

Standard Heats of Formation, $\Delta H_{\text{formation}}$

Reaction ¹	Substance	ΔH_f (kJ/mol)
$\text{Al}(s) \rightarrow \text{Al}(s)^2$	$\text{Al}(s)$	0
$2 \text{Al}(s) + 3/2 \text{O}_2(g) \rightarrow \text{Al}_2\text{O}_3(s)^3$	$\text{Al}_2\text{O}_3(s)$	-1670
$\text{C}(s, \text{graphite}) \rightarrow \text{C}(s, \text{graphite})$	$\text{C}(s, \text{graphite})$	0
$\text{C}(s, \text{graphite}) \rightarrow \text{C}(s, \text{diamond})$	$\text{C}(s, \text{diamond})$	2
$\text{C}(s) + 1/2 \text{O}_2(g) \rightarrow \text{CO}(g)$	$\text{CO}(g)$	-111
$\text{C}(s) + \text{O}_2(g) \rightarrow \text{CO}_2(g)$	$\text{CO}_2(g)$	-394
$\text{C}(s) + 2 \text{H}_2(g) \rightarrow \text{CH}_4(g)$	$\text{CH}_4(g)$	-75
$\text{C}(s) + 2 \text{H}_2(g) + 1/2 \text{O}_2(g) \rightarrow \text{CH}_3\text{OH}(l)$	$\text{CH}_3\text{OH}(l)$	-239
$2 \text{C}(s) + \text{H}_2(g) \rightarrow \text{C}_2\text{H}_2(g)$	$\text{C}_2\text{H}_2(g)$	227
$2 \text{C}(s) + 2 \text{H}_2(g) \rightarrow \text{C}_2\text{H}_4(g)$	$\text{C}_2\text{H}_4(g)$	52
$2 \text{C}(s) + 3 \text{H}_2(g) \rightarrow \text{C}_2\text{H}_6(g)$	$\text{C}_2\text{H}_6(g)$	-85
$2 \text{C}(s) + 3 \text{H}_2(g) + 1/2 \text{O}_2(g) \rightarrow \text{C}_2\text{H}_5\text{OH}(l)$	$\text{C}_2\text{H}_5\text{OH}(l)$	-278
$\text{Ca}(s) + 1/2 \text{O}_2(g) \rightarrow \text{CaO}(s)$	$\text{CaO}(s)$	-636
$\text{Ca}(s) + \text{O}_2(g) + \text{H}_2(g) \rightarrow \text{Ca}(\text{OH})_2(s)$	$\text{Ca}(\text{OH})_2(s)$	-987
$\text{Ca}(s) + \text{S}(s) + 2 \text{O}_2(g) \rightarrow \text{CaSO}_4(s)$	$\text{CaSO}_4(s)$	-1433
$\text{Cl}_2(g) \rightarrow \text{Cl}_2(g)$	$\text{Cl}_2(g)$	0
$2 \text{Fe}(s) + 3/2 \text{O}_2(g) \rightarrow \text{Fe}_2\text{O}_3(s)$	$\text{Fe}_2\text{O}_3(s)$	-822
$\text{H}_2(g) \rightarrow \text{H}_2(g)$	$\text{H}_2(g)$	0
$\text{H}_2(g) + \text{Cl}_2(g) \rightarrow \text{HCl}(g)$	$\text{HCl}(g)$	-92
$\text{H}_2(g) + 1/2 \text{O}_2(g) \rightarrow \text{H}_2\text{O}(g)$	$\text{H}_2\text{O}(g)$	-242
$\text{H}_2(g) + 1/2 \text{O}_2(g) \rightarrow \text{H}_2\text{O}(l)$	$\text{H}_2\text{O}(l)^4$	-286
$\text{H}_2(g) + \text{S}(s) \rightarrow \text{H}_2\text{S}(g)$	$\text{H}_2\text{S}(g)$	-20.
$\text{H}_2(g) + \text{S}(s) + 2 \text{O}_2(g) \rightarrow \text{H}_2\text{SO}_4(l)$	$\text{H}_2\text{SO}_4(l)$	-811
$\text{Hg}(l) \rightarrow \text{Hg}(g)$	$\text{Hg}(g)$	59
$\text{Mg}(s) + 1/2 \text{O}_2(g) \rightarrow \text{MgO}(s)$	$\text{MgO}(s)$	-602
$1/2 \text{N}_2(g) + 3/2 \text{H}_2(g) \rightarrow \text{NH}_3(g)$	$\text{NH}_3(g)$	-46
$1/2 \text{N}_2(g) + 3/2 \text{H}_2(g) + 1/2 \text{Cl}_2(g) \rightarrow \text{NH}_4\text{Cl}(g)$	$\text{NH}_4\text{Cl}(s)$	-315
$\text{N}_2(g) (g) + 1/2 \text{O}_2(g) \rightarrow \text{N}_2\text{O}(g)$	$\text{N}_2\text{O}(g)$	82
$2\text{N}_2(g) (g) + 2 \text{O}_2(g) \rightarrow \text{N}_2\text{O}_4(g)$	$\text{N}_2\text{O}_4(g)$	10
$\text{Na}(s) + 1/2 \text{Cl}_2(g) \rightarrow \text{NaCl}(s)$	$\text{NaCl}(s)$	-411
$\text{Na}(s) + 1/2 \text{O}_2(g) + 1/2 \text{H}_2(g) \rightarrow \text{NaOH}(s)$	$\text{NaOH}(s)$	-427
$\text{O}_2(g) \rightarrow \text{O}_2(g)$	$\text{O}_2(g)$	0
$\text{S}(s) \rightarrow \text{S}(s)$	$\text{S}(s)$	0
$\text{S}(s) + \text{O}_2(g) \rightarrow \text{SO}_2(g)$	$\text{SO}_2(g)$	-297
$\text{S}(s) + 3/2 \text{O}_2(g) \rightarrow \text{SO}_3(g)$	$\text{SO}_3(g)$	-395

¹ This column of equations where the substances is never included in Thermodynamic tables because it assumed that the person reading the table can knows that the substances listed are being made from their elements.

² The common room temp form of the ΔH_f the elements are not normally listed in tables because by definition the ΔH_f would be zero. I've included them in this table however as an instructional aid.

³ Fractional coefficients are used in the formation reactions because ΔH_f values are for one mole of the single product.

⁴ Note that you release 44 kJ of extra potential energy when hydrogen and oxygen form water, $\text{H}_2\text{O}(l)$, compared to just making steam, $\text{H}_2\text{O}(g)$. This is often brought up in AP Chemistry questions.