Dinitrogen trioxide

Dinitrogen trioxide is the <u>chemical</u> <u>compound</u> with the <u>formula</u> N_2O_3 . This deep blue <u>solid</u>^[1] is one of the simple <u>nitrogen oxides</u>. It forms upon mixing equal parts of <u>nitric oxide</u> and <u>nitrogen</u> <u>dioxide</u> and cooling the mixture below $-21 \, ^{\circ}\text{C} \, (-6 \, ^{\circ}\text{F})$:^[2]

Molar mass Appearance	76.01 g/mol deep blue tinted gas
<u>Density</u>	1.447 g/cm ³ , liquid 1.783 g/cm ³ (gas)
Melting point	-100.7 ^[1] °C
	(-149.3 °F; 172.5 K)
Boiling point	3.5 °C (38.3 °F;
	276.6 K)
	(dissociates ^[1])
Solubility in water	very soluble
<u>Solubility</u>	soluble in <u>ether</u>
<u>Magnetic</u> <u>susceptibility</u> (χ)	-16.0·10 ⁻⁶ cm ³ /mol

Structure

Molecular shape planar, C_s

<u>Dipole moment</u> 2.122 D

Thermochemistry

Heat capacity (C) 65.3 J/mol K

Std molar $314.63 \text{ J K}^{-1} \text{ mol}^{-1}$ entropy (S^{Θ}_{298})

Std enthalpy of formation ($\Delta_f H^{\mathbb{Z}}_{298}$)

+91.20 kJ/mol

Hazards

EU classification (DSD) (outdated)

Highly toxic (T+)

verify (what is *2) Infobox references

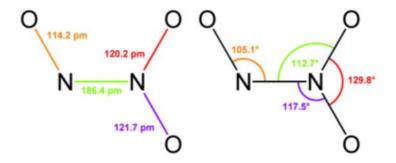
$$NO + NO_2 \Rightarrow N_2O_3$$

Dinitrogen trioxide is only isolable at low temperatures, i.e. in the liquid and solid <u>phases</u>. At higher temperatures the equilibrium favors the constituent gases, with $\underline{K}_{diss} = 193 \text{ kPa} (25 \,^{\circ}\text{C}).^{[3]}$

Structure and bonding

Typically, N-N bonds are similar in length to that in <u>hydrazine</u> (145 pm). Dinitrogen

trioxide, however, has an unusually long N-N bond at 186 pm. Some other nitrogen oxides do also possess long N-N bonds, including dinitrogen tetroxide (175 pm). The N_2O_3 molecule is planar and exhibits Cs symmetry. The dimensions displayed below come from microwave spectroscopy of low-temperature, gaseous N_2O_3 :[2]



It is the <u>anhydride</u> of the unstable <u>nitrous</u> <u>acid</u> (HNO₂), and produces it when mixed into water. An alternative structure might be anticipated for the true anhydride, i.e. O=N-O-N=O, but this isomer is not observed. If the nitrous acid is not then used up quickly, it decomposes into nitric oxide and nitric acid. Nitrite salts are sometimes produced by adding N_2O_3 to solutions of bases:

$$N_2O_3 + 2 NaOH \rightarrow 2 NaNO_2 + H_2O$$

References

ONIDES OF NITROGEN

Nitrogen sesquioxide N2O3

 N_2O_3 can only be obtained at low temperatures. It can be made by condensing equimolar amounts of NO and NO₂ together, or by reacting NO with the appropriate amount of O₂. This gives a blue liquid or solid, which is unstable and dissociates into NO and NO₂ at -30°C.

$$\begin{array}{c} NO + NO_2 \rightarrow N_2O_3 \\ 4NO + O_2 \rightarrow 2N_2O_3 \end{array}$$

It is an acidic oxide and is the anhydride of nitrous acid HNO₂. With alkali it forms nitrites.

$$N_2O_3 + H_2O \rightarrow 2HNO_2$$

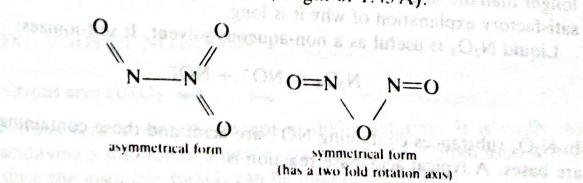
 $N_2O_3 + NaOH \rightarrow 2NaNO_2 + H_2O$

N₂O₃ reacts with the concentrated acids, forming nitrosyl salts:

$$N_2O_3 + 2HCIO_4 \rightarrow 2NO[CIO_4] + H_2O$$

 $N_2O_3 + 2H_2SO_4 \rightarrow 2NO[HSO_4] + H_2O$

The oxide exists in two different forms. These may be interconverted by irradiation with light of the appropriate wavelength. The N—N bond length from microwave spectra is 1.864 Å in the asymmetrical form. This is exceptionally long and thus the bond is exceptionally weak compared with the N—N bond found in hydrazine (length of 1.45 Å).



NOOT + NH, NO: - NH, CT