

SL & HL Answers to Bond enthalpies questions

- The sum of the strong O—H and C=O bonds being formed is always greater than the sum of the weaker C—H, O=O (and C=C or C≡C where relevant) bonds being broken.
- Energy in to break N≡N and 3H—H = $945 + (3 \times 436) = 2253$ kJ
 Energy out to form 6N-H = $6 \times 391 = 2346$ kJ
 ΔH for reaction (which involves formation of two mol of NH₃) = $2253 - 2346 = -93$ kJ
 Enthalpy of formation of ammonia = $\frac{1}{2} \times -93 = -46.5$ kJ mol⁻¹
- Energy in to break C—H and Cl—Cl = $414 + 242 = 656$ kJ
 Energy out to form C—Cl and H—Cl = $324 + 431 = 755$ kJ
 $\Delta H = 656 - 755 = -99$ kJ mol⁻¹
- Energy in to break C=C and H—H = $614 + 436 = 1050$ kJ
 Energy out to form C—C and 2C—H = $346 + (2 \times 414) = 1174$ kJ
 $\Delta H = 1050 - 1174 = -124$ kJ mol⁻¹
 - Extra energy in to vaporize cyclohexene to gaseous state = 33.5 kJ mol⁻¹
 Extra energy out to condense cyclohexane gas to liquid state = 32.0 kJ mol⁻¹
 Enthalpy of hydrogenation of cyclohexene = $-124 + (33.5 - 32.0) = -122.5$ kJ mol⁻¹
 Difference with literature value = $[(122.5 - 121.9) / 121.9] \times 100 = 0.49\%$.
 This difference is because average bond enthalpies were used for the C-C, C-H and C=C bonds which are not the exact values for the bonds in cyclohexene and cyclohexane.
 - Energy in to vaporise C₆H₆ and to break 3C=C and 3H—H = $33.9 + (3 \times 614) + (3 \times 436) = 3184$ kJ
 Energy out to form 3C—C and 6C—H and condense C₆H₁₂ = $(3 \times 346) + (6 \times 414) + 32.0 = 3554$ kJ
 Enthalpy of hydrogenation of benzene = $3184 - 3554 = -370$ kJ mol⁻¹
 - Benzene is expected to give out either 370 kJ mol⁻¹ (calculated from average bond enthalpies) or 366 kJ mol⁻¹ (calculated from 3 x literature value) when hydrogenated. In fact it only gives out 210 kJ mol⁻¹ as the difference in energy has been used to overcome the delocalization energy. Hence the delocalization energy of benzene is approximately 160 kJ mol⁻¹.