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Determination of Direct Reaction Contribution by Reducing the Cross Section Fluctuation Data to the Polarization-like Ones

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The normalized variance (NV) of the differential cross section (CS) equals 1)

 $(\langle 6^2 \rangle - \langle 6 \rangle^2)/\langle 6 \rangle^2 = (1 - y_d^2)/N$, (1) where $y_d = 6_d/\langle 6 \rangle$ is the contribution of direct reaction CS 6_d into summarized energy-averaged CS $\langle 6 \rangle = 6_d + \langle 6_{fl} \rangle$ with $\langle 6_{fl} \rangle$ being the average fluctuating CS and N being the number of independent channels. The formula (1) reflects the well-known problem: The NV of the CS gives only the relation between y_d and N but does not allow to determine each quantity separately. Probability distribution (PD) of the CS also is practically not sensible to y_d and N values for the NV of CS (1) being fixed. In this contribution a method is proposed for the determination of both y_d and N values.

Suppose we have the excitation function $\mathcal{G}(E)$ for nuclear reaction of the arbitrary spin structure $S_1+S_2\rightarrow S_3+S_4$ and let $\mathcal{G}(E)$ be measured with the high energy resolution $\Delta E \ll \gamma$, where γ is the energy coherence length ¹. Let us take for each value of $\mathcal{G}(E)$ the energy-shifted one $\mathcal{G}(E+e)$ with $e \gg \gamma$. Then $\mathcal{G}(E)$ and $\mathcal{G}(E+e)$ are uncorrelated random functions. Now turn from $\mathcal{G}(E)$ and $\mathcal{G}(E+e)$ to the new variables

s(E) = f(E) + f(E+e), r(E) = (f(E) - f(E+e))/s(E). (2) Taking into account the stationarity of the random process f(E) and using the PD of the CS¹ it is easy to obtain the simultaneous PD of $y(E) = s(E)/\langle s \rangle$ and r(E)

$$F_{2N}(\mathbf{r},\mathbf{y}) = 2N^{2}(1-y_{d})^{-2}y_{d}^{1-N}y^{N}\exp(-2N(y_{d}+\mathbf{y})/(1-y_{d}))(1-\mathbf{r}^{2})^{(N-1)/2}$$

$$I_{N-1}(2Ny_{d}^{1/2}y^{1/2}(1+\mathbf{r})^{1/2}/(1-y_{d}))I_{N-1}(2Ny_{d}^{1/2}y^{1/2}(1-\mathbf{r})^{1/2}/(1-y_{d})), \qquad (3)$$

where $I_{N-1}(x)$ stands for the Bessel function of an imaginary argument. Using (3) we can investigate the statistical correlation (SC) between y and r. The simple analysis shows that for the NV of the CS being fixed the greater y_d the stronger this correlation (Fig.1). Hence the study of SC between y and r do makes possible the unique determination of both y_d and N.

To test the SC method we use ${}^{28}\text{Si}(\vec{p},p)$ data at $\text{E}_{p}=12-18$ MeV, $\vartheta=160^{\circ}$ with $\vartriangle \text{E}_{p}=50$ keV energy steps 2 . Using of the polarized beam in this case results in the one-channel situation, N=1 for the partial CS

 $6 \pm 6A$ (A is the analyzing power (AP)). Hence, from (1) it is easy to determine the y_d value for both partial CS. The results of SC analysis after trend-reducing by averaging over 650 keV interval ²) are shown in Fig.2. The ability of method proposed to increase the initial statistics was used in the histograms derivation. Namely, the sets of s and r were taken which differed by energy shifts $e \leq 1$ MeV. The values of e have been chosen from the condition of uncorrelation between $5(E) \pm 5(E)A(E)$ and $5(E+e) \pm 6(E+e)A(E+e)$ on the 10% level of statistical accuracy.

The possibility to determine y_d and N values arises due to the reduction of CS fluctuation data to polarization-like ones. Indeed, the analysis proposed is analogous to the investigation of SC between AP and CS for the reaction of $1/2+J_1+J_2+J_3$ type ³) with direct reaction AP being zero. Here s(E) takes part of the CS and r(E) corresponds to the AP. At the same time the analysis ³) is based principally on the uncorrelation condition of compound-nucleus phases, which does not hold in the presence of direct reactions ⁴). The method proposed above is more general. Indeed, the compound-nucleus phase correlations naturally effect(δ_{fl}) and hence y_d and N values, but it is expected that their influence on the PD of the CS and simultaneous PD (3) is negligible. Note as a conclusion that investigation of the stability of the PD and correlations between s and r under the variation of an interval e can be used for the identification of the intermediate structure in nuclear reactions.

Fig.1. $\langle r^2(y) \rangle$ Correlation depen- 0.2 dences $\langle r^2(y) \rangle$ for different y_d and N values with NV 0.1 of the CS being fixed and equal to 0.1.



Fig.2. Experimental (histograms) and theoretical correlation dependences <r²(y)> for ²⁸Si(p,p) scattering.

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