

# Measurement of the damping mechanisms of the GDR at high excitation energy using the GARFIELD and HECTOR setup

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## I. INTRODUCTION

The damping mechanisms of the Giant Dipole Resonance (GDR) in the high temperature region (high excitation energy) are still not well understood [1].

The two main relaxation mechanisms giving rise to the measured damping width which have been identified are the collisional damping and the thermal large-amplitude fluctuations of the nuclear shape. The phenomenon of collisional damping is essentially connected with the coupling of the giant modes with the quantal fluctuations of the nuclear surface. Various models of the collisional damping width exist some predicting a constant behaviour with temperature (refs [2-4]), and one predicting an increase with temperature [5]. The second important mechanism at work in the breaking of the strength of the GDR is the coupling of the vibration to the large amplitude fluctuations (shape fluctuations) of the nuclear surface which are induced by temperature [6,7].

In fusion-evaporation reactions at low energies corresponding to  $T \leq 2$  MeV, the width of the GDR built on excited states is observed to increase rapidly with bombarding energy (by approximately a factor of 2 by going from 0 to 2 MeV) due to increasing spin-induced deformation, and increasing thermal shape fluctuations [8]. In this regime the global systematics of the GDR width are reasonably well described by calculations made within the thermal shape fluctuation model and using the  $T=0$  value of the collisional damping width (namely the intrinsic width of the photo-absorption line shape) [6,7]. Beyond the bombarding energy at which the maximum angular momentum that the nucleus can sustain without fissioning saturates, one expects that the GDR width would increase more slowly.

At excitation energy larger than  $E^* \geq 100$  MeV there is the open question whether the width of the GDR saturates or increases. This has been newly addressed particularly in relation to the problem of how well one can determine the excitation energy of the nucleus on which the GDR is built. Up to now a large number of the performed measurements are essentially inclusive and therefore at the highest

excitation energies more exclusive measurement are necessary for the understanding of this problem.

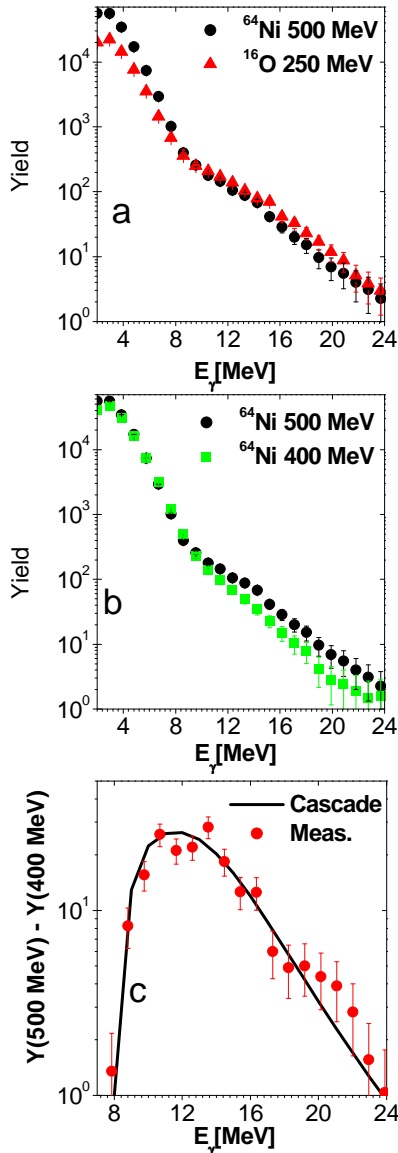
Recently, measurements were made using the HECTOR [9] and GARFIELD [10] setup, in which both high energy gamma-rays and charged particle emission in coincidence with evaporation residues were recorded. In order to evaluate the role of the preequilibrium emission we have chosen two different reactions leading to the same excitation energies, but at different value of the bombarding energies. The measured reactions were the most symmetric and asymmetric reactions in the entrance channel, namely  $^{64}\text{Ni} + ^{68}\text{Zn}$  at  $E_{\text{beam}} = 500$  and  $400$  MeV (corresponding to 200 and 150 MeV of nominal excitation energy) and  $^{16}\text{O} + ^{116}\text{Sn}$  at  $E_{\text{beam}} = 250$  MeV (with  $E^* = 200$  MeV, as deduced from kinematics for complete fusion).

## II. THE EXPERIMENT

The experiment was made using the GARFIELD set up combined with the  $\text{BaF}_2$  detectors of the HECTOR array. The GARFIELD vacuum scattering chamber was equipped with one of the two drift chambers of the GARFIELD apparatus (gaseous microstrips coupled with  $\text{CsI(Tl)}$  crystals) from  $\theta = 30^\circ$  to  $90^\circ$ , while the  $\text{BaF}_2$  were positioned at backward angles. In the forward direction, between  $4^\circ$  and  $12^\circ$ , two couples of PSPACs were positioned symmetrically with respect to the beam. Some  $\text{SiLi}$  detectors were positioned between each PSPAC couple at the larger angles. So far we have concentrated in the measurements of  $^{132}\text{Ce}$  at two energies (corresponding to  $E^* = 150$  and  $200$  MeV), the higher measured using the two different reactions  $^{64}\text{Ni} + ^{68}\text{Zn}$  and  $^{16}\text{O} + ^{116}\text{Sn}$ . The pulsed beam was provided by the TANDEM+ALPI accelerator system. With the system GARFIELD + HECTOR we could measure both  $\gamma$ -rays and charged particles in coincidence with residual nuclei detected in the PSPAC detectors.

### III. PRELIMINARY RESULTS OF THE MEASUREMENTS

Some preliminary results of the measurements are presented in figure 1. In that figure the high-energy gamma-ray spectra measured using the reaction  $^{64}\text{Ni} + ^{68}\text{Zn}$  ( $E_{\text{beam}} = 400, 500 \text{ MeV}$ ) and  $^{16}\text{O} + ^{116}\text{Sn}$  ( $E_{\text{beam}} = 250 \text{ MeV}$ ) are shown.



**Fig.1:** Panel a: gamma-ray spectra measured for the reactions  $^{64}\text{Ni} + ^{68}\text{Zn}$  at  $E_{\text{beam}} = 500 \text{ MeV}$  and  $^{16}\text{O} + ^{116}\text{Sn}$  at  $E_{\text{beam}} = 250 \text{ MeV}$ .

Panel b : gamma-ray spectra measured for the reaction  $^{64}\text{Ni} + ^{68}\text{Zn}$  at  $E_{\text{beam}} = 400$  and  $500 \text{ MeV}$ .

Panel c: Spectrum obtained as a difference of spectra in panel b in comparison with a statistical model calculation.

The spectra are in coincidence with the heavy recoiling nuclei identified with the PSPAC detectors at forward angles. There are two important features that can

be observed in these spectra. The first is that in case of the  $^{64}\text{Ni} + ^{68}\text{Zn}$  reaction one sees an increase with excitation energy of the production yield of high energy gamma-rays which is consistent with that predicted by the statistical model and assuming 100 % of the Energy Weighted Sum Rule of the giant dipole mode.

This is shown in panel b and c of figure 1, and in particular panel c shows the difference spectrum (obtained by subtracting the data corresponding to the reactions  $^{64}\text{Ni} + ^{68}\text{Zn}$  at  $E_{\text{beam}} = 400, 500 \text{ MeV}$ ) in comparison with the statistical model prediction made with the code CASCADE [11].

The second feature observed in the data concerns the reaction dependence of the gamma-ray yield. In fact, the data at the highest excitation energy for  $^{64}\text{Ni} + ^{68}\text{Zn}$  are found to be different than those obtained with the reaction  $^{16}\text{O} + ^{116}\text{Sn}$  in spite of the fact that kinematically these two reactions should lead to the same compound nucleus and excitation energy. In particular one can note in panel a of figure 1 that the spectral shapes are different mainly in the region of the rapidly exponential decrease. Concerning the particle emissions we have observed sizable differences for the two reactions at forward angles but we need to complete the calibration and analysis work, presently in progress, on the particle detectors before we can infer something on the possible pre-equilibrium contributions. We expect that a relevant step forward in the understanding of nuclear structure of finite temperature should be made by completing these analyses.

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