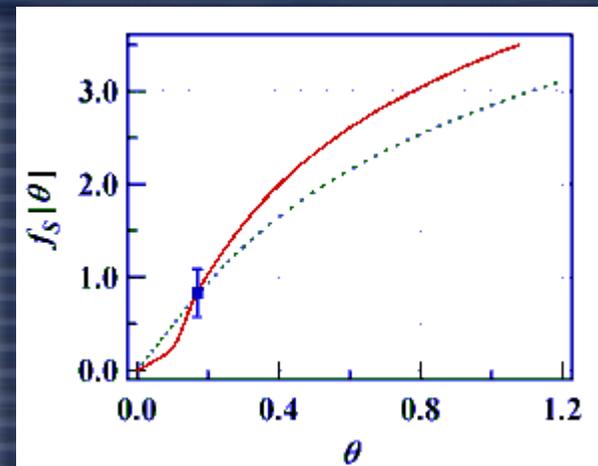
Helmholtz free energy



Chemical potential



Entropy

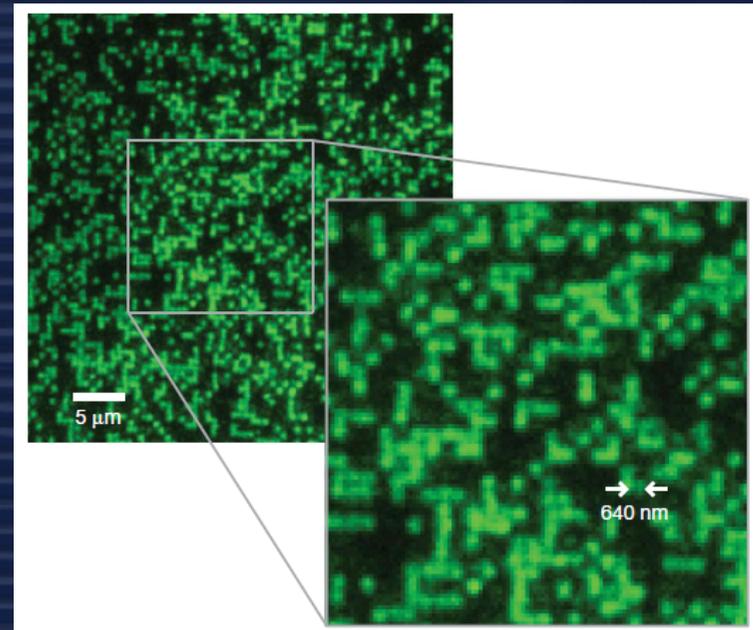
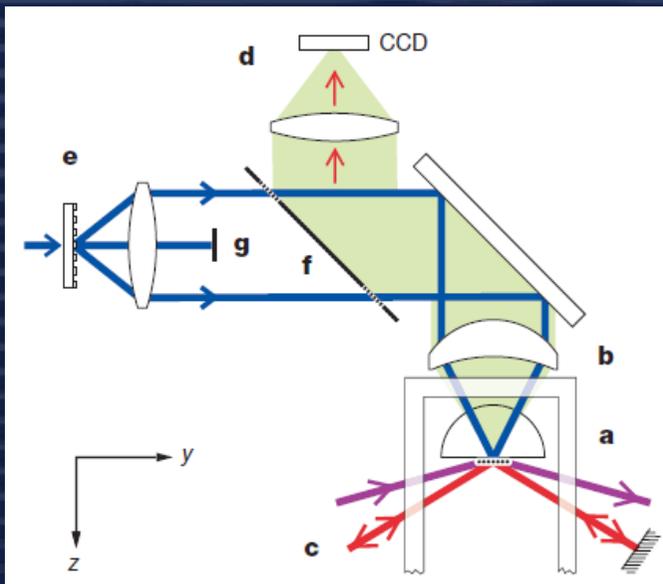


In the case of unitary gas, equation of state $p(\mathbf{r}) = 2\varepsilon(\mathbf{r})/3$ is available (exceptional case !!) which enable us to measure **local thermodynamic quantities**.

Then, how can we determine local thermodynamic quantities without help of equation of state ?



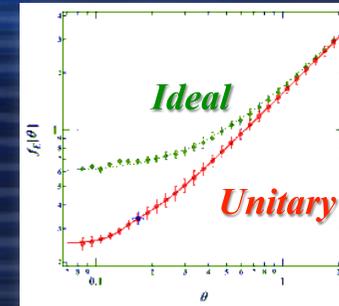
- Box potential
- High resolution local probe



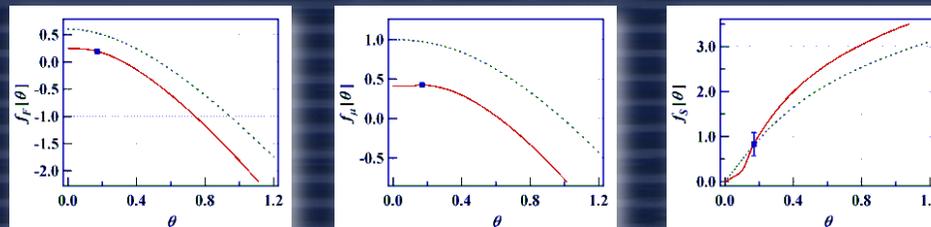
W. S. Bakr et al. Nature 462, 74 (2009).

Summary

- *The universal function of the internal energy was determined at the unitarity limit*

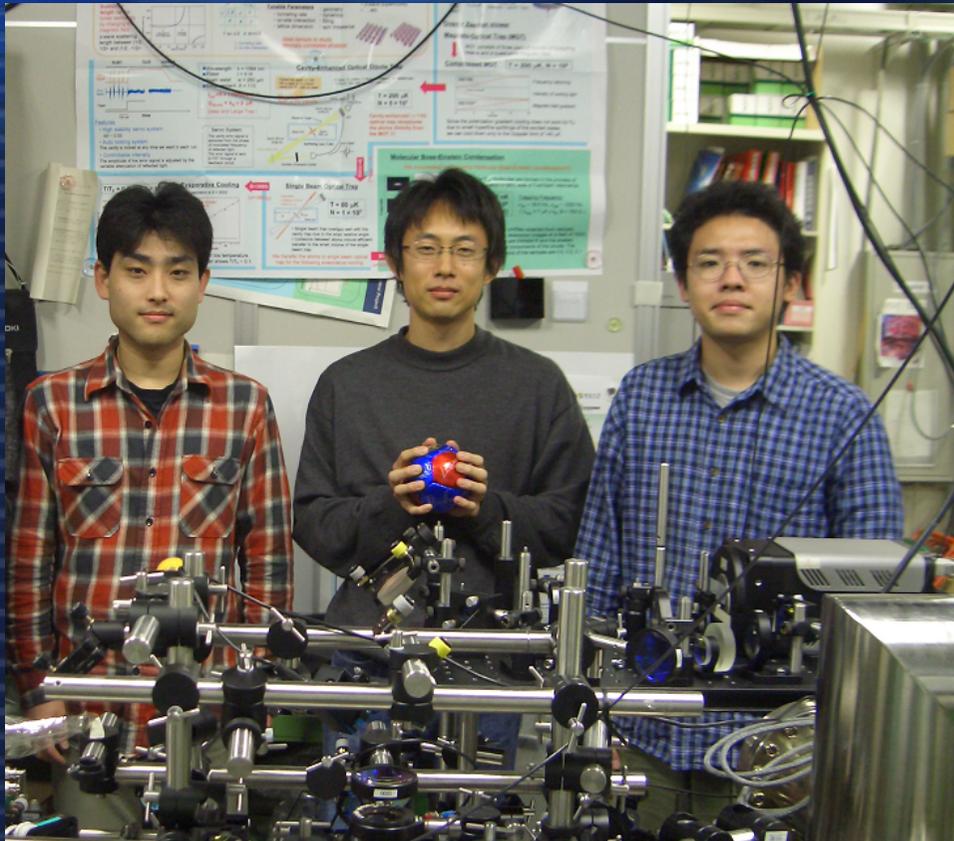


- *The other thermodynamic functions were derived from the thermodynamic relationship*
- *The critical parameters were determined at the superfluid transition temperature*



M. Horikoshi, S. Nakajima, M. Ueda and T. Mukaiyama, Science, **327**, 442 (2010).

The team (ERATO project)



T. Mukaiyama
(Group leader)

M. Horikoshi
(Postdoc)

S. Nakajima
(Ph.D student)

Unitary gas

Efimov physics



Masahito Ueda
(project leader)

Equation of state for a unitary Fermi gas

$$\Delta E = \Delta N E_F(n) f_E(T/T_F)$$

$$\Delta S = \Delta N k_B f_S(T/T_F)$$

$$E_F(n) = \frac{\hbar^2}{2m} (6\pi^2 n)^{2/3}$$

$$p = - \left(\frac{\partial(\Delta E)}{\partial(\Delta V)} \right)_{\Delta N, \Delta S}$$

$$\Delta S = \text{一定} \quad \longrightarrow \quad T/T_F = \text{一定}$$

$$p = - \left(\frac{\partial(\Delta E)}{\partial(\Delta V)} \right)_{\Delta N, T/T_F} = -\Delta N f_E(T/T_F) \left(\frac{\partial(E_F(n))}{\partial(\Delta V)} \right)_{\Delta N, T/T_F}$$

$$= \frac{2}{3} n E_F(n) f_E(T/T_F)$$

$$= \frac{2}{3} \varepsilon(n)$$

