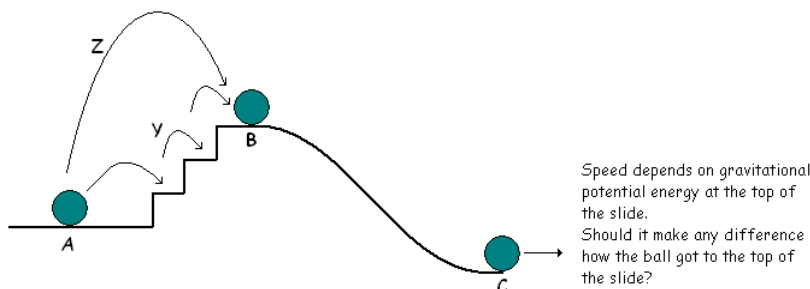


State Variables X

$\Delta X = X_f - X_i$ It does not matter how one gets from initial to final state.

For example, consider the following example concerning gravitational potential energy:



The speed that the ball achieves at the bottom of the slide depends on the potential energy that it has at the top.

The change in potential energy E_{gp} from point A to point B is $\Delta E_{gp} = E_{gpB} - E_{gpA}$.

$$E_{gpB} = \Delta E_{gp} + E_{gpA}$$

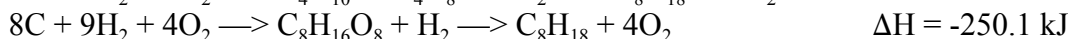
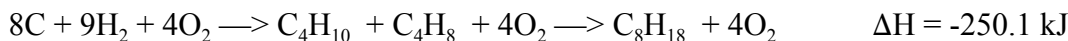
If ΔE_{gp} depends on how the ball gets from point A to B (route Y or Z, in this example), then the ball will achieve a different speed at the bottom of the slide depending on how it got to the top. Does this make any sense?

If not, then any route from point A to B must have the same ΔE_{gp} .

In other words: $\Delta E_{gp} = E_{gpF} - E_{gpI}$

It must also be the case for chemical reactions. If $\Delta H = H_f - H_i$, then it should not matter what sort of reaction path a reaction follows.

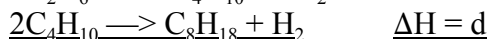
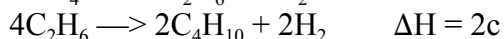
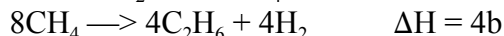
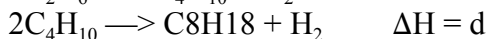
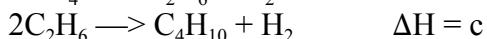
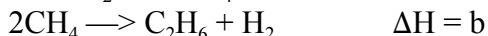
Both of the following processes must have the same overall ΔH , since they have the same initial and final states:



Why is this important? Because if any reaction of interest (target reaction) can be represented by the sum of a series of steps, then the ΔH for the target reaction must equal the sum of the ΔH values for each of the steps.

Example: target reaction is $8C + 9H_2 \longrightarrow C_8H_{18}$ $\Delta H = ?$

Known Reactions



Hess' Law: $\Delta H_{\text{target}} = \Sigma \Delta H_{\text{known}}$

Why is Hess' Law important.

- any reaction can be represented by a series of other reactions
- many target reactions cannot be measured using calorimetry because:
 - they are too slow
 - they are too dangerous
 - they are too difficult (other products are made instead)