# A Direct Construction of the Nuclear Effective Interaction from Scattering Phase Shifts

Kenneth S. McElvain UC Berkeley and Lawrence Berkeley National Laboratory

Work Done in Collaboration with Wick Haxton

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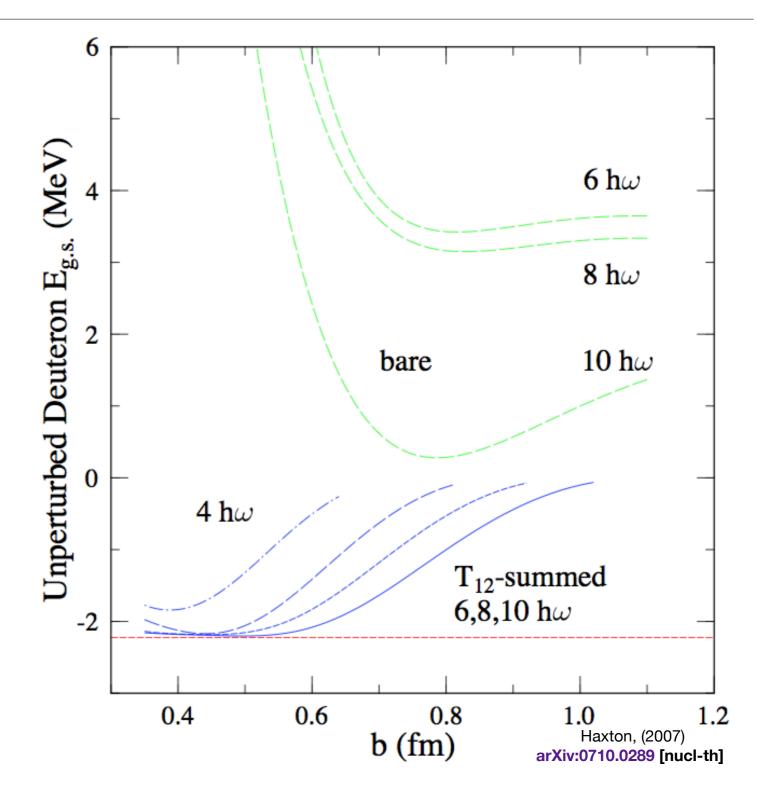
## Background: Interaction From a Potential

 $H_{eff} = P \left( H + H \frac{1}{E_i - OH} QH \right) P \psi_i = E_i P \psi_i$ Bloch-Horowitz:  $H_{eff} = P \frac{E}{E - TO} \left| T - T \frac{Q}{E} T + V + V \frac{1}{E - OH} V \right| \frac{E}{E - OT} P$ Haxton-Luu Form:  $\frac{E}{E-QT}^{P}$  acts only on edge state, restoring the long range waveform. 3 R<sub>6,0</sub>(r) The first terms in H<sub>eff</sub> perform a complete sum of scattering by QT. 0  $P \frac{E}{E-TQ} V \frac{1}{E-QH} V \frac{E}{E-QT} P$  is short range and is 3 5 r(fm) replaced by a contact gradient expansion which can be easily fit by taking matrix elements of the known potential.

## Deuteron Binding Energy Convergence

With QT summed to all orders, shrinking the HO length scale b enables the capture of the important part of V with no ET terms in a very small basis.

Shows the power of summing the IR contribution.

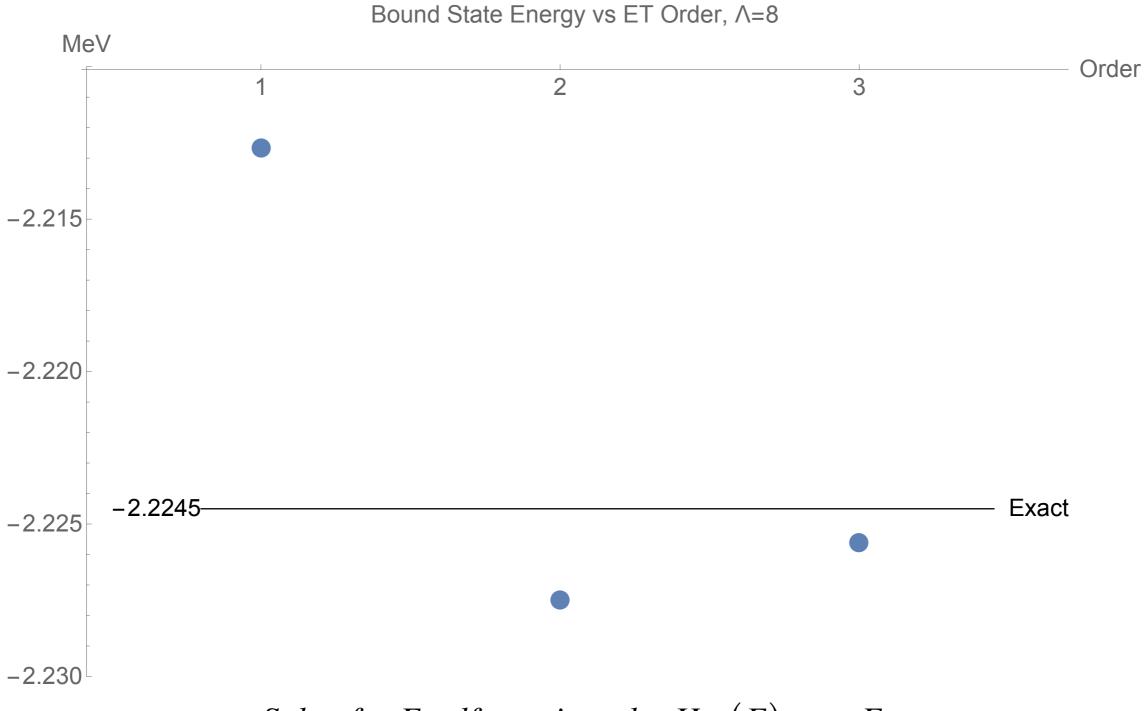


## Fitting to Continuum Scattering Phase Shifts

Same H<sub>eff</sub> 
$$H_{eff} = P \frac{E}{E - TQ} \left[ T - T \frac{Q}{E} T + V + V \frac{1}{E - QH} V \right] \frac{E}{E - QT} P$$
  
But  $\dots \frac{E}{E - QT} P$  is no longer  
unique. The boundary  
condition at infinity is a  
periodic boundary condition  
corresponding to the phase  
shift.

The contact gradient expansion coefficients are used to satisfy the self consistent energy constraint at a set of continuum energies. Then we predict ...

#### Practice Problem: S Channel Only, Hard Core + Well



Solve for E self consistently:  $H_{eff}(E)\psi_P = E\psi_P$ 

## Coupled Channel Deuteron Binding Energy

Phase shifts generated from Argonne v18 potential.

Eight parameters fit through NNLO to produce self consistent energies.

	alo3S1	anlo3S1	anloSD	$N^2$	<b>Binding Energy</b>
Reference	-1.7589	-0.02221	-0.1271	yes	-2.22
Reference-	-1.7589	-0.02221	-0.1271	no	-2.08
Fit	-1.7803	-0.02167	-0.1268	no	-2.27

The short range part of the interaction comes from the phase shifts. The IR contribution comes from the transform of the edge state, yielding a sum of the scattering by QT.

# Summary

- Key Ideas for generalization to continuum
  - Match phase shifts at each energy by constraining Greens' function
  - Fit contact gradient coefficients to produce self consistent energies across a range of energies
- The result is an accurate small basis interaction
  - Due to separation of UV and IR and complete sum of IR contributions
  - Only need long range part of V to be accurate. The rest of the potential contribution is captured from phase shifts

#### See me at the poster session for questions!

Thank You!

# NN ET Operators

$${}^{3}S_{1} \leftrightarrow {}^{3}S_{1} : a_{LO}^{3S1}\delta(\vec{r}) + a_{NLO}^{3S1}\left(\bar{\nabla}^{2}\delta(\vec{r}) + \delta(\vec{r})\bar{\nabla}^{2}\right) + \cdots$$

$${}^{3}S_{1} \leftrightarrow {}^{3}D_{1} : a_{NLO}^{SD}\left(\bar{D}^{0}\delta(\vec{r}) + \delta(\vec{r})\bar{D}^{0}\right) + \cdots$$

$${}^{3}D_{1} \leftrightarrow {}^{3}D_{1} : a_{NNLO}^{3D1}\left(\bar{D}^{2} \cdot \delta(\vec{r})\bar{D}^{2}\right) + \cdots$$

$$\vec{D}^{0} = \left[\left(\sigma(1) \otimes \sigma(2)\right)^{(2)} \otimes \left(\vec{\nabla} \otimes \vec{\nabla}\right)^{(2)}\right]_{0}^{0} \sim S_{12}$$

$$\vec{D}_{M}^{2} = \left[\vec{\nabla} \otimes \vec{\nabla}\right]_{M}^{2}$$