

Table 2.5

Electron affinities of the elements (kJ mol <sup>-1</sup> ) <sup>a</sup>			Electron affinities of molecules <sup>a</sup>		
Z	Element	Value	Z	Element	Value
1	H	72.775	34	Se → Se <sup>1-</sup>	194.980
2	He	0		Se <sup>1-</sup> → Se <sup>2-</sup>	-410 <sup>b</sup>
3	Li	59.63	35	Br	324.6
4	Be	0	36	Kr	0
5	B	26.7	37	Rb	46.887
6	C	153.89	38	Sr	0
7	N → N <sup>1-</sup>	7	39	Y	29.6
	N <sup>1-</sup> → N <sup>2-</sup>	-673 <sup>b</sup>	40	Zr	41.1
	N <sup>2-</sup> → N <sup>3-</sup>	-1070 <sup>b</sup>	41	Nb	86.1
8	O → O <sup>1-</sup>	140.986	42	Mo	71.9
	O <sup>1-</sup> → O <sup>2-</sup>	-744 <sup>b</sup>	43	Tc	53
9	F	328.0	44	Ru	101.3
10	Ne	0	45	Rh	109.7
11	Na	52.871	46	Pd	53.7
12	Mg	0	47	Ag	125.6
13	Al	42.5	48	Cd	0
14	Si	133.6	49	In	28.9
15	P → P <sup>1-</sup>	72.02	50	Sn	107.3 <sup>c</sup>
	P <sup>1-</sup> → P <sup>2-</sup>	-468 <sup>b</sup>	51	Sb	103.2
	P <sup>2-</sup> → P <sup>3-</sup>	-886 <sup>b</sup>	52	Te	190.16
16	S → S <sup>1-</sup>	200.42	53	I	295.18
	S <sup>1-</sup> → S <sup>2-</sup>	-456 <sup>b</sup>	54	Xe	0
17	Cl	349.0	55	Cs	45.509
18	Ar	0	56	Ba	0
19	K	48.387	57	La	48
20	Ca	0	58-71	Ln	50
21	Sc	18.1	72	Hf	0
22	Ti	7.62	73	Ta	31.06
23	V	50.6	74	W	78.63
24	Cr	64.26	75	Re	14.47
25	Mn	0	76	Os	106.1
26	Fe	15.7	77	Ir	151.0
27	Co	63.7	78	Pt	205.3
28	Ni	111.5	79	Au	222.76
29	Cu	118.4	80	Hg	0
30	Zn	0	81	Tl	19.2
31	Ga	28.9	82	Pb	35.1
32	Ge	119.0 <sup>c</sup>	83	Bi	91.2
33	As → As <sup>1-</sup>	78	84	Po	183.3
	As <sup>1-</sup> → As <sup>2-</sup>	-435 <sup>b</sup>	85	At	270.1
	As <sup>2-</sup> → As <sup>3-</sup>	-802 <sup>b</sup>	86	Rn	0

<sup>a</sup> Unless otherwise noted, all values are from Hotop, H.; Lineberger, W. C. *J. Phys. Chem. Ref. Data* 1985, 14, 731.

<sup>b</sup> Pearson, R. G. *Inorg. Chem.* 1991, 30, 2856-2858.

<sup>c</sup> Miller, T. M.; Miller, A. E. S.; Lineberger, W. C. *Phys. Rev. A* 1986, 33, 3558-3559.

Table 2.6

Electron affinities of molecules <sup>a</sup>			
Molecule	Experimental (kJ mol <sup>-1</sup> ) <sup>b</sup>	Molecule	Experimental (kJ mol <sup>-1</sup> ) <sup>b</sup>
CH <sub>3</sub>	752	OCN	340
C≡CH	285	SiH <sub>3</sub>	140
C <sub>2</sub> H <sub>5</sub>	165	PH <sub>2</sub>	150
C <sub>6</sub> H <sub>5</sub>	100	PtF <sub>5</sub>	630
C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub>	85	PtF <sub>6</sub>	770
CN	365	SH	223
N <sub>3</sub>	266	SO <sub>2</sub>	107
NH <sub>2</sub>	74	SO <sub>3</sub>	160
NO	232	SCN	205
NO <sub>2</sub>	220	SF <sub>5</sub>	290
NO <sub>3</sub>	375	SF <sub>6</sub>	101
O <sub>2</sub>	42	Cl <sub>2</sub>	230
O <sub>3</sub>	203	Br <sub>2</sub>	240
OH	176	TeF <sub>5</sub>	430
OCH <sub>3</sub>	155	TeF <sub>6</sub>	320
O- <i>i</i> -C <sub>4</sub> H <sub>9</sub>	184	I <sub>2</sub>	240
O- <i>neo</i> -C <sub>5</sub> H <sub>11</sub>	183	WF <sub>6</sub>	330
OC <sub>6</sub> H <sub>5</sub>	220	UF <sub>2</sub> O <sub>2</sub>	325
O <sub>2</sub> H	104	UF <sub>6</sub>	540

<sup>a</sup> Lias, S. G.; Bartmess, J. E.; Liebman, J. F.; Holmes, J. L.; Levin, R. D.; Mallard, W. G. *J. Phys. Chem. Ref. Data* 1988, 17, Supplement 1, 1-86.

<sup>b</sup> Uncertainty is approximately ±20 except for numbers given to three significant digits.

## Problems

- Calculate the  $r$  value in pm at which a radial node will appear for the 2s orbital of the hydrogen atom.
- Which quantum numbers reveal information about the shape, energy, orientation, and size of orbitals?
- How many orbitals are possible for  $n = 4$ ? Which of these may be described as *gerade*?
- How many radial nodes do 3s, 4p, 3d and 5f orbitals exhibit? How many angular nodes?
- Make a photocopy of Fig. 2.8. Draw two lines, one along the  $z$  axis, and one at a 45° angle away from the  $z$  axis. Along one of these lines measure the distance from the origin (nucleus) to each contour line and plot the value of the contour line at that distance ( $r$ ). Do this for all contours on both lines. Compare your drawing with Fig. 2.4.
- Determine the maximum number of electrons that can exist in a completely filled  $n = 5$  level. Give four possible quantum numbers for a 5f electron of the hydrogen atom.
- The signs of the unsquared wave functions are usually shown in plots of the squared functions. Why do you think this practice exists?
- Sometimes 2p orbitals are drawn as shown below:

