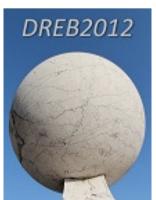
## Open Questions and Perspectives in DREB Physics (from a theoretician point of view)







### Direct reactions (from Wikipedia)

An intermediate energy projectile transfers energy or picks up or loses nucleons to the nucleus in a single quick ( $10^{-21}$  second) event.

Energy and momentum transfer are relatively small.

These are particularly useful in experimental nuclear physics, because the reaction mechanisms are often simple enough to calculate with sufficient accuracy to probe the structure of the target nucleus.

## Role of theory (shopping list)

Standard models with better structure input
Check of selectivity of input information
Adjustment of standard model to accomplish new
situations (e.g. weak binding)

Consistence of reaction models with structure models
Clarification of reaction mechanism

New genuine models (inspired by new development in
nuclear structure)

Quoting Daphne Jackson: "In order to make theoretical predictions which may be compared with the experimental data on direct reactions, it is necessary to introduce a number of simplifying assumptions to reduce the many-body problem to a tractable form"

Traditional "pulling horses" in the description of direct reactions:

Optical potentials
Collective formfactors for inelastic modes
Single-particle formfactors for transfer reactions
Coupled channels
DWBA

Semiclassical approximations (for heavy-ion scattering)

Use of structure information is maximized by the introduction of microscopic potentials and formfactors, fully exploiting the knowledge of the nuclear many-body functions

Example of "simple" process from the reaction mechanism point of view: radiative capture (one-step process induced by pure e.m. field) e.g.  $^{7}$ Be(p. $^{7}$ ) $^{8}$ B

Improvements come from the nuclear structure, entering at different levels, both for the bound and the continuum states

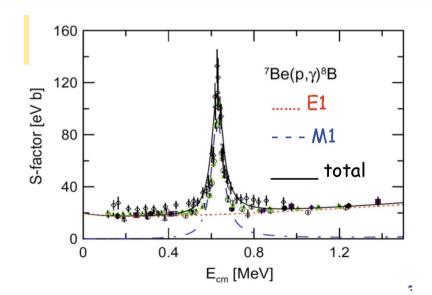
## Simplest model uses bound and continuum single-particle wave functions

$$\int_0^\infty r^\lambda \psi^l(E_i, r) \psi^{l'}(E_f, r) dr$$

simply defined by a potential (e.g. Woods-Saxon) generated by the inert <sup>7</sup>Be core

Schrödinger equation:

$$-\frac{\hbar^2}{2\mu}\frac{d^2}{dr^2}\psi^l(r) + \left[V(r) + \frac{\hbar^2}{2\mu}\frac{l(l+1)}{r^2}\right]\psi^l(r) = E\psi^l(r)$$



Bertulani Z. Phys. A356, 293 (1996)

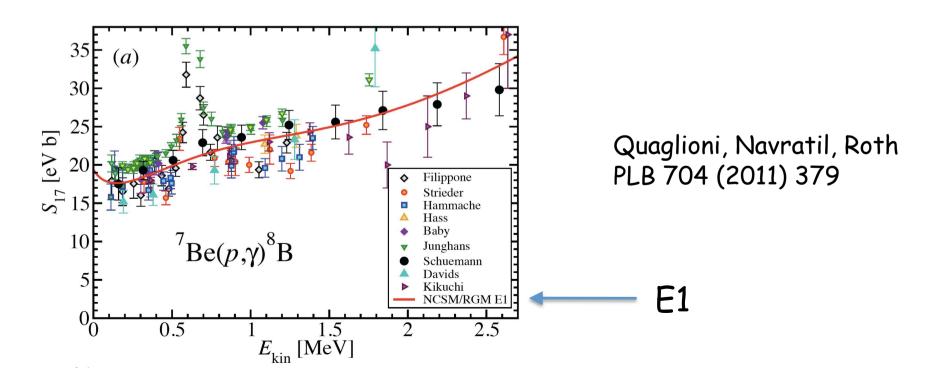
e.g., Woods-Saxon

# But now other models use more sophisticated many-body wave functions for both the bound and the scattering channels

Examples: Cluster, ab initio, no-core shell model, no-core shell model, no-core shell model plus RGM

(Realistic interactions H = T +  $\Sigma$ vij +  $\Sigma$ vijk, accurate wave functions of <sup>7</sup>Be, <sup>8</sup>B to deduce the ANC of <sup>8</sup>B)

OBS: need for an equivalent treatment of both bound and unbound channels



Other "simple" process: "safe" Coulomb excitation

Only structure information needed (or vice versa extracted): transition  $B(E\lambda)$  and multipole moments Reaction description: Coupled channels

Leftover problem: choice of the relevant model space and technical management of the coupled channel calculation, with delicate minimization procedures

Obs: correction to standard non-relativistic and relativistic Coulomb field

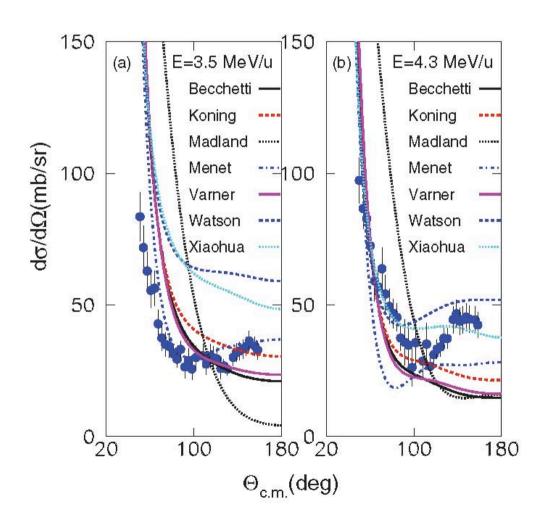
## Elastic scattering and optical potentials ("mean field" domain)

Basic tool: folding potentials of densities and NN interaction (M3Y, JLM, ....). Energy dependence. Novel interactions (NN+NNN+ ..., tensor force, ..)

Polarization part
(Feshbach theory, leading order, BWP, Sakuragi, ....).
Choice of the model space in CC formalim. Coupling to continuum

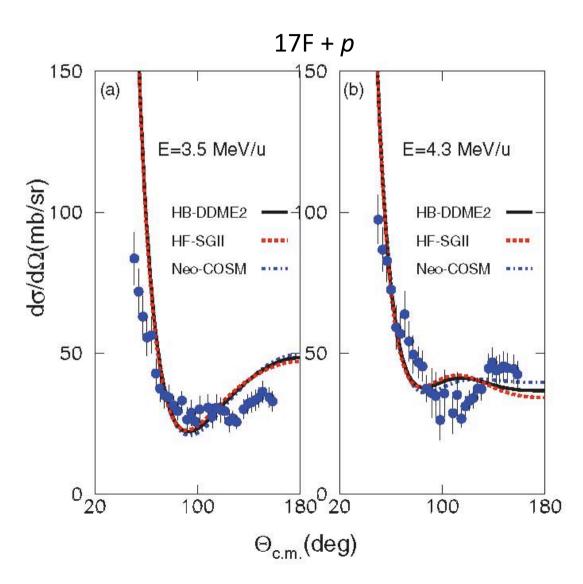
"universal" potentials, eg AW, São Paulo

Example of sensitivity (or not) to different microscopic inputs Elastic scattering  $p+^{17}F$  at LNL (Pakou et al)



Using different optical potentials on the market

Now using JLM folding procedure with different structure options for the <sup>17</sup>F density: rather similar results, hence not great resolving power in this case



JLM Parameters at E=3.5 (4.3)

COSM (cluster orbital shell model, Masui, Kato, and Ikeda)  $\lambda_{V} = 0.8 \ (0.93) \quad \lambda_{W} = 0.75 \ (0.86)$ 

DDME2 (Relativistiv Mean Field, G. Lalazissis,)  $\lambda_{\text{V}} = 0.85 \ (0.99) \ \lambda_{\text{W}} = 0.73 \ (0.65)$ 

HF-SGII (Hartree Fock, AV)  $\lambda_{V} = 0.86 (0.90) \quad \lambda_{W} = 0.66 (0.66)$ 

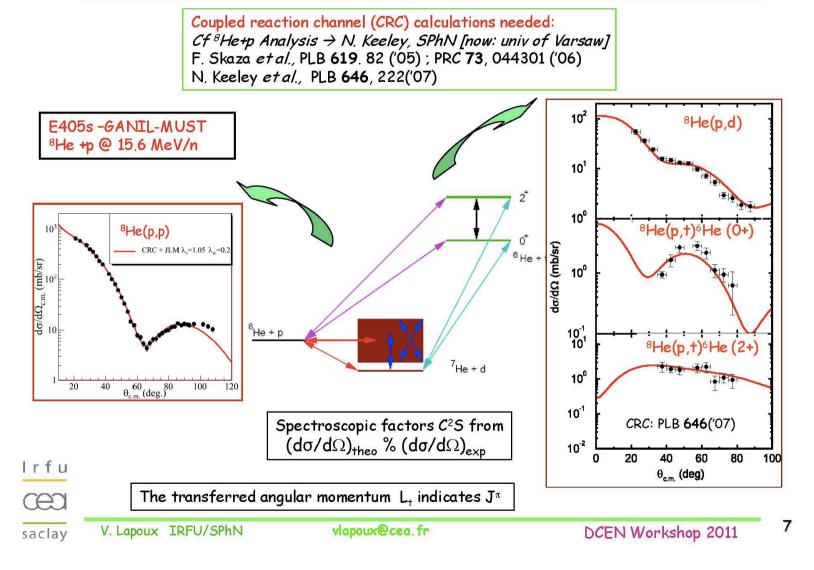
Basic problem: consistent treatment of all relevant diagonal (potentials) and non diagonal (formfactors) matrix elements in a sufficiently large model space (to include explicitly all strongly coupled channels, but not too large to be numerically hard to treat). Transfer channels obviously included, with all consequent problems

OBS problems more explicit in the case of weakly-bound systems (coupling to continuum states .....)

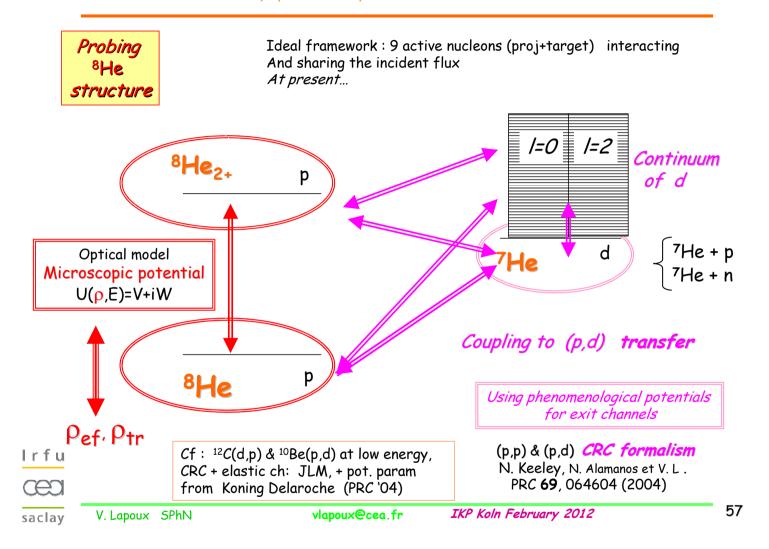
Rather complicate affair .....

## Example: 8He + p

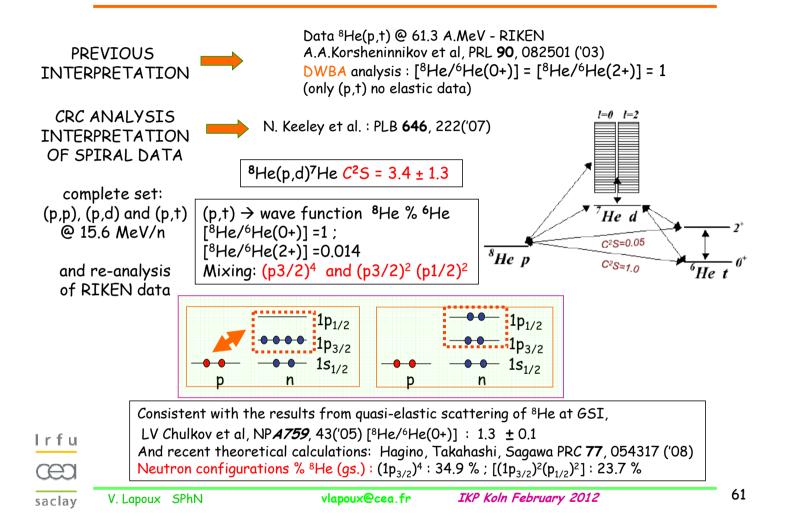
Interpretation of direct reactions: ex of <sup>8</sup>He+p @ 15.6 MeV/nucleon



Courtesy of Valerie Lapoux



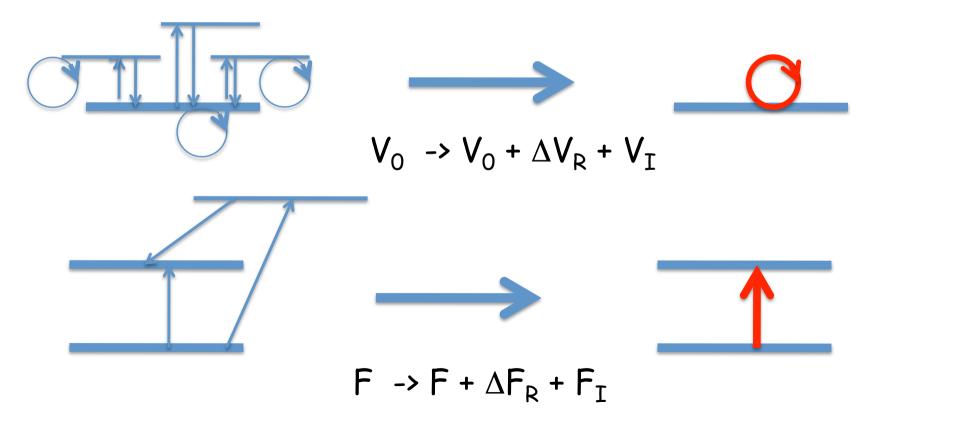
### Courtesy of Valerie Lapoux



Polarization contribution for optical potential and formfactors (due to the reduction of model space). Construction of effective potentials and couplings

Need for simple and reliable prescription for the exploitation of the Feshbach prescription, beyond the Coulomb excitation case and/or perturbative situations (eg Broglia, Pollarolo, Winther).

Complex bare couplings? Weakly-bound systems and continuum-continuum: Repulsive polarization contribution?



## Inelastic scattering (the traditional domain of "collective modes")

Need for microscopic formfactors
(for collective and non-collective excitations)
Role of higher-order couplings (beyond standard DWBA)

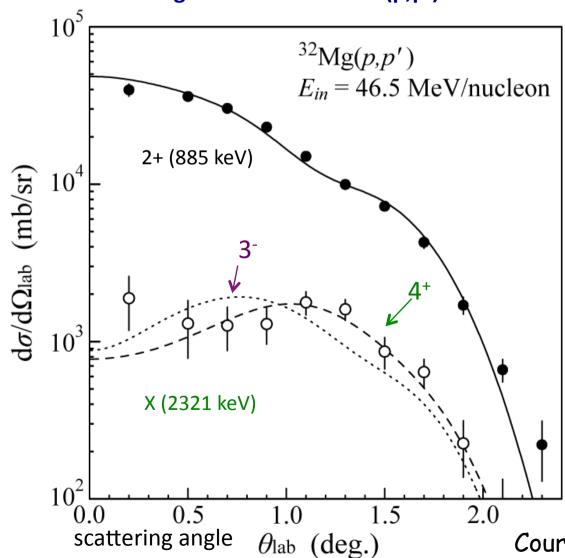
Polarization contribution to formfactors (imaginary part?)

Excitation to unbound states

### Example of inelastic scattering in a strongly coupled system

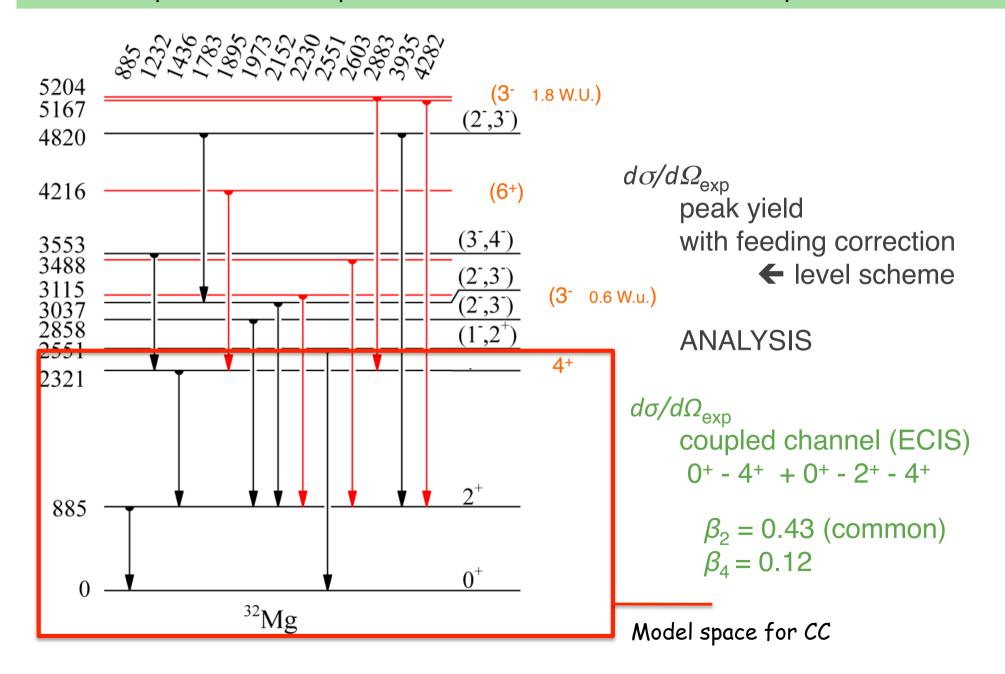
<sup>32</sup>Mg(p,p') in inverse kinematics at DALI2 (RIKEN)- the 2332 keV state

### Angular distribution of (p,p')

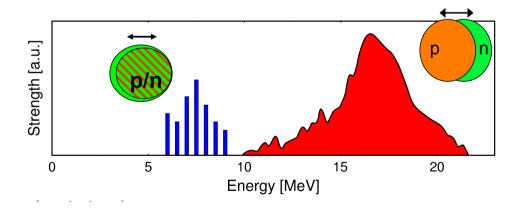


Courtesy of Tohru Motobayashi

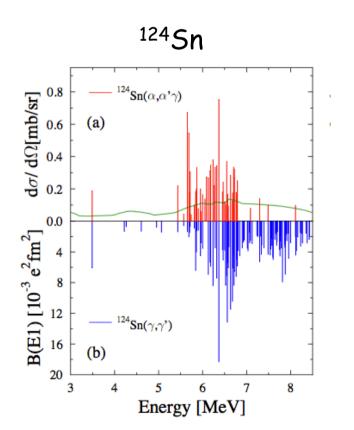
## <sup>32</sup>Mg(p,p') - Coupled channel analysis with collective formfactors Need for microscopic formfactors? Truncation of model space? Nuclear and Coulomb equal deformation parameters? Can we learn more from experiment?

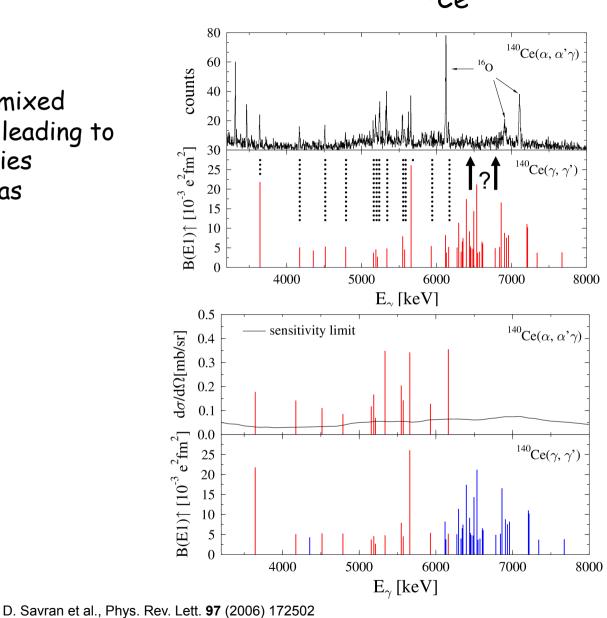


An example where the use of microscopic formfactors is compulsory: excitation of PDR (Pygmy Dipole States) in neutron-rich nuclei



Low-lying dipole states have a mixed isoscalar/isovector character, leading to different excitation probabilities tested with different probes, as  $(\alpha, \alpha')$  or  $(\gamma, \gamma')$ 

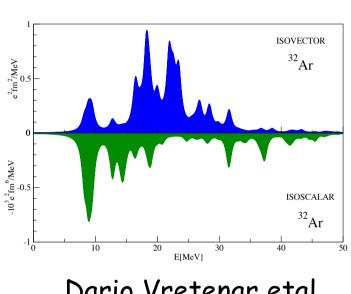




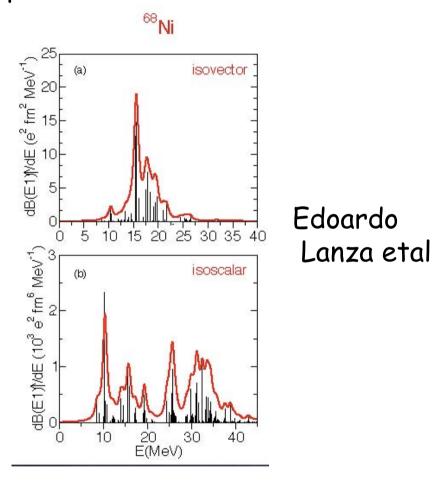
J. Endres et al., Phys. Rev. C 80 (2009) 034203

J. Endres, et al., Phys. Rev. Lett. 105 (2010) 212503

## from theory one gets response to isovector and (leading-order) isoscalar dipole operator



Dario Vretenar etal



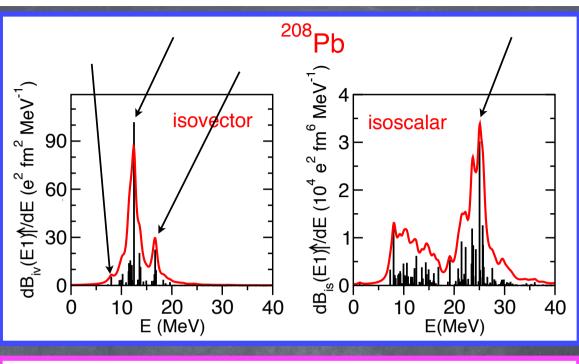
But how the cross sections for reactions as  $(\alpha, \alpha')$  depend on the (isoscalar) dipole response? They are connected, but not proportional. One has to pass through the explicit construction of microscopic formfactors using transition densities that must be proveded by structure calculations

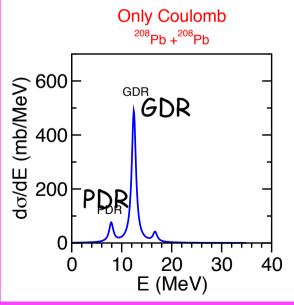
Calculations of inelastic cross section for four selected states

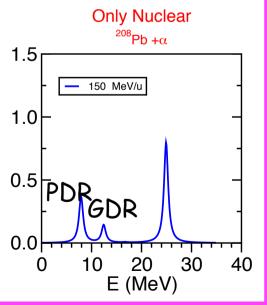
Different relative population of PDR and GDR with different probes

Microscopic FF for nuclear excitation have been obtained from microscopic RPA transition densities

Edoardo Lanza etal







One-particle transfer and knock-out (the traditional domain of "single-particle degree of freedom")

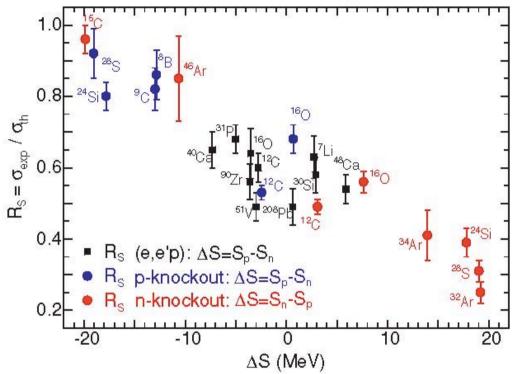
Hot issue: the spectroscopic strength and its deviation from the "bare" single-particle picture

This implies solving quite a number of points, arising from

1. structure models but also

2. reaction models and associated parameters (optical potentials, mean-field potentials generating single-particle wf, higher-order processes, etc)

OBS All problems blow up in the case of weakly-bound systems



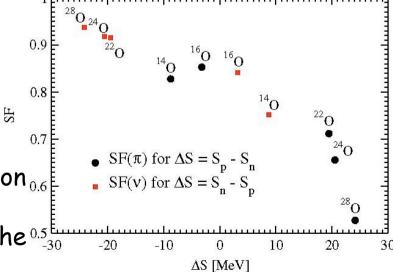
Systematic behavior: spectroscopic factors as a function of the difference In the proton and neutron Fermi energy (a measure of proton/neutron mass asymmetry)

From Alexandra Gade

Quenching of spectroscopic Factors for Proton Removal in Oxygen Isotopes Ø. Jensen etal, 2011

Example of "sophisticated" structure calculation 0.6 Coupled-cluster calculation N3LO including coupling to scattering states above the 0.5

including coupling to scattering states above the  $0.5\frac{E_{111}}{30}$  neutron separation threshold



Quoting Alan Wuosmaa (Helios collaboration) for the light nuclei

Despite concerns about the reliability of reaction theory, good absolute agreement with *ab-initio* form-factor calculations can be achieved at the 20-30% level for (d,p), even for unbound states.

Other reactions are problematic: e.g. absolute (d,t) and (d,3He) results show a strong dependence on optical-model parameters

In all cases, it seems that relative spectroscopic factors can be reproduced very well by *ab-initio* or SM calculations in many (but not all) cases.

Interesting to extend studies further to unbound states to test structure calculations

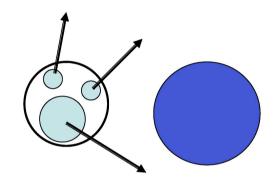
A wise opinion (John Schiffer)

Simple ways of treating data can allow us to extract essential structural information, even if reaction theorists are not completely happy.

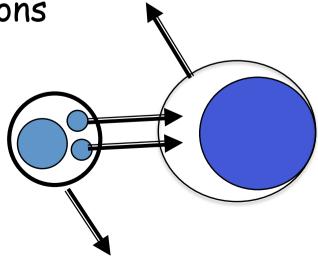
One hopes that a new generation will be able to start from what was Learned, and build on it, and not get bogged down with the fascinating sophistries of reaction theories. We need not have to rediscover all the blind alleys (of two-step processes, coupled channels ... etc) that obscured simple underlying information

## Reactions involving pairs of nucleons (the traditional domain of "nuclear correlations")

- 1. two-nucleon knockout reactions
- 2. two-nucleon break-up reactions



3. two-nucleon transfer reactions



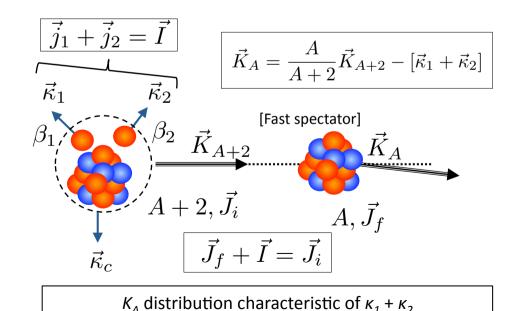
#### 1. two-nucleon knockout reactions

Basic information arise from the momentum distribution, which turns out to be depending on the structure and the correlations in initial and final states

The process can be described in terms of two "correlated" knock-out of single neutrons

(Jeff Tostevin etal, Edward Simpson etal)

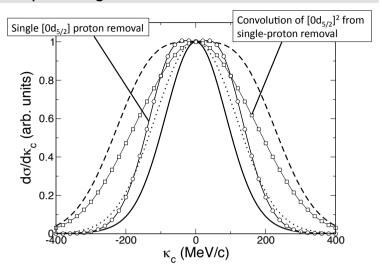
#### Momentum distributions



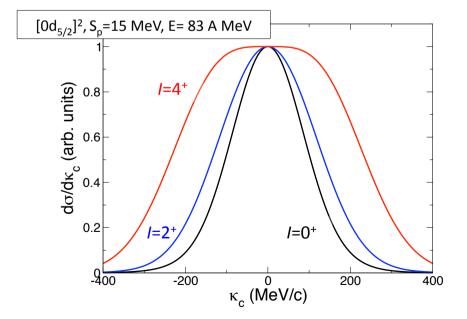
From a presentation by Edward Simpson

Dependence of momentum distribution on the angular momentum coupling of the two nucleons and on the specific orbitals

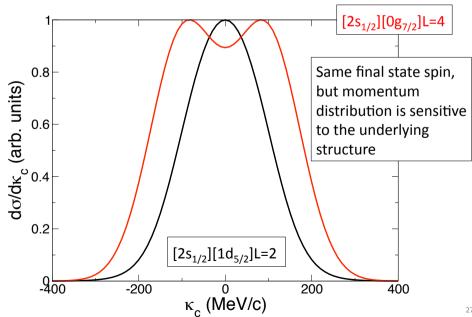
#### Example I: Single/Uncorrelated





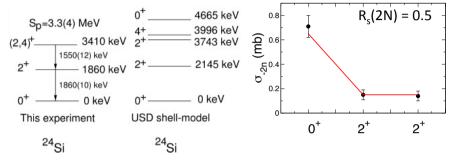


### Example: $^{208}\text{Pb}(-2p) \rightarrow ^{206}\text{Hg}(J_f=3^+)$



#### <sup>26</sup>Si(-2n): Cross section results

#### Results



Shell model (sd-shell, USD) two-nucleon amplitudes

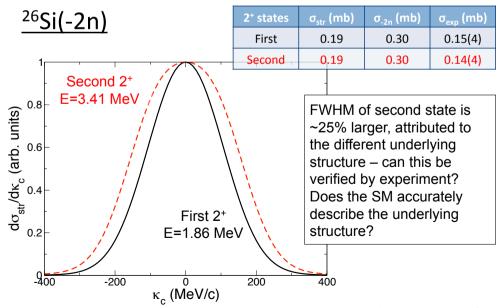
| State       | [0d <sub>5/2</sub> ] <sup>2</sup> | [0d <sub>5/2</sub> ,0d <sub>3/2</sub> ] | [0d <sub>3/2</sub> ] <sup>2</sup> | [1s <sub>1/2</sub> ,0d <sub>3/2</sub> ] | [1s <sub>1/2</sub> ,0d <sub>5/2</sub> ] |
|-------------|-----------------------------------|-----------------------------------------|-----------------------------------|-----------------------------------------|-----------------------------------------|
| 2+ (First)  | -0.70074                          | 0.43499                                 | 0.00594                           | -0.00188                                | -0.02781                                |
| 2+ (Second) | -0.38021                          | -0.12354                                | -0.12945                          | -0.15876                                | -0.58292                                |
| 4+ (First)  | 1.57469                           | 0.41519                                 | -                                 | -                                       | -                                       |

Yoneda et al., PRC 74, 021303(R) (2006)

An example of detailed "microscopic" calculation based on microscopic correlated WF's

Requirement for full success: absolute cross section and momentum distribution

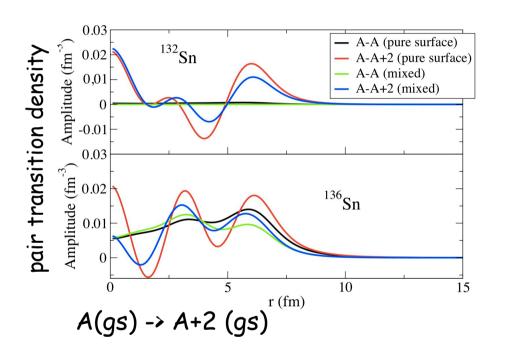
#### Structure Sensitivity

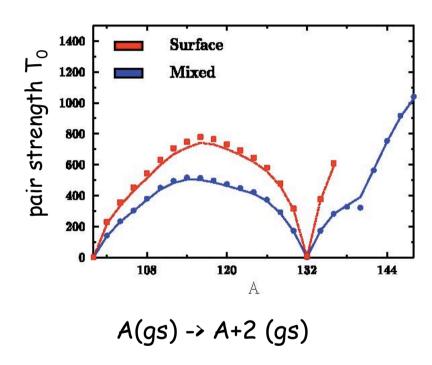


### 2. two-nucleon transfer reactions

It is widely accepted that pairing correlations strongly effect (and enhance) two-particle transfer reactions. But the quantitative connection is not obvious. Will cross section scale with the square of the two-particle transfer matrix elements? Or the radial dependence of the two-particle transition densities contain more information? And how this information enter into the reaction mechanism?

Example of structure investigation of pairing correlations: can we discriminate among different forms of pairing interactions? Example: can we distinguish pure surface pairing interaction from mixed (volume plus surface) in HFB calculations? (from Grasso, Lacroix, AV)





How the different behavior in the tail enter in the reaction mechanism? Is only the integrated value  $T_0$  relevant?

BASIC PROBLEM: The reaction mechanism

Variety of models on the market

The fully microscopic approach is based on sequential two-step process (each step transfers one particle)

Microscopy: Pairing enhancement comes from the coherent interference of the different paths through the different intermediate states in (a-1) and (A+1) nuclei, due to the correlations in initial and final wave functions

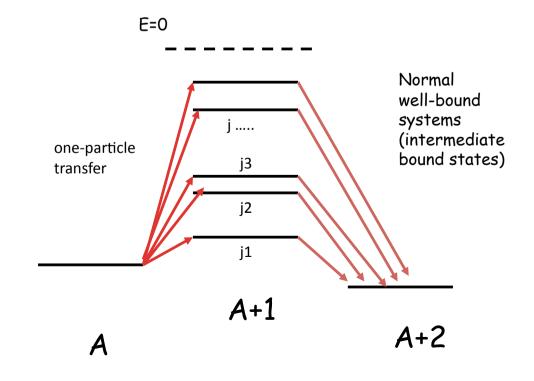
Building blocks: single-particle formfactors and wf's

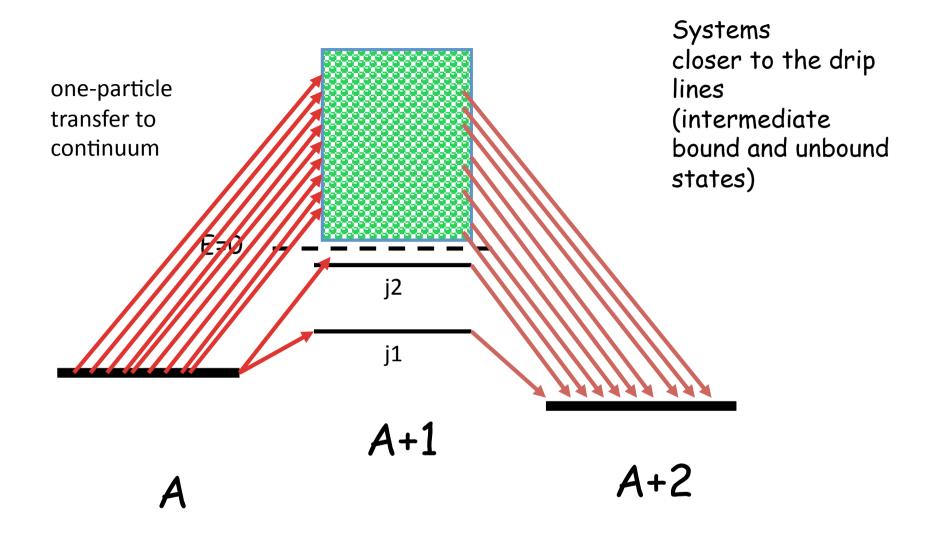
Problems: quantal calculations rather complex (taking into account full recoil), semiclassical more feasible (but approximate treatment of recoil)

All microscopy and nuclear structure information are contained in the two-particle transfer amplitudes (from correlated initial and final wave functions), which give the weight of each two-step path, and in the single particle transfer formfactors, which need single particle wavefunctions in target and projectile

Obs: Basic idea: dominance of mean field, which provides the framework for defining the single-particle content of the correlated wave functions

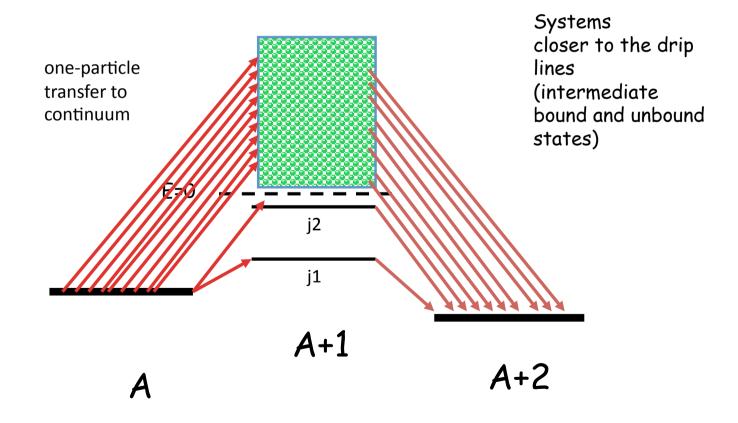
Examples in this meeting by Ricardo Broglia and Rituparna Kanungo

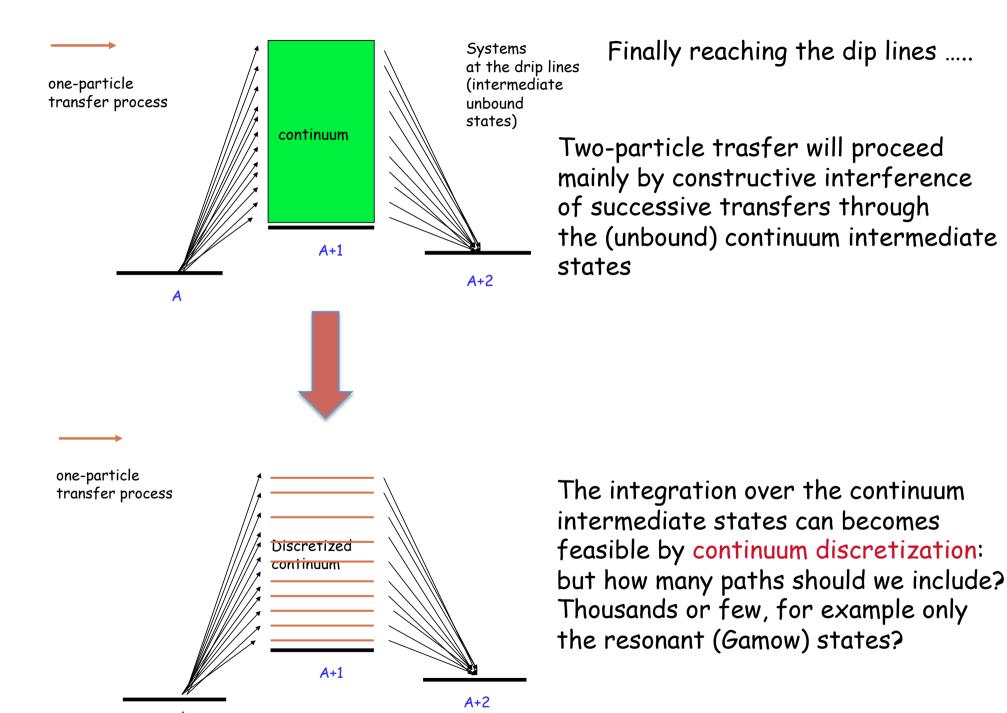




$$|A=2> = \{\sum_{i} X_{i} [a_{i}^{+} a_{i}^{+}]_{0}^{+} \int dE X(E) [a^{+}(E)a^{+}(E)]_{0} \} |A>$$

But moving from the stability line ......





## 3. Two-particle break-up processes

Break-up of a two-particle halo system is a rather complex 4-body process. To make it simpler let us consider an one-dimensional case, that should hopefully display the essential features of the three-dimensional case

## One-dimensional three-body model



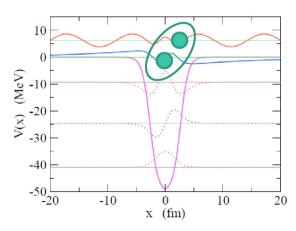
Two interacting neutrons in a one-dimensional potential well:

$$H = -\frac{\hbar^2}{2m} \frac{d^2}{dx_1^2} + V(x_1) - \frac{\hbar^2}{2m} \frac{d^2}{dx_2^2} + V(x_2) + v_{nn}(x_1, x_2)$$

density-dependent contact interaction:

$$v_{nn}(x, x') = -g \left(1 - \frac{1}{1 + e^{(|x| - R)/a}}\right) \delta(x - x')$$

Hagini, Vitturi, Sagawa, PerezBernal 2011



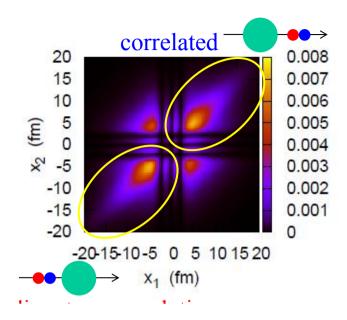
$$\Psi_{gs}(x_1, x_2) = \sum_{n \le n'} \alpha_{nn'} \Psi_{nn'}(x_1, x_2)$$

$$\Psi_{nn'}(x_1, x_2) \propto \mathcal{S}[\phi_n(x_1)\phi_{n'}(x_2)] \times |S = 0\rangle$$

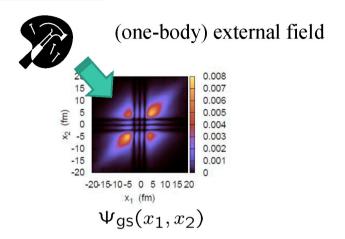
•S = 0 state: symmetric for the spatial part of wf

•n, n': the same parity

two-particle density:  $|\Psi_{gs}(x_1, x_2)|^2$ 



#### **Nuclear Breakup Process**



Time-dependent two-particle Schroedinger equation:

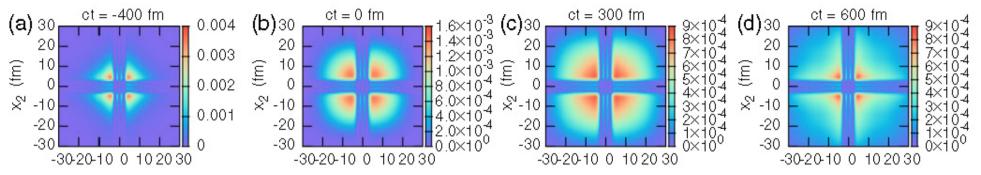
$$i\hbar \frac{\partial}{\partial t} \Psi(x_1, x_2, t) = [H + V_{\text{ext}}(x_1, x_2, t)] \Psi(x_1, x_2, t)$$

$$V_{\text{ext}}(x_1, x_2, t) = \sum_{i=1,2} V_c e^{-t^2/2\sigma_t^2} e^{-(x_i - x_0)^2/2\sigma_x^2}$$

 $V_{\rm c} = 3 {\rm MeV}, \, \sigma_{\rm t} = 2.1 {\rm \ hbar/MeV}, \, x_0 = 0$ 

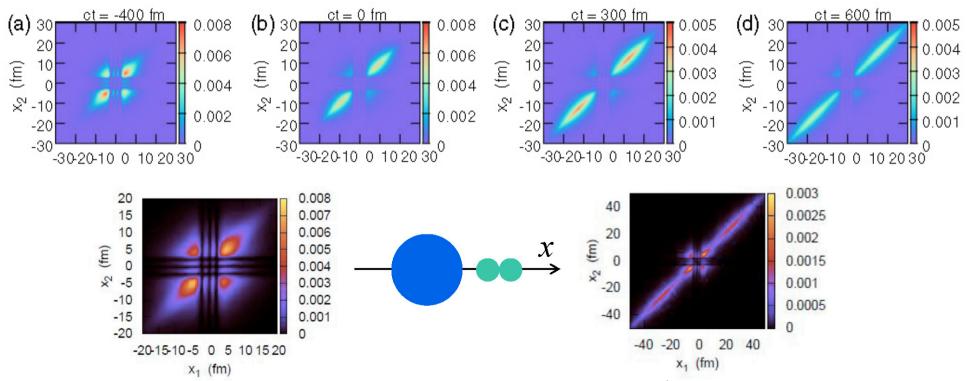
The perturbing interaction (that produces the break-up) is a one-body field (i.e. acting individually on each of the two particles). The enhanced two-particle break-up originates from the correlation in the two-particle wave function, and not from a forced two-neutron cluster reaction mechanism

## Time evolution (uncorrelated case)



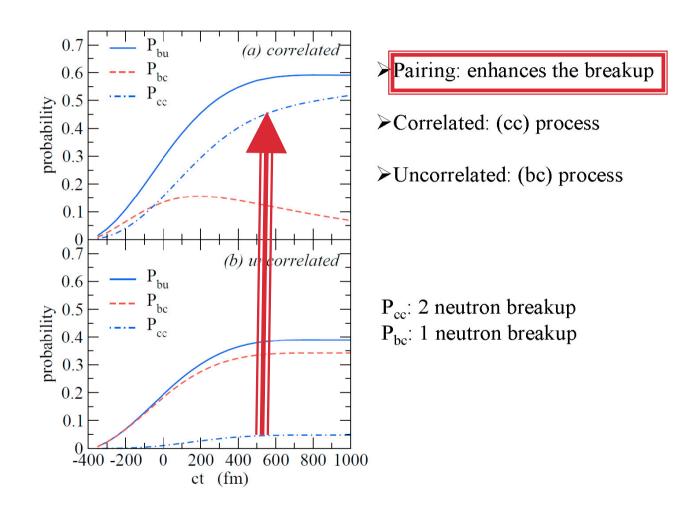
OBS: Large component with only one-particle in the continuum

## Time evolution (correlated case)



Large di-neutron emission

As a net result, the two-particle break-up process is enhanced by the correlation (although the external perturbing field is of one-body nature)



## Sub-barrier heavy-ion fusion

Strictly speaking this is not a "direct" process (on the contrary), but strongly effected by the competing direct channels

Interplay of different coupled channels (elastic, inelastic, transfer)

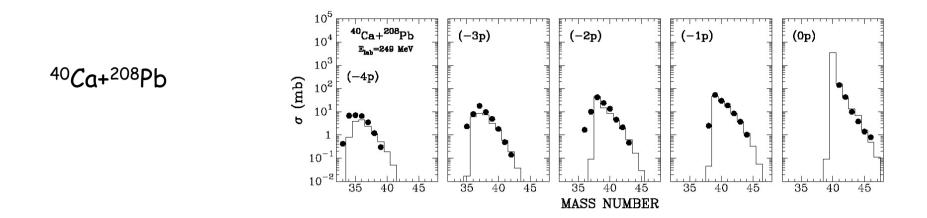
For weakly-bound systems still interesting issue of the effect of the coupling to continuum (enhancement or suppression?)

## Multi-nucleon transfer reactions

Again, not a "genuine" direct process, but a key case as test of pairing modes in the "vibrational multiphonon-like" and in the "rotor-like" pairing cases. In the case of population of excited states is fundamental in describing the transition from grazing reactions to more central deep-inelastic collisions)

OBS: instrumental for structure studies with  $\gamma$ -spectroscopy for systems far from stability, but this is another story ....

#### Example of multi-nucleon transfers at Legnaro

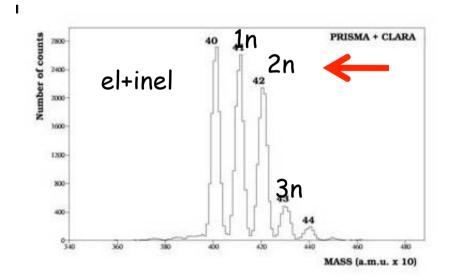


Obs: transfer of particles on both directions

Transition from direct to deep inelastic (cf Q distributions)

Example: Neutron transfer channels

(odd-even transfer effect? Structure effect?)



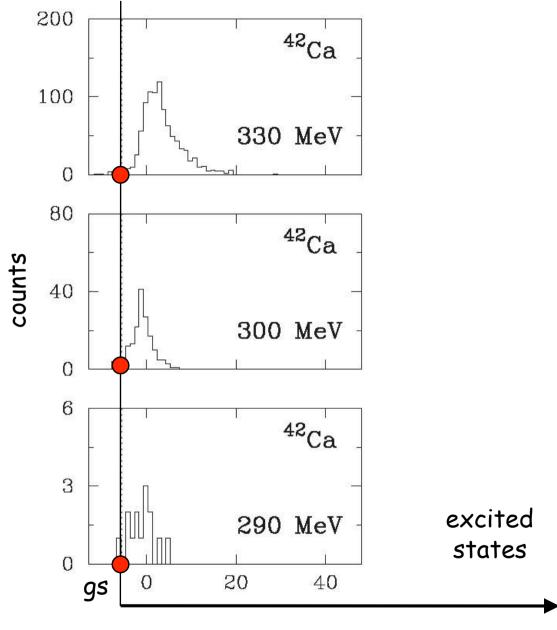
## Experimental evidence

Negligible transitions to GS due to Q-value effects.
What information on pairing correlations?

Example

<sup>96</sup>Zr+<sup>40</sup>Ca

Selecting final <sup>42</sup>Ca mass partition

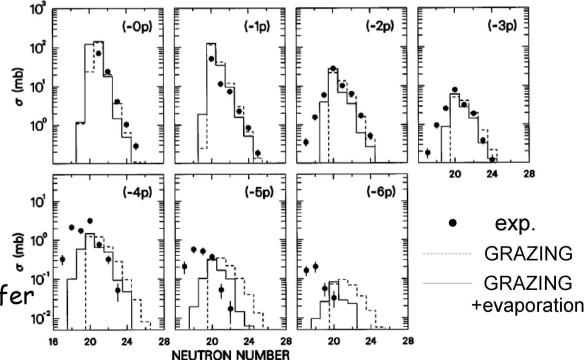


Total kinetic energy loss (MeV)

# Basic and most popular approach: "Grazing model" (Nanni Pollarolo and Aage Winther)

- semiclassical description of trajectory
- single-particle transfers
- two-particle transfers (double counting?)
- collective inelastic excitations
- sufficient phase-space for multi-transfer?
- "bare" ion-ion potential?
- structure information?
- excellent for "average" behavior.
   Specific cases?
   Weakly-bound systems and treatment of continuum?
- collective vs non-collective transfer (and non L=0 pairs)

<sup>40</sup>Ca+<sup>124</sup>Sn El<sub>ab</sub>=170 MeV



Other approach: TDHF with particle number projection technique (Cedric Simenel), with reduction of computational time by a factor 100 (still in fieri and under discussion the range of validity: maximum number of transferred particles? Residual correlations? In particular pairing correlations? Reaction mechanism beyond mean field?)

Example:  ${}^{40}Ca+{}^{124}Sn E_{lab}=174 MeV$ 

Sekizawa and Yabana SLy5

OBS: Overall agreement is good when transferred proton number is small

