According to Table 2.3, 1 gram of H₂O gives up 2260 J when it condenses. Since each
gram of ice requires 333.5 J to melt, we get

2260 J 1g 333 5 16.78 grams of ice that can be melted.

- 2.76. , H (prods) , H(rcts) (2 mol)(26.5kJ/mol) 0 0 53.0kJ
- 2.78. H 60, H°[CO₂(g)] H°[C₁₀₀(s)] 60, H°[O₂(g)] 26367kJ/mole 60(393,51kJ/mol) x 0 26367kJ/mole
- 2.80. The reactions are:

,H "[C_{so}(s)] 2757kJ/mole

$$\begin{split} 2[\text{NaHCO}_3(s) \ \text{Na}(s) + \frac{1}{2} \text{H}_2(g) + \text{C}(s) + \frac{3}{2} \text{O}_2(g)] & 2 \text{--}_1 H = +1901.62 \text{ kJ} \\ 2 \text{ Na}(s) + \text{C}(s) + \frac{3}{2} \text{O}_2(g) \ \text{Na}_2 \text{CO}_3(s) & _1 H = -1130.77 \text{ kJ C} \\ (s) + \text{O}_2(g) \ \text{CO}_2(g) & _1 H = -393.51 \text{ kJ} \\ \text{H}_2(g) + \frac{1}{2} \text{O}_2(g) \ \text{H}_2 \text{O}(l) & _1 H = -285.83 \text{ kJ} \end{split}$$

This yields the overall reaction (you can verify that), and the overall $_{mn}H$ is the sum of the values on the right: $_{mn}H = 91.51 \text{ kJ}$.

- 2.82. The reaction is $2 \text{ A1}(s) + \text{Fe}_2\text{O}_3(s) \text{ Al}_2\text{O}_3(s) + 2 \text{ Fe}(s)$. The $_{\text{ran}}H$ is (using data from the appendix): (2 mol)(0 kJ/mol) + (1 mol)(-1675.7 kJ/mol) (1 mol)(-825.5 kJ/mol) (2 mol)(0 kJ/mol) = -850.2 kJ.
- 2.84. In this case, since the combustion occurs in open atmosphere in (assumed) constant pressure, this time q₀ = H = -31,723 J. If the expansion of gases is done against a constant atmospheric pressure of 1 atm, to determine work we need to know the net volume of gas produced. From exercise 2.55, we found that there was a net change of -0.00491 mol of gas produced (ignoring the volume of the benzoic acid itself). Thus,
- V nRT (0.0049 mol)(0.08205 L atm/mol K)(24.6.273.15) 0.120 L less gas volume

Using this change in volume to calculate work: $w p = V (1atm)(-0.120 L) \frac{101.32 J}{1L} 12.1 J$

Finally, to calculate U:

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