

# Assessment of the $^{16}\text{O}(\text{d},\text{p})^{17}\text{O}$ and $^{16}\text{O}(\text{d},\alpha)^{14}\text{N}$ cross sections

A. Gurbich

*Institute of Physics and Power Engineering*

*Obninsk, Russia*

The  $^{16}\text{O}(\text{d},\text{p}_1)^{17}\text{O}$  reaction is known to be very popular for oxygen analysis. A relatively wide plateau in the 800-900 keV energy region provides favorable conditions for IBA and so the cross section in this region was measured in a number of works. Most of the results were obtained at  $150^\circ$  (see Fig. 1) and therefore additional measurements are needed in a wide interval of angles.

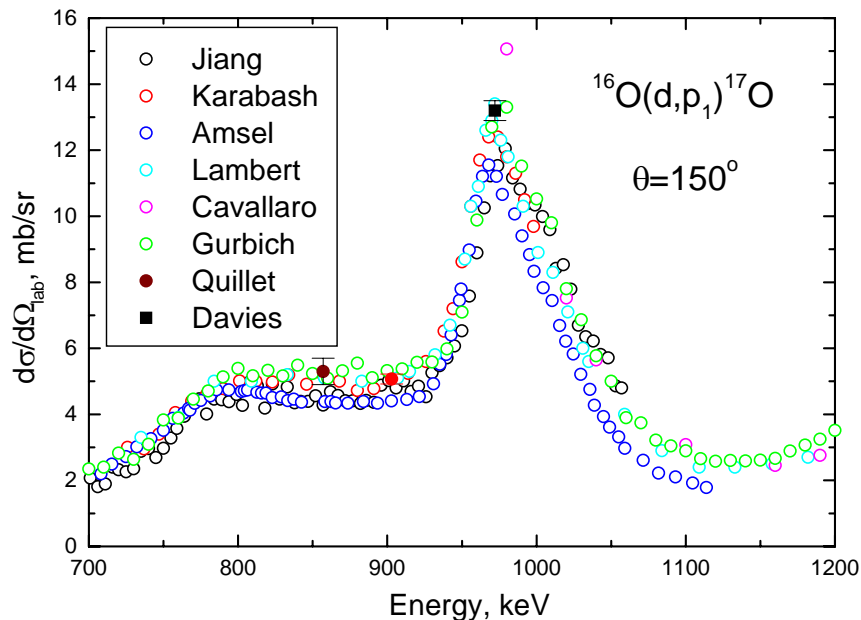


Fig. 1. Experimental data available for the  $^{16}\text{O}(\text{d},\text{p}_1)^{17}\text{O}$  reaction in the energy range from 700 to 1200 keV at  $150^\circ$ . Solid points represent the data reported in a numerical form.

An ambiguity should be noted concerning the  $^{16}\text{O}(\text{d},\text{p}_1)^{17}\text{O}$  cross section obtained by Amsel et al. and presented by the authors in graphical form in five original publications [1-5]. There is every indication that the data are the same in all the figures. Strange enough, but none of the papers contains a description of the cross section measurements. Thus nothing is known about experimental conditions at which the data were obtained. The scattering angle is  $165^\circ$  lab in the figures in all the papers except for [2] where the scattering angle is not explicitly indicated at all. However it is the figure from [2] that is reproduced in Handbooks [6-7], the cross section being attributed to the angle of  $150^\circ$ . This angle was mentioned in the paper [2], but on another occasion.

There are at least two papers [3,8] where Amsel et al. demonstrate an application of the  $^{16}\text{O}(d,p_1)^{17}\text{O}$  cross section to particular studies, with the experimental setup being presented in the figures. The detector is fixed at  $165^\circ$  in both cases. Amsel's data are compared in Fig.2 with the data sets obtained for  $150^\circ$  [9] and  $164^\circ 15'$  [10] and the agreement is much better for the  $164^\circ 15'$  case in the plateau region. The decrease of the cross section with increasing angle for backward angles corresponds to the angular distribution shown in Fig. 3.

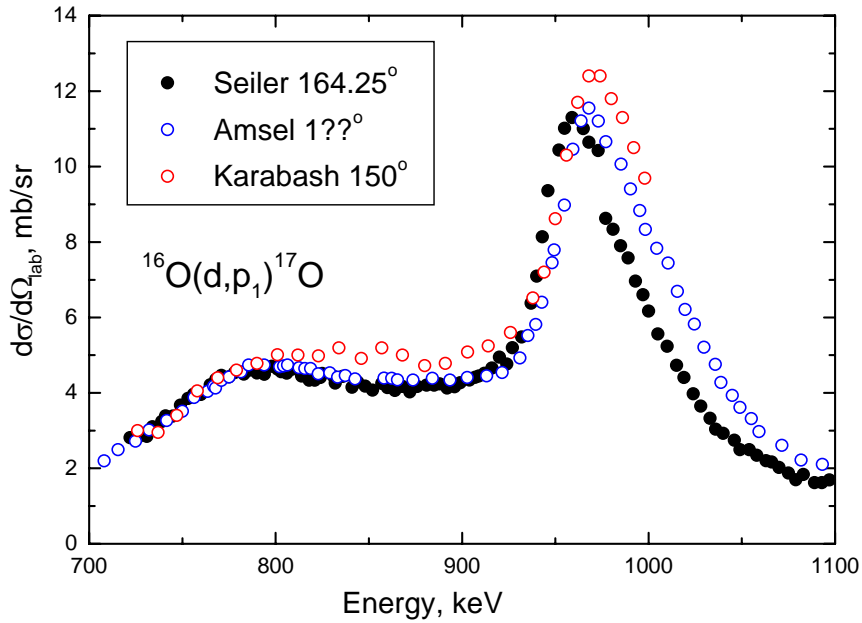


Fig. 2. Comparison of Amsel's data with results obtained for  $150^\circ$  and  $165^\circ$ .

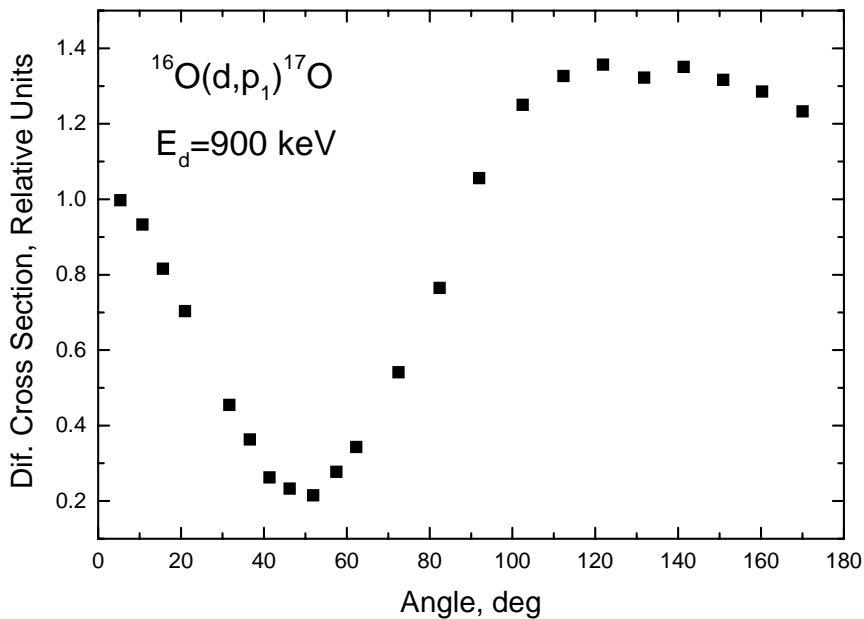


Fig. 3. Angular distribution for the  $^{16}\text{O}(d,p_1)^{17}\text{O}$  reaction at 900 keV [11].

Summing up it seems likely that the angle in Handbooks [6-7] was assigned to the data by mistake, however Prof. Amsel tends to believe that the good number for the scattering angle is  $150^\circ$  [12]. His reasons are derived from speculations about preferable experimental conditions for application of the reaction rather than grounded on some notes or reminiscences concerning the cross section measurements.

Special efforts were applied for absolute calibration of the  $^{16}\text{O}(d,p_1)^{17}\text{O}$  cross section in Refs. [13-16]. The obtained results along with the absolute data from [9] and [17] published in tabular form are compared in Table 1.

Table 1. Absolute values for the  $^{16}\text{O}(d,p_1)^{17}\text{O}$  cross section

Energy, keV	Cross section, mb/sr	Target	Reference
857	$5.3 \pm 0.4$	Ta <sub>2</sub> O <sub>5</sub>	Quillet
903	$5.07 \pm 0.15$	Al <sub>2</sub> O <sub>3</sub>	Karabash
972	$13.6 \pm 0.4$	Ta <sub>2</sub> O <sub>5</sub>	Lennard89
972	$13.3 \pm 0.4$	Ta <sub>2</sub> O <sub>5</sub>	Davies80
972	$13.2 \pm 0.3$	Ta <sub>2</sub> O <sub>5</sub>	Davies83
857	$4.28 \pm 0.11$	SiO <sub>2</sub>	Jiang
969	$11.22 \pm 0.45$		
974	$11.53 \pm 0.46$		
979	$12.05 \pm 0.48$		

All the results except for [17] are in a good agreement. As is seen from Table 1 the peak in the cross section [17] is shifted by 7 keV and the values are lower both at the plateau and for the peak. The peak to plateau ratio is 2.82 in [17] versus 2.57 in average for the other works.

At higher energies the data were measured at various angles and comparison is difficult (Fig. 4). The significant difference between the data from [18] ( $150^\circ$ ) and [19] ( $142.2^\circ$ ) near 1.6 MeV in Fig. 4 can be caused by the cross section resonance behavior.

Scarce information is available for the  $^{16}\text{O}(d,p_0)^{17}\text{O}$  and  $^{16}\text{O}(d,\alpha_0)^{14}\text{N}$  cross sections. The comparison between the  $^{16}\text{O}(d,p_0)^{17}\text{O}$  data from [18] and [19] demonstrates reasonable agreement (Fig. 5).

Cumulative information on the studied cross sections is presented in Table 2. Some ambiguity should be mentioned concerning Ref. [10]. The conversion of the angle from the laboratory system into the centre-of-mass one depends on the energy. So it is impossible to assign the same c.m. angle to all points of the excitation function as is done in the paper. There is no

indication whether the correct c.m. angle was applied in conversion of the measured yield into the cross section. It worth noting that the dependence of the angle conversion rate on energy is actually small.

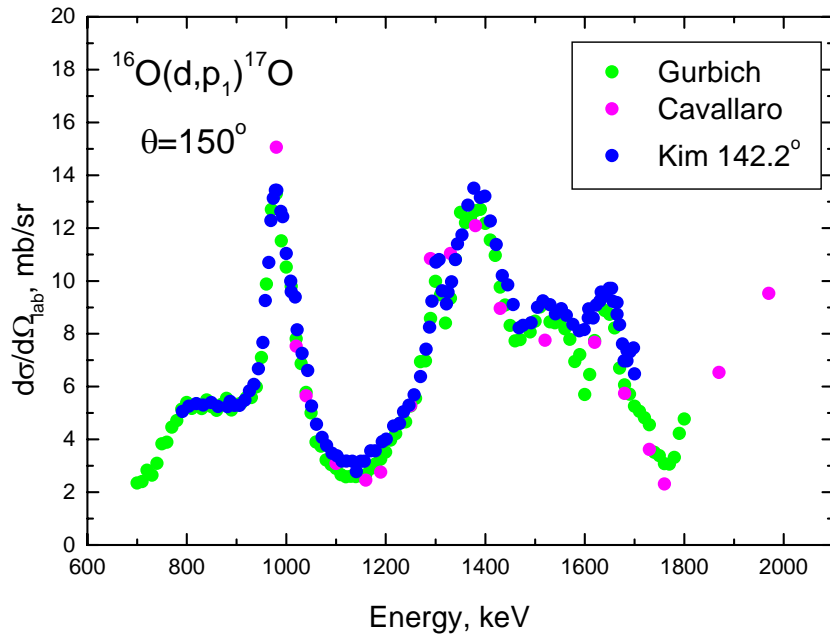


Fig. 4. Comparison of different data for the  $^{16}\text{O}(d,p_1)^{17}\text{O}$  reaction in a wide energy region (Cavallaro's excitation function was constructed from angular distributions presented in [20]).

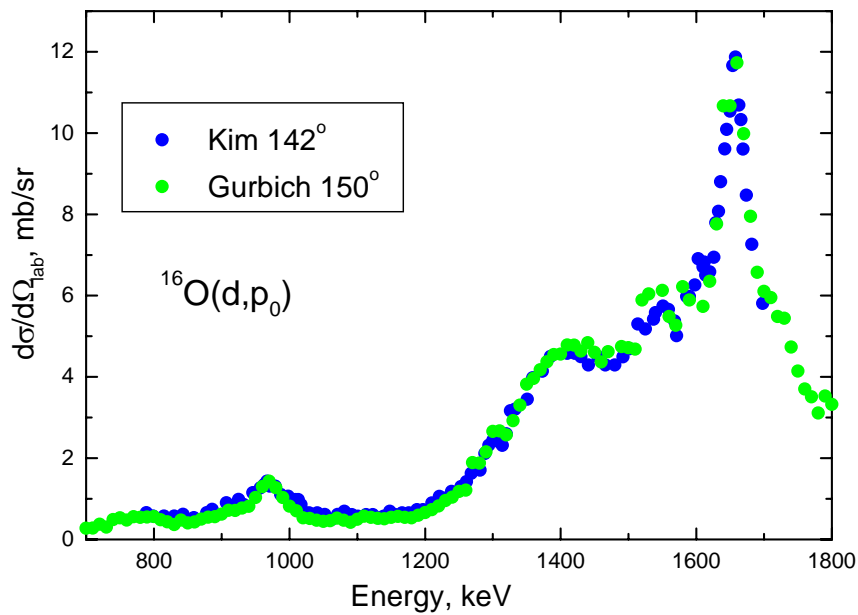


Fig. 5. Comparison of different data for the  $^{16}\text{O}(d,p_0)^{17}\text{O}$  reaction.

Table 2. Cumulative information on the deuteron induced reactions for  $^{16}\text{O}$ .

Energy range (MeV)	Reaction	Target	The energy (MeV) of angular distribution measurement	The angle of excitation function measurement	Error	Data presentation	Notes	Ref.
0.8-1.7	(d,p <sub>0</sub> ), (d,p <sub>1</sub> ), (d,α <sub>0</sub> )	Gas		51.4, 66.9, 86.7, 127.7, 142.2, 164.3	5%	Graph	Added to IBANDL	Kim
0.7-1.0	(d,p <sub>1</sub> )	Al <sub>2</sub> O <sub>3</sub> , 62.8 μg/cm <sup>2</sup>		150	3-5%	Table		Karabash
0.98-1.97	(d,d <sub>0</sub> ), (d,p <sub>0</sub> ), (d,p <sub>1</sub> ), (d,α <sub>0</sub> )	Gas	0.98, 1.02, 1.04, 1.10, 1.16, 1.19, 1.25, 1.29, 1.34, 1.38, 1.43, 1.52, 1.62, 1.68, 1.73, 1.76, 1.87, 1.97	0.98, 1.02, 1.04, 1.10, 1.16, 1.19, 1.25, 1.29, 1.34, 1.38, 1.43, 1.52, 1.62, 1.68, 1.73, 1.76, 1.87, 1.97	6%	Graph	Excitation function for 150° derived from angular distributions was added to IBANDL	Cavallaro
0.857	(d,p <sub>1</sub> )	Ta <sub>2</sub> O <sub>5</sub> , 361·10 <sup>15</sup> cm <sup>-2</sup>		150	7.5%	Value		Quillet
0.972	(d,p <sub>1</sub> )	Ta <sub>2</sub> O <sub>5</sub>		150	2%	Value	Added to IBANDL	Davies80, Davies83
0.7-1.8	(d,p <sub>0</sub> ), (d,p <sub>1</sub> )	Al <sub>2</sub> O <sub>3</sub> , 60 μg/cm <sup>2</sup>		150	7.5%	Graph, IBANDL	Mistakes were corrected in the IBANDL files	Gurbich
0.972	(d,p <sub>1</sub> )	Ta <sub>2</sub> O <sub>5</sub>		150		Value	Added to IBANDL	Lennard89
0.7-1.2	(d,p <sub>1</sub> )	Ta <sub>2</sub> O <sub>5</sub>		150	5%	Table	Added to IBANDL	Lennard91
0.7-1.06	(d,p <sub>1</sub> ), (d,α <sub>0</sub> )	SiO <sub>2</sub>		150	4%	Table, Graph		Jiang
0.55-0.66	(d,p <sub>0</sub> )	Ta <sub>2</sub> O <sub>5</sub>		150	10%	Graph		Berty

Table 2 continued

0.65-2.0	(d,p <sub>0</sub> ), (d,p <sub>1</sub> ), (d,α <sub>0</sub> )	Gas		164.25	5%	Graph	Added to IBANDL (EXFOR data converted from c.m. to lab.) instead of data from NDT.	Seiler
0.5-3.0	(d,p <sub>0</sub> ), (d,p <sub>1</sub> ), (d,α <sub>0</sub> )	SiO <sub>2</sub> , Ta <sub>2</sub> O <sub>5</sub>		135	12%	Graph	Added to IBANDL (EXFOR data) instead of Jarjis' data.	Debras
0.8-2.0	(d,p <sub>0</sub> ), (d,p <sub>1</sub> ), (d,α <sub>0</sub> )	Ta <sub>2</sub> O <sub>5</sub>	0.900, 0.950, 0.986, 1.013, 1.040, 1.067, 1.069, 1.145, 1.206, 1.266, 1.299, 1.310, 1.385	90, 135, 165 (d, α); 10, 87, (d,p <sub>0</sub> ,1)		Graph		Amsel64
0.42-1.12	(d,p <sub>1</sub> )	Presumably Ta <sub>2</sub> O <sub>5</sub>		150(?), 165(?)		Graph		Amsel [1-5]
0.84-1.02	(d,α <sub>0</sub> )	SiO <sub>2</sub>		160		Graph		Picraux
0.76-0.95	(d,p <sub>1</sub> ), (d,α <sub>0</sub> )	SiO <sub>2</sub>		145		Graph		Turos

## References

1. G. Amsel, D. Samuel, *Physics. Chem. Solids* 23 (1962) 1707.
2. G. Amsel, D. Samuel, *Anal. Chem.* 39 (1967) 1689.
3. G. Amsel, G. Bernager, B. de Gelas, P. Lacombe, *J. Appl. Phys.* 39 (1968) 2246.
4. G. Amsel, D. David, G. Beranger, P. Boisot, *Rev. Phys. Appl.* 3 (1968) 373.
5. G. Amsel, J.P. Nadai, E. d'Artemare, D. David, E. Girard, J. Moulin, *Nucl. Instr. and Meth.* 92 (1971) 481.
6. J.W. Mayer, E. Rimini, eds., *Ion Beam Handbook for Material Analysis*, Academic Press, New York, 1977.
7. J.R. Tesmer, M. Nastasi, eds., *Handbook of Modern Ion Beam Materials Analysis*, MRS, Pittsburg, PA, 1995.
8. G. Amsel, D. David, G. Beranger, P. Boisot, B. de Gelas, P. Lacombe, *J. Nucl. Materials*, 29 (1969) 144.
9. V.A Karabash, A.N. Sosnin, V.S Shorin, *Voprosy At. Nauky Techn. Ser.: Yad. Const. No. 3* (1988) 31.
10. R.F. Seiler, C.H. Jones, W.J. Anzick, D.F. Herring, K.W. Jones, *Nucl. Phys.* 45 (1963) 647 (reproduced by H.J.Kim, W.T.Milner and F.K.McGowan *Nuclear Data Tables v.A3* (1967) 123).
11. G. Amsel, *Ann. Phys.* 9 (1964) 197.
12. G. Amsel, private communications.
13. V. Quillet, F. Abel, M. Schott, *Nucl. Instr. and Meth. B* 83 (1993) 47.
14. J.A. Davies, P.R. Norton, *Nucl. Instr. and Meth.* 168 (1980) 611.
15. J. A. Davies, T.E. Jackman, H. Plattner, I. Bubb, *Nucl. Instr. and Meth.* 218 (1983) 141.
16. W.N. Lennard, S.Y. Tong, I.V. Mitchel, G.R. Massoumi, *Nucl. Instr. and Meth. B*43 (1989) 187.
17. W. Jiang, V. Shutthanandan, S. Thevuthasan, D.E. McCrready, W.J. Weber, *Nucl. Instr. and Meth. B* 207 (2003) 453.
18. A.F. Gurbich, S.L. Molodtsov, *Nucl. Instr. and Meth. B*226 (2004) 637.
19. H.C. Kim, R.F. Seiler, D.F. Herring, K.W. Jones, *Nucl. Phys.* 57 (1964) 526.
20. S. Cavallaro, A. Cunsolo, R. Potenza, A. Rubbino, *Il Nuovo Cimento*, 14A (1973) 692.
21. W.N. Lennard, G.R. Massoumi, P.F.A. Alkemade, I.V. Mitchell, S.Y. Tong, *Nucl. Instr. and Meth. B*61 (1991) 1.
22. M.Berty and A.V.Drigo, *Nucl. Instr. and Meth.* 201 (1982) 476.

23. S.T. Picraux, Nucl. Instr. and Meth. 149 (1978) 289.
24. A. Turos, L. Wielunski, J. Olenski, Phys. Stat. Sol. A 16 (1973) 211; A. Turos, L. Wielunski, A. Barcz, Nucl. Instr. and Meth. 111 (1973) 605.
25. G. Debras and G. Deconnink, J. Rad. Chem. 38 (1977)193 (reproduced by Jarjis).