IUPAC Task Group on Atmospheric Chemical Kinetic Data Evaluation

– Data Sheet AQ\_OH\_60

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**HCHO(aq) + H2O → CH2(OH)2(aq) (1)**

**HO(aq) + CH2(OH)2(aq) → CH(OH)2 + H2O (2)**

*(product distribution given by Chin and Wine (1994))*

**Rate coefficient data**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| k/ L mol-1 s-1 | T/K | pH | I/ mol L-1 | Reference | Technique/ Comments | |
| *Relative Rate Coefficients* | | | | | | |
| 1.3 109 | 291 | 1 | - | Merz and Waters, 1949 | Fenton-mechanism (a) |
| 1.3 109 | 294 | - | - | Hart et al., 1964 | Steady-state method (b) |
| (7.7 ±1.2)  108 | 297 | 1.5 – 5.7 | - | Chin and Wine, 1994 | LFP-LPA (d1) |
| 2.53  1010 exp[-(1040 ± 70)/T] | 279 – 319 | 1.5 – 5.7 | - |  | LFP-LPA (d2) |

The equilibrium constant for the hydration (1) is recommended as K298 K = 2000 by Doussin and Monod (2013).

*GR* (aq): Aqueous phase thermochemical data not available. As well, gas phase thermochemical data *R* (g) are not available.

**Comments**

(a) Merz and Waters are giving a relative rate constant *k* = 3.0, relative to the reference reaction (HO + Fe2+) without stating a specific rate constant; product analysis by colorimetric determination; for the recalculation of this value, the selected rate coefficient for the reference reaction *k*(HO + Fe2+) = 4.3  108 M‑1s‑1 has been used.

(b) The rate constant has been determined from the plotted data, as there was no value given in the paper (Figure 1; Hart et al., 1964); reference reaction: HO + H2O2, with *k*(HO + H2O2) = 4.5  107 M‑1s‑1 as determined by Schwarz (1962); rate coefficients have been recalculated using the selected values for the reference reactions (2.97 × 107 M‑1s‑1). It has been stated by Hart et al., that due to the lower reactivity, nearly exclusively methanediol is expected to be part of the reaction, rather than molecular formaldehyde; as no exact temperature is given, T = 294 K is assumed for room temperature.

(c) Reference reaction: HO + SCN‑ with temperature-dependent rate coefficient given as ln(*k*) = 28.7665 – (1655 / T) M‑1 s‑1; *c*(SCN‑) = (0.5 – 5)  10‑5 M; The error includes an estimated ±10% uncertainty in the H2C(OH)2 concentration (c1); *k*(T) data are given relative to the reference reaction (Table 1; Chin and Wine, 1994), errors are 1σ; *k*297(HO + SCN-) = 1.18  1010 M‑1s‑1 (calculated from given data) (c2); rate coefficients have been recalculated using the selected values for the T dependent reference reaction (Zhu et al., 2003).

**Preferred Values**

|  |  |  |
| --- | --- | --- |
| **Parameter** | **Value** | ***T*/K** |
|  |  |  |
| *k* / L mol-1 s-1 | 1.12  109 | 298 |
|  |  |  |

*Reliability*

|  |  |  |
| --- | --- | --- |
| Δ log *k* | ±0.3 | 298 |
|  |  |  |

*Comments on Preferred Values*

For the room temperature rate constant, the mean value of the data by Merz and Waters (1949), Hart et al. (1964) and Chin and Wine (1994) is suggested. The estimated uncertainty for that rate ceofficient is ±50% or Δ log *k* = 0.3. While the two determinations at room temperature agree well with each other after the recalculation, the discrepancy with the temperature dependent data is significant. A reliable Arrhenius expression is rather not to be determined and recommended.

**References**

Chin, M. and Wine, P. H.: Aquat. Surf. Photochem., 85-96, 1994.

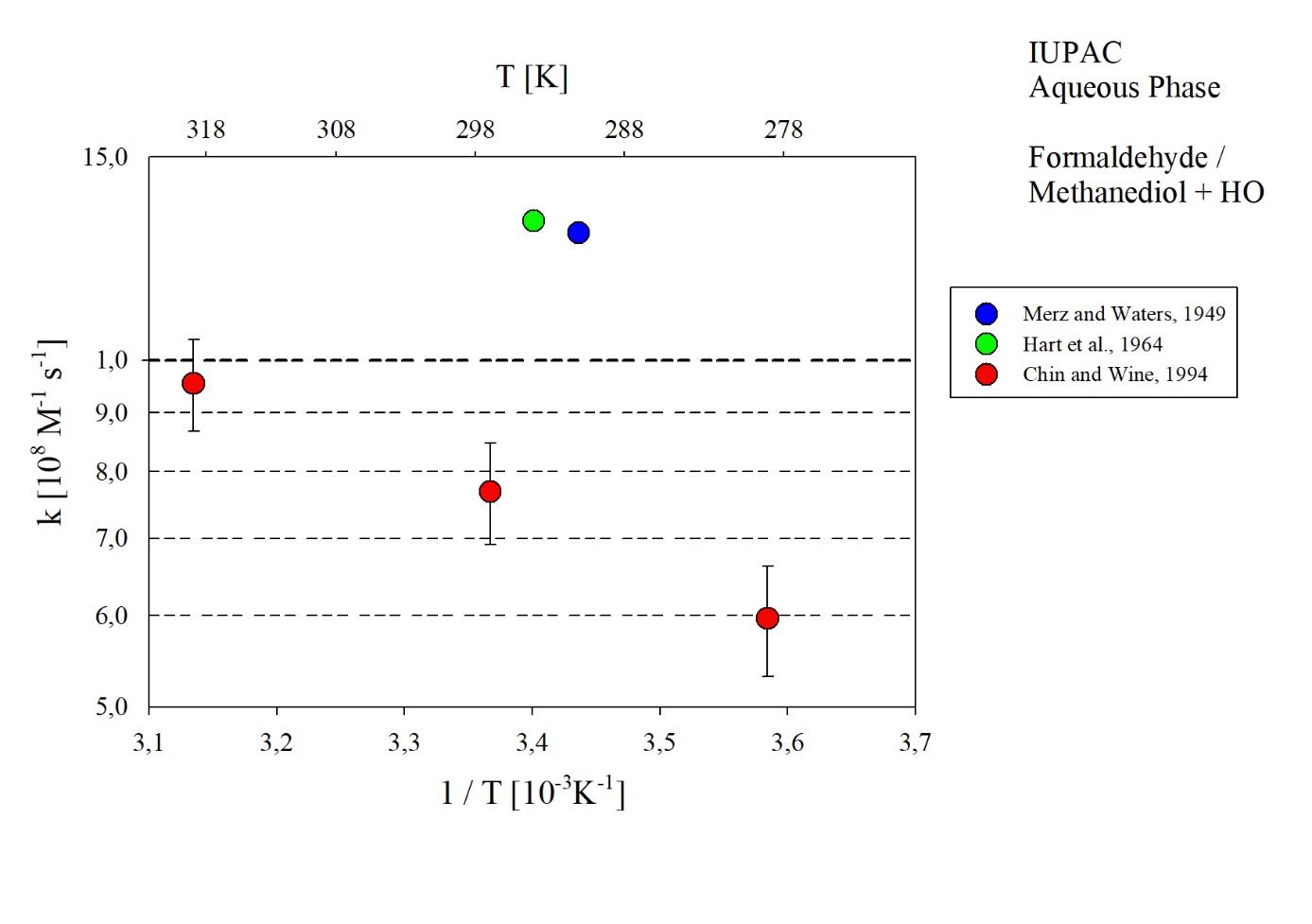
Doussin, J. F. and Monod, A.: Atmos. Chem. Phys., 13(23), 11625-11641, 2013.

Hart, E. J., Thomas, J. K. and Gordon, S.: Radiat. Res. Suppl., 4, 74-88, 1964.

Merz, J. H. and Waters, W. A.: Faraday Soc., 2(1), 179-191, 1947.

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T-dependent rate coefficients for the reaction of Formaldehyde / Methanediol with HO in aqueous solution. All data given in the plot have been considered for evaluation.