
Equilibrium constants for hydrolysis and associated equilibria in critical compilations

Titanium(III)

| Equilibrium reactions | lgK at infinite dilution and T = 298 K | | |
|---|--|-----------------------|------------------------|
| | Perrin, 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 |

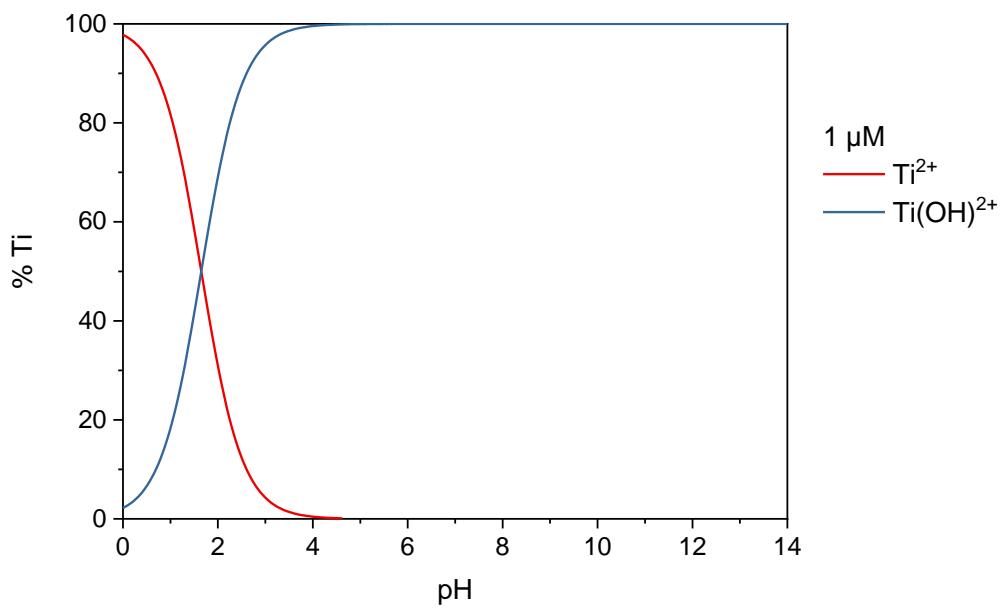
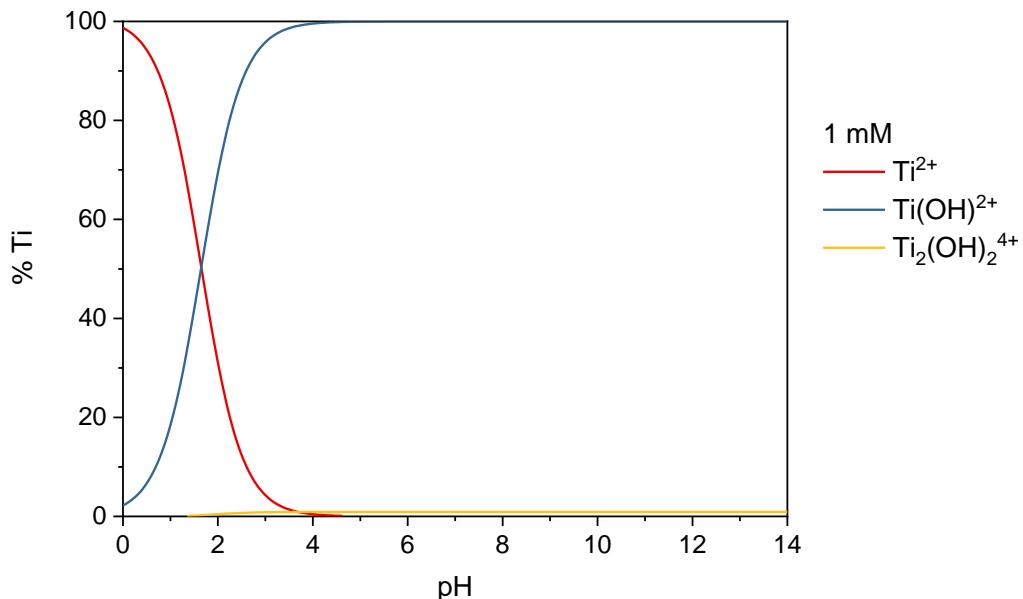
C.F. Baes and R.E. Mesmer, *The Hydrolysis of Cations*. Wiley, New York, 1976, p. 151.

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 mM = 1×10^{-3} mol L⁻¹ and 1 µM = 1×10^{-6} mol L⁻¹) with the ‘best’ equilibrium constants above (in green). Calculations assume $T = 298$ 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 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 |
| $\text{TiO}_2(\text{s}) + 4 \text{H}^+ \rightleftharpoons \text{Ti}^{4+} + 2 \text{H}_2\text{O}$ | | -3.56 ± 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.

Distribution diagrams

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

