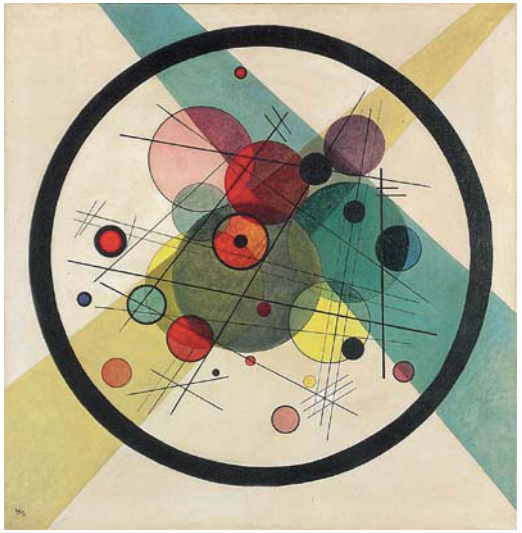
Two-particle transfer reactions: a key tool for the study of phase transitions in nuclei

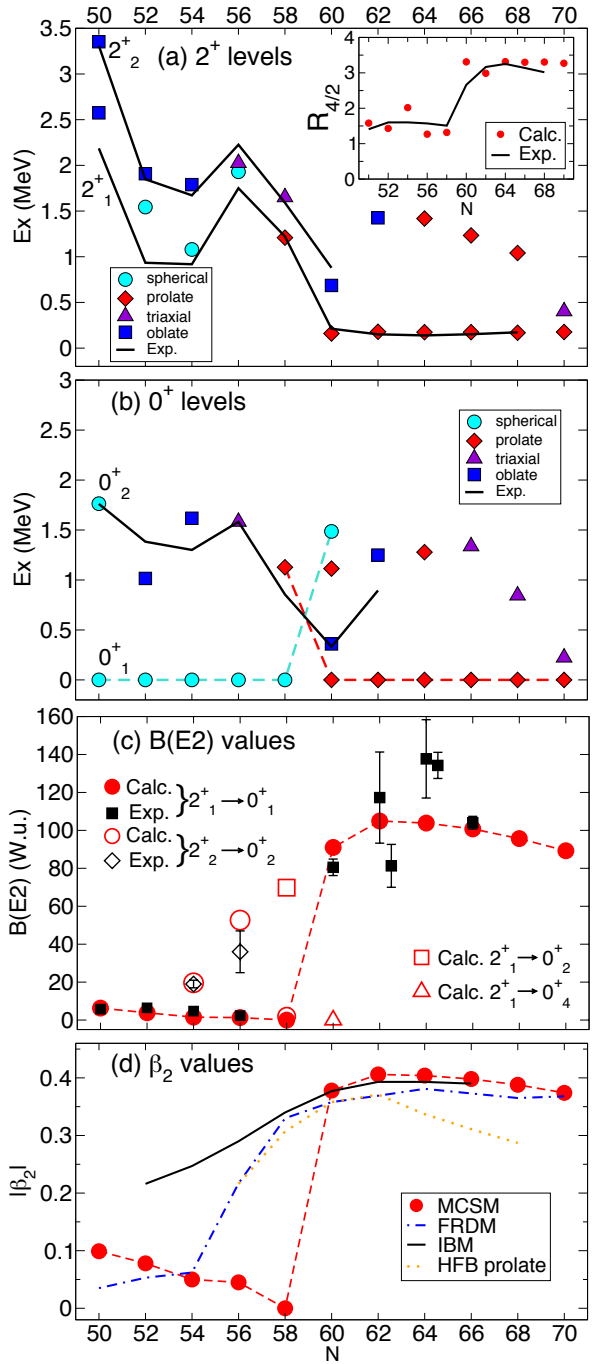
Andrea Vitturi (Padova)



Basic point to discuss: how the nuclear behavior of the pairing degree of freedom can provide an additional and complementary clear-cut signature of the occurrence and on the nature of the phase transition in nuclear systems.

This dynamical source of information should be complementary (but as important) to the one associated to other properties (as for example, in the case of even-even nuclei, the energy of the first 2+ state, the ratio *E4/E2* and the magnitude of the electromagnetic E2 transition connecting ground state and the first excited 2+ state)

How to single out phase transitions?



The main road to use dynamics to study pairing effects along phase transitions is clearly provided by the study of those processes where a pair of particles is involved, e.g. transferred from/to another nucleus (two-particle transfer) or ejected into the continuum (two-particle break-up or
two-particle knock-out). Clearly the probabilities for such processes must be influenced by the particle-particle correlations, but these will depend on the specific "shape phase" of the system. So they will be sensitive to any change in the status of the system, for example along an isotope chain.

The essential quantity to characterize the system from the pairing point of view is given by the "pairing response", namely all the T_0 values of the square of the matrix element of the pair creation (or removal) operator

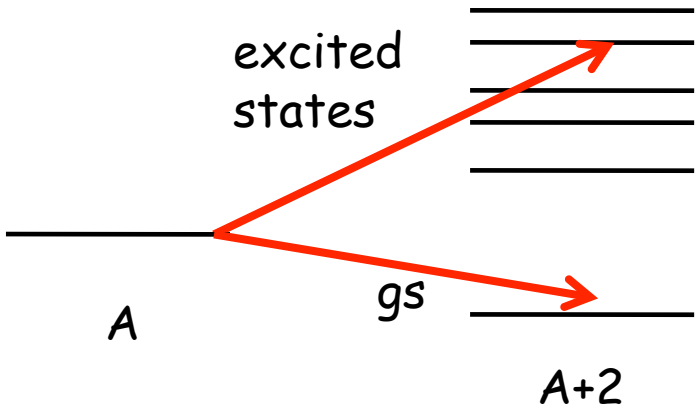
$$P^+ = \sum_j [a^+_j a^+_j]_{00} \qquad \text{(and similarly for } P^-\text{)}$$

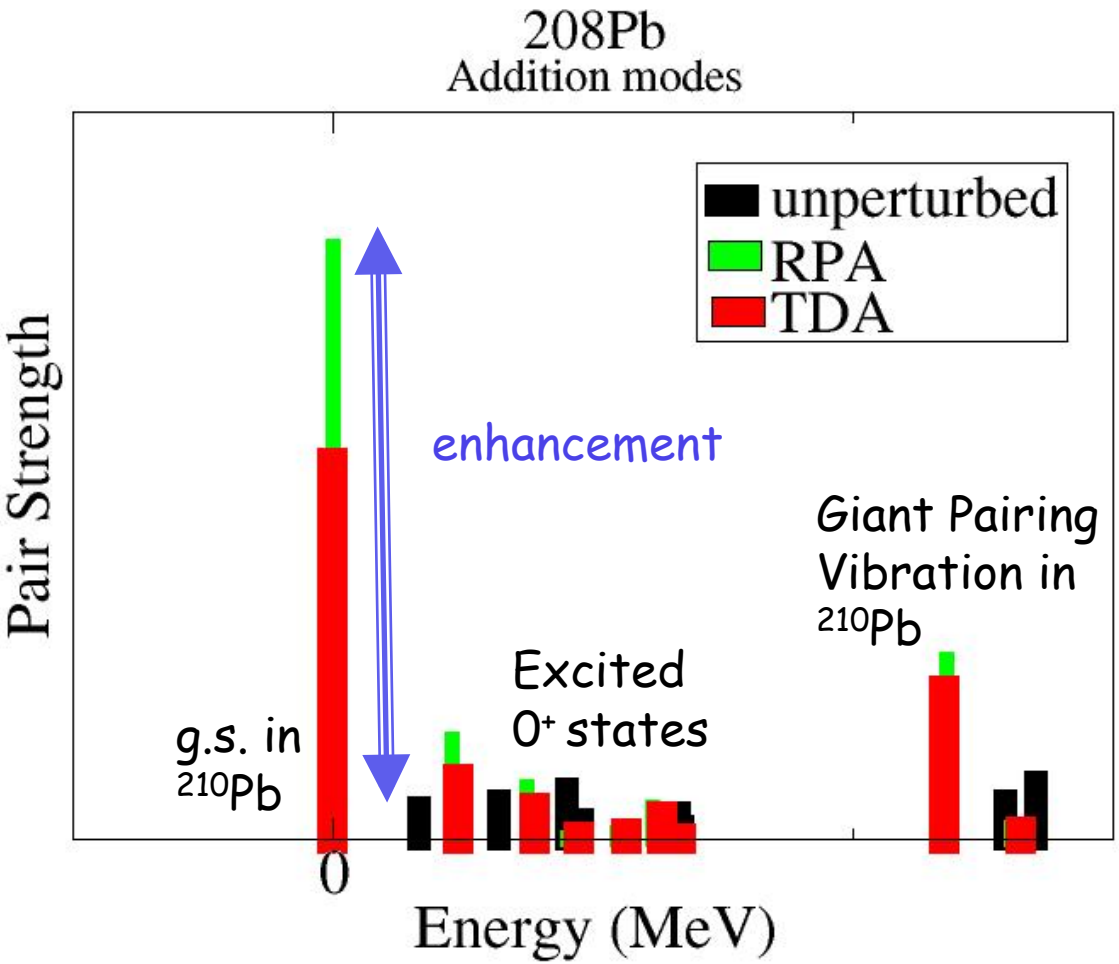
connecting the ground state of nucleus N with all 0+ states of nucleus A+2 (or A-2). It is often assumed that the cross section for two-particle transfer just scale with T_0.

The traditional way to define and measure the collectivity of pairing modes is to compare with single-particle pair transition densities and matrix elements to define some "pairing" single-particle units and therefore "pairing" enhancement factors.

Obs: We discuss here monopole T=1 pairing modes, i.e. 0+ states, but similar arguments would apply to T=0 neutron-proton pairs.

Typical "pairing" response



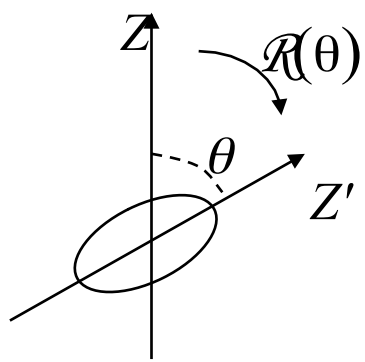
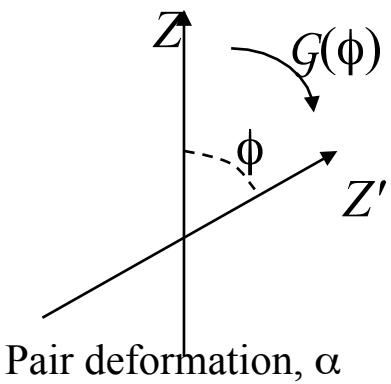


Sofia, Dasso, Vitturi

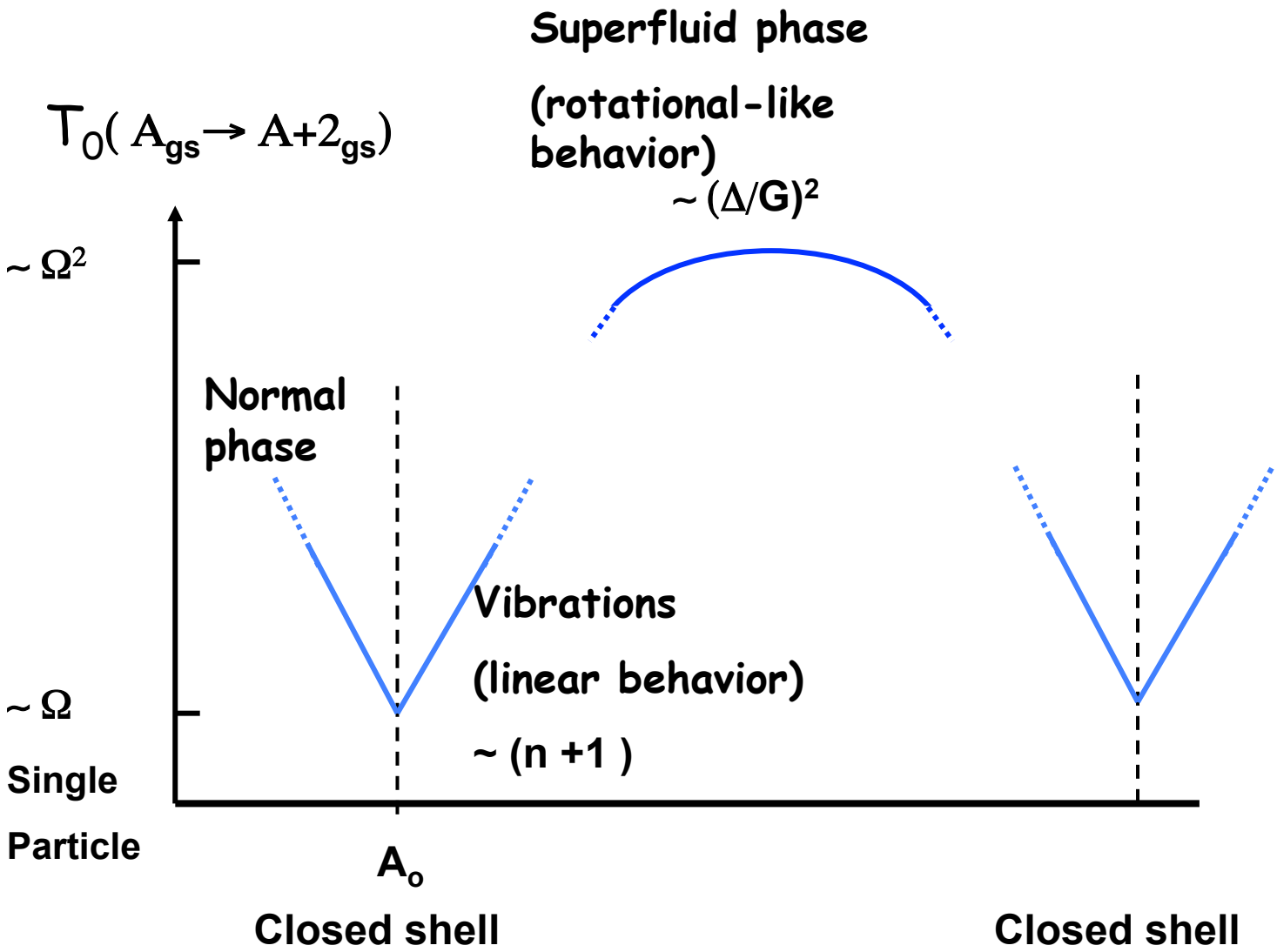
The pairing response is characterized by the pairing phase (normal or superfluid) and by the shape phase (e.g. spherical or deformed). Therefore it will be a clear signature of phase transitions (in addition to the standard signatures, as E_4/E_2, B(E2), etc) in both the

shape degree of freedom

pairing degree of freedom

Shape Transitions	Pairing Transitions
$\mathcal{R}(\theta)=\exp(-iI\theta)$	$\mathcal{G}(\phi)=\exp(-i\mathcal{N}\phi)$
Angular Momentum, I	Particle Number, $\mathcal{N}$
	
β, γ, Euler angles θ	Pair deformation, α Gauge angle, ϕ
Violation of spherical symmetry	Violation of particle number
Physical space	Abstract "gauge" space

Phase transition from "normal" to "superfluid" phases: characteristic behavior of the pair transfer matrix element



OBS: Similar behavior as a function of temperature or angular momentum

PHYSICAL REVIEW C **85**, 034317 (2012)

Pair-transfer probability in open- and closed-shell Sn isotopes

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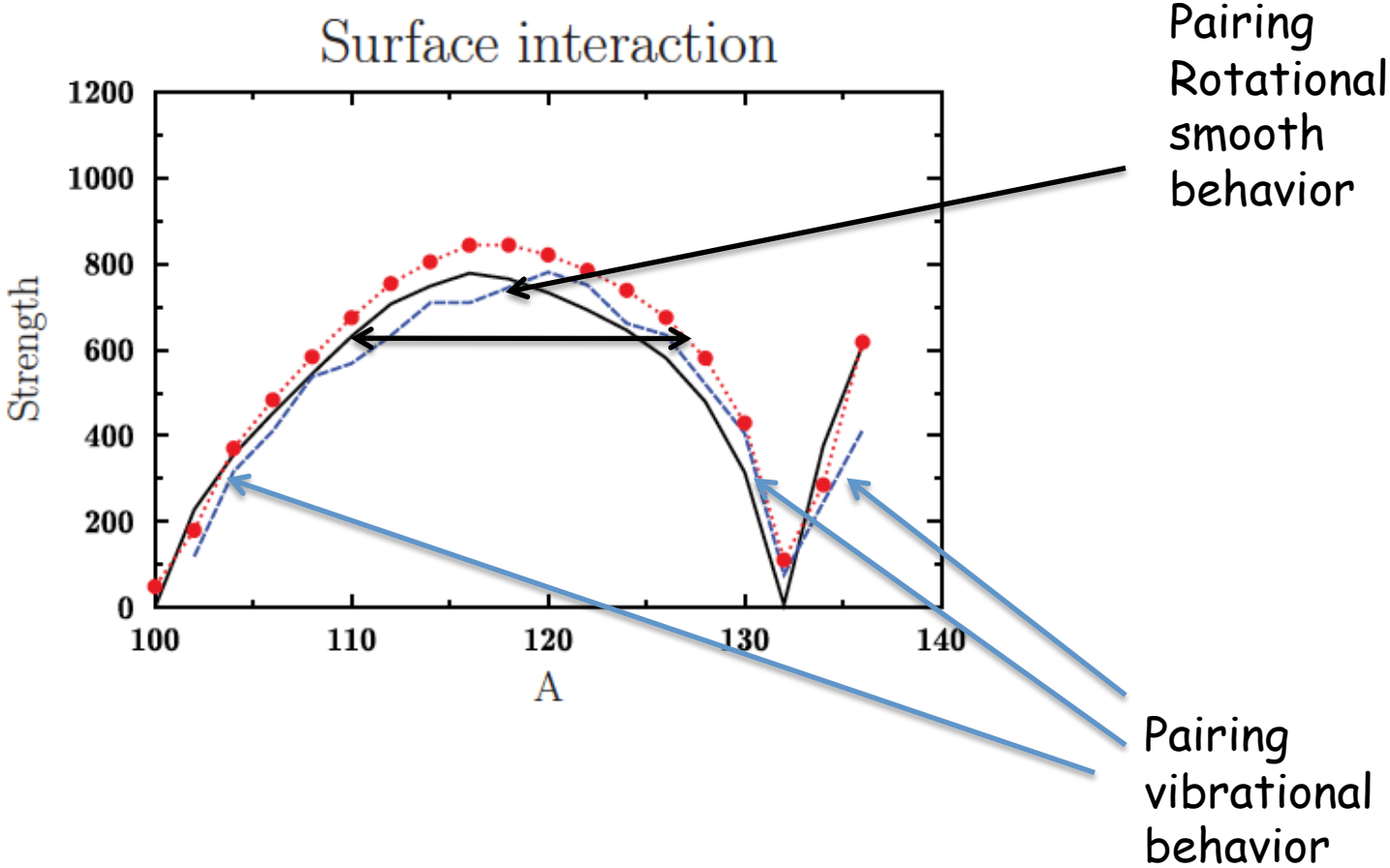
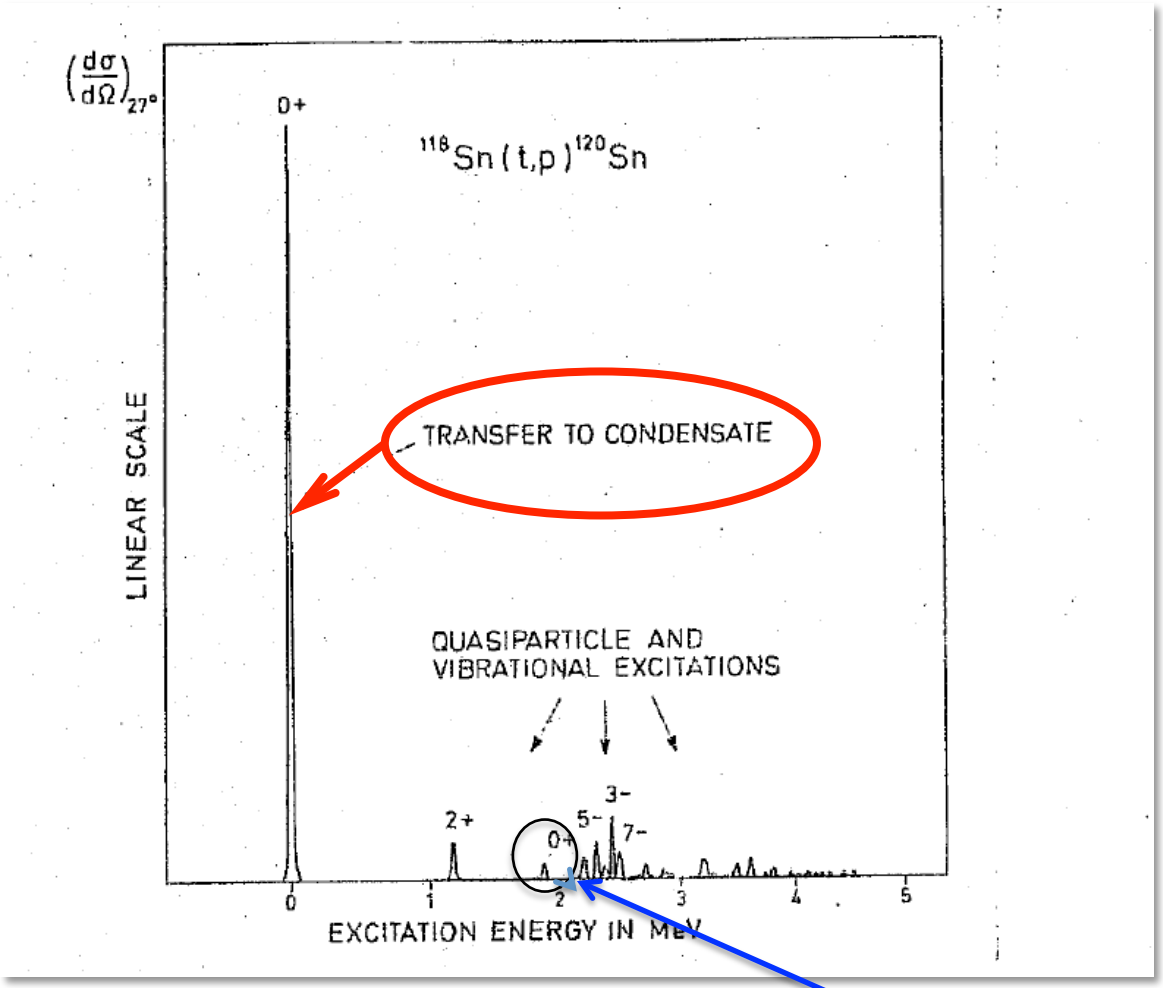


An example of a "superfluid" nucleus (pairing rotations), which shows a characteristic pairing response



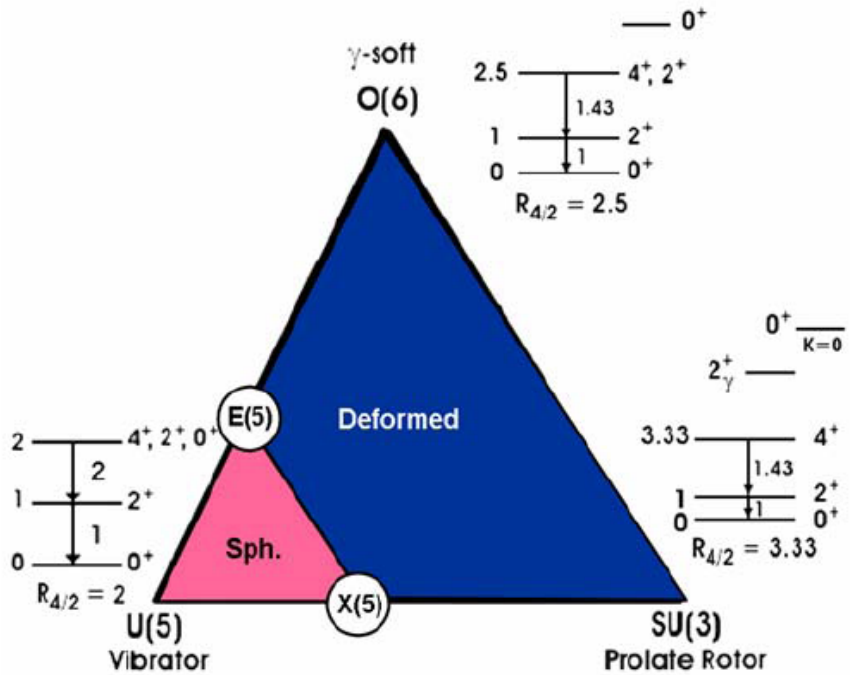
Practically **all** pairing strength goes to the ground state

J.H.Bjerregaard *et al.* NPA 110 **1** (1968)

excited 0+ state

In a similar way pair-transfer probabilities show characteristic behaviors in correspondence of shape phase transitions

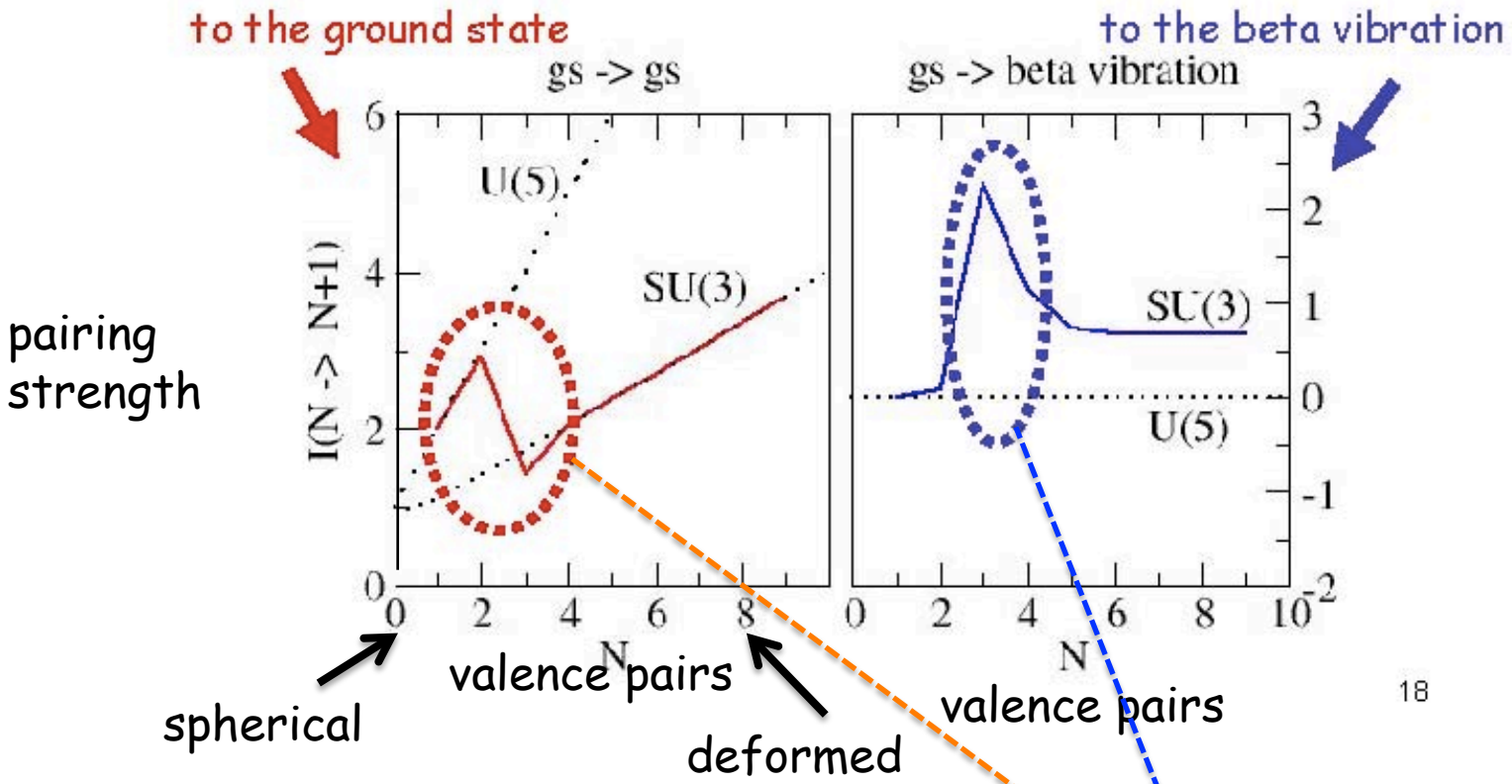
For simplicity we move within the framework of the Interacting Boson Model, but the results are similar within other microscopic models



The IBM does not explicitly use the fermion degrees of freedom. From mapping procedure the "form" of the two-particle addition operator is simply assumed as s^+, neglecting higher-order terms, as s^+s^+s or $[d^+d^+]_0 s$ or $[d^+s^+d]_0$ etc ……

OBS: See OAI mapping

Example: L=0 pair transfer in a phase transition from spherical to axial deformation (from U(5) to SU(3) in algebraic language)

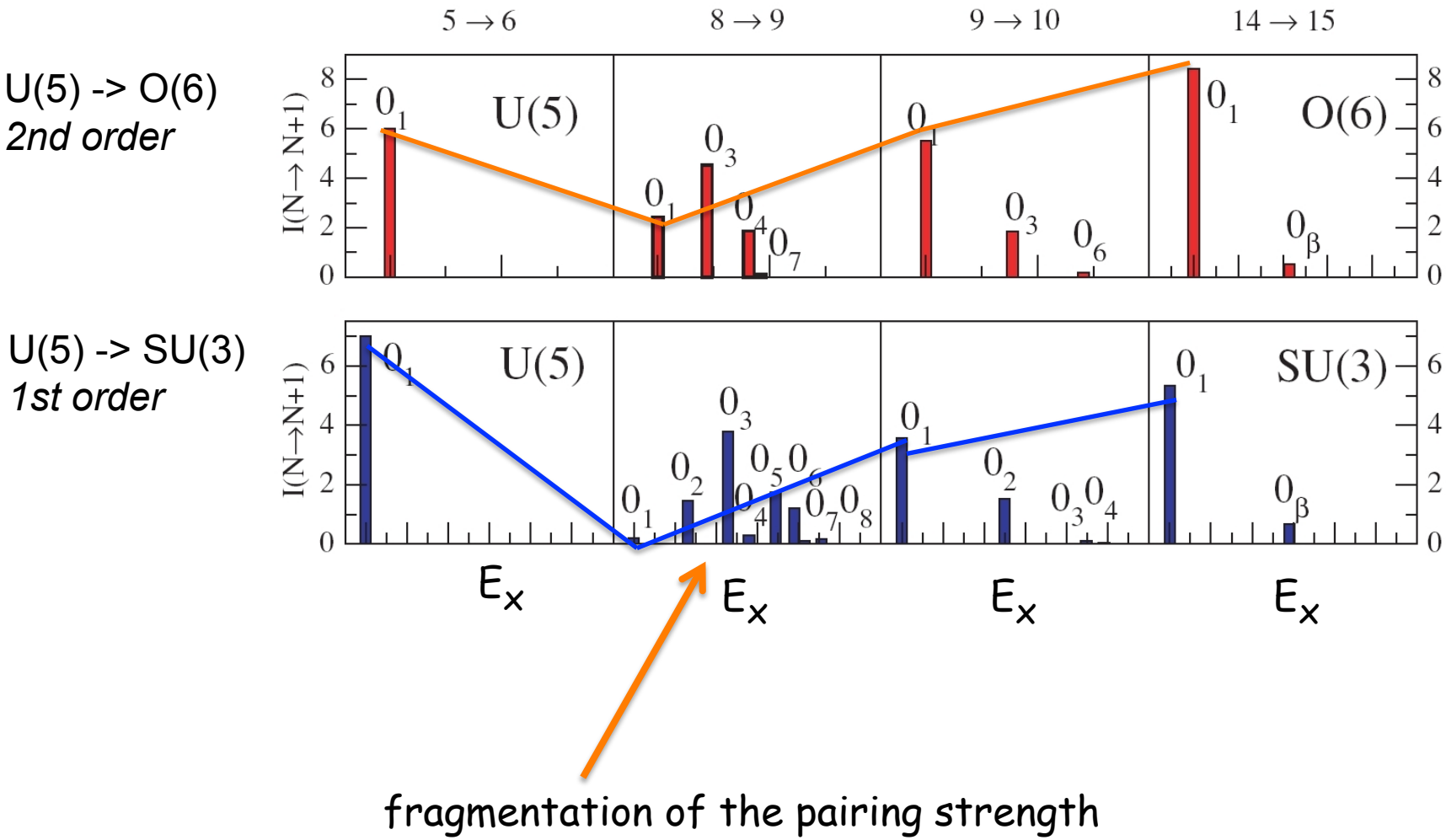


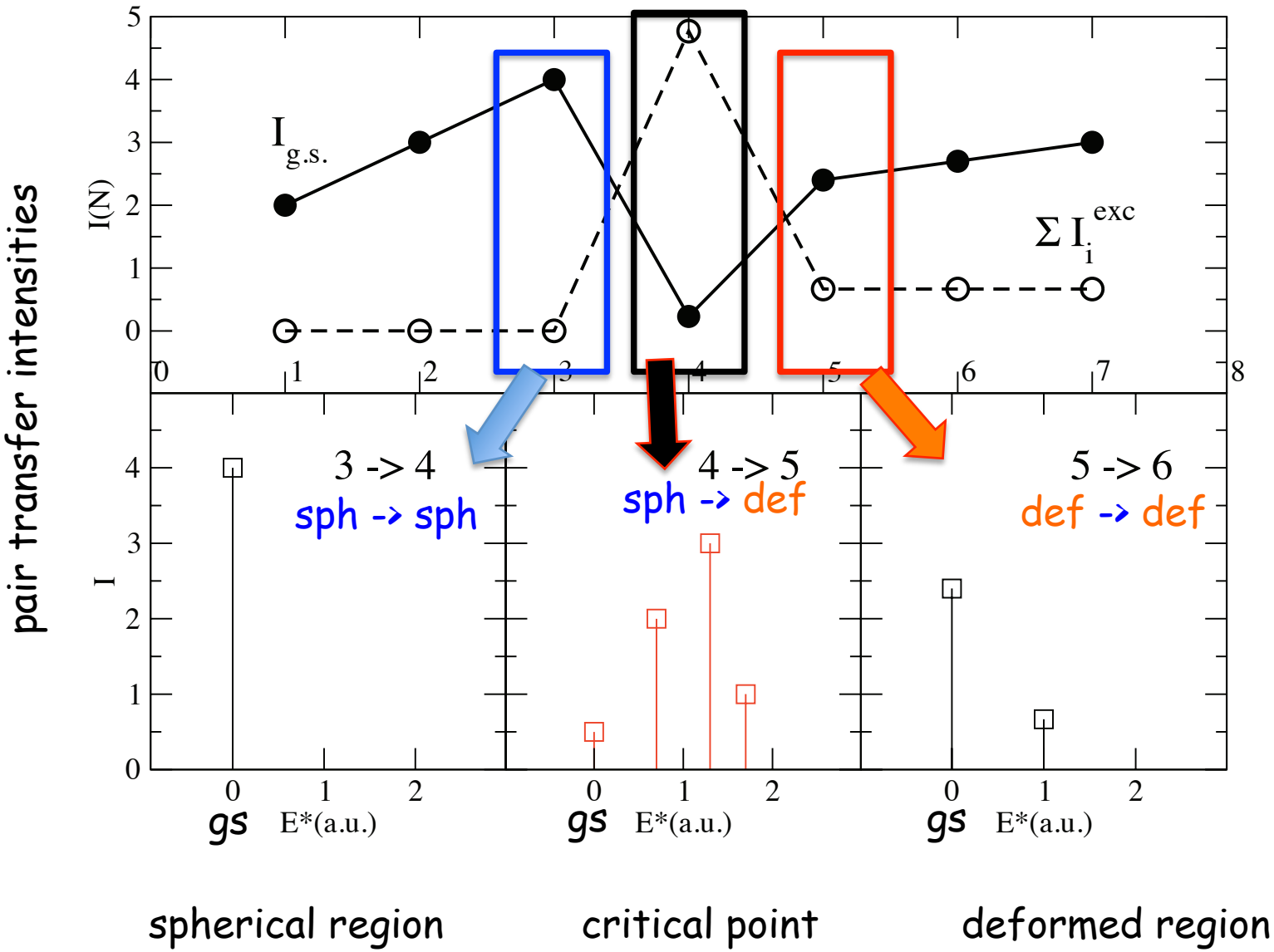
Fossion, Alonso,Arias,Fortunato and A. V.
Cf. also Zhang and Iachello

There is a clear signal at the phase transition

Obs: fragmentation of the pairing strength in correspondence to phase transitions along an isotope chain (in this case chosen to take place at N=8)

Increasing number of valence pairs




pair transfer intensities
I(N)
$I_{g.s.}$
ΣI_i^{exc}
3 -> 4
sph -> sph
4 -> 5
sph -> def
5 -> 6
def -> def
I
gs
E*(a.u.)
spherical region
critical point
deformed region

Another scenario of phase transition:
shape co-existence, for example of a spherical and a deformed state within the same nucleus

$$|0^+_{gs},N\rangle = \alpha\ |N\rangle_{U(5)} + \beta\ |N+2\rangle_{SU(3)}$$

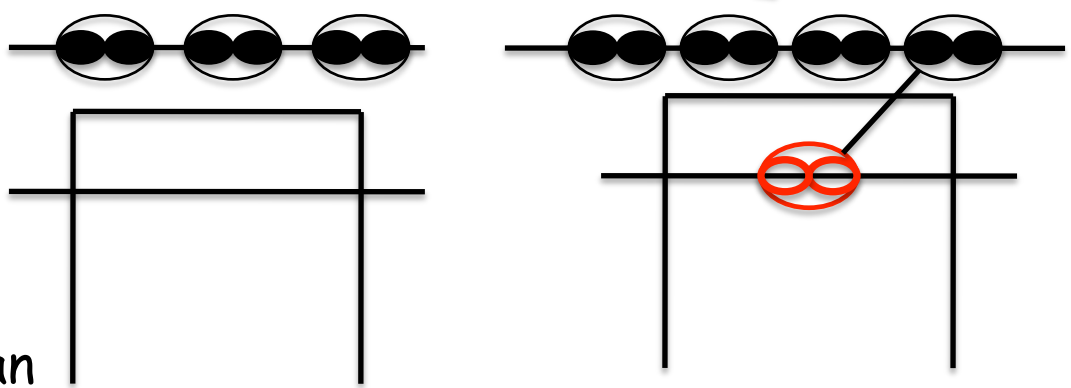
$$|0^+_{exc},N\rangle = -\ \beta\ |N\rangle_{U(5)} + \alpha\ |N+2\rangle_{SU(3)}$$

Mixing of two configurations, with mixture changing along the isotope chain

N particle pairs

0 hole pairs

U(5) hamiltonian (spherical)



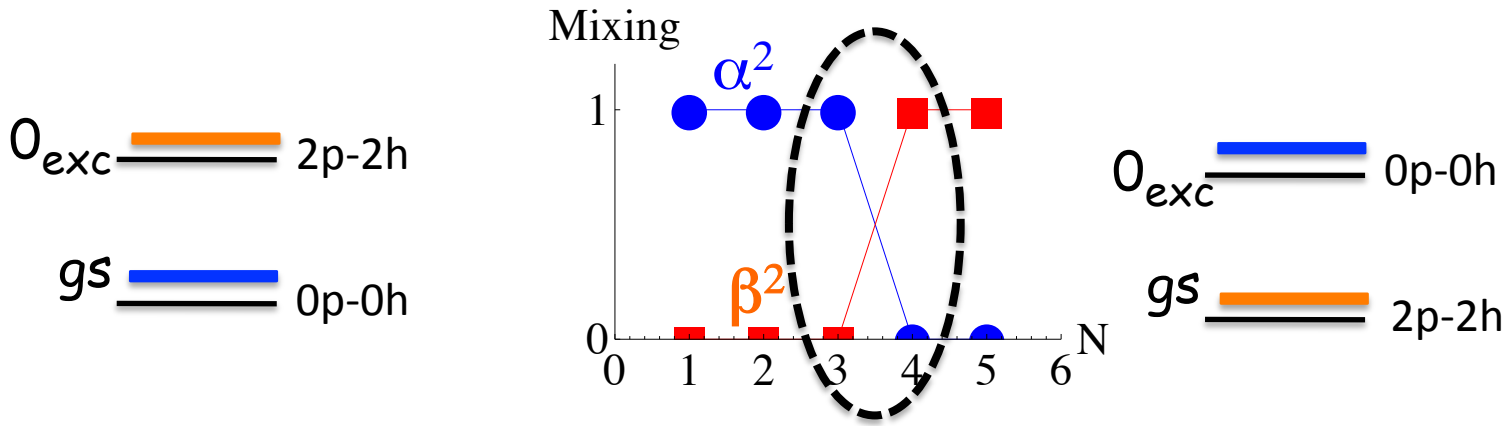
$|N\rangle_{U(5)}$

$|N+2\rangle_{SU(3)}$

N+1 particle pairs,
1 hole pair (2p-2h exc): total N+2 pairs

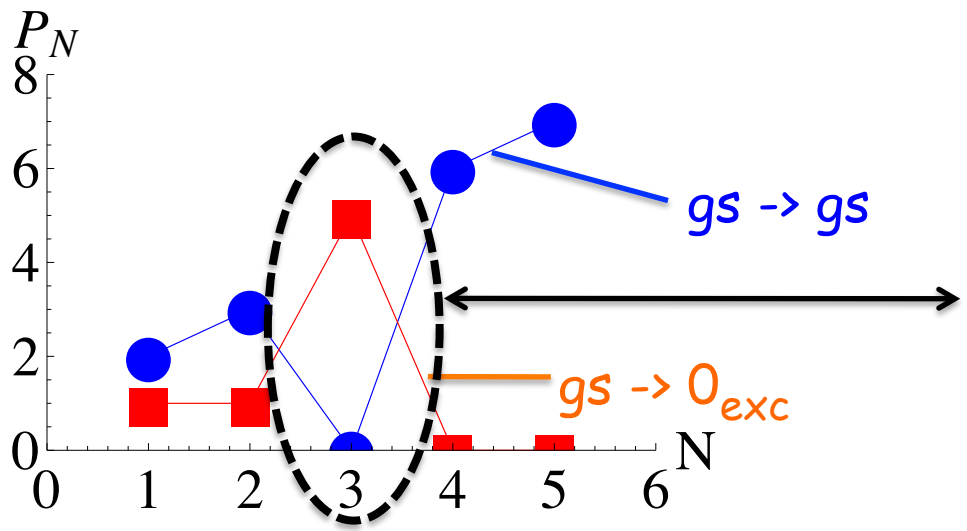
SU(3) hamiltonian (deformed)

A simple model: along the isotope chain a sharp inversion of the structure



Transfer operator in now more complex: s^+ + s (one can create a particle pair or destroy a hole pair)

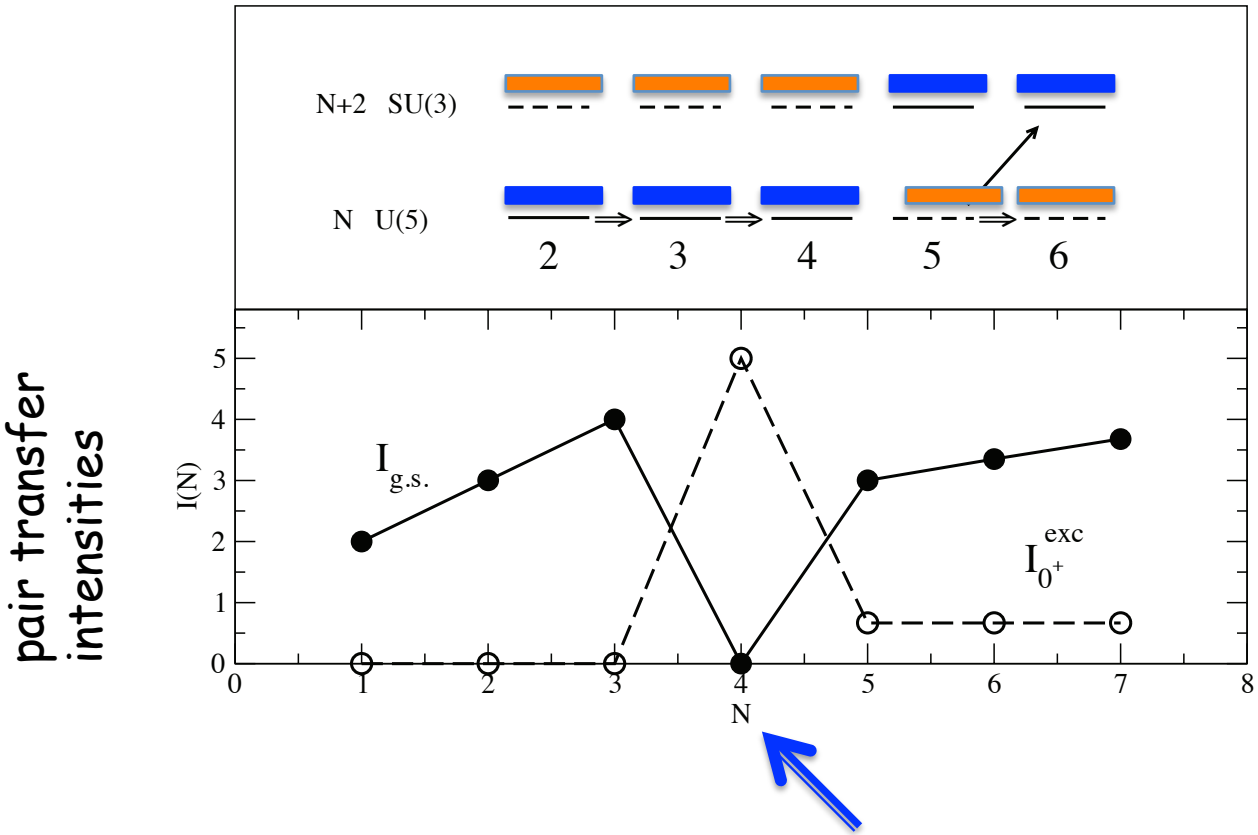
Pair creation probability N -> N+2



There is a clear signal at the phase transition

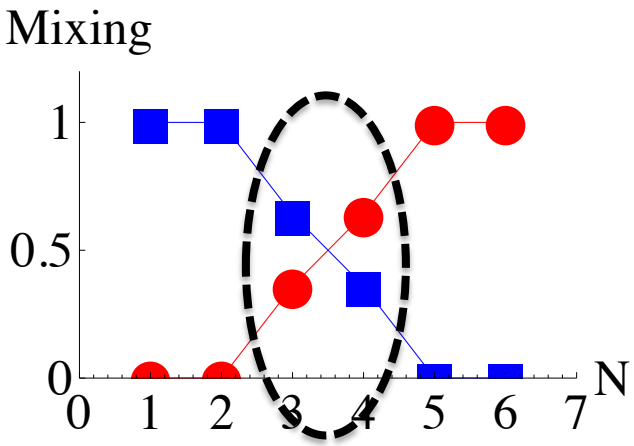
deformed

spherical



As in the previous situation a clear discontinuity appears at the critical point. However, at variance with the previous case, the pair strength is always practically concentrated in a single state, without the fragmentation illustrated in the previous case

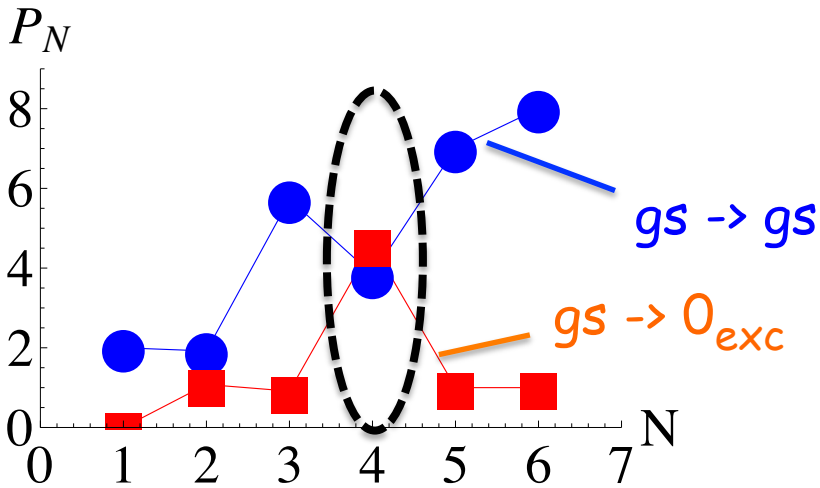
Another case: shape-coexistence with a smoother transition



OBS: Cf. E0 transitions between the two 0+ states

Transfer operator for pair removal : $s + s^+$ (one can destroy a particle pair or create a hole pair)

Pair removal probability N <- N+2



So far we have considered matrix elements of the pair operator: but what about pair transfer cross sections?

Unfortunately, at variance, for example, from low-energy one-step Coulomb excitation, where the excitation probability is directly proportional to the B(Eλ) values, the reaction mechanism associated with pair transfer is rather complicated and the possibility of extracting spectroscopic information on the pairing field is not obvious. The situation is actually more complicated even with respect to other processes (as inelastic nuclear excitation) that may need to be treated microscopically, but where the reaction mechanism is somehow well established.

We expect an correlation between cross sections and square of the pair operator. But if the qualitative behavior may be clear, the quantitative aspects require a proper treatment of the reaction mechanism. All approaches, ranging from macroscopic to semi-microscopic and to fully microscopic, try to reduce the actual complexity of the problem, which is a four-body scattering (the two cores plus the two transferred particles), to more tractable frameworks.

Two models are most popular:
A, Successive single-particle transfer
B. Cluster transfer

A

Sequential two-step process: each step transfers one particle

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

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

Expansion to second-order in the transfer potential

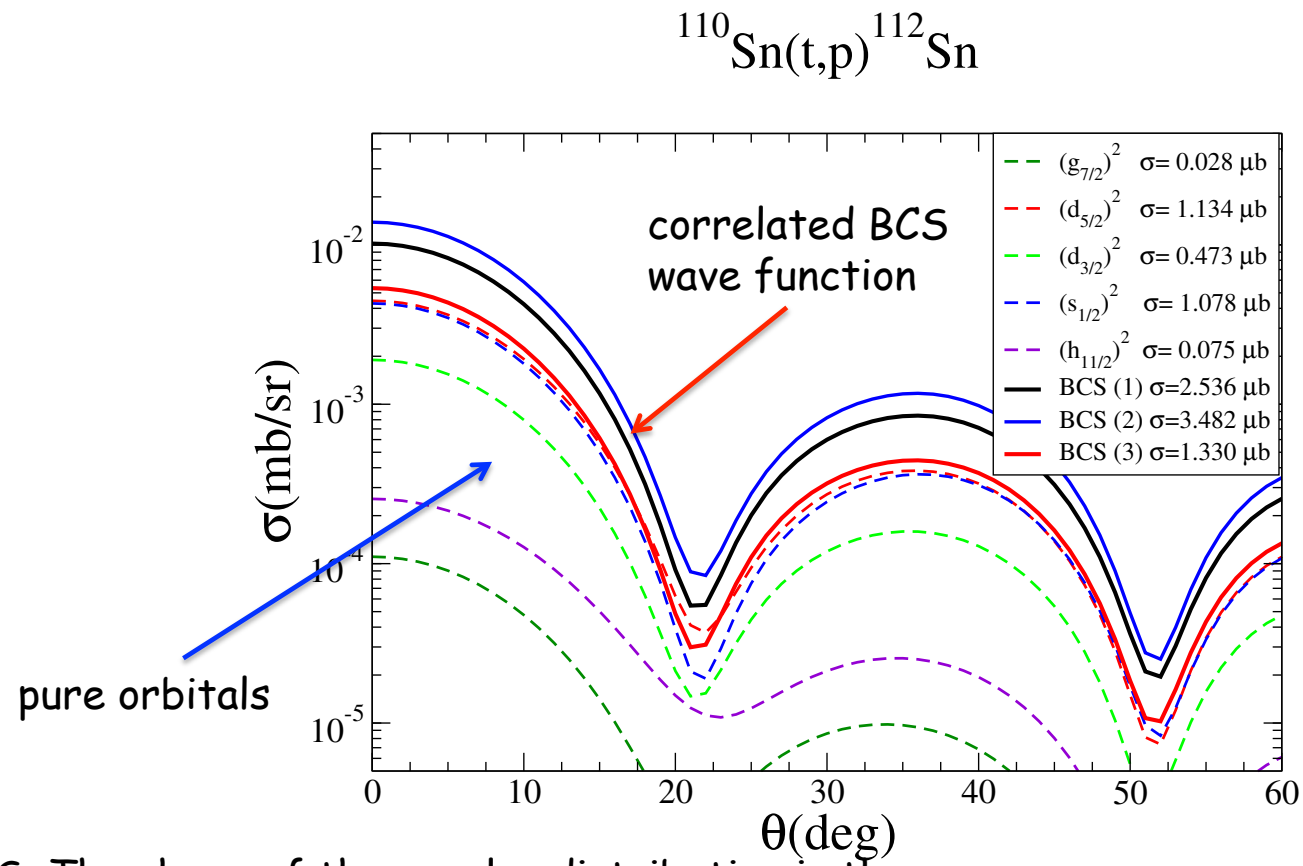
Simultaneus	+	Sequential	+	not-orthogonality
(first-order)		(second-order)		(second-order)

this is <u>not</u> the cluster contribution

these two terms may approximately cancel each other

Effect of kinematical conditions

The transfer probabilities **vary strongly** with the involved orbital. In addition whether the final wave function only involves a "pure" orbital, or whether it is correlated



OBS: The shape of the angular distribution is the same, being associated with the L=0 transfer

But the dependence on the microscopy also arises from the reaction mechanism associated with each specific two-particle transfer process

If we consider the same case as before, i.e. the transfer of two neutrons from ^{110}Sn to ^{112}Sn (0+; gs), but using different reactions, e.g. (14C,12C) or (18O,16O) the ranking of the cross sections associated to the different orbitals changes.

projectile

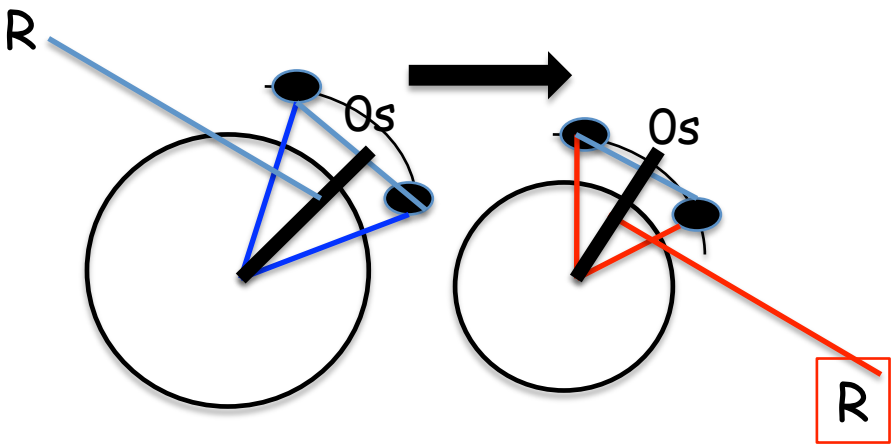
single-particle orbital in the target

^{112}Sn	(t,p)	(^{14}C,^{12}C)				(^{18}O,^{16}O)		
		$(0p_{1/2})^2$	$(1s_{1/2})^2$	$(0p_{3/2})^2$	$(0d_{5/2})^2$	$(0d_{5/2})^2$	$(1s_{1/2})^2$	Conf_1
$(0g_{7/2})^2$	2.80E-5	1.73E-5	1.19E-4	7.09E-4	9.00E-4	1.19E-3	2.01E-4	1.24E-3
$(1d_{5/2})^2$	1.13E-3	3.00E-2	4.71E-3	5.54E-3	1.18E-3	3.55E-3	1.13E-2	1.19E-2
$(2s_{1/2})^2$	1.08E-3	1.53E-2	5.38E-3	7.05E-3	1.16E-3	5.02E-3	1.42E-2	1.59E-2
$(1d_{3/2})^2$	4.73E-4	1.34E-3	2.79E-3	9.87E-3	4.14E-3	1.26E-2	6.62E-3	1.83E-2
$(0h_{11/2})^2$	7.50E-5	7.77E-4	5.29E-5	1.05E-4	7.65E-5	1.10E-4	9.06E-5	1.88E-4
Conf_A	2.54E-3	8.77E-2	2.26E-2	3.77E-2	1.21E-2	8.60E-3	1.95E-2	7.53E-2

BCS

B

Cluster-transfer model (suggested by the close radial correlation of the pairs)



Initial and final cluster wave functions are obtained by taking the overlap between the two-particle wave functions and a 0s wave function for the relative motion

Also in this case the resulting cross section depends on the specific single-particle orbitals (via the Talmi-Moshinsky brackets), but the dependence is different from the one associated with the sequential transfer (!!!)

The preference to either model may depend on the colliding systems and on kinematical conditions.

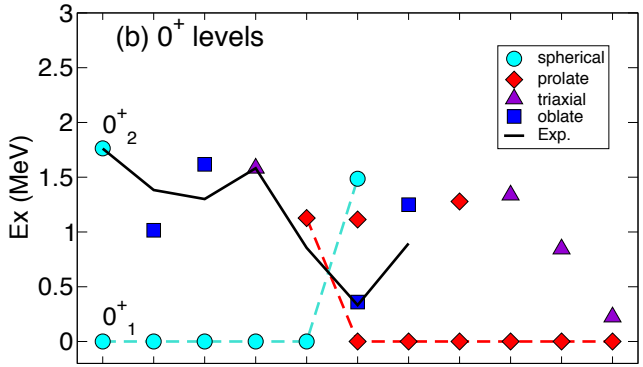
The proper approach will depend on the competition between the two colliding single-particle mean fields and the residual two-body interaction (for relatively weak interaction the mean fields will prevail, while in the other extreme of infinite pairing correlation the cluster structure will take over).

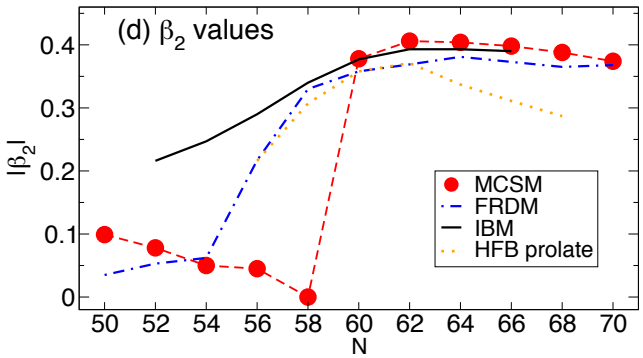
One case in more details (with full microscopic wave functions):

Shape phase transition in Zr isotopes

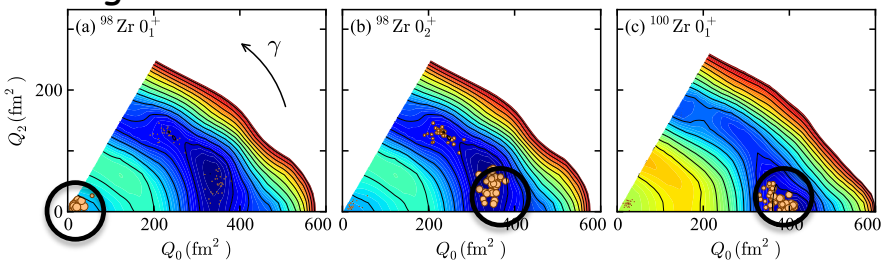
Other examples: Mg isotopes, Ni isotopes

First example: Shape phase transition in Zr isotopes between N=58 and 60





0_{gs} in ^{98}Zr 0_2 in ^{98}Zr 0_{gs} in ^{100}Zr



similar shape

Quantum Phase Transition in the Shape of Zr isotopes

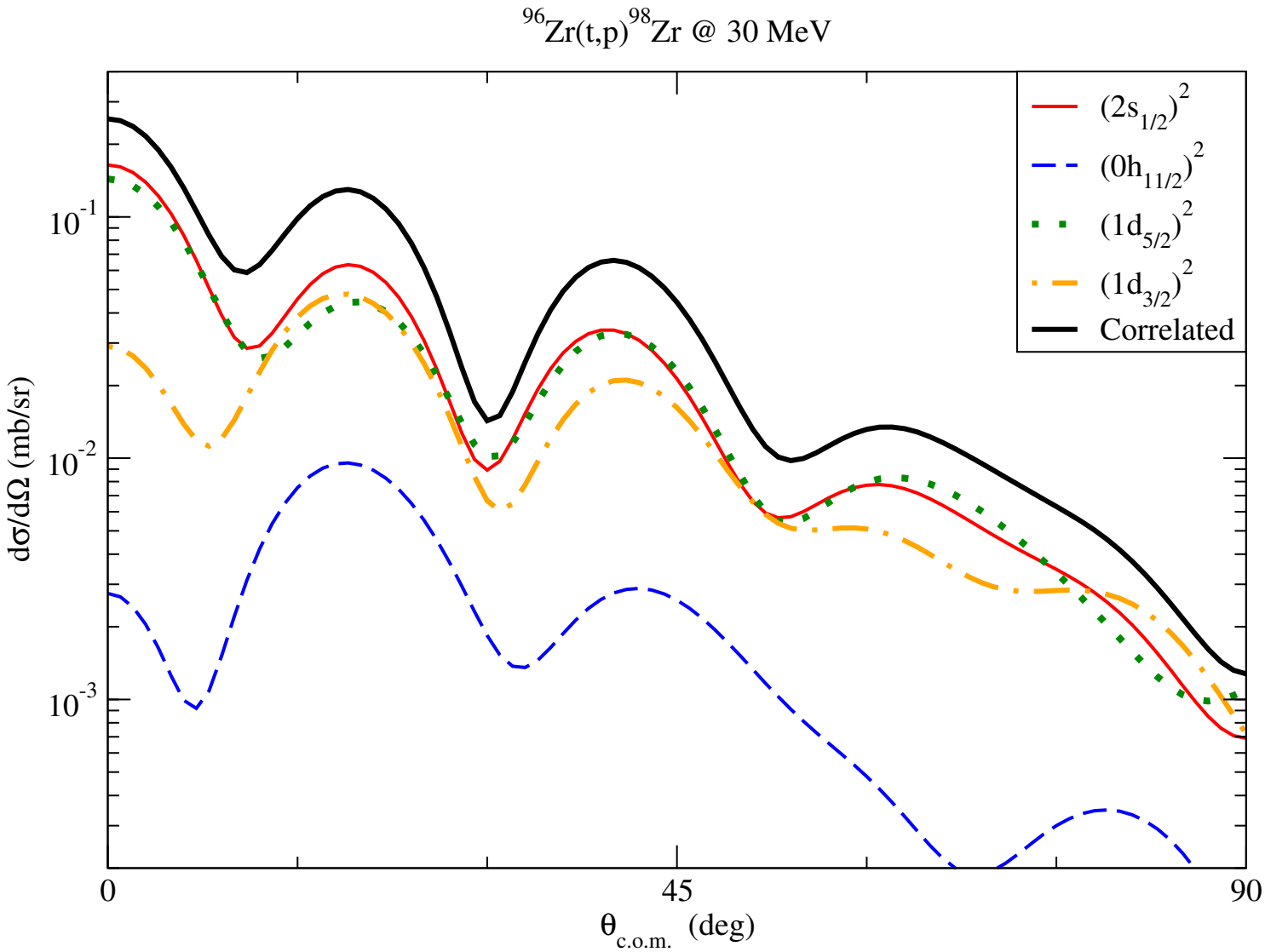
Tomoaki Togashi[1], Yusuke Tsunoda[1], Takaharu Otsuka[1,2,3,4] and Noritaka Shimizu[1]

relevant 2-particle spectroscopic amplitudes

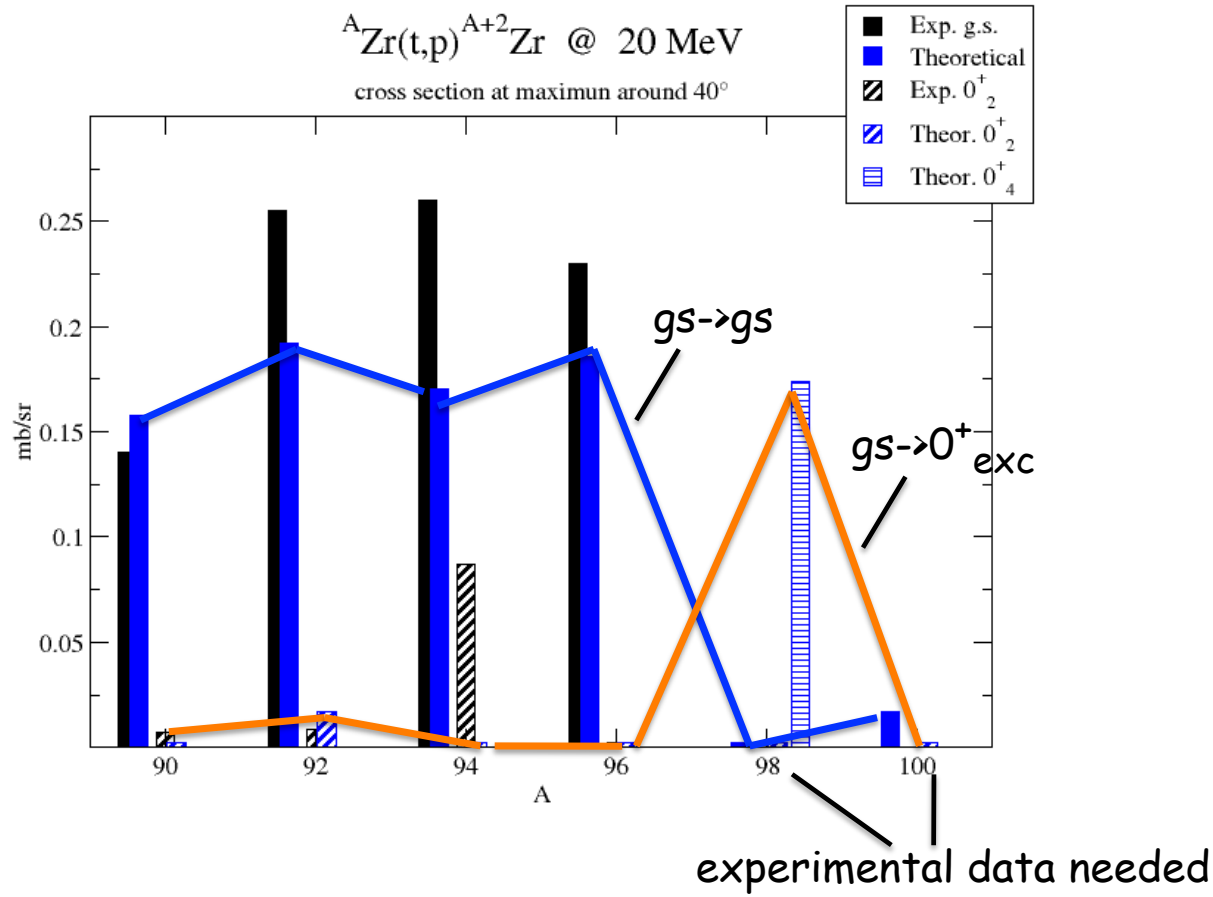
	90>92gs	92>94gs	94>96gs	96>98gs	98>100gs	98>100 ($0+_4$)	100>102gs
d5/2	**0.74**	**0.86**	**0.86**	0.13	0.0	0.16	0.08
S1/2	0.10	0.08	0.10	**0.90**	0.0	0.16	0.05
d3/2	0.13	0.18	0.16	0.07	0.0	**0.90**	0.04
h11/2	0.22	0.20	0.19	0.08	0.0	0.14	**0.55**

$d_{5/2}$ $s_{1/2}$ $d_{3/2}$ $h_{11/2}$

Cross sections for pure configurations



Calculation of two-particle transfer reactions using:
sequential model for the reaction mechanism
one- and two-particle spectroscopic amplitudes from the Tokyo group



Conclusions:

Pairing response (tested in two-particle transfer reactions but also in other dynamical processes involving pairs of particles) gives strong constrains on nuclear wave functions. The effect is amplified in correspondence of critical situations associated with shape phase transitions, with "abnormal" population of excited 0+ states and weakening of the ground state transition.

Further data on two-particle transfer reactions are definitely needed

Thanks to
Jose' Antonio Lay and Lorenzo Fortunato (Padova/Sevilla)
Tomoaki Togashi, Yusuke Tsunoda and Takaharu Otsuka (Tokyo)