

Titanium(III)

Equilibrium reactions	lgK at infinite dilution and $T = 298\text{ K}$		
	Perrin et al., 1969	Baes and Mesmer, 1976	Brown and Ekberg, 2016
$\text{Ti}^{3+} + \text{H}_2\text{O} \rightleftharpoons \text{TiOH}^{2+} + \text{H}^+$	−1.29	−2.2	−1.65 ± 0.11
$2\text{Ti}^{3+} + 2\text{H}_2\text{O} \rightleftharpoons \text{Ti}_2(\text{OH})_2^{4+} + 2\text{H}^+$		−3.6	−2.64 ± 0.10

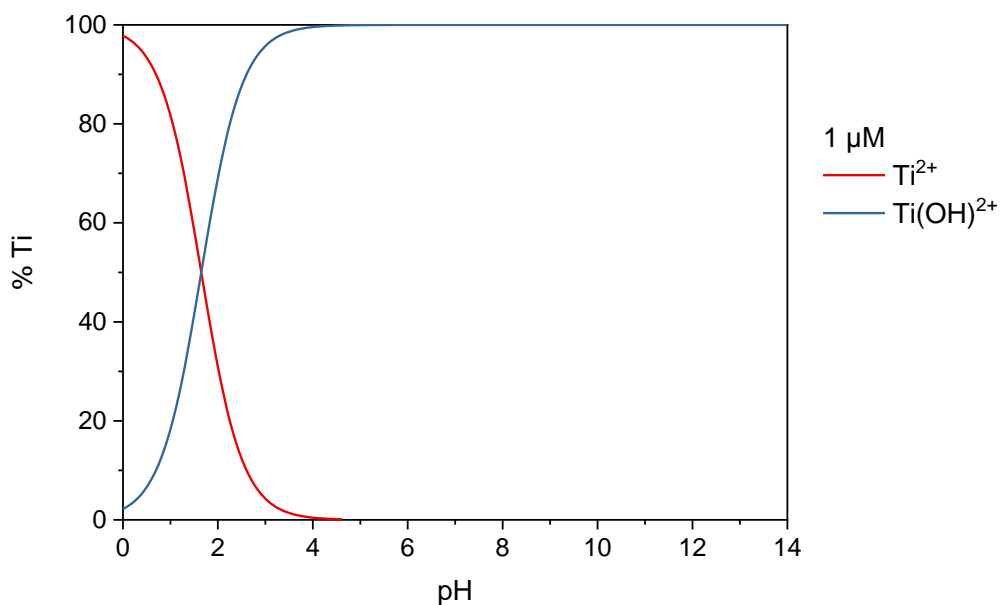
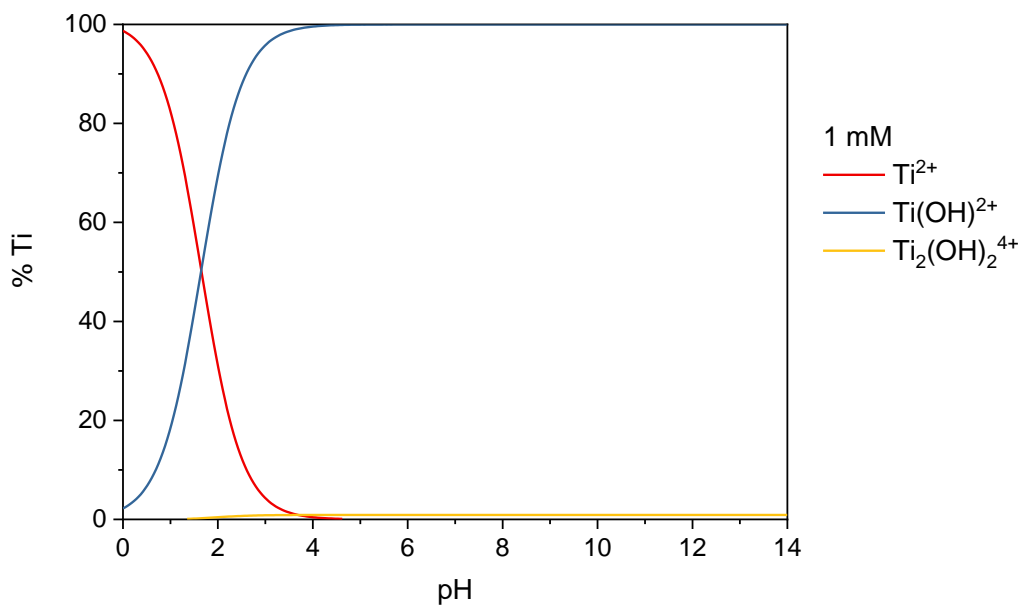
C.F. Baes and R.E. Mesmer, *The Hydrolysis of Cations*. Wiley, New York, 1976, p. 151.

P.L. Brown and C. Ekberg, *Hydrolysis of Metal Ions*. Wiley, 2016, pp. 433–442.

D.D. Perrin, *Dissociation Constants of Inorganic Acids and Bases in Aqueous Solutions*. International Union of Pure and Applied Chemistry. Commission on Electroanalytical Chemistry. Butterworths, 1969, pp. 208.

Distribution diagrams

These diagrams have been computed at two Ti(III) concentrations ($1 \text{ mM} = 1 \times 10^{-3} \text{ mol L}^{-1}$ and $1 \text{ }\mu\text{M} = 1 \times 10^{-6} \text{ mol L}^{-1}$) with the 'best' equilibrium constants above (in green). Calculations assume $T = 298 \text{ K}$ for the limiting case of zero ionic strength (*i.e.*, even neglecting plotted ions).



Equilibrium constants for hydrolysis and associated equilibria in critical compilations

Titanium(IV)

Equilibrium reactions	lgK at infinite dilution and $T = 298 \text{ K}$	
	Baes and Mesmer, 1976	Brown and Ekberg, 2016
$\text{Ti(OH)}_2^{2+} + \text{H}_2\text{O} \rightleftharpoons \text{Ti(OH)}_3^+ + \text{H}^+$	≤ -2.3	
$\text{Ti(OH)}_2^{2+} + 2 \text{H}_2\text{O} \rightleftharpoons \text{Ti(OH)}_4 + 2 \text{H}^+$	-4.8	
$\text{TiO}^{2+} + \text{H}_2\text{O} \rightleftharpoons \text{TiOOH}^+ + \text{H}^+$		-2.48 ± 0.10
$\text{TiO}^{2+} + 2 \text{H}_2\text{O} \rightleftharpoons \text{TiO(OH)}_2 + 2 \text{H}^+$		-5.49 ± 0.14
$\text{TiO}^{2+} + 3 \text{H}_2\text{O} \rightleftharpoons \text{TiO(OH)}_3^- + 3 \text{H}^+$		-17.4 ± 0.5
$\text{TiO(OH)}_2 + \text{H}_2\text{O} \rightleftharpoons \text{TiO(OH)}_3^- + \text{H}^+$		-11.9 ± 0.5
$\text{TiO}_2(\text{c}) + 2 \text{H}_2\text{O} \rightleftharpoons \text{Ti(OH)}_4$	~ -4.8	
$\text{TiO}_2(\text{s}) + \text{H}^+ \rightleftharpoons \text{TiOOH}^+$		-6.06 ± 0.30
$\text{TiO}_2(\text{s}) + \text{H}_2\text{O} \rightleftharpoons \text{TiO(OH)}_2$		-9.02 ± 0.02
$\text{TiO}_2 \times \text{H}_2\text{O} \rightleftharpoons \text{Ti(OH)}_2^{2+}[\text{OH}^-]$		
$\text{TiO}_2(\text{s}) + 4 \text{H}^+ \rightleftharpoons \text{Ti}^{4+} + 2 \text{H}_2\text{O}$		-3.56 ± 0.10

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Distribution diagrams

These diagrams have been computed at two Ti(IV) concentrations ($1 \text{ mM} = 1 \times 10^{-3} \text{ mol L}^{-1}$ and $1 \text{ }\mu\text{M} = 1 \times 10^{-6} \text{ mol L}^{-1}$) with the 'best' equilibrium constants above (in green). Calculations assume $T = 298 \text{ K}$ for the limiting case of zero ionic strength (*i.e.*, even neglecting plotted ions).

