Gold fragmentation induced by stopped antiprotons

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A natural gold target was irradiated with the antiproton beam from the low-energy antiproton ring at CERN. Antiprotons of 200 MeV/*c* momentum were stopped in a thick target, products of their annihilations on Au nuclei were detected using the off-line γ -ray spectroscopy method. In total, yields for 114 residual nuclei were determined, providing a dataset to deduce the complete mass and charge distribution of all products with *A* \approx 20 from a fitting procedure. The contribution of evaporation and fission decay modes to the total reaction cross section as well as the mean mass loss were estimated. The fission probability for Au absorbing antiprotons at rest was determined to be equal to $(3.8\pm0.5)\%$, in good agreement with an estimation derived using other techniques. The mass-charge yield distribution was compared with the results obtained for proton and pion induced gold fragmentation. On average, the energy released in \bar{p} annihilation is similar to that introduced by \approx 1 GeV protons. However, compared to proton bombardment products, the yield distribution of antiproton absorption residues in the *N*-*Z* plane is clearly distinct. The data for antiprotons exhibit also a substantial influence of odd-even and shell effects.

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I. INTRODUCTION

The large energy of almost 2 GeV released in nucleonantinucleon annihilation has brightened hopes to observe some unique nuclear reactions induced in this way. Especially energetic antiprotons were presumed to reach the deep interior of the nucleus. Exotic processes, such as phase transitions to quark-gluon plasma and explosive decay of that hot system had been expected to occur $[1,2]$, but were not verified in experiments performed shortly after the commissioning of the low-energy antiproton ring (LEAR) at CERN [3,4]. Nevertheless, the character of the reactions starting with antiproton absorption in nuclei is quite unique in comparison with reactions induced by protons or heavy ions. Whereas the excitation energy carried in by postannihilation mesons is quite large, the linear and angular momentum transfer as well as the matter compression are reduced, particularly in the stopped antiproton case that is preceded by the exotic atom phase. Hence, one may investigate a clear thermal reaction aspect with suppressed collective motion complications.

Such phenomena were intensively studied during the LEAR era for stopped and energetic antiprotons. The spectra for neutrons and light charged particles $[5-10]$, mass yield distributions $[11–16]$, and characteristics of the fission fragments $[17-19]$ were measured for a wide range of targets. The mean excitation energy derived from these studies, \approx 150 MeV and \approx 300 MeV for heavy nuclei absorbing stopped and 1.2 GeV antiprotons, respectively, compares well with the average values obtained for protons that have about 1 GeV larger energies, i.e. approximately the nucleon rest mass.

One of the most distinct features observed in reactions with stopped antiprotons is the large probability (≈ 0.1) of very small energy transfer, when the target nucleus loses only one nucleon in annihilation and is left excited below the nucleon separation energy. Nuclear spectroscopy studies of the relative yield of both types of these residual $(A_T - 1)$ nuclei (a neutron or a proton lost in such soft antiproton absorptions) were used to establish a new and powerful method of probing the nuclear periphery composition $[23 26$.

The irradiation of the heavy, gold target gave us a chance to study the competition between evaporation and fission induced by antiproton absorption. The yield distribution of heavy residual nuclei complements the results obtained from on-line measurements of neutron and charged particle spec- *Electronic address: piotr@camk.edu.pl tra [8]. Since gold is a commonly used target, there was also

Yield distributions of residual nuclei were studied as soon as more intense \bar{p} beams were provided by LEAR. Many targets were irradiated with antiprotons, mainly at rest energy, but only few of them were examined in detail: natCu [14], $92,95$ Mo [11], 98 Mo [12], $natAg$ [15], $natBa$ [13], 165 Ho [12], and 181 Ta [16]. Average quantities such as the mean mass removed from the target as well as individual yield features, e.g., isomeric ratios, were investigated. Clear oddeven effects in the mass and charge yield distribution were observed. Theoretical calculations, based on intranuclear cas- $\text{cade} + \text{evaporation models}, \text{were able to reproduce only the}$ gross features and failed to predict yield dependence on the detailed N and Z [13]. The odd-even phenomenon, although postulated to be present and used to model the yield distribution for years $[20,21]$ and sometimes reaching very large values $[22]$, still seems to be almost unexplored theoretically in a more quantitative way.

the opportunity to compare these data with a rich set of information gathered from irradiations with energetic protons or pions. In particular, the yield distribution after the reaction of 1 GeV protons with Au has been extensively studied in older and recent γ -ray spectroscopy measurements [27,28] and also with a new method using the mass-charge spectrometry for inverse kinematic reactions [29,30]. Our preliminary data of the reaction \overline{p} + Au at rest have already been published $[31]$; this work presents the complete results obtained after fitting the *Y*(*A*,*Z*) yield distribution.

The paper is organized as follows. In Sec. II we briefly present some details of the experiment; in Sec. III data analysis is described together with the yield fitting method. Experimental results are presented and initially discussed in Sec. IV. In Sec. V we compare the results of this experiment with those obtained with other projectiles impinging on Au and with data coming from studies of antiproton absorption in various targets. Finally, our main conclusions are presented in Sec. VI.

II. EXPERIMENT

A thick target of natural gold was irradiated with the antiproton beam of 200 MeV/*c* momentum from LEAR facility. The target of the total thickness of 549 mg/cm² was composed of ten foils of 80, 30, 30, 30, 2, 30, 30, 37, 80, and 200 mg/cm^2 , starting from the beam side. The initial energy of the antiprotons, equal to about 21 MeV, was reduced in the scintillation counter (from pilot B) and in some additional moderators (mylar, silicon) to about 6.5 MeV at the target surface. Such an arrangement assured that the majority of antiprotons were stopped in few central Au foils. The very central and extremely thin foil of 2 mg/cm² was applied to monitor the x-ray activity, while the last and thickest one (200 mg/cm^2) was used to check the secondary reactions level.

Two scintillation counters, *S*1 and *S*2, were used to control beam intensity and transverse dispersion. The first anticounter *S*1 had a hole of 10 mm diameter and the active area of counter *S*2 was a disc of the same diameter. Consequently the signal *S*1*S*2 indicated particles going towards the target. The irradiation lasted 15 min with the total number of antiprotons equal to $(9.25 \pm 0.35) \times 10^8$ ($\overline{S1}S2$ number).

Monitoring of the target activity started 13 min after irradiation and continued at CERN for one week; afterwards the spectra were collected in Warsaw, the last one was taken more than a year after target activation. Two HPGe detectors were used at CERN, a γ -ray counter of 15% relative efficiency for all foils and an x-ray counter for the thinest one. In Warsaw two more efficient γ -ray detectors were applied, of 20% and of 60% relative efficiency, and a third x-ray detector for the thin foil.

All collected spectra were analyzed with the program $ACTIV$ [32,33]. Gamma-ray lines were identified by their energies, half-lives, and intensity ratios. The decay data were taken from the eighth edition of the Table of Isotopes $[34]$.

Experimental yields for all detected residual nuclei, normalized to 1000 $\frac{3}{p}$ stopped in the target, are listed in Table I. The independent yields represent the total number of nuclei, summed over all isomers. Cumulative yields include also the yields of all β -decay precursors of a given isotope. Besides that, we present partial yields for some isomers not representing the whole production for a given (*A*,*Z*) pair as well as some production limits for Hg nuclei. Mercury may be produced from gold after \overline{p} absorption in a charge exchange reaction, when one of the annihilation π^+ , π^0 pions exchanges charge with a target neutron. Such a phenomenon was observed for some targets irradiated with stopped antiprotons $[12,13,15,16]$, where nuclei of target charge plus one were produced at a level ranging from 0.5 to 5 per $1000\overline{p}$. On the other hand, for some other targets, studied in the neutron halo project, rather low upper limits $(0.5–2%)$ were given [35]. Our data obtained for Au, except for the 195 Hg isomers with low γ intensity, indicate that such an effect should occur very rarely.

Initially, the distribution of the activity induced in individual target foils was estimated several hours after the irradiation with the use of the measurement of the 186 Ir(*g*) yield in each foil. Five inner foils $(30, 30, 2, 30, 30 \text{ mg/cm}^2)$ gathered about 90% of the total activity and only these foils were monitored later. This reduced the γ -ray self-absorption effect. The distribution of the target activity was determined more precisely afterwards on the basis of yields obtained for six evaporation residues: 186 Ir, 184 Ir, 183 Os, 181 Re, 157 Dy, and 152Dy. On average, five inner foils stopped (89.7 \pm 2.1)% of the whole number of antiprotons. These data, together with the results for 196 Au and 192 Au, were used also to estimate the yield introduced by secondary reactions with particles (mainly pions and neutrons) produced after antiproton annihilation on target nuclei. It was done by comparing the number of given nuclei produced per foil thickness unit, averaged for three inner foils, with a similar result obtained for the last, thickest foil. The upper limit for the secondary reactions leading to 196 Au is equal to 3%, the limit for 192 Au is about 2.6% and for the rest of quoted isotopes it does not exceed 2%, i.e., in all cases it is below the yield uncertainties. The negligible influence of the secondary reactions on our results is additionally confirmed by very small upper limit given in Table I for 198 Au, a (n, γ) reaction product.

III. DATA ANALYSIS

The method based on off-line γ -ray spectroscopy has some limitations. The most important is the necessity to use some phenomenological model to reconstruct the yields of unobservable products $[12,20,21,36,37]$. In similar experiments with other projectiles the data are sparsely spread over the *N*-*Z* plane. In this case less detailed models may be applied and the results obtained are only a first-order approximation of the true yield distribution $Y(A,Z)$, even when cumulative cross sections are involved in the fitting procedure. Moreover, the precision of tabulated absolute γ -ray intensities sometimes leaves much to be desired and, finally, the decomposition of spectra with a few hundreds of lines, as they are measured for heavier targets, becomes a challenge for the persistence of the evaluator.

TABLE I. Experimental and fitted yields of residual nuclei from gold fragmentation induced by stopped antiprotons. Yield type: *I*: independent, *C*: cumulative, in brackets: total yield fitted for isotopes for which only one isomeric state was observed.

Nuclide	$T_{1/2}$	Experiment $\lceil N/1000p \rceil$	Fit	Type	Nuclide	Half-life	Experiment $\lceil N/1000p \rceil$	Fit	Type
$^{198}\mathrm{Au}^\mathrm{g}$	2.7d	< 0.6		\boldsymbol{I}	$^{165}\mathrm{Tm}$	$30.1\ \mathrm{h}$	14.4 ± 0.8	14.2	$\cal C$
$^{196}\mathrm{Au}^{\mathrm{m2}}$	$9.7\ \mathrm{h}$	1.52 ± 0.18		I	$^{163}\mathrm{Tm}$	$1.81\ \mathrm{h}$	10.1 ± 0.6	10.1	$\cal C$
$^{196}\mathrm{Au}$	$6.2\ \mathrm{d}$	75.0 ± 3.4		I	$^{161}\mathrm{Tm}$	$33.0\ \mathrm{min}$	8.6 ± 1.1	8.6	$\cal C$
$^{195}\mathrm{Hg^m}$	$41.6\ \mathrm{h}$	< 3.1		Ι	^{161}Er	3.2h	2.69 ± 0.67	2.73	Ι
$^{195}\mathrm{Hg^{8}}$	$9.9\ \mathrm{h}$	< 11.7		Ι	^{160}Er	$26.6\ \mathrm{h}$	9.8 ± 0.8	9.9	\boldsymbol{C}
$^{195}\mathrm{Au}$	$186\ \mathrm{d}$	38.7 ± 3.1		Ι	$^{159}\mathrm{Er}$	$36.0\ {\rm min}$	8.2 ± 0.8	8.2	\mathcal{C}_{0}^{0}
$^{194}\mathrm{Au}$	38.0 _h	11.8 ± 0.6		Ι	$\mathrm{^{157}Dy}$	8.2h	8.0 ± 0.4	7.9	\boldsymbol{C}
193 Hg ^m	$11.8\ \mathrm{h}$	< 0.5		Ι	$^{155}\mathrm{Dy}$	9.9h	6.2 ± 0.3	6.2	$\cal C$
$^{193}Hg^8$	$3.8~\mathrm{h}$	< 2.8		Ι	$\mathrm{^{155}Tb}$	$5.3\,\mathrm{d}$	0.27 ± 0.09	0.30	\boldsymbol{I}
$^{193}\mathrm{Au}$	$17.7\ \mathrm{h}$	9.9 ± 2.2		Ι	154Ho8	$11.8~\mathrm{min}$	2.93 ± 0.20	(3.16)	$\cal C$
192 Hg	4.9 h	< 1.3		Ι	$^{153}\mathrm{Tb}$	2.3d	4.77 ± 0.24	4.78	$\cal C$
192Au	4.9 h	8.5 ± 0.6	8.37	Ι	$\rm ^{153}Gd$	242 d	0.13 ± 0.04	0.12	I
$^{192}\text{Ir}^8$	73.8 d	1.61 ± 0.07	2.6	Ι	$^{152}\mathrm{Dy}$	$2.4\ \mathrm{h}$	3.60 ± 0.11	3.60	$\cal C$
$^{191}Hg^m$	$51.0~\mathrm{min}$	< 1.7		Ι	$\rm ^{152}Tb$	$17.5\ \mathrm{h}$	0.67 ± 0.13	0.67	\boldsymbol{I}
$^{195}\mathrm{Hg^{8}}$	49.0 min	< 2.8		Ι	$\rm ^{151}Tb$	$17.6\ \mathrm{h}$	3.55 ± 0.20	3.55	$\cal C$
$^{191}\mathrm{Au}$	3.2h	6.7 ± 0.4	6.8	Ι	$\rm ^{151}Gd$	$124~\mathrm{d}$	0.27 ± 0.08	0.24	\boldsymbol{I}
^{191}Pt	2.9d	7.9 ± 0.8	8.6	Ι	150 Dy	$7.2\,$ min	1.95 ± 0.34	1.95	$\cal C$
$^{190}\mathrm{Hg}$	$20.0\ \mathrm{min}$	< 2.2		Ι	$^{150}\mathrm{Tb}^{\mathrm{g}}$	$3.5\ \mathrm{h}$	2.85 ± 0.27	(2.9)	$\cal C$
$^{190}\mathrm{Au}$	42.8 min	5.0 ± 0.4	5.0	Ι	$\rm ^{149}Gd$	9.3 d	3.12 ± 0.20	3.13	$\cal C$
190 Ir ^{m2}	$3.3\ \mathrm{h}$	2.04 ± 0.15		Ι	$\rm ^{148}Tb^g$	1.0 _h	1.13 ± 0.16	(1.73)	$\cal C$
190 Ir	11.8d	1.95 ± 0.10	4.0	Ι	$\rm ^{147}Gd$	$38.1~\mathrm{h}$	2.9 ± 0.2	2.9	$\cal C$
189Pt	$10.9\ \mathrm{h}$	12.5 ± 1.1	12.2	\boldsymbol{C}	$^{147}\mathrm{Eu}$	24.1 d	0.44 ± 0.20	0.51	\boldsymbol{I}
189 Ir	13.2 d	5.1 ± 0.8	4.8	\boldsymbol{I}	$\rm ^{146}Gd$	48.3 d	2.24 ± 0.13	2.23	$\cal C$
$188p_t$	10.2 d	13.2 ± 0.4	10.6	\boldsymbol{C}	$^{146}\mbox{Eu}$	4.6d	0.73 ± 0.08	0.71	$\cal C$
^{188}Ir	41.5 h	5.6 ± 0.4	5.8	\boldsymbol{I}	$\rm ^{145}Eu$	5.9 d	2.15 ± 0.34	2.19	$\cal C$
187Pt	2.4 h	9.8 ± 1.5	10.2	\boldsymbol{C}	$^{143}\mathrm{Pm}$	265d	2.0 ± 0.4	1.9	\boldsymbol{C}
^{187}Ir	10.5 _h	7.3 ± 1.0	6.8	Ι	$^{139}\mathrm{C}$	138 d	1.2 ± 0.1	1.2	\boldsymbol{C}
$186p_1$	$2.0\ \mathrm{h}$	10.4 ± 1.8	8.8	\boldsymbol{C}	135 Ce	$17.7~\mathrm{h}$	0.81 ± 0.21	0.78	$\cal C$
186 Ir ^m	$2.0\ \mathrm{h}$	$<$ 4.0		Ι	$^{132}\rm{La}$	$4.8~\mathrm{h}$	0.32 ± 0.08	0.34	Ι
$^{186}\mathrm{Ir^g}$	$16.6\ \mathrm{h}$	7.8 ± 0.8	$7.8\,$	Ι	$^{131}\mbox{Ba}$	$11.5~\mathrm{d}$	0.63 ± 0.24	0.62	$\cal C$
^{185}Ir	$14.4~\mathrm{h}$	16.3 ± 1.1	16.5	\boldsymbol{C}	$^{129}\mathrm{Cs}$	32.1 h	0.60 ± 0.20	0.56	I
185 Os	93.6 d	5.2 ± 0.8	4.1	Ι	$^{127}\mathrm{Xe}$	36.4 d	0.51 ± 0.05	0.52	$\cal C$
184Pt	17.3 min	8.2 ± 0.8	9.0	\boldsymbol{C}	124 I	$4.2\ \mathrm{d}$	0.16 ± 0.07	0.15	\boldsymbol{I}
$^{184}\mathrm{Ir}$	3.1 _h	8.3 ± 0.5	9.6	$\cal C$	$^{121}\mathrm{Te}$	$16.8\,\rm\,d$	0.39 ± 0.11	0.38	$\cal C$
183 Ir	58.0 min	14.2 ± 2.3	16.4	$\,$	103 Ru	39.3 d	0.48 ± 0.07	0.48	$\,c\,$
^{183}Osm	9.9 h	11.4 ± 0.8)		$\cal C$	$^{96}\mathrm{Tc}$	4.3 d	0.23 ± 0.05	0.23	\boldsymbol{I}
183Os ^g	$13.0\ \mathrm{h}$	9.7 ± 0.6	23.0	\boldsymbol{C}	95Nb	$35.0\ \mathrm{d}$	$0.65\!\pm\!0.07$	0.65	\boldsymbol{I}
$^{183}\mathrm{Re}$	70.0 d	2.22 ± 0.54	2.23	\boldsymbol{I}	^{95}Zr	64.0 d	0.16 ± 0.03	0.16	$\cal C$
^{182}Ir	15.0 min	11.8 ± 0.6	14.8	\boldsymbol{C}	$\rm{^{93}Mo^m}$	6.9h	0.23 ± 0.04	(0.23)	I
^{182}Os	$22.1\ \mathrm{h}$	9.8 ± 1.0	9.9	I	${}^{89}Zr$	3.3d	0.58 ± 0.05	0.59	$\cal C$
181 Re	19.9 h	24.9 ± 1.4	25.5	\boldsymbol{C}	${}^{88}\mathrm{Zr}$	83.4 d	0.30 ± 0.04	0.30	$\cal C$
180 _{Os}	21.7 min	19.9 ± 0.8	19.9	$\cal C$	88 Y	$107~\mathrm{d}$	0.81 ± 0.11	0.83	I
179 Re	19.5 min	20.8 ± 2.0	20.3	$\cal C$	^{87}Y	3.3d	0.75 ± 0.04	0.74	$\cal C$
$^{178}\mathrm{Re}$	13.2 min	15.5 ± 2.4	15.7	$\cal C$	86Y	14.7 h	0.38 ± 0.04	0.39	\boldsymbol{C}
177W	2.3h				${}^{85}Sr$	$64.8\,\mathrm{d}$			
$^{177}\mathrm{Ta}$	56.6h	20.8 ± 1.2 3.4 ± 1.1	21.2 $2.1\,$	$\cal C$	$\rm ^{84}Rb$	32.8 d	0.81 ± 0.11 0.78 ± 0.12	0.83	\boldsymbol{C}
$^{176}\mathrm{Ta}$			26.4	Ι	83 Rb			0.81 0.85	I
$^{175}\mathrm{Ta}$	8.1 h	26.5 ± 2.0		$\cal C$	${}^{82}Rb^m$	86.2 d	0.89 ± 0.16		\boldsymbol{C}
$^{175}\mathrm{Hf}$	$10.5\ \mathrm{h}$	21.2 ± 1.5	21.6	\boldsymbol{C}		$6.5~\mathrm{h}$	0.40 ± 0.11	(0.43)	\boldsymbol{I}
	$70.0\,\,{\rm d}$	2.3 ± 0.6	1.6	\boldsymbol{I}	${}^{82}\text{Br}$	$35.3~\mathrm{h}$	0.31 ± 0.08	0.30	\boldsymbol{I}

Nuclide	$T_{1/2}$	Experiment $\lceil N/1000\overline{p}\rceil$	Fit	Type	Nuclide	Half-life	Experiment $\lceil N/1000p \rceil$	Fit	Type
173 Ta	3.1 _h	16.1 ± 2.7	15.5	\mathcal{C}_{0}^{0}	75 Se	112d	0.36 ± 0.05	0.36	\mathcal{C}_{0}^{0}
173 Hf	23.6h	5.8 ± 0.7	5.7	I	^{74}As	17.8 _d	0.51 ± 0.08	0.51	I
173 Lu	1.37 y	0.54 ± 0.13	1.3	Ι	^{72}As	26.0 h	0.22 ± 0.07	0.22	I
172 Ta	36.8 min	8.1 ± 0.7	8.1	\mathcal{C}_{0}^{0}	^{72}Ga	14.1 _h	0.27 ± 0.07	0.27	I
172 Hf	1.9 _v	12.5 ± 1.4	12.3	\boldsymbol{I}	${}^{69}Zn^m$	13.8 _h	0.34 ± 0.04	(0.32)	I
172 Lu	6.7d	1.34 ± 0.20	1.36	I	59Fe	44.5 d	0.36 ± 0.07	0.38	$\mathcal{C}_{0}^{(n)}$
171 Lu	8.24d	20.8 ± 1.0	20.9	\mathcal{C}_{0}^{0}	48 _V	16.0 _d	0.19 ± 0.03	0.19	\mathcal{C}_{0}^{0}
170 Hf	16.0 _h	16.1 ± 1.4	16.1	\mathcal{C}_{0}^{0}	46 Sc	83.8 d	0.24 ± 0.07	0.24	I
170 Lu	2.0 d	3.8 ± 0.7	3.9	Ι	^{41}Ar	1.8 _h	0.22 ± 0.05	0.22	\mathcal{C}_{0}^{0}
169 Lu	34.1 h	16.0 ± 0.6	16.0	\mathcal{C}_{0}^{0}	^{24}Na	15.0 _h	0.27 ± 0.07	0.27	\mathcal{C}_{0}^{0}
169 Yb	32.0 _d	2.15 ± 0.67	2.18	I	^{22}Na	2.6 y	< 2.7	0.31	\mathcal{C}
166 Yb	56.7 h	15.5 ± 0.8	15.5	\mathcal{C}_{0}^{0}	7Be	53.3 d	< 3.5		\mathcal{C}_{0}^{0}

TABLE I. (Continued.)

Targets irradiated with antiproton beams, much less intense than the proton beams, have a rather low activity level. In particular, yields obtained for the fission fragments were close to our detection limit. Keeping in mind one of our goals, the estimation of the probability for antiproton induced fission, the data analysis needed special care and some feedback. A primary set of yield results obtained from the spectrum analysis served as an input for the model distribution fitting at its early stage, when the best approach was searched for. This relates to the choice of the final formula as well as to the division of the data to subsets assuring the lowest total χ^2 . Afterwards, a modeling procedure was applied to check, confirm or eliminate some doubtful experimental yields. For some mass regions it appeared necessary to apply an additional or separate evaluation, and we describe it at the end of this section.

A. Fitting procedure

The formula, used to describe the yield distribution was rather complex in order to be as universal as possible and to test various models. This complexity mainly arose from the aim of taking into account cumulative yields and from introducing the (*N*,*Z*) evenness corrections. The general formula was factorized into two components, mass and charge distributions, Y_A and Y_{ZP} , respectively,

$$
Y(A,Z) = Y_A Y_{ZP} . \tag{1}
$$

The distribution over the mass (the main distribution ridge) was modeled with the exponential of a fourth-order polynomial with parameters $a_1 - a_5$,

$$
Y_A = e^{(a_1 + a_2A + a_3A^2 + a_4A^3 + a_5A^4)}.
$$
 (2)

This was useful for testing the fit in broader mass regions, where the ridge shape may change more rapidly.

The form of the second factor in Eq. (1) , Y_{ZP} , was much more complex: the charge distribution was multiplied by the odd-even corrections F_p . When needed, an additional component, containing the sum of yields of the decay predecessors of the given (*A*,*Z*) isotope, was added here

$$
Y_{ZP} = \sum_{k=0}^{5} F_P(Z + ck, N - ck) e^{-(Z + ck - Z_p)^{w}/2\sigma^2},
$$
 (3)

where the term with $k=0$ corresponds to the independent $Y_{ZP}(A, Z)$ yield and the terms with $k=1-5$ stand for the precursors contributions. The upper limit of the sum over *k* was set to 5, because the charge distribution for given *A* is rather narrow and neglecting $k > 5$ did not change the sum by more than 1%.

The most probable charge path Z_p and the charge distribution width σ were expressed as third-order polynomials of *A*, with parameters $a_6 - a_9$ and $a_{10} - a_{13}$, respectively,

$$
Z_p = a_6 + a_7A + a_8A^2 + a_9A^3,
$$
 (4)

$$
\sigma = a_{10} + a_{11}A + a_{12}A^2 + a_{13}A^3. \tag{5}
$$

The value of the factor *c* in the sum of the yield cumulation for a given *A* depended on the side of the stability valley on which the given isotope lies,

$$
c = \begin{cases} 1, & EC, \beta^+ \quad \text{decay (neutron-deficient nuclei)}, \\ -1, & \beta^- \quad \text{decay (neutron-rich nuclei)}. \end{cases}
$$
 (6)

The power index w in the exponent argument in Eq. (3) was allowed to be different for $Z>Z_p$ and $Z \leq Z_p$,

$$
w = \begin{cases} a_{16}, & Z > Z_p \\ a_{17}, & Z \le Z_p, \end{cases}
$$
 (7)

where a_{16} was always set to 2 and $a_{17}=2$ or $a_{17}=1.5$ was used in order to test the asymmetric charge distribution in the latter case.

Finally, the odd-even correction was assumed to be a simple factor depending on the *Z* and *N* evenness combination

 $a_{14}a_{15}$, $Z - \text{odd}, N - \text{odd}.$ A division of the whole *A*-*Z* plane into subregions may be experimental data.

treated as yet another model parameter. To have control over it, we have plotted positions of all detected nuclei in the *N*-*Z* plane; this appeared to be very helpful for a preliminary determination of the Z_p path, especially for regions with many data. The final mass region division is illustrated in Fig. 1 by the solid lines showing Z_p fitted for seven data regions. Mass range limits were fixed to get the smallest total χ^2 for the whole data range and to possibly simplify the model for the course of the Z_p path. We have tested many alternative divisions, especially for the region of the heavy evaporation residues ($143 \leq A \leq 183$). The fitting applied to broader *A* ranges

FIG. 1. The path of the most probable atomic number Z_p corresponding to the best fit values for the parameters $a_6 - a_8$ in seven mass regions (solid lines). Positions of nuclei, for which a cumulative or an independent yield was determined, are denoted by circles and triangles, respectively. Dashed line - valley of stability approximated by the relation $Z_{\beta} = A/(1.98 + 0.0155A^{2/3})$ [38]. Open square indicates the target nucleus, vertical and horizontal dotted crosses are plotted for the magic $N=82$ and the closed shell $Z=64$ and for the magic $N = 50$ and the closed shell $Z = 40$.

than those listed in Table II, resulted in at least one order of magnitude larger χ^2 values, mainly due to rapid changes in the Z_p course at $A=162$ and $A=150$. For three separate regions of the lighter products with $A=121-139$, 82–103, and 24–75 the limits were defined by the grouping of the

As can be seen from Table II, presenting the final parameters, sometimes the best fit is obtained when the number of parameters exceeds the number of data. This was done by fixing some parameters when the others were fitted, and vice versa. Various combinations and order of fixing (or releasing) of parameters as well as their total number were tested. The shape of the mass ridge could be parametrized with a maximum of four parameters, the most probable *Z* path was approximated via a parabola except for two cases and the charge distribution width was constant or changed linearly with mass. Odd-even corrections were applied only for three heaviest mass regions, where a larger number of points and

TABLE II. Best fit parameters obtained for the yield distribution model applied to six mass regions.

Parameter		Mass range						
	$163 - 182$	$150 - 161$	$143 - 149$	$121 - 139$	$82 - 103$	$24 - 75$		
Y_A a_1	$-93.98(2)$	$-90.63(3)$	$-479.2(1)$	18.13(9)	$-30.97(5)$	$-2.07(11)$		
a ₂	1.0951(1)	1.060(2)	6.486(1)	$-0.207(1)$	0.690(1)	$-0.0059(18)$		
a_3	$-0.00310(1)$	$-0.00302(2)$	$-0.02192(3)$	$-0.000706(5)$	$-0.00390(6)$	0.00029(3)		
a_4			$-0.3(2)E-7$	$0.884(4)E-5$				
Z_p a_6	96.26(4)	136.6(1)	$-3.06(9)$	$-7.2(3)$	$-142.1(1)$	0.77(10)		
a ₇	$-0.6681(2)$	$-1.201(1)$	0.5051(6)	0.525(3)	4.886(1)	0.456(2)		
a_8	0.003081(1)	0.00485(5)	$-0.000329(4)$	$-0.00029(2)$	$-0.0456(1)$	$-0.000329(2)$		
a ₉				$0.8(14)E-7$	0.000153(1)			
σ a_{10}	$-0.82(2)$	1.17(4)	1.12(8)	0.91(9)	1.15(4)	2.12(11)		
a_{11}	0.0105(2)					$-0.013(2)$		
F_P a_{14}	0.67(3)	0.82(5)	0.58(7)	1	1			
a_{15}	0.74(3)	0.91(6)	0.90(8)		1			
$w a_{16}$	2	2	2	\mathfrak{D}	\overline{c}	2		
a_{17}	1.5	1.5	1.5	1.5	\mathfrak{D}	\mathfrak{D}		
χ^2/NDF	0.045	0.006	0.164	0.131	0.082	0.020		

FIG. 2. Normalized *Z* distributions for six mass ranges. The fitted function (line) is the exponent $\exp(-|(Z-Z_p)/\sigma|^w)$, where *w* is the charge distribution power index, Z_p is the most probable Z , and σ is the *Z* distribution width. Experimental values (\circ - independent, \blacksquare - cumulative) are normalized as described in the text. All distributions have the same normalized width equal to unity (note the asymmetric shapes for four distributions of heavier isotopes, with different slope for both sides).

smaller relative errors of the experimental data allowed one to get a reliable fit. The shape of the charge distributions was modeled better by using an asymmetric form for the evaporation residues, lying further from the stability valley (see Fig. 1). For lighter, fission products the Z_p path goes closely along the valley of stability and here a symmetric Gaussian shape was more adequate.

The normalized charge distributions for six mass intervals are plotted in Fig. 2 against the normalized charge difference $(Z - Z_p)/\sigma$. This reduces the distributions to the same width in the case where σ is not constant in the given region. The fitted function is the simple exponent $exp(-|(Z-Z_p)/\sigma|^{w})$, then, for comparison, experimental data $Y(A,Z)_E$ are normalized with three factors coming from the fit

$$
Y_Z = \frac{Y(A,Z)_E}{Y(A)F_P(N,Z)f_I}.\tag{9}
$$

Here, the mass distribution $Y(A)$ is as in Eq. (1), the oddeven correction $F_P(N,Z)$ follows Eq. (8), and the factor f_I corresponds to the independent yield fraction in the case of the cumulative yield

$$
F_P(Z,N) \int_{Z-0.5}^{Z+0.5} e^{-(z-Z_p)^w/2\sigma^2} dz
$$

$$
\sum_{k=0}^{S} \left[F_P(Z+c k, N-c k) \int_{Z+c k-0.5}^{Z+c k+0.5} e^{-(z+c k - Z_p)^w/2\sigma^2} dz \right].
$$
(10)

*f ^I*5

Sometimes charge distributions are normalized to unity integral over *Z* to get the total yield for a given *A* equal directly to Y_A [36,39]. However, when odd-even corrections are used, the distribution of the total yield cannot be described by a simple continuous function as in Eq. (2) . Also the generalization of the charge distribution shape with the twovalued (or released) index w leads to problems in obtaining an analytical form of the normalization factor for this function. Hence our distributions for six mass regions are normalized only to the same width $\sigma=1$, not to the same integral.

B. Treatment of the heaviest residues

It is a well known fact that no simple phenomenological model can properly describe the charge distribution of yields for isotopes lying near the target nuclide $[21,36]$. The main reason for this is the asymmetric, non-Gaussian shape of the charge distribution, with the most probable *Z* and width rapidly changing with mass. Such a phenomenon is clearly seen when one uses a longer section of the *Y*(*A*,*Z*) yield, along constant *Z* value instead of constant *A*. Figure 3 presents isotopic $Y_A(A)$ distributions obtained in this experiment for elements ranging from Au to Lu. Also isotopic distributions obtained for the heaviest element after stopped antiproton absorption on $176Yb$, $148Nd$, and $130Te$ targets [35] exhibit such a behavior: a steep and narrow distribution for the target element, Z_T , a flat and broad distribution for the Z_T -1 element and deformed quasi-Gaussian shapes for some smaller *Z*, with the deformation on the heavy mass side decreasing with increasing distance from Z_T . Even though the low mass side for all elements but Z_T may be described with the same slope, the slope at the higher mass side changes rapidly and cannot be fitted well with a fixed isotopic distribution asymmetry, i.e. with the unique, constant a_{17} parameter. As a consequence, the heaviest elements should be excluded from the global fit and their $Y_A(A)$ yields have to be fitted separately for a given *Z*.

The method of the yield completion for the heaviest elements is recursive: at the beginning we estimate the lacking yields of the lighter Au isotopes. With the use of these results, cumulative, experimental yields for Pt are converted to independent ones and the isotopic distribution for this element is evaluated. Then, a similar procedure is applied for Ir, Os, and so on. The method was applied down to Ta and Hf elements, where the yields for $A \ge 175$ were corrected. Finally, the summed $Y(A)$ yields for $A \ge 175$ presented with a line in Fig. 4 are the combination of results of both evalua-

FIG. 3. Isotopic yield distributions for the nine heaviest residual elements produced in the fragmentation of gold with stopped antiprotons. Dashed lines show yields obtained from the fit applied to the region $163 \leq A \leq 182$; triangles, open circles, and dots represent the experimental yields: cumulative, cumulative transformed to independent, and independent, respectively. Solid lines illustrate the corrections of the isotopic yield distribution for the heaviest isotopes of the elements from Pt to Hf, and for the lightest Au isotopes (see Sec. III for discussion).

tions: the $163 \leq A \leq 182$ region global fit and the procedure described above.

The platinum distribution was the most laborious case, due to the lack of radioactive isotopes above mass 191 and owing to the strong odd-even *N* effect (up to 30%), observed for this even-*Z* element produced after low-energy absorption of antiprotons. The overall shape of the isotopic distribution was assumed to be similar to that observed for Tm residues after Yb fragmentation with antiprotons $|35|$, with an increasing enhancement of yields for three heaviest nuclei, which for Pt are those with the mass numbers 194, 195, and 196. The number for 196 Pt obtained in this way $(16/1000\bar{p})$ was compared with the result of another estimation, based on the so-called halo factor dependence on separation energy S_n of the neutron from target nucleus [25]. For heavy nuclei with S_n close to 8 MeV, the halo factor is of the order of 4–5, hence, using ¹⁹⁶Au yield the $Y(^{196}Pt)$ should be between 14 and 23 per 1000 \bar{p} . These two estimations are consistent. For the lightest Pt isotopes, we assumed that the steep slope coming from the $163 \leq A \leq 182$ region fit is a good approximation. This assumption was used also for consecutive, lower *Z* elements (we have checked the justification

FIG. 4. Mass distribution of residual nuclei produced by the reaction of stopped \bar{p} with Au. The cumulative sum of experimentally observed yields for a given mass is shown with crosses, the corresponding fitted $Y(A)$ yield for all input data (averaged over Z for given *A*) is represented by full circles. The line shows the sum of fitted yields completed with interpolated yields for mass intervals not fitted. For the heaviest isotopes $(A \ge 175)$ an additional correction of the mass yield data was performed, as described in Sec. III.

of this approach using the lightest mass data points of these isotopes in an additional test fit). At last, on the basis of observed changes of Pt yield for odd and even *N* isotopes an appropriate correction was applied.

C. Region of the α -decay

There is a narrow bump at $A=147$ in the mass yield distribution, a feature observed for Au $[27,28]$ and Ta $[16,40]$ target fragmentation after reactions with protons, heavy ions and stopped antiprotons. The enhancement of the cross section in this region was suggested to be the result of α decay of nuclei above the $N=82$ shell [27], but the authors did not estimate quantitatively this effect due to the lack of charge dispersion curves that could not be fitted for limited experimental data. We have done such an estimation for our data performing a preliminary fit for isotopes not affected by α decay, i.e., for $154 \leq A \leq 161$, adding ¹⁵⁰Dy with an independent yield. Taking into account a charge dispersion yield *Y*(*A*,*Z*) obtained in this way, we have calculated appropriate decay corrections for experimentally measured yields of 153Tb (+2.9% correction), 153Gd (+2.8%), 152Dy (+7.2%), ¹⁵²Tb (+6.0%), ¹⁵¹Tb (+14.8%), ¹⁵¹Gd (+13.6%), ¹⁵⁰Tb (+32.5%), ¹⁴⁹Gd (+5.5%), ¹⁴⁷Gd (-13.7%), ¹⁴⁷Eu $(+32.5\%),$ ^{149}Gd $(+5.5\%),$ ^{147}Gd $(-13.7\%).$ (-12.1%) and ¹⁴⁶Gd (-23.7%). The experimental results listed in Table I are the corrected ones and were used for fitting in two mass ranges affected by this effect. As can be seen from Fig. 4 the corrections obtained are too small to remove the local yield maximum at $A=147$ (crosses show the yield before correction, circles after that). Therefore, α decay alone cannot explain fully such a feature and the observed yield enhancement in this region should be partially ascribed to the closed $N=82$ shell influence.

IV. RESULTS

The experimental data are presented in Table I, together with the fit results for isotopes representing full yield for a given pair of *A* and *Z*. Results are normalized to yield per $\frac{1000\overline{p}}{1000\overline{p}}$ with the total number of antiprotons stopped in the target (9.25 \times 10⁸). The final mass yield distribution is presented in Fig. 4. The cumulative sum of all yields observed for a given mass number *A* is here compared with the total yield *Y*(*A*) obtained from the fit via summation of all fitted *Y*(*A*,*Z*) values over *Z*, or from the interpolation between fitted mass regions. The global curve of the fitted *Y*(*A*) yield, when compared with the summed experimental yields, forms its exact skyline in almost the whole region of the evaporation residues. A deviation from this rule is observed for three mass ranges: the heaviest, with $A > 176$, a few mass numbers around $A=147$ and all fission fragments ($A<120$). Except for the second region (affected by the α decay), this is the result of prevailing accumulation of the isobaric yield by nondetectable isotopes. The depression of the observed yield of the heaviest evaporation residues is narrow but deep, with a maximum decrease to about 40% of the fitted *Y*(*A*) for $A \approx 192$, and comes mainly from the stable Pt isotopes produced. For fission products, where the Z_p path goes over the stability valley, the observed yield is strongly suppressed and its outline reaches only about 20–50 % of the fitted yield.

Leaving out two heaviest masses, the maximal *Y*(*A*) yield is reached at mass 180 but the largest individual $Y(A,Z)$ production is fitted for 176W. The small yield peaks observed for some even masses $(A=180,176,170,...)$ are due to the strongest odd-even effect for some even *Z* lying almost on the Z_p path. On the other hand, the global mass yield minimum appears between $A=105$ and $A=120$. As numerous γ lines of strongly populated heavier nuclei covered this region, no valuable production limits can be given for this region and we have to stick to the interpolated curve.

After evaluation of the mass yield curve it is possible to estimate the relative yields for different reaction channels. The fission fragments mass range should be treated with some care as their multiplicity is equal to 2 or greater when one takes into account any multifragmentation process. Assuming that all residues with $40 \leq A \leq 120$ are binary fission products (i.e. two heavy residues per antiproton) and neglecting the lightest $A < 40$ masses, we have obtained the summed fission yield. Comparing this number with the total *Y*(*A*) integral in the mass limits from 40 to 196, we have extracted the probability of gold fission induced by stopped antiprotons to be (3.8 ± 0.5) %. The lighter mass region $(A=10)$ -40), not taken into account in fission due to possible multiplicity $>$ 2 and/or the not fully negligible chance to have a fission partner in the $A > 120$ region, constitutes additionally less than 0.9% of the total yield (the error quoted above takes this into account). Our result compares well with the fission probability of $3.1(3)\%$ obtained in an experiment where fission fragment yields were measured with PIN diodes $[41]$ and is substantially larger than the value of 1.5% derived from another experiment using also on-line technique $[42]$.

FIG. 5. Comparison of the mass yield distribution obtained for protons (right axis) and stopped antiprotons (left axis). Curves for protons at 0.49 GeV, 1.0 GeV, and 3.0 GeV adopted from Ref. [27], for the inverse kinematic reaction of Au on H at 800 MeV from Refs. [29,30]. The yield axis for antiprotons was normalized with a factor of 1.75.

V. DISCUSSION

A. Antiprotons versus other projectiles

The properties of reactions induced by stopped antiproton absorption can be investigated by comparison with yield distributions obtained for other, more ''classical'' projectiles. We have confined this comparison to the gold target as the literature is quite rich here $[27,28,36,39,43-45]$. The two other popular neighbor-mass targets, Pb and Ta, represent rather different decay scenarios, with, respectively, more and less pronounced fission channel.

1. Mass yield curve

First, we present a rather qualitative comparison with the yield curve shapes extracted for protons. Figure 5 shows the summed isobaric *Y*(*A*) yield obtained for stopped antiprotons plotted together with yield distributions resulting from Au fragmentation by 0.49, 0.8, 1.0, and 3.0 GeV protons $[27,29,30]$. Since the yields for stopped antiprotons and protons are measured in different units, we have normalized our yield axis with an arbitrary factor equal to 1.75, providing the concordance between \overline{p} 's and 1 GeV protons results in the 150–170 mass range. It should be stated here that the yield curve presented for fission residues in the case of 1 GeV protons was fitted with only 5 mass points $[27]$.

The most striking differences between the curves shown in Fig. 5 are seen for the fission region. The fission probability for gold excited by protons, estimated as in Sec. IV, is equal to $\approx 6.5\%$, $\approx 3.7\%$, and $\approx 3.3\%$ for 1 GeV, 0.8 GeV, and 0.49 GeV protons, respectively. Then, 800 MeV protons seem to correspond to stopped antiprotons, but fission takes only a small part of the total yield and the comparison of distribution shapes in the evaporation region is much more adequate. Such inspection leads to the conclusion that stopped antiprotons match protons at 1 GeV.

Besides the level and width of the fission hump, a major feature distinguishing the mass yield shapes observed for

TABLE III. Characteristics of the mass yield distribution after the reaction of different projectiles with gold nuclei.

Projectile	Energy (GeV)	ΔA	Reference
$\pi^{+,-}$	0.0	7.2 ± 1.1	$[48]$
$\pi^{+,-}$	0.1	8.7 ± 1.0	[44]
$\pi^{+,-}$	0.18	10.4 ± 1.2	[44]
$\pi^{+,-}$	0.3	12.2 ± 1.3	$[44]$
\boldsymbol{p}	0.2	8.6 ± 1.1	$[27]$
\boldsymbol{p}	0.49	14.9 ± 1.8	$\lceil 27 \rceil$
\boldsymbol{p}	0.8 ^a	17.0 ± 1.4	$\lceil 45 \rceil$
\boldsymbol{p}	0.76	17.8 ± 2.3	$[28]$
\boldsymbol{p}	0.8	18.0 ± 2.3	$[28]$
\boldsymbol{p}	1.0	20.7 ± 2.8	$\lceil 27 \rceil$
\boldsymbol{p}	1.2	23.2 ± 3.2	$[28]$
\boldsymbol{p}	1.6	26.5 ± 3.4	$[28]$
\boldsymbol{p}	2.6 ^b	30.1 ± 1.9	$[36]$
\boldsymbol{p}	3.0	30.6 ± 4.3	$[27]$
\boldsymbol{p}	11.5	30.3 ± 4.2	$[43]$
\boldsymbol{p}	800.0	26.1 ± 3.7	$[39]$
\bar{p}	0.0 ^b	17.3 ± 1.3	$[37]$
\bar{p}	0.0	20.0 ± 0.8	this work

a Inverse kinematic reaction.

^bFit for generalized formula.

stopped antiprotons and protons is the distribution for the heaviest masses close to the target. Here the experimental situation is much better than in the fission case: more reliable nuclear spectroscopy data is additionally confirmed by the results of inverse kinematic measurements. The yield distribution for protons is rather unchanged in energy range from 0.5 to 1 GeV and forms a plateau between $A \approx 175$ and *A* \approx 194. On the contrary, for antiprotons in this mass range, not only for the gold target $[15,16,35]$, the yield slowly decreases from $A \approx (A_T - 20)$ to $A \approx (A_T - 3)$ and then strongly rises, reaching an absolute maximum at $A=(A_T-1)$. The enhancement of yield for few masses closest to the target may be explained by two mechanisms. The first one is the soft antiproton absorption, where almost all annihilation pions miss the rest of the target nucleus. Then A_T ⁻¹ nuclei are left with very low excitation energy. Only after antiproton absorption on nucleons occupying a deeper states $[46]$ rearrangement of nucleon configurations results in a mass loss of one or two additional units. The probability of the production of A_T ⁻¹ nuclei is quite large, about 10% for targets used in our nuclear periphery studies $[25,26]$ and the results obtained for gold are also of this order of magnitude (cf. Table I). The second mechanism leading to low excitation energies is the class of all processes where the annihilation pions escape unabsorbed by the target nucleus, but due to the sizable total π -nucleus cross section excite this nucleus enough to emit few nucleons. The probability of such a kind of quasielastic meson escapes may be quite large when the nuclear diffuseness and partial opacity are considered $[47]$.

FIG. 6. Average mass removal from the Au targets irradiated with ∇ - pions, \bullet - protons (references cited in Table III) and stopped antiprotons (pion data are shifted right by the π rest mass). The hatched band shown for antiproton data reflects the error in ΔA and a possible range of energy deposition for stopped \bar{p} annihilation with one of the target nucleons.

A quantitative comparison between antiprotons and other projectiles is presented in Table III and Fig. 6. For this purpose we have calculated the average mass removed from the target, ΔA , defined as

$$
\Delta A = A_T^* - \frac{\int_{A_{min}}^{A_{max}} Y(A)A dA}{\int_{A_{min}}^{A_{max}} Y(A) dA},
$$
\n(11)

where A_T^* equals A_T for protons and pions or A_T -1 for antiprotons. A_{min} is the lower integration limit adjusted to get all single heavy residues and A_{max} is equal to A_T for protons and pions or to A_T –2 for antiprotons. The residual mass A_T ⁻¹ is ignored in the integration of the reaction yields for antiprotons since it attests no reaction (soft \overline{p} absorption).

To get consistent and comparable results, for each data set taken from the literature we have applied a uniform method to determine ΔA . Only experimental data representing the highest (approximately the whole) cumulative yield for a given *A* were used to construct the mass yield distribution curve. The absolute errors of the quantities presented in Table III were estimated to be of the order of 10–20%, but the relative errors should be smaller. In addition to proton data, results for pions absorbed by the gold target are presented; their energy range is limited as compared to the rest of data but coincides with the kinetic energy of pions emitted in antiproton annihilation. The average mass removal from the target nucleus smoothly correlates with the projectile energy. As can be seen from Table III and Fig. 6, the average mass removed from the gold target by stopped antiprotons is only slightly lower than ΔA obtained for 1 GeV protons.

2. Charge distribution

The information on the reaction mechanism, obtained from the investigation of mass yield distributions, may be

FIG. 7. (Upper part) Distance of the fitted most probable *Z* from the beta stability line: protons (or heavy ions) compilation at saturation limit $[36]$, antiprotons at rest energy compilation $[37]$, Au at 800 MeV on H $[29,30]$, protons at 11.5 and 300 GeV $[43]$, protons at 3 and 6 GeV $[27]$, and fission for protons at 1 GeV $[27]$. (Lower part) Charge distribution widths as functions of residue mass. The results of the same evaluations as in the upper part are presented (with unchanged notation), except for protons at 1 GeV and at 3 and 6 GeV, for which the width parameter is equal to that obtained for 11.5 and 300 GeV [27].

enriched by examining other features of the *Y*(*A*,*Z*) yield topography. Two such properties were compared for results of gold fragmentation by antiprotons and other projectiles: the course of the fitted Z_p path and the charge dispersion width. Figure 7 illustrates such an inspection for some cases quoted in Table III. To bring differences into prominence, we have recalculated $Z_p(A)$ to its distance δ_{stab} from the line modeling the beta stability valley Z_β (defined as in the caption to Fig. 1). Such a presentation was earlier applied to study the distribution of products of gold projectile fragmentation on C and Al targets $[49]$. To compare with our results we present δ_{stab} obtained for the inverse kinematic Au + p reaction at 800 MeV $[29,30]$ and for energetic protons [27,43]. Besides this, we have also plotted curves derived for the Au target from a general formula describing the $Z_p(A)$ path for products of various medium and heavy target fragmentations induced with protons at the fragmentation limit [36] and with stopped antiprotons [37]. The lower part of Fig. 7 shows the dependence of the charge distribution width on the product mass.

As can be seen from Fig. 7, the relative course of Z_p and Z_β for stopped antiprotons changes now much more dramatically than in Fig. 1. The δ_{stab} calculated for more general $Y(A,Z)$ models [36,37] are smooth due to the broad *A* range fitting. Z_p fitted for 11.5 and 300 GeV protons [43] lies very close to these curves in the evaporation residues region. Surprisingly, Z_p derived for the inverse reaction at smaller energy extends further towards the neutron-deficient nuclei for heavy products. In the fission region the situation is reversed except for the lightest products. The curve plotted for 3 and 6 GeV protons lies closer to the valley of stability, for 1 GeV protons only the fission region is represented as there are no fit parameters given in Ref. $[27]$.

When *Y*(*A*,*Z*) models are applied to shorter mass ranges a rather noncontinuous behavior with segments of rapidly changing position and orientation is observed. This happens both for protons $[39]$ and heavy ions $[50–52]$ investigated with γ -ray spectroscopy technique and for heavy ion reactions on Au studied with the inverse kinematic technique [49]. Obviously, this situation cannot be ascribed merely to the uncertainties in the experimental data, even in the worst cases. In our case, the experimental data distribution in the *N*-*Z* plane shown in Fig. 1 strongly favors the segmentation of the evaporation region in fitting.

Generally, lower-energy reactions lead to Z_p running closer to the valley of stability, with passages to the neutronrich side for fission fragments. Antiproton data show a peculiar tendency: although Z_p lies quite away from the valley of stability for evaporation residues, it does not reach such a neutron-deficient region as the energetic protons. Such a behavior may be partially explained by the influence of shell effects observed in antiproton distribution for *N*/*Z* crossings at $(106,76)$, $(82,64)$, and $(50,40)$. The *Y*(*A*,*Z*) yield reaches local maxima in these regions, the most probable *Z* goes towards the more neutron-deficient nuclei and the charge dispersion becomes broader, as can be seen from Fig. 7.

The width of the *Z* distribution, σ , was found to decrease smoothly with decreasing *A* when a generalized approach is used for protons $[36]$ or antiprotons $[37]$. On the contrary, results of fitting within shorter mass regions are again inconsistent with compilations using broad mass regions, with quite small widths for evaporation residues and with a large scatter of σ for fission fragments [39,51,52]. For antiprotons stopped in Au the charge width is rather small in the evaporation region, especially in comparison with 800 MeV/ nucleon Au on H data. On the other hand, products of fission induced by stopped antiprotons are distributed quite broadly, similarly to the proton reaction products.

From a methodological point of view, results on $Z_p(A)$ and $\sigma(A)$ fitted in different ways are not consistent, even for protons at similar energies. The *Y*(*A*,*Z*) modeling applied for wide mass regions may be reasonable for limited experimental data and for generalization purposes; however, this approach washes out any possible feature of more discrete nature. Hence, the division of input data to some *A* subregions should work better in detailed studies, especially for lower excitation reactions.

B. Antiprotons stopped in various targets

There were many other targets irradiated with low-energy antiprotons from LEAR $[12–16,35]$. A review of some results of these experiments will allow us to look closely at antiproton induced reactions. Mass-charge yield models were

TABLE IV. Characteristics of mass yield distribution after antiproton stopping in different targets.

Target	ΔA	Reference
$\mathrm{nat} \mathrm{Cu}$	13.9 ± 1.8	$[14]$
96 Ru	15.8 ± 1.9	$[35]$
96Zr	16.2 ± 2.0	$[35]$
$^{98}\rm{Mo}$	16.2 ± 2.1	$\lceil 12 \rceil$
106 Cd	16.4 ± 2.3	$[35]$
$natAg$	17.7 ± 1.8	$\lceil 15 \rceil$
$^{130}\mathrm{Te}$	18.9 ± 1.6	$[35]$
$n^{nat}Ba$	17.9 ± 2.1	$\lceil 13 \rceil$
$^{144}\mathrm{Sm}$	17.5 ± 1.9	$\lceil 35 \rceil$
${}^{148}\mathrm{Nd}$	19.4 ± 2.0	$[35]$
$\rm ^{160}Gd$	19.2 ± 2.2	$[35]$
$^{165}\mathrm{Ho}$	21.7 ± 2.5	$\lceil 12 \rceil$
176 Yb	21.1 ± 2.0	$[35]$
$natTa$	22.4 ± 2.2	$\lceil 16 \rceil$
$\mathrm{nat}_{\mathrm{Au}}$	20.0 ± 0.8	This work

fitted only for a part of these targets; the parameters of the mass yield distributions for the rest were evaluated on the basis of summed direct experimental *Y*(*A*) yields. However, either for the former or the latter results, the yields for the heaviest nuclei, close to the target, are underestimated since a significant part of the total $Y(A)$ is hidden in nondetectable isotopes. To take this effect into account, we recalculated ΔA values obtained for other targets in the way as it was done for Au (see Sec. V A). The results are listed in Table IV. The removed mass increases with increasing target mass, as illustrated in Fig. 8. Such a behavior is consistent with the simple geometrical picture of an excitation energy proportional to the number of participating nucleons $[8]$, hence to the volume of the nuclei bombarded with annihilation mesons.

Using the ΔA value obtained for the Au target we may estimate the mean thermal excitation energy of the decaying system. The compilation of the measured particle emission

FIG. 8. Average mass removed from different targets after stopped antiproton absorption. Results from earlier experiments studying mass yield curves $[12–16,35]$ are recalculated after yield correction for the heaviest products.

 $[5,6,8]$ gives 5.4 nucleons ejected in the cascade +preequilibrium stages through *n*, *p*, *d*, *t*, ³He, and ⁴He ejectiles. Hence we have, on average, 14.6 evaporated nucleons and assuming 8 MeV separation energy and 3 MeV kinetic energy per nucleon [8] leads to 161 ± 23 MeV stored in the thermalized system. Such a result compares nicely with the value of 183 ± 21 MeV derived from the measurements of the spectra of neutrons and of light charged particles $[8]$.

C. Odd-even effects

Data on odd-even and shell effects observed in the *Y*(*A*,*Z*) yield distribution are rather rarely discussed in the yield modeling context $[20,21,37]$. Their influence on the yields is difficult to observe if the experimental dataset is limited and errors are of the order of the possible odd-even correction. The conclusion of Rudstam $[20]$, looking for a general formula predicting cross sections for p and α induced reactions, was that there is no need to introduce such a correction as the experiment to model yield ratios for various *N* and *Z* combinations do not show any clear correlation with the nucleon number evenness. Later, Silberberg and Tsao $[21]$ found a moderate effect, modeled with factors equal to 1.25, 0.9, 1.0 and 0.85 for even-even, odd-*N*, odd-*Z*, and odd-odd (*N*,*Z*) pairs, respectively.

Since off-line nuclear spectroscopy was applied to study the *Y*(*A*,*Z*) distribution, the odd-even effect was observed in reactions induced with stopped antiprotons $[12,13]$. Results for lighter targets $(^{92,95,98}\text{Mo}, ^{nat}\text{Ba})$ are consistent, with a 18–26% correction for odd-*Z* nuclei and a 32–34% correction for odd- N nuclei (and the sum of these values in the odd-odd case). Corresponding values fitted for 165 Ho [12] are not so evident, the yield of odd-*N* nuclei is strongly reduced (by $\approx 66\%$) whereas there is no need to correct the odd-*Z* results. Because these fits were made simultaneously for the whole heavy residue region, no dependence on the emitted number of nucleons (hence excitation) was studied, also no indication for any shell effects was reported.

Using the heavy and fissionable gold nuclei to absorb antiprotons, we have the opportunity to investigate the oddeven and shell effects in a wide evaporation and fission products mass range. In Fig. 9 we present three fitted yield distributions, as functions of mass, charge, and neutron number of the products. Corrections fitted for odd-*N* and odd-*Z* yield were shown in Table II with parameters a_{14} and a_{15} , respectively. They seem to be the largest for the medium ΔA region $(161< A < 182)$, since amongst the heavier residues only the odd-*N* Pt isotopes exhibit a clear yield reduction of about 30%. A small odd-*N* effect ($\leq 10\%$) seems also to appear for the other heavy even-*Z* products. Lighter evaporation residues are produced more uniformly over changing nucleon numbers evenness, although for products close to the closed $N=82$ shell (143 $\lt A \lt 149$) the even-*N* isotopes are strongly favored. We have not been able to study oddeven effects for the lightest evaporation and all fission products because of their small cross sections with large relative uncertainties and because of scarcity of experimental data. It should be stated that the correction for odd-odd nuclei used here slightly differs from that applied before for stopped an-

FIG. 9. Yield of residues after gold fragmentation by stopped antiprotons: (a) as a function of the atomic mass A , (b) as a function of the neutron number N , and (c) as a function of the atomic number *Z*.

tiprotons $[12,13]$ as we use the multiplicative form $[Eq. (3)]$ instead of a correction factor equal to $1-P_n-P_p$, where P_n and P_p are parameters fitted for odd- N and odd- Z nuclei, respectively. Since we observe that the odd-even effect is stronger when both *N* and *Z* are odd or even, the multiplication is more adequate.

The use of odd-even corrections for stopped antiproton reactions strongly improves the fit, with χ^2 reduction by more than one order of magnitude. Thus it cannot be treated as a trivial improvement via adding another parameter. Moreover, since corrections obtained for different mass regions are consistent, therefore, taking into account previous results obtained for lighter targets $[12,13]$ the inclusion of such a component is unavoidable in the correct modeling of the data coming from experiments with stopped antiprotons. Its strength, more pronounced than in the corresponding energetic proton data at 800 MeV $[29,30]$, is one of the most distinct features of the antiproton absorption induced reactions.

VI. CONCLUSIONS

The independent and/or cumulative yields for 114 isotopes produced after absorption of stopped antiprotons in gold were measured by using the off-line γ -ray spectroscopy technique. On this basis, with the help of a phenomenological model, the whole yield distribution was extracted for residues ranging from the target mass minus one down to the light fission products with mass ≥ 20 . The fission probability was estimated to be $(3.8\pm0.5)\%$, in agreement with the results of measurements using on-line techniques.

An average thermal excitation energy, gained by the Au nucleus after \bar{p} annihilation, was shown to be quite similar to that of 1 GeV protons, although the fission probability for such protons is almost twice as large. Moreover, the inspection of the yield distribution over the *A*-*Z* plane indicates a fairly peculiar character of the reaction induced by lowenergy antiprotons. The most probable *Z* course is quite different, lying closer to the stability line and exhibiting a more complex shape. Furthermore, the charge dispersion over *Z* does not compare with that observed for 0.8 GeV protons, being almost twice as narrow.

The average mass removal observed for various targets reacting with stopped antiprotons rises linearly with increasing target mass. This behavior is consistent with in-beam studies of the light particle emission. ΔA derived from mass yield data helps to complete such measurements, unable to detect charged particles of the lowest energy.

A clear odd-even and some shell effects distinguish evidently the reaction with stopped antiproton from those induced by energetic protons. The dependence of this phenomenon on the residue mass was studied for the first time. The strength of such effects seems to diminish with the excitation energy, although for long evaporation chains and fission products it may be unobserved due to scarce and uncertain data.

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