1980
a) 2 points

$$
1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{3}
$$

b) 2 points
for the two electrons in the $4 \mathrm{~s}: 4,0,0,+1 / 2$ and $4,0,0,-1 / 2$
for the three electrons in the $4 \mathrm{p}: 4,1,-1,+1 / 2 ; 4,1,0,+1 / 2$ and $4,1,+1,+1 / 2$
c) 2 points

Paramagnetic. It has three unpaired electrons.
d) 2 points
$\mathrm{Na}_{3} \mathrm{As}$ - each Na gives up one electron to the As, the As has a complete octet and the sodium atoms are ionically bonded to the arsenic
$\mathrm{AsCl}_{3}$ - the three chlorines each have one half-filled orbital and the aresenic has three. So three covalent bonds are created and the As has one non-bonding pair to make a pyramidal structure.
$\mathrm{AsF}_{5}$ - fluorine is so eletronegative that it draws the two electrons of the non-bonding pair of $\mathrm{AsCl}_{3}$ into bonding. A 4 d orbital is involved in the $\mathrm{sp}^{3} \mathrm{~d}$ hydridization, yielding a trigonal bipyramidal shape.

## 1981

a) 1 point

Quantized energy levels or discrete energies or wave properties of electron produce discrete energy states in a gas.
b) 2 points

The excited state atoms can relax to several lower energy states (see diagram in c).
Each final state energy level produces a separate series.
c) 2 points

d) 3 points

Emission spectra are photons emitted from excited state systems as they drop to lower energy states.
Absorption spectra result from the absorption of electromagnetic radiation. Electrons are excited to a highter energy state.

Hydrogen atoms are in the lowest electronic energy state at $25^{\circ} \mathrm{C}(\mathrm{n}=1)$ so absorptions will be $\mathrm{n}=1$ to $n=2,3,4$, etc.

## 1987

Special rules for this question:
An ambiguous 2 pt . response receives only 1 pt
An incorrect statement in an otherwise correct 2 pt response will result in a score of 1 pt The answers labeled (i) below received two points; (ii) received one point.
a) two points

The radii of the alkali metal ions increase with increasing atomic number because:
(i) the principle quantum number (or shell or energy level) increases
(ii) there is an increase in shielding (or the number of orbitals increases)
b) two points

The chloride ion is larger than the chlorine atom because:
(i) electron- electron repulsion increases (or shielding increases or the electron-proton ratio increases or the effective nuclear charge decreases)
(ii) an extra electron generally increases the size
c) two points

The ionization energy of Mg is greater than that for Al because:
(i) the 3 p orbital is at a higher energy than the 3 s orbital (or the electron in Al is shielded from the nucleus more completely by the 3 s electron than the 3 s electrons shield one another from the nucleus)
(ii) a 3p electron is easier to remove than a 3s electron
d) two points

The much greater difference between the 2nd and 3rd ionization energies in Mg (relative to the difference between the 1st and 2nd) is due to the 3rd electron being removed from the 2 p subshell after the first 2 were removed from the 3 s subshell.

## 1987

a) three points

It is impossible to determine (or measure) bothe the position and momentum of any particle (or object, or body) simutaneously.

OR
The more exactly the position of a particle is known, the less exactly the position or velocity of the particle is known.

OR
([delta] x) ([delta] p) [greater than or equal to] h-bar (or h / 4pi)
$\mathrm{h}=$ Planck's constant
[delta] $\mathrm{x}=$ uncertainty in position
[delta] $\mathrm{p}=$ uncertainty in momentum

## Notes:

1 point is given for the notion of simultaneous determination. (A number of students give the first sentence but omit the word simultaneously. (They got 2 out of 3.) If the second or third versions of the answer are given, simultaneity is understood.
If they give the equation, they must have a > sign, not just an = sign, or they lose 1 pt .
A student who gives a correct answer but adds erroneous material gets one point deducted.
b) five points

Bohr postulated that the electron in a H atom travels about the nucleus in a circular orbit and has a fixed angular momentum. With a fixed radius of orbit and a fixed momentum (or energy), ([delta] x) ([delta] p) $<\mathrm{h} / 4 \mathrm{pi}$
and violates the uncertainty principle.
Students receive 2 pts total for the above. If they say only "the electron travels in circular orbits", they get 1 point. They also get credit for saying elliptical orbits. To receive full credit, they must describe an aspect of the Bohr theory.

The following portion of the answer is worth three points.
The wavelength of a particle is given by the deBroglie relation:
[gamma] = h/mv

For masses of macroscopic objects, $\mathrm{h} / \mathrm{m}$ is so small for any v that [gamma] is so small as to be undetectable. For an electron, $m$ is so small that $\mathrm{h} / \mathrm{mv}$ yields a detectable [gamma].

OR
They may say the product of the uncertainties in postion and velocity depends on $h / m$ and since $h$ is so small ( $\mathrm{h}=6.63 \times 10^{-34} \mathrm{~J} \bullet \mathrm{~s}$ ) unless m is very small, as for an electron, the product of the uncertainties is too small to be detected.

Students may discuss the fact that measuring the position and momentum requires having a photon strike the particle. The photon has an energy comparable to that of an electon but small compared to that of a macroscopic object. They must stress mass rather than size as the important distinction to get full credit.
A student who says [gamma] or ([delta] x) ([delta] p) depends on "size" and not mass gets 2 points if discussion is otherwise okay.
6.


The diagram shows the first ionization energies for the elements from Li to Ne. Briefly (in one to three sentences) explain each of the following in terms of atomic structure.
(a) In general, there is an increase in the first ionization energy from Li to Ne .
(b) The first ionization energy of $B$ is lower than that of Be .
(c) The first ionization energy of O is lower than that of N .
(d) Predict how the first ionization energy of Na compares to those of Li and of Ne . Explain.
(a) Across the period from $L i$ to Ne the number of protoas is inoreasing in the nucleus hence the nuclear charge is increasing with a consaquently stronger attraction for electrons and. an increase in. I.E.
(b). The electron ionized in the case of. Be is a 2 s electron … ....(1) .. whereas.. in the case of..B it is.-a $2 p$-electron.....
2p..electrons are higher in energy than $2 s$ electrons becuuse of ...(1) lesser...penctration of the core... .......
The electron ionized in $O$ is paired with another elactron in. The sane orbital, whereas in $N$ the electuon comes from a singly-ocengied orbbital. The-ioniestim anegy of the $O$ electron is less becuuse of the repulsion betreen two electrons.. . in the same ambital.
(d). The -ionization energy of Na will be less than that of both ... (1) $L_{i}$ and $N_{e}$ because. the electhon reaured-comes from-an orbital which is...farther from the naclens, therafone less $\qquad$ Hightly held.
\# 6 ( 8 pts)
(3) (4.) Electron configuration of $\mathrm{Na}: \mathrm{igg}$ ( 1 point) $\left\{\begin{array}{l}\text { Energy difference explanation between } \mathrm{Na}: \mathrm{mg} \\ \text { Size difference explanation between } \mathrm{Na}: m g\end{array}\right\} \begin{aligned} & \text { any } \\ & \text { ipo in }\end{aligned}$ Note:
If only Na or Mg is used 1 point can be earned by showing the respective electron configuration and using one of the other explanations.

Shizding/effective nuclear charge discussion
(1) (B.) Correct direction and explanation of any one of the following:
shielding differences
energy differences
\# proton / / election differences
(2) (c. (i) Ni unpaired electrons/paramaguatic
(ii) In paired electrons/diamagutic
(iii) $N$ i unpaired electrons / In paired elections
(iiii) Ni paramagnetic, In diamagnetic orbital discussion/Hend's Rule/Diagrams $\qquad$
(2) (D.) Expanded octet or $s p^{3} d$ hybrid of phosphorous (1 pair Lack of d orbitals in nitrogen (1 point)
nitrogen is too or $\begin{aligned} & \text { OR all } 5 \\ & 5\end{aligned}$ Bonding site o.

## STANDARDS

## Question 9

(a) $\mathrm{Ca}^{2+}$ has fewer electrons, thus it is smaller than Ca
(1 pt.)
The outermost electron in Ca is in a 4 s orbital, whereas the outermost electron in $\mathrm{Ca}^{2+}$ in in a $3 p$ orbital

Note: The first point is earned for indicating the loss of electrons, the second point for indicating the outermost electrons are in different shells ... must account for the magnitude of the size difference between Ca and $\mathrm{Ca}^{2+}$.
(b) $U$ for CaO is more negative than $U$ for $\mathrm{K}_{2} \mathrm{O}$, so it is more ( pt .) difficult to break up the CaO lattice (stronger bonds in CaO ).
$\mathrm{Ca}^{2+}$ is smaller than $\mathrm{K}^{+}$, so internuclear separations (between cations and $O^{-}$) are less,
OR,
(I pt.)
$\mathrm{Ca}^{2+}$ is more highly charged than $\mathrm{K}^{+}$, thus cation- $\mathrm{O}^{2-}$ bonds are stronger

Note: understanding what "lattice energy" is earns 1 point; size or charge explanation needed for the second point. Responses that use Lewis structures or otherwise indicate molecules rather than ionic lattice earn no points.
(c) (i) Ca has more protons and is smaller. The outermost electrons are more strongly held by the nuclear charge (l pt.) of Ca .
(ii) The outermost electrons in Ca are in the $4 s$, which is a higher energy orbital (more shielded) than the $3 p$ electron in $K$.

Note: for (i), the idea of attraction between nucleus and electrons must be present; for (ii), a
"noble-gas configuration" argument must be tied to an energy argument in order to earn credit.
(d) The highest energy (outermost) electron in Al is in a $3 p$ orbital, whereas that electron in Mg is in a $3 s$ orbital. (1 pt.)

The $3 p$ electron in Al is of higher energy (is more shielded) than is the $3 s$ electron in Mg .

Note: noting that different orbitals are involved earns the first point; a correct energy argument earns the second point.
Responses that attribute the greater stability of $C a$ over $K$ (or $\mathrm{K}^{+}$over $\mathrm{Ca}^{+}$, or Mg over Al ) to the stability of a completely filled (vs. half or partially filled) orbital earn NO credit.

## Question 6

(a) Response must contain a cogent discussion of the forces between the nucleus and the outermost (or "ionized") electron. For example, a discussion of "the outermost electron on $\mathrm{K} . .$. " should include one of the following:
i. it is farther from nucleus than the outermost electron on Li
ii. it is more shielded from the nucleus (or "experiences a lower effective nuclear charge") than the outermost electron on Li

2 points
for any one
iii. it is in a higher energy orbital (4s) than the outermost electron on $\mathrm{Li}(2 s)$."

Notes: " K is larger than Li " earns 1 point.
No points earned for " K electron is easier to remove" (or some other restatement).
(b) Nitrogen has one less proton than oxygen

Nitride and oxide ions are isoelectronic
1 point
or,
In nitride ion the electron/proton ratio is greater, causing more repulsion; thus, nitride is the larger ion.
(c) A Zn atom has more protons (10 more) than an atom of Ca

Electrons in $d$ orbitals of Zn have a lower principal quantum number; thus, they are not in orbitals that are farther from the nucleus.
(d) Correct identification of the orbitals involved ( $2 s$ versus $2 p$ )

Clear statement that the two orbitals have different energies 1 point
Note: Arguments that "the $2 p$ orbital is farther out than the $2 s$ orbital", or that "the Be atom has a filled subshell, which is a more stable configuration" earn no explanation point.

General note: For all parts (a) through (d), discussions of position in the periodic table earn no points.

# AP ${ }^{\circ}$ CHEMISTRY 1999 SCORING GUIDELINES <br> <br> Question 2 

 <br> <br> Question 2}

9 points
One point deduction for mathematical error (maximum once per question)
One point deduction for error in significant figures* (maximum once per question)
*number of significant figures must be correct within +/- one digit.
(a) (i)

$$
v=\frac{c}{\lambda}=\frac{3.00 \times 10^{17} \mathrm{~nm} / \mathrm{sec}}{495 \mathrm{~nm}}\left(\text { or },=\frac{3.00 \times 10^{8} \mathrm{~m} / \mathrm{sec}}{495 \times 10^{-9} \mathrm{~m}}\right)=6.06 \times 10^{14} \mathrm{sec}^{-1}
$$

(ii) $E=h v=\left(6.626 \times 10^{-34} \mathrm{~J} \mathrm{sec}\right)\left(6.06 \times 10^{14} \mathrm{sec}^{-1}\right)=4.02 \times 10^{-19} \mathrm{~J}$
(iii) $\left(4.02 \times 10^{-19} \mathrm{~J}\right)\left(6.022 \times 10^{-23} \mathrm{~mol}^{-1}\right)(0.00100 \mathrm{~kJ} / \mathrm{J})=242 \mathrm{~kJ} / \mathrm{mol}$

Note: No units required if answers are numerically same as above. No penalty if answers are correct with different units and units are explicitly indicated (e.g., for part (ii), $4.02 \times 10^{-22} \mathrm{~kJ}$ is acceptable)
(b) (i) Energy is emitted.

The $n=6$ state is at a higher energy than the $n=2$ state. Going from a high energy state to a low energy state means that energy must be emitted.

Note: The key idea is that the energy of the $n=6$ state is higher (more excited) than the lower (less excited) $n=2$ state. The argument that " $e^{-}$is farther away" at the $n=6$ level is not accepted.
(ii) $E_{2}=\frac{-2.178 \times 10^{-18} \mathrm{~J}}{2^{2}}=-5.45 \times 10^{-19} \mathrm{~J}, \quad E_{6}=\frac{-2.178 \times 10^{-18} \mathrm{~J}}{6^{2}}=-6.05 \times 10^{-20} \mathrm{~J}$

$$
\Delta E=E_{6}-E_{2}=-6.05 \times 10^{-20} \mathrm{~J}-\left(-5.45 \times 10^{-19} \mathrm{~J}\right)=4.84 \times 10^{-19} \mathrm{~J}
$$

OR,

$$
\Delta E=2.178 \times 10^{-18}\left(\frac{1}{2^{2}}-\frac{1}{6^{2}}\right) \mathrm{J}=2.178 \times 10^{-18}(0.2222) \mathrm{J}=4.84 \times 10^{-19} \mathrm{~J}
$$

Note: Point earned for determining the energy of transition.
Negative energies acceptable.

## AP ${ }^{\circ}$ CHEMISTRY <br> 1999 SCORING GUIDELINES

## Question 2 (cont.)

$$
E=\frac{h c}{\lambda} \Rightarrow \lambda=\frac{h c}{E}
$$

OR,

$$
v=\frac{E}{h}=\frac{4.84 \times 10^{-19} \mathrm{~J}}{6.626 \times 10^{-34} \mathrm{Jsec}}=7.30 \times 10^{14} \mathrm{sec}^{-1}
$$

Note: Point earned for writing or using $E=\frac{h c}{\lambda}$, or for calculating the frequency, $v$
$\lambda=\frac{c}{v}=\frac{3.00 \times 10^{8} \mathrm{~m} \mathrm{sec}^{-1}}{7.30 \times 10^{14} \mathrm{sec}^{-1}} \times \frac{1 \mathrm{~nm}}{10^{-9} \mathrm{~m}}=411 \mathrm{~nm}$
OR,
$\lambda=\frac{\left(6.626 \times 10^{-34} \mathrm{~J} \mathrm{sec}\right)\left(3.00 \times 10^{17} \mathrm{~nm} \mathrm{sec}^{-1}\right)}{4.84 \times 10^{-19} \mathrm{~J}}=411 \mathrm{~nm}$
Note: Point earned for correct wavelength; $\lambda=4.11 \times 10^{-7} \mathrm{~m}$ accepted.
Negative wavelength not accepted.
(iii) The positive charge holding the electron is greater for $\mathrm{He}^{+}$, which has a $2+$ nucleus, than for H with its $1+$ nucleus. The stronger attraction means that it requires more energy $\mathbf{1} \boldsymbol{p t}$ for the electron to move to higher energy levels. Therefore, transitions from high energy states to lower states will be more energetic for $\mathrm{He}^{+}$than for H .

Note: Other arguments accepted, such as, " $E$ is proportional to $Z^{2}$. Since $Z=2$ for $\mathrm{He}^{+}$ and $Z=1$ for H , all energy levels in $\mathrm{He}^{+}$are raised (by a factor of 4)." Other accepted answers must refer to the increased charge on the $\mathrm{He}^{+}$nucleus, and NOT the mass.

One point deduction for mathematical error (maximum once per question)
One point deduction for error in significant figures* (maximum once per question)
*number of significant figures must be correct within +/- one digit

# AP ${ }^{\circledR}$ Chemistry 2000 - Scoring Standards 

## Question 7

(8 points)
(a) The isotopes have the same number (34) of protons,

1 pt. but a different number of neutrons.

1 pt.

- No comment about the number of electrons is necessary
(b) $\quad 1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 4 s^{2} 3 d^{10} 4 p^{4}$

$$
1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6} 3 d^{10} 4 s^{2} 4 p^{4} \quad \text { lpt. }
$$

- No point is earned for $[\mathrm{Ar}] 4 s^{2} 3 d^{10} 4 p^{4}$, because the question specifically asks for a complete electron configuration.

Since there are three different $4 p$ orbitals, there must be two unpaired electrons. $\mathbf{1 p t}$.


Notes: The second part should have some explanation of Hund's rule, and may include a diagram. The second point can still be earned even if the first point is not IF the electron configuration is incorrect, but the answer for the second part is consistent with the electron configuration given in the first part.
(c) (i) The ionized electrons in both Se and Br are in the same energy level, but Br has more protons than Se , so the attraction to the nucleus is greater.

1 pt.
Note: There should be two arguments in an acceptable answer -- the electrons removed are from the same ( $4 p$ ) orbital and Br has more protons (a greater nuclear charge) than Se .
(ii) The electron removed from a Te atom is in a $5 p$ orbital, while the electron 1 pt. removed from an Se atom is in a $4 p$ orbital. The $5 p$ orbital is at a higher energy than the $4 p$ orbital, thus the removal of an electron in a $5 p$ orbital requires less energy.

# AP ${ }^{\circledR}$ Chemistry 2000 - Scoring Standards 

## Question 7

(continued)
(d)

1 pt.


Notes: One point earned for a correct Lewis diagram and a sketch. The Lewis diagram and the molecular structure may be combined into one sketch if both aspects (electron pairs and structure) are correct. Dots, lines, or a mixture of both can be used in the Lewis diagram. The lone pair of electrons need not be shown in the sketch -- just the atomic positions. No credit earned for just a verbal description of molecular geometry ("see-saw", "saw-horse", or something "distorted"), because the question clearly asks the student to "sketch the molecular structure".

The $\mathrm{SeF}_{4}$ molecule is polar, because the polarities induced by the bonds and
1 pt. the lone pair of electrons do not cancel.

