

# Thermal effects on nuclear matter properties



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# Outlook

- Why finite-temperature EOSs in NS physics?
- Nuclear Hamiltonian
  - Correlated basis functions – effective interaction
- Generalisation to finite temperature
- Thermal effects on single-particle properties
  - Fermi distribution, spectrum and effective mass
- Average quantities: exact calculation VS fit
- Conclusions and perspectives

# Hot EOS

- Merger and post-merger phases of a BNS coalescence
- Proto-NS evolution
- Supernova modelling
- Important phenomena involved
  - Thermal contribution to the pressure
  - Bulk viscosity
  - Neutrino emission and absorption

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# Nuclear Many-Body Theory (NMBT)

- The basis for Nuclear Many-Body Theory is the Hamiltonian

$$H = \sum_{i=1}^A \frac{\mathbf{p}_i^2}{2m} + \sum_{j>i=1}^A v_{ij} + \sum_{k>j>i=1}^A V_{ijk}$$

where, in general,  $v_{ij} = \sum_p v^p(r_{ij}) O_{ij}^p$

- Nucleon-nucleon (NN) potential: Argonne  $v_6'$  (AV6P)
- Three-nucleon (3N) potential: Urbana IX (UIX)
- AV6P+UIX results are very close to AV18+UIX [Lovato+ (2022), arXiv:2202.10293]
  - Repulsive part of UIX  $\rightarrow$  additional repulsion  $\rightarrow$  lower equilibrium density of SNM

# CBF Effective Interaction

- NN forces are strongly repulsive at short distance → standard many-body perturbation theory cannot be used
- The CBF effective interaction is defined through

$$\langle H \rangle = \langle \Psi_0 | H | \Psi_0 \rangle = T_F + \langle \Phi_0 | \sum_{i < j} v_{ij}^{\text{eff}} | \Phi_0 \rangle$$

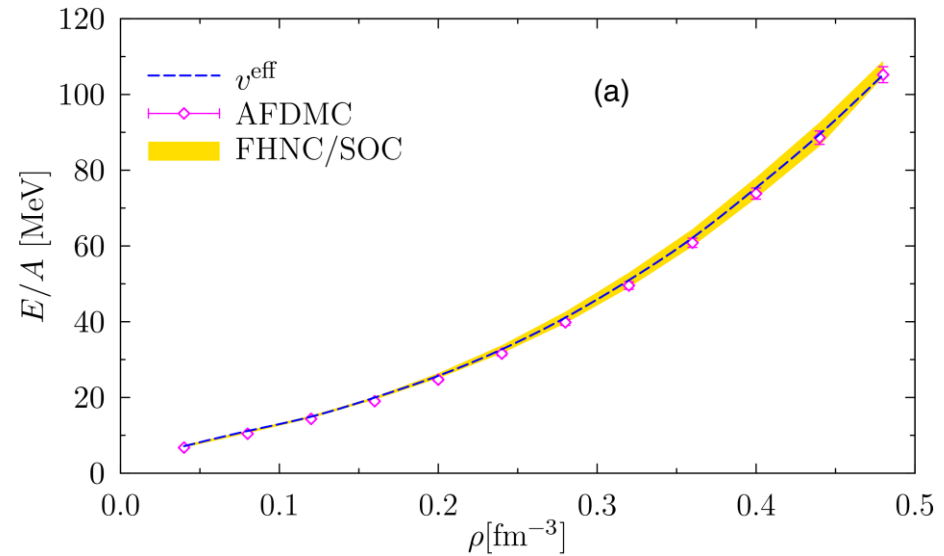
Correlated basis Fermi gas basis

- Include NN and 3N forces and correlation effects

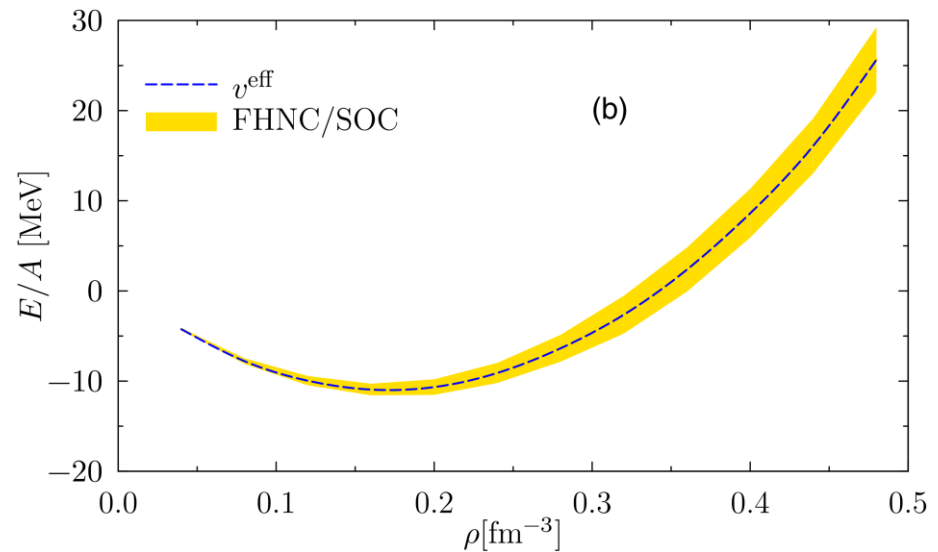
$$\langle H \rangle = T_F + \sum_n (\Delta E)_n = T_F + \langle \Phi_0 | \sum_{i < j} v_{ij}^{\text{eff}} | \Phi_0 \rangle$$

# CBF Effective Interaction - Results

PNM



SNM



Lovato & Benhar, **PRC** 96 054301 (2017)

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# Generalisation to finite temperature

- Our basic assumption: at  $T \neq 0$  and  $T \ll m_\pi \approx 140$  MeV, the Hamiltonian is largely unaffected by thermal effects
- Thermodynamic consistency is not easily achieved in finite-temperature perturbation theory, which can be tested by comparing the two, a priori equivalent, definitions of pressure

$$P = \rho \left( \mu - \frac{F}{N} \right) \qquad P = - \frac{\partial F}{\partial V} = \rho^2 \frac{\partial F}{\partial \rho} \frac{1}{N}$$

- Variational approach: minimisation of a trial grand canonical potential

# Variational principle

- The minimisation yields the solution for the Fermi distribution

$$n(k, T) = \{1 + e^{\beta[e(k, T) - \mu]}\}^{-1}$$

where the single-particle energy  $e(k, T)$  is given by

$$e(k, T) = \frac{k^2}{2m} + U_k + \delta e$$

where

$$U_k = \sum_{k'} \langle kk' | v | kk' \rangle_A n(k', T)$$

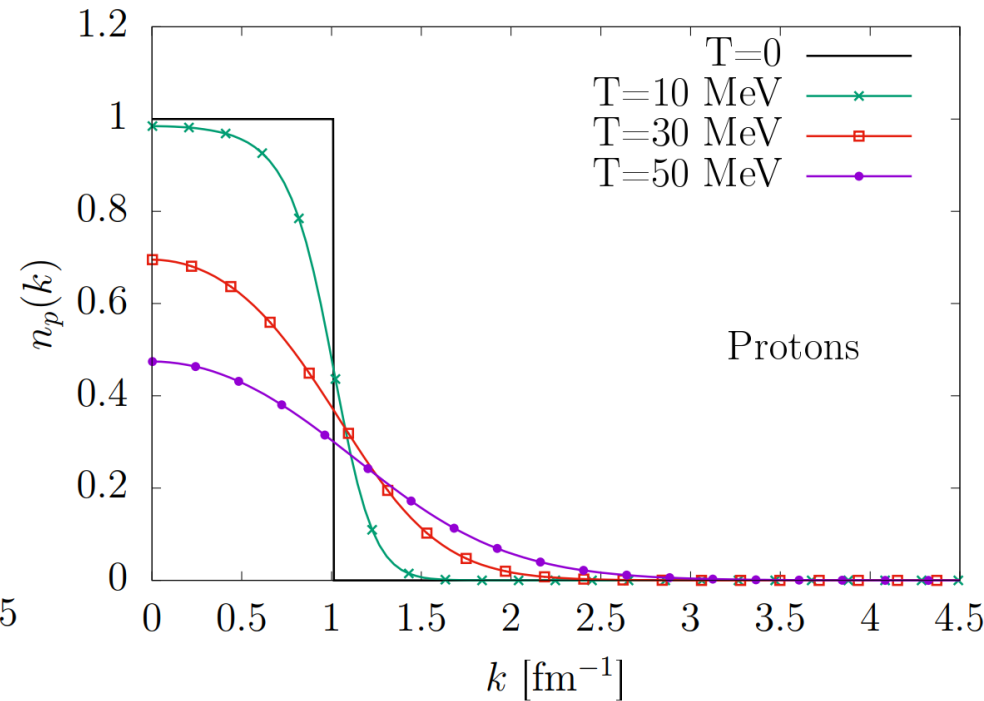
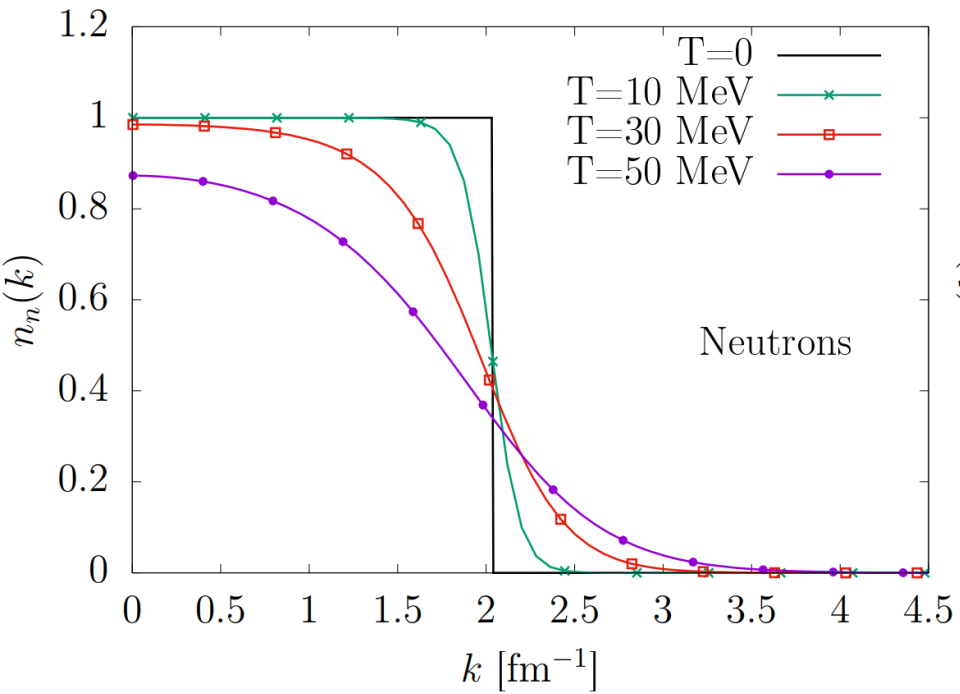
$$\delta e = \frac{1}{2} \sum_{k, k'} \langle kk' | \frac{\partial v}{\partial \rho} | kk' \rangle_A n(k, T) n(k', T)$$

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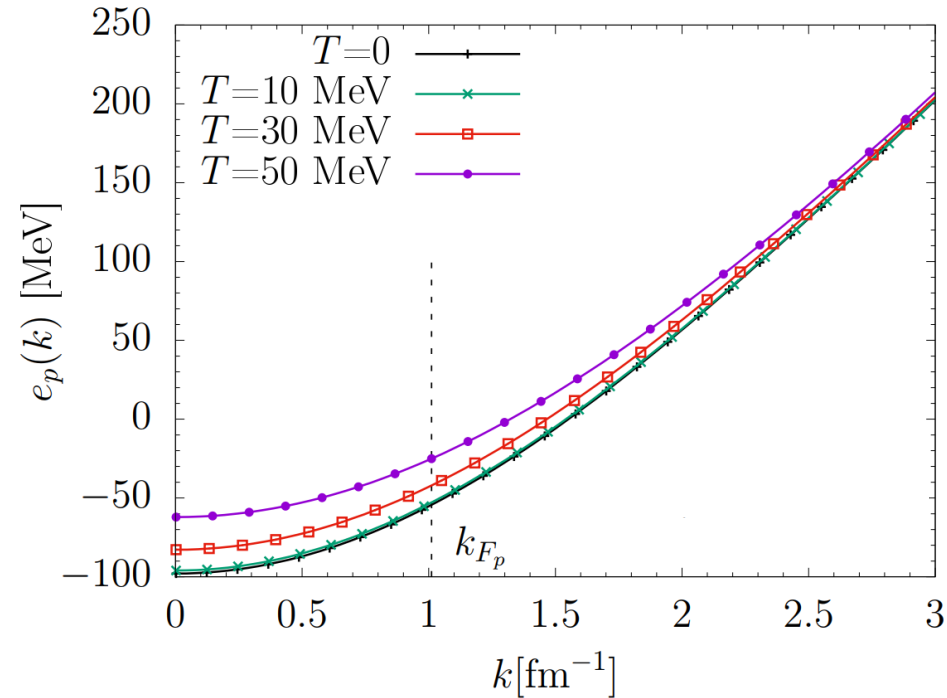
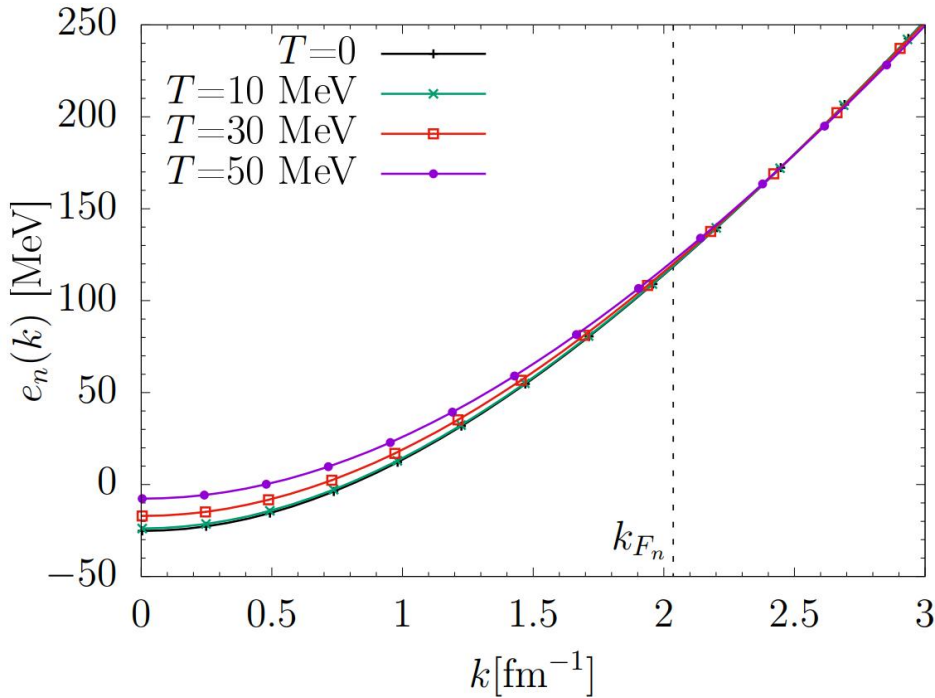
# Fermi distributions

↳ Contain all thermodynamic information



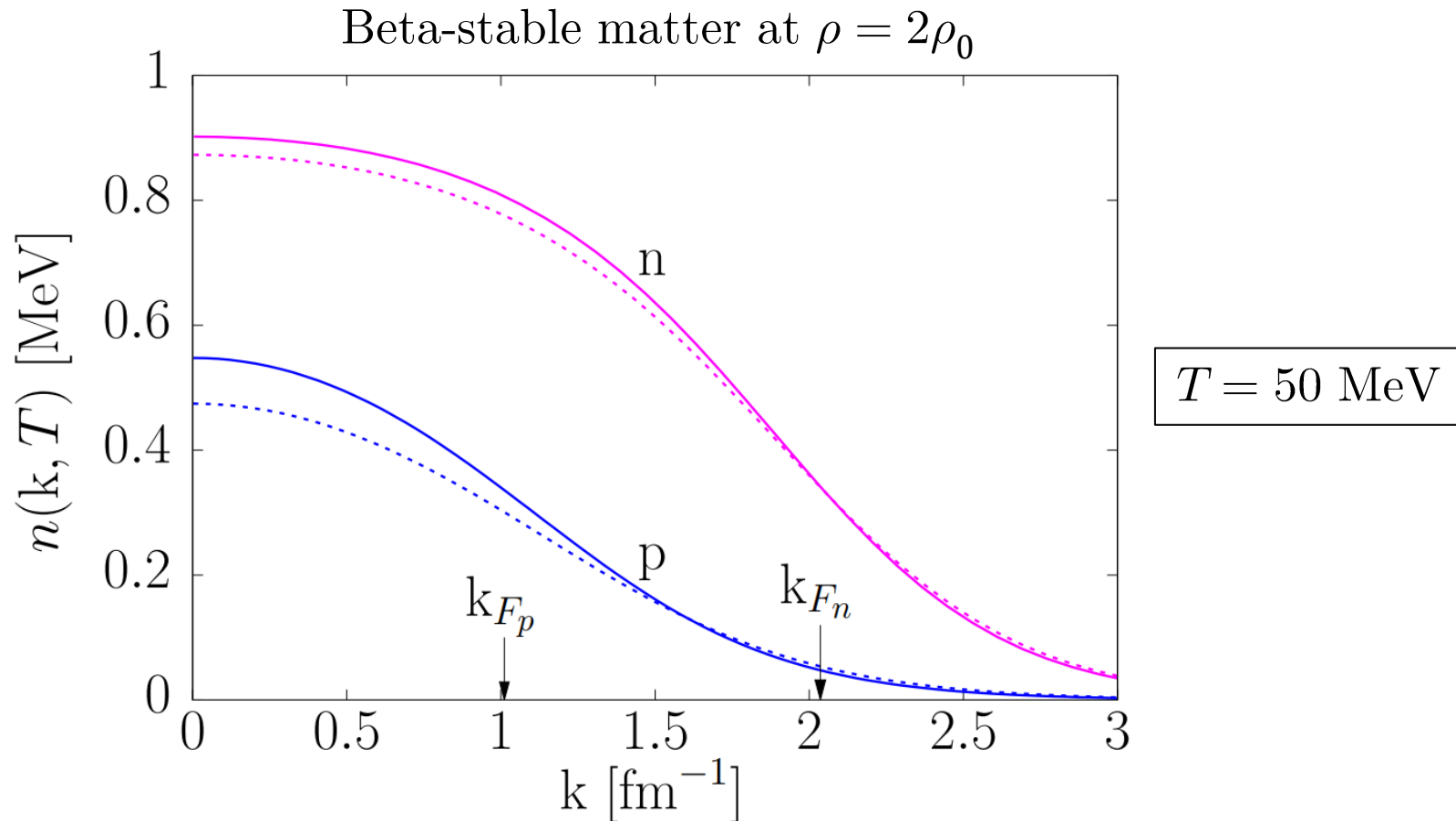
Beta-stable matter at  $\rho = 2\rho_0$

# Single-particle energies (spectrum)



Beta-stable matter at  $\rho = 2\rho_0$

# Spectrum $\rightarrow$ Fermi distribution

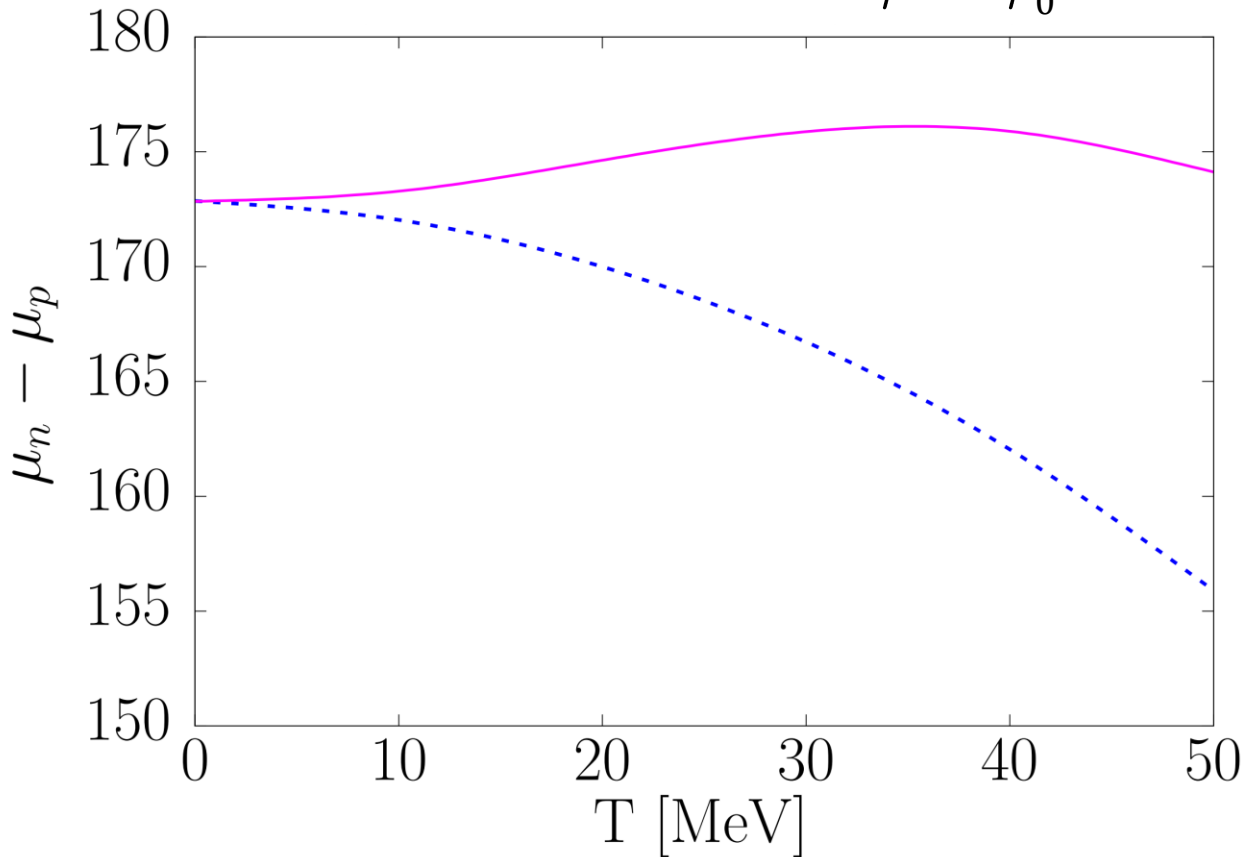


Solid line: Fermi distribution assuming  $e(k, T)$

Dashed line: Fermi distribution assuming  $e(k, T = 0)$

# Spectrum $\rightarrow$ Chemical potential

Beta-stable matter at  $\rho = 2\rho_0$



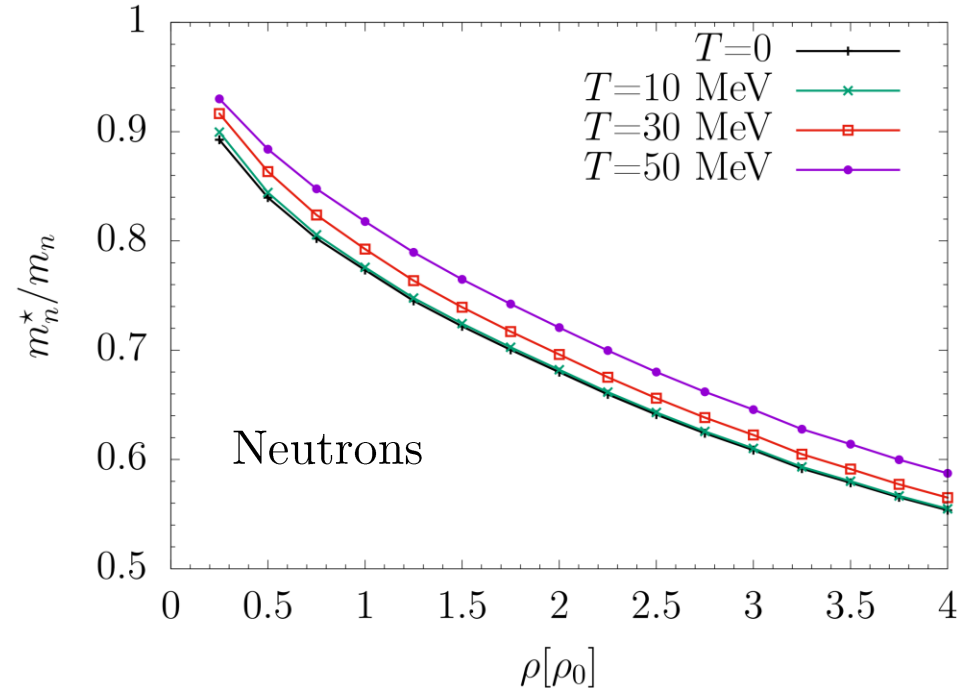
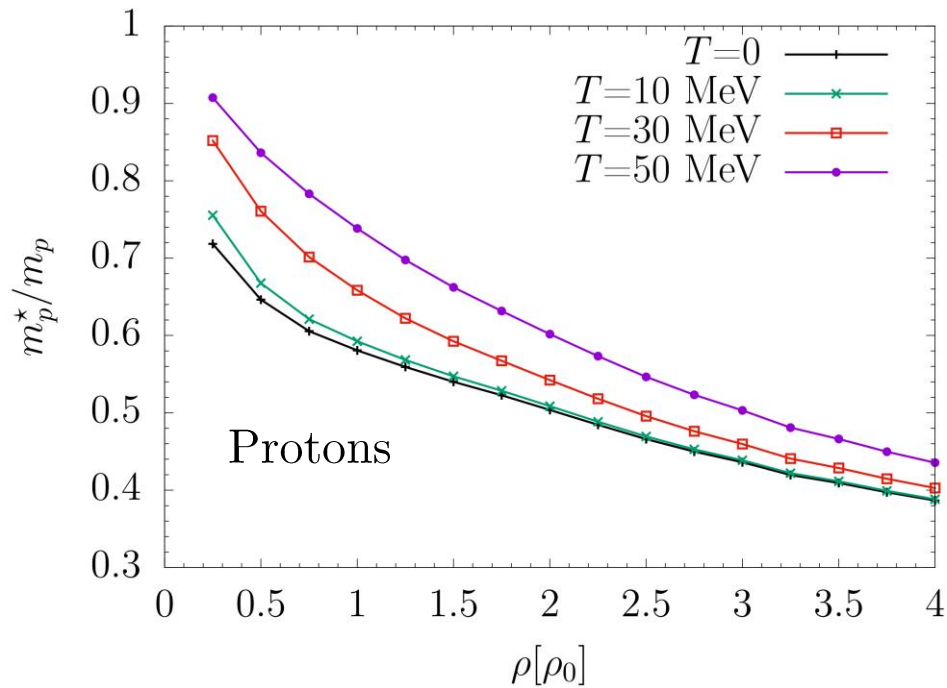
Changes of chemical composition in  $\beta$ -stable matter

Dashed line: assuming  $e(k, T)$

Solid line: assuming  $e(k, T=0)$

# Effective mass

↳ Determines the nucleon dispersion relation  $\rightarrow \frac{1}{m^*(k, T)} = \left( \frac{1}{k} \frac{de(k, T)}{dk} \right)_{k=k_F}$



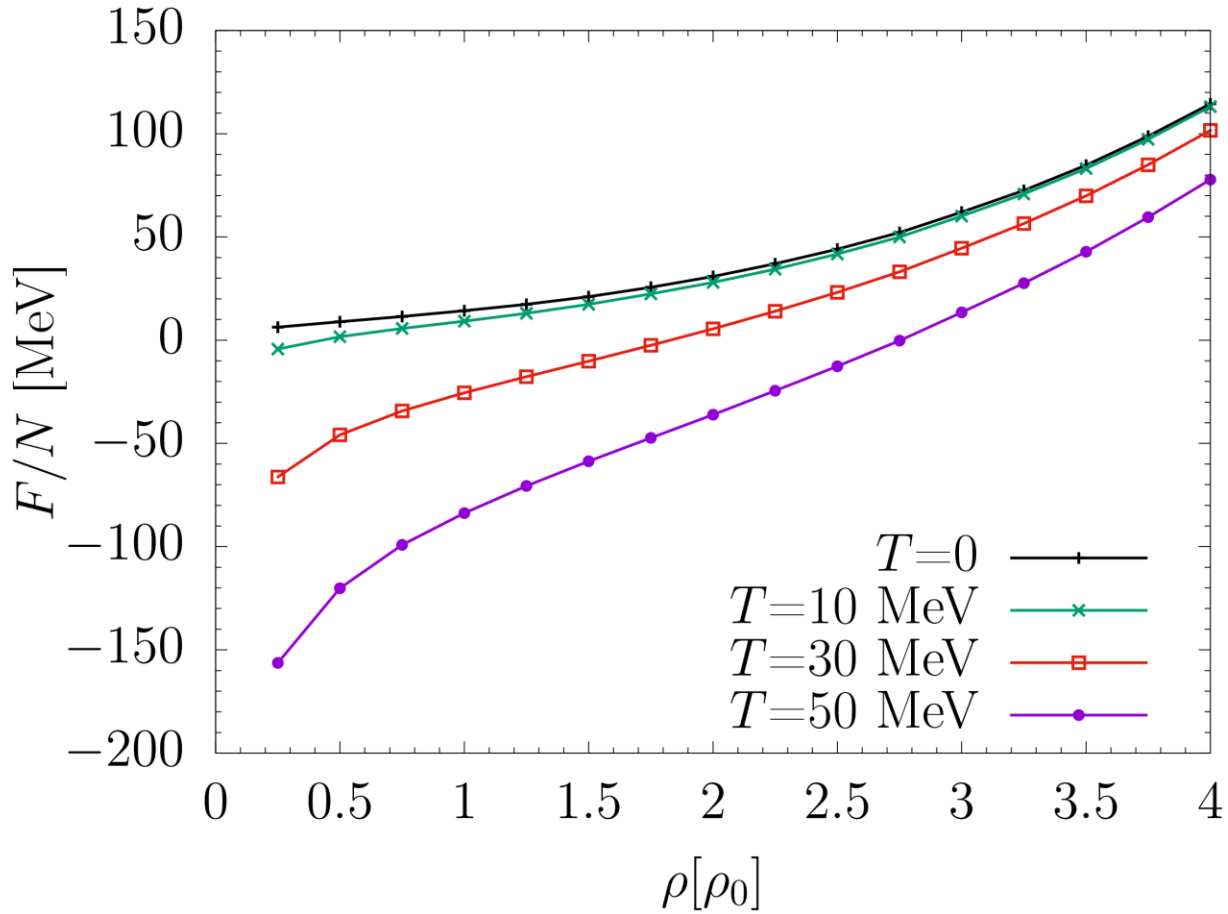
Neutrons  $\rightarrow$  larger density  $\rightarrow$  weaker temperature effects



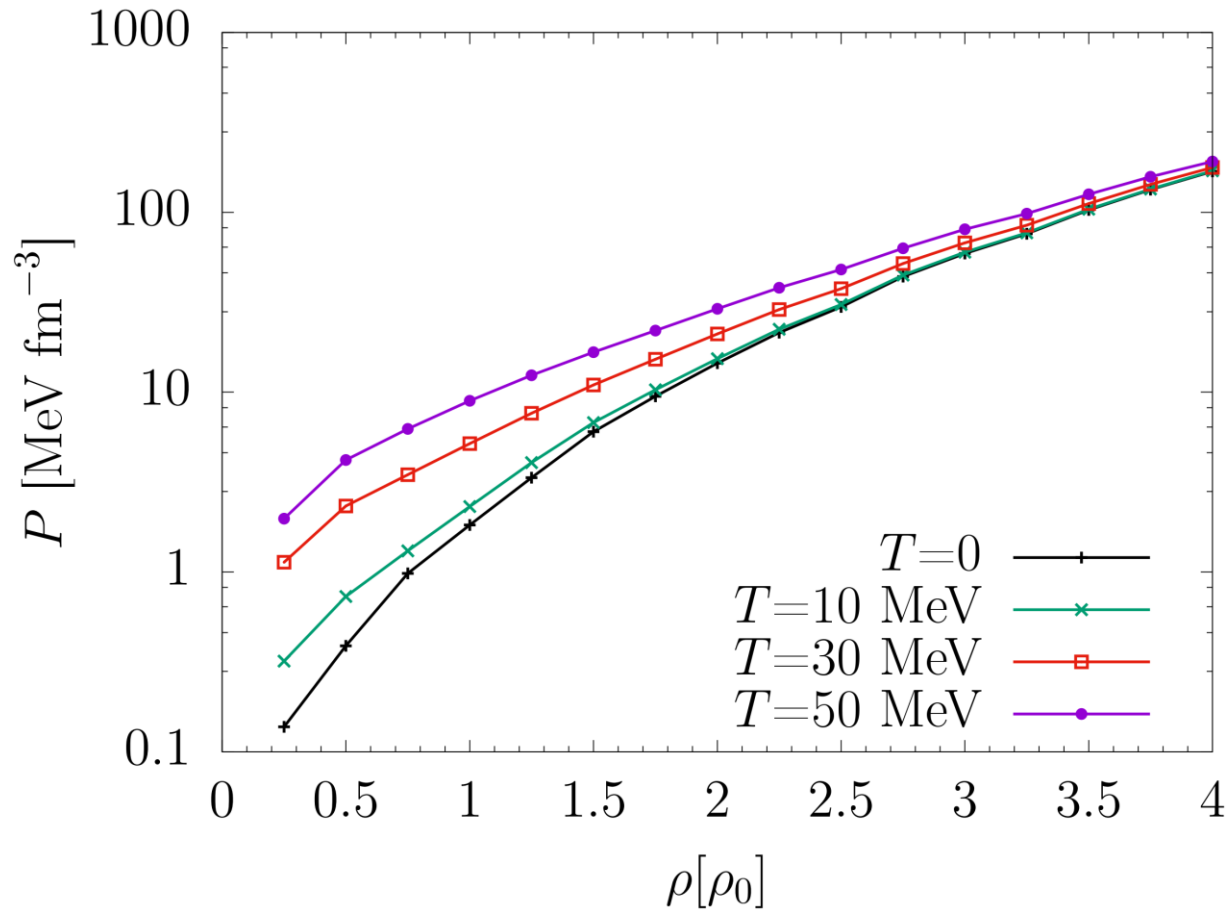
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# Free energy



# Pressure



Clearly at high densities the thermal contribution becomes less relevant

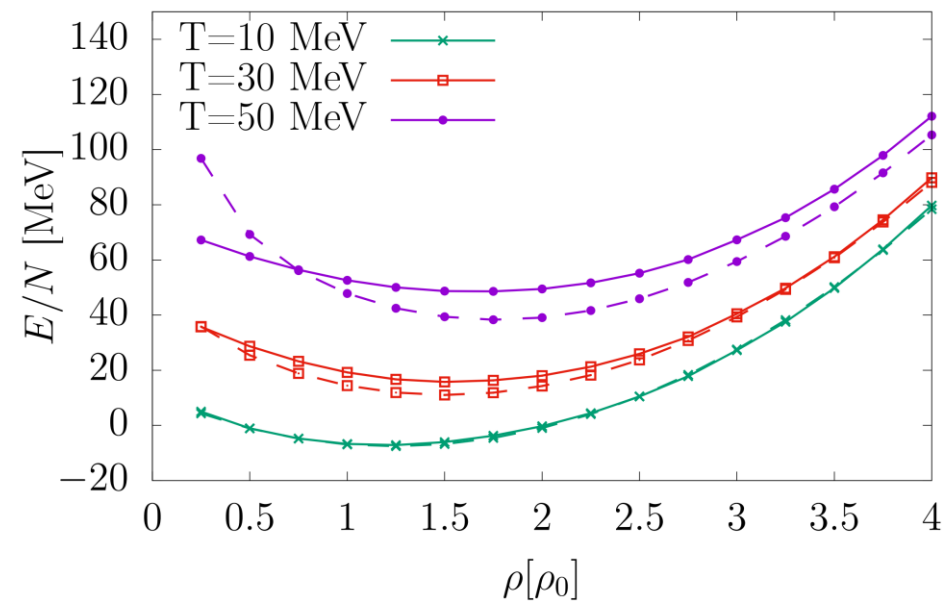
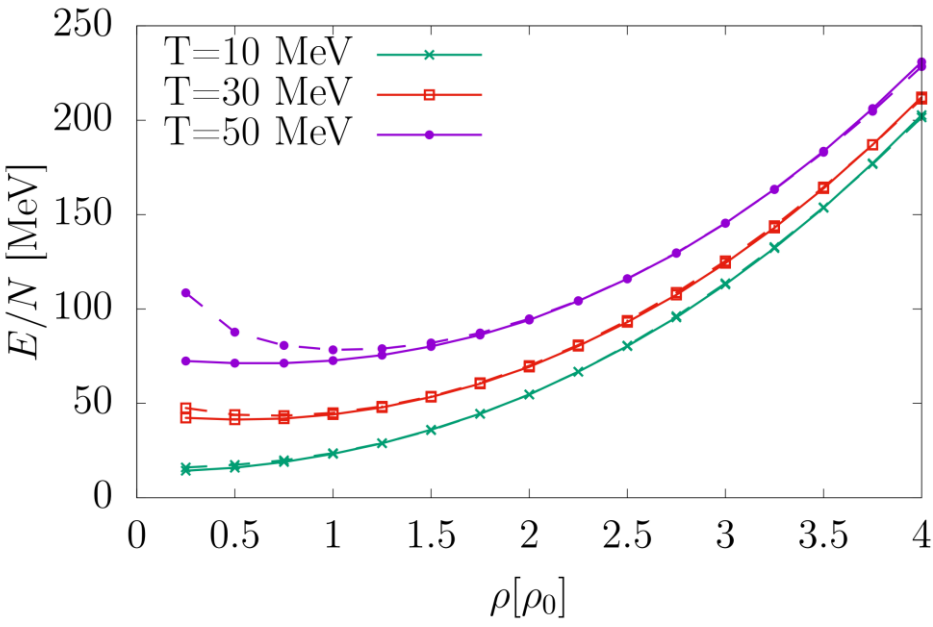
# Exact results VS fit

- Lack of hot EOSs: resort to parametrisations that generalise the thermal contribution to an arbitrary  $T = 0$  EOS
- Widely used model: thermal contribution added as an ideal fluid at  $T$
- Recently Raithel et al. (2019) have proposed a parametrisation to simulate thermal effects
  - Intermediate densities: ideal fluid
  - High densities: degenerate Fermi gas (Sommerfeld expansion)

# Exact results VS Raithel+ (2019)

PNM

SNM



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# Conclusions and perspectives

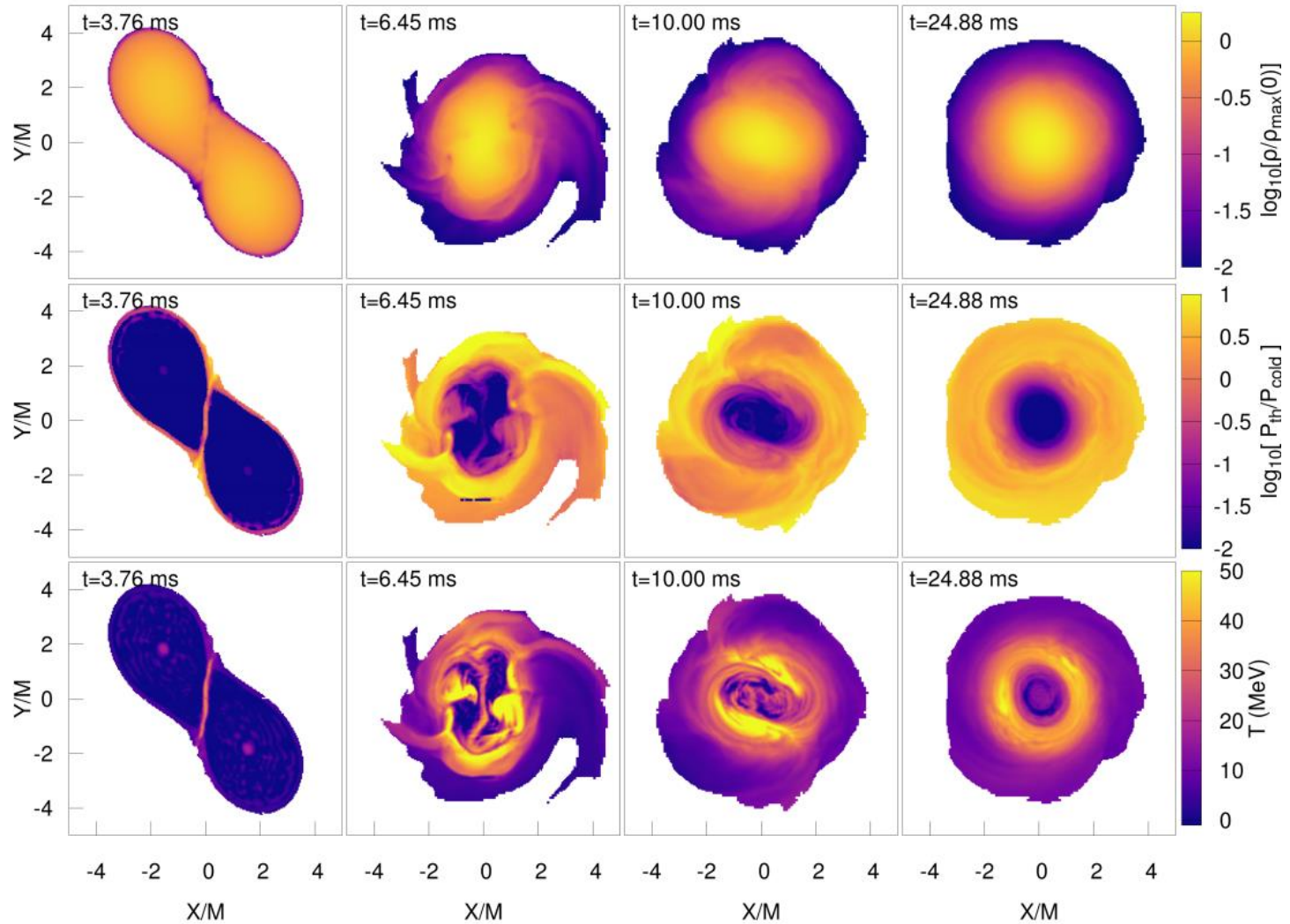
- **Robust finite-temperature EOSs** are fundamental to multimessenger astrophysics
- **Thermal effects** are more important to **protons**, due to their **lower density**
- **Temperature** also affects the **chemical composition** of NS matter
- Changes in **single-particle energies** and effective masses → relevant to every **nucleon collision process**
- The application to the calculation of bulk viscosity is being carried out
- A physically sound parametrisation of the thermal component is being studied

# Backup



# Merger simulation

# Simulations



Raithel+, **PRD** 104 063016 (2021)

# CBF Effective Interaction

# CBF Effective Interaction

- The effective interaction can be written as

$$v_{ij}^{\text{eff}} = \sum_{p=1}^6 v^{\text{eff},p}(r_{ij}) O_{ij}^p$$

- And the correlated ground state as

$$|\Psi_0\rangle \equiv \frac{F|\Phi_0\rangle}{\langle\Phi_0|F^\dagger F|\Phi_0\rangle^{1/2}}$$

- The correlation operator is written in the form

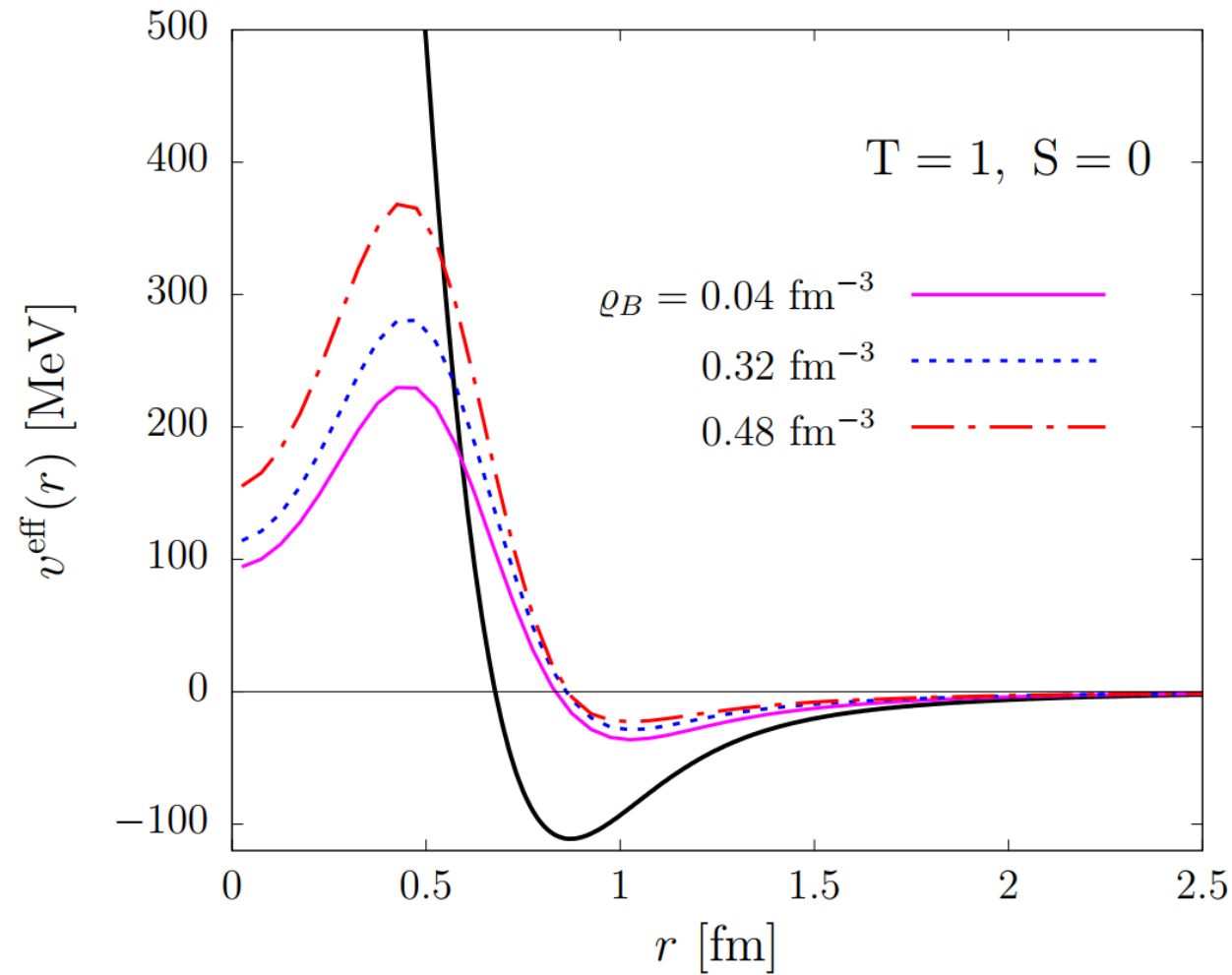
$$F(1, \dots, N) = \mathcal{S} \prod_{j>i=1}^N f_{ij}$$

# CBF Effective Interaction

- The two-body correlation functions reflect the complexity of the NN potential and can be conveniently expressed as

$$f_{ij} = \sum_{p=1}^6 f^p(r_{ij}) O_{ij}^p$$

# CBF Effective Interaction



Radial dependence of the CBF effective potential in the  $S = 0, T = 1$  channel

# Finite-temperature perturbation theory

# Finite-temperature perturbation theory

- Our basic assumption: at  $T \neq 0$  and  $T \ll m_\pi \approx 140$  MeV, the Hamiltonian is largely unaffected by thermal effects
- All thermodynamic functions of a system in equilibrium at temperature  $T$  can be obtained from the grand canonical potential

$$\Omega = -\frac{1}{\beta} \ln Z$$

where the partition function  $Z$  is

$$Z = \text{Tr } \Phi$$

with

$$\Phi = e^{-\beta(H - \mu N)}$$



# Finite-temperature perturbation theory

- The basis for the derivation of finite-temperature perturbation theory is the Bloch equation

$$-\frac{\partial\Phi}{\partial\beta} = (H - \mu N)\Phi$$

- To find the perturbative expansion of  $Z$  we can explore the similarity between this equation and Schroedinger's
- Rewriting the Hamiltonian as  $H = H_0 + H_I$ ,

$$\begin{aligned} -\frac{\partial\Phi}{\partial\beta} &= [(H_0 - \mu N) + H_I]\Phi \\ &= (H'_0 + H_I)\Phi \end{aligned}$$

# Finite-temperature perturbation theory

- $H_0$  and  $H_I$  are written as

$$H_0 = \sum_k e_k a_k^\dagger a_k$$

$$H_I = \frac{1}{2} \sum_{k,k',q,q'} \langle k'q' | v | kq \rangle a_{k'}^\dagger a_{q'}^\dagger a_q a_k - \sum_k U_k a_k^\dagger a_k$$

where  $e_k$  is the single-particle energy, which plays an important role in determining thermal effects.

$$e_k = \frac{\mathbf{k}^2}{2m} + U_k = t_k + U_k$$

# Finite-temperature perturbation theory

- It can be shown that, at first order,  $\Omega = \Omega_0 + \Omega_1$  can be written as

$$\begin{aligned}\Omega_0 &= -\frac{1}{\beta} \ln Z_0 = -\frac{1}{\beta} \sum_k \ln (1 + e^{-\beta(e_k - \mu)}) \\ &= \sum_k (e_k - \mu) n_k + \frac{1}{\beta} \sum_k [n_k \ln n_k + (1 - n_k) \ln(1 - n_k)]\end{aligned}$$

$$\Omega_1 = -\frac{1}{\beta} \ln Z_1 = \frac{1}{2} \sum_{kk'} \langle kk' | v | kk' \rangle_A n_k n_{k'} - \sum_k U_k n_k$$

# Finite-temperature perturbation theory

- It can be shown that, at first order,  $\Omega = \Omega_0 + \Omega_1$  can be written as

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$$n_k = [1 + e^{\beta(e_k - \mu)}]^{-1} \quad \text{Fermi distribution}$$

$$\Omega_1 = -\frac{1}{\beta} \ln Z_1 = \frac{1}{2} \sum_{kk'} \langle kk' | v | kk' \rangle_A n_k n_{k'} - \sum_k U_k n_k$$

# Variational calculation at finite $T$

# Hartree-Fock grand canonical potential

$$\begin{aligned}\tilde{\Omega} = & \sum_k t_k n_k + \frac{1}{2} \sum_{k,k'} \langle kk' | v | kk' \rangle_A n_k n_{k'} \\ & + \frac{1}{\beta} \sum_k [n_k \ln n_k + (1 - n_k) \ln(1 - n_k)]\end{aligned}$$

- The given potential is totally valid in a variational context
- It recovers the HF approximation at  $T=0$
- The formalism is thermodynamically consistent by construction