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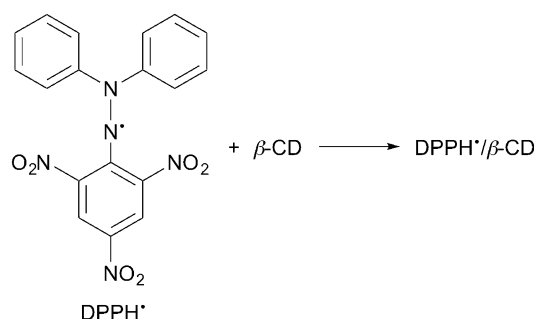
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Solubilisation of a 2,2-diphenyl-1-picrylhydrazyl radical in water by β -cyclodextrin to evaluate the radical-scavenging activity of antioxidants in aqueous media†

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A 2,2-diphenyl-1-picrylhydrazyl radical (DPPH[•]) was successfully solubilised in water by β -cyclodextrin (β -CD). DPPH[•]/ β -CD thus obtained was demonstrated to be a powerful tool to evaluate the antioxidative activity of water-soluble antioxidants, such as ascorbate and Trolox, in aqueous buffer solutions.

A relatively stable radical, 2,2-diphenyl-1-picrylhydrazyl (DPPH[•]) (Scheme 1), is frequently used as a reactivity model of reactive oxygen species (ROS) to evaluate the radical-scavenging activity of antioxidants.^{1–6} DPPH[•] shows a characteristic absorption band at around 520 nm, which disappears upon addition of compounds with radical-scavenging activity. However, alcoholic cosolvents, such as methanol and ethanol, are required to use DPPH[•] in aqueous systems due to its little solubility in water.^{7–9} In such a case, concentrated buffer solutions cannot be used to control the pH of the reaction systems because buffer salts are precipitated in the alcoholic reaction media. Cyclodextrins (CDs) are cyclic oligosaccharides that have a hydrophobic internal cavity and a hydrophilic external surface. Thus, CDs form inclusion complexes



Scheme 1 Incorporation of DPPH[•] into β -cyclodextrin (β -CD).

with a wide range of hydrophobic molecules and solubilise them in water.^{10,11} We report herein the solubilisation of DPPH[•] in water using β -cyclodextrin (β -CD; Scheme 1), which consists of 7 glucopyranoside units. The scavenging reaction of β -CD-solubilised DPPH[•] (DPPH[•]/ β -CD) by water-soluble antioxidants in phosphate buffer solution (0.1 M, pH 7.4) demonstrated that DPPH[•]/ β -CD would be a powerful tool to evaluate the antioxidative activity in aqueous media without alcoholic cosolvents.

15 mL of boiling water (Milli-Q) or a phosphate buffer solution (0.1 M, pH 7.4) was added to the mixture containing DPPH[•] (0.23 mmol) and β -CD (0.35 mmol), and the suspension was cooled to room temperature. The filtration of the suspension using a membrane filter (pore size: 0.22 μ m) gave a deep violet solution. This solution showed an absorption band at 527 nm, which is diagnostic of DPPH[•] (Fig. 1). Thus, DPPH[•] could be solubilised in water by β -CD. A significant red shift of the band due to DPPH[•]/ β -CD as compared to those of free DPPH[•] in *n*-hexane (509 nm), MeOH (516 nm), EtOH (517 nm) and acetonitrile (519 nm) suggests that the >N–N•– moiety of DPPH[•] may exist outside of the β -CD cavity and strongly interact with water. The concentration of DPPH[•] was estimated to be 5.9×10^{-5} M by using the ϵ value of $11\,000\text{ M}^{-1}\text{ cm}^{-1}$ determined for DPPH[•] in a 1 : 1 ethanol–buffer solution.⁹ When β -CD was replaced by α - or γ -CD, which consists of 6 or 8 glucopyranoside units and thus has a smaller or bigger hydrophobic cavity than β -CD,

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† Electronic supplementary information (ESI) available: Experimental details and EPR spectra at room temperature (Fig. S1). See DOI: 10.1039/c5cc02236c



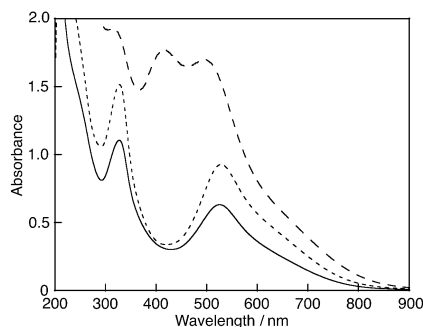


Fig. 1 UV-vis spectra of extracts of the mixture of DPPH• (0.23 mmol) and β-CD (0.45 mmol) in 15 mL of boiling phosphate buffer (0.1 M, pH 7.4) (—), acetate buffer (50 mM, pH 4.4) (---) and borate buffer (14 mM, pH 9.1) (— · —).

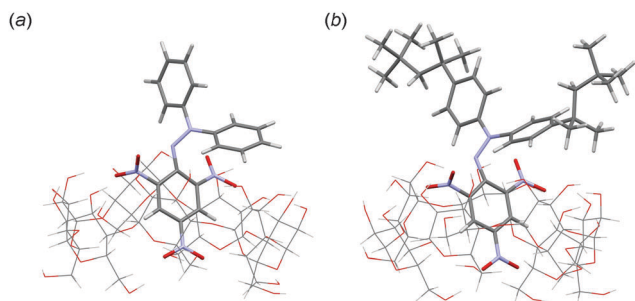


Fig. 2 Optimised structures of the inclusion complex of (a) DPPH• with β-CD and (b) DOPPH• with β-CD calculated by DFT (UB3LYP/3-21G:C-PCM solvation model parameterised for water).

DPPH• was not solubilised in water. The 2,2-di(4-*tert*-octylphenyl)-1-picrylhydrazyl radical (DOPPH•) could not be solubilised in water by β-CD in the same manner, either. Fig. 2a shows an optimised structure of the inclusion complex of DPPH• with β-CD calculated by the density functional theory (DFT) (UB3LYP/3-21G:C-PCM solvation model parameterised for water) (see the ESI†). The picryl moiety of DPPH• is incorporated into the hydrophobic cavity of β-CD. DOPPH• is also incorporated into β-CD as shown in Fig. 2b. The calculated association energy between DPPH• and β-CD ($-31 \text{ kcal mol}^{-1}$) by DFT is significantly less negative than that between DOPPH• and β-CD ($-44 \text{ kcal mol}^{-1}$). These theoretical results suggest that DOPPH• may not be solubilised by β-CD due to the hydrophobic *tert*-octyl groups. DPPH• solubilised by β-CD in water or the phosphate buffer solution (0.1 M, pH 7.4) is stable at least for several days at room temperature. When a boiling acetate buffer solution (50 mM, pH 4.4) was used instead of the phosphate buffer, DPPH• could also be solubilised by β-CD (Fig. 1). On the other hand, a brown solution with absorption bands at 416 and 505 nm was obtained using the boiling borate buffer solution (14 mM, pH 9.1) as shown in Fig. 1. This suggests that DPPH• is unstable under basic conditions as reported previously.⁹ In fact, addition of 0.75 mL of a borate buffer solution (0.1 M, pH 9.1) to DPPH•/β-CD in water (Milli-Q) (2.3 mL) resulted in a gradual decrease in the absorption band at 527 nm, accompanied by an increase in the band at 412 nm with clear isosbestic points at 252, 304, 346 and 491 nm as shown in Fig. 3. The one-electron reduced DPPH• (DPPH^{•-}) is reported to have an

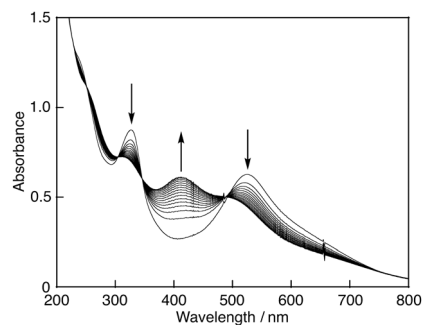


Fig. 3 Spectral change observed for DPPH•/β-CD ($5.8 \times 10^{-5} \text{ M}$) in borate buffer (25 mM, pH 9.1) at 298 K. Interval: 20 min.

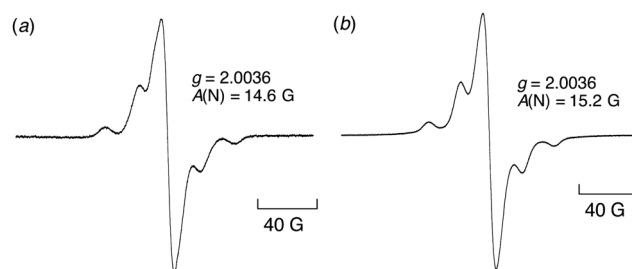
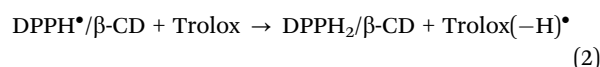


Fig. 4 EPR spectra of (a) DPPH•/β-CD ($1.0 \times 10^{-5} \text{ M}$) in distilled water at 77 K and (b) DPPH• ($1.0 \times 10^{-5} \text{ M}$) in MeOH at 77 K.

absorption band at 426 nm in a 1:1 ethanol-buffer solution,⁹ suggesting that DPPH^{•-} may be included in the products. However, the detailed reaction mechanism of DPPH• under basic conditions is under investigation and will be reported elsewhere.

The EPR spectrum of DPPH•/β-CD observed in water at room temperature has the same *g* value (2.0036) and hyperfine coupling constant (7.8 G) as those of DPPH• in MeOH (2.0036 and 7.9 G, respectively) (see the Fig. S1, ESI†). A slightly small hyperfine coupling constant (14.6 G) was observed in water as compared to that in MeOH (15.2 G) for the EPR spectra recorded at 77 K (Fig. 4).

When ascorbic acid (AscH₂) was added to the phosphate buffer solution (0.1 M, pH 7.4) of DPPH•/β-CD, the band at 527 nm disappeared immediately with clear isosbestic points at 320, 338 and 431 nm as shown in Fig. 5a. Since the *pK_a* value of AscH₂ is reported to be 4.1,¹² AscH₂ undergoes deprotonation and exists in its anionic form, AscH⁻, in phosphate buffer solution (0.1 M, pH 7.4). Thus, this spectral change indicates that AscH⁻ efficiently scavenged DPPH• in phosphate buffer [eqn (1)]. When AscH⁻ was replaced by Trolox, a water-soluble analogue of α-tocopherol (vitamin E), a similar spectral change was observed due to the scavenging reaction of DPPH• by Trolox [Fig. 5b and eqn (2)].¹³



Spectral titrations (insets of Fig. 3a and b) show the same stoichiometry with both antioxidants, the DPPH•/antioxidant molar ratio being 2:1.⁷ The decay of the absorbance at 527 nm monitored by a stopped-flow technique obeyed pseudo-first-order kinetics,



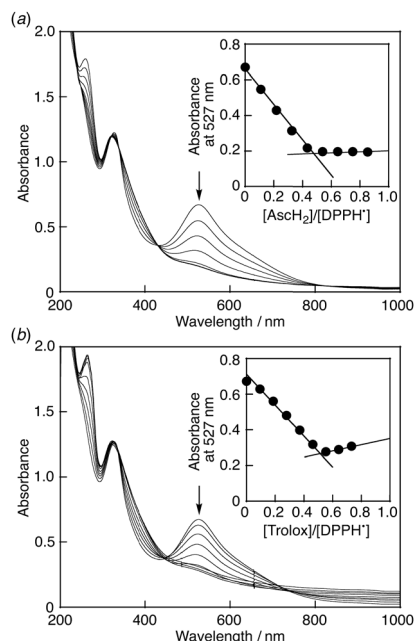


Fig. 5 Spectral changes observed upon addition of (a) AsCH_2 ($0\text{--}5.4 \times 10^{-5}$ M) or (b) Trolox ($0\text{--}4.6 \times 10^{-5}$ M) to $\text{DPPH}^\bullet/\beta\text{-CD}$ (6.3×10^{-5} M) in phosphate buffer (0.1 M, pH 7.4). Insets: plots of the absorbance at 527 nm vs. (a) $[\text{AsCH}_2]/[\text{DPPH}^\bullet]$ and (b) $[\text{Trolox}]/[\text{DPPH}^\bullet]$.

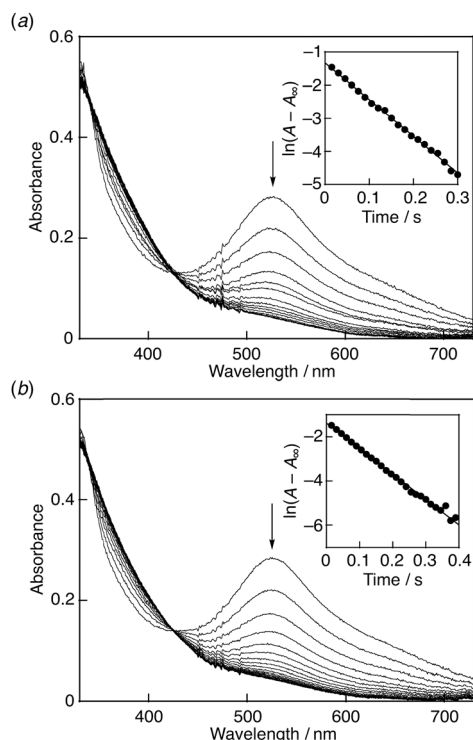


Fig. 6 Spectral changes (interval: 25 ms) observed during reactions of (a) AsCH_2 (1.4×10^{-3} M) with $\text{DPPH}^\bullet/\beta\text{-CD}$ (2.6×10^{-5} M) and (b) Trolox (6.3×10^{-4} M) with $\text{DPPH}^\bullet/\beta\text{-CD}$ (3.0×10^{-5} M) in phosphate buffer (0.1 M, pH 7.4) at 298 K. Insets: the first-order plots of the absorbance at 527 nm.

when the concentration of AsCH_2 ($[\text{AsCH}_2]$) was maintained at more than a 10-fold excess of $\text{DPPH}^\bullet/\beta\text{-CD}$ concentration (Fig. 6a).

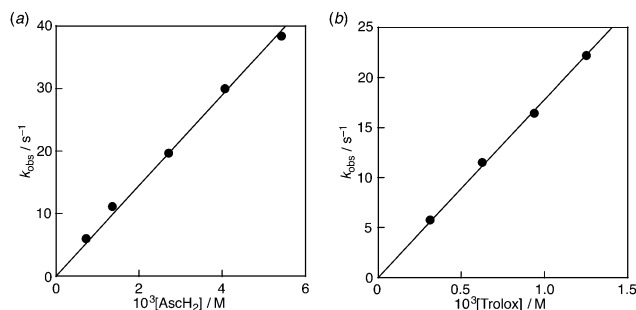


Fig. 7 Plots of k_{obs} vs. (a) $[\text{AsCH}_2]$ and (b) $[\text{Trolox}]$.

The pseudo-first-order rate constant (k_{obs}) linearly increased with increasing $[\text{AsCH}_2]$ (Fig. 7a). From the slope of the linear plot the second-order rate constant (k) for the scavenging of $\text{DPPH}^\bullet/\beta\text{-CD}$ by AsCH_2 was determined in a phosphate buffer (0.1 M, pH 7.4) to be $7.2 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$. The k value for Trolox was also determined in the same manner to be $1.8 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$ (Fig. 6b and 7b), which is exactly the same as that measured in a 1:1 ethanol-buffer (pH 7.4) solution.^{9,14} Thus, $\beta\text{-CD}$ does not inhibit the reaction of DPPH^\bullet with the antioxidants, while the stability and reactivity of nitroxyl radicals were reported to be significantly changed by the complexation with cyclodextrins.¹⁵

In summary, $\beta\text{-CD}$ -solubilised DPPH^\bullet in water has been demonstrated to be a powerful tool to evaluate the antioxidative activity of antioxidants in aqueous media, especially in highly concentrated buffer solutions without precipitation of buffer salts.

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Notes and references

- M. S. Blois, *Nature*, 1958, **181**, 1199.
- (a) K. U. Ingold and D. A. Pratt, *Chem. Rev.*, 2014, **114**, 9022; (b) G. Litwinienko and K. U. Ingold, *Acc. Chem. Res.*, 2007, **40**, 222.
- (a) E. van Wenum, R. Jurczakowski and G. Litwinienko, *J. Org. Chem.*, 2013, **78**, 9102; (b) M. Musialik, R. Kuzmich, T. S. Pawłowski and G. Litwinienko, *J. Org. Chem.*, 2009, **74**, 2699; (c) P. Mulder, G. Litwinienko, S. Lin, P. D. MacLean, L. R. Barclay and K. U. Ingold, *Chem. Res. Toxicol.*, 2006, **19**, 79; (d) G. Litwinienko and K. U. Ingold, *J. Org. Chem.*, 2005, **70**, 8982; (e) M. Musialik and G. Litwinienko, *Org. Lett.*, 2005, **7**, 495; (f) G. Litwinienko and K. U. Ingold, *J. Org. Chem.*, 2004, **69**, 5888; (g) G. Litwinienko and K. U. Ingold, *J. Org. Chem.*, 2003, **68**, 3433.
- (a) R. Amorati, S. Menichetti, C. Viglianisi and M. C. Foti, *Chem. Commun.*, 2012, **48**, 11904; (b) M. C. Foti, *Int. J. Chem. Kinet.*, 2012, **44**, 524; (c) V. D. Kancheva, L. Saso, S. E. Angelova, M. C. Foti, A. Slavova-Kasakova, C. Daquino, V. Enchev, J. Firuzi and M. Nechev, *Biochimie*, 2012, **94**, 403; (d) M. C. Foti, C. Daquino, G. A. DiLabio and K. U. Ingold, *Org. Lett.*, 2011, **13**, 4826; (e) M. C. Foti, R. Amorati, G. F. Pedulli, C. Daquino, D. A. Pratt and K. U. Ingold, *J. Org. Chem.*, 2011, **75**, 4434; (f) M. C. Foti, C. Daquino, I. D. Mackie, G. A. DiLabio and K. U. Ingold, *J. Org. Chem.*, 2008, **73**, 9270; (g) M. C. Foti and C. Daquino, *Chem. Commun.*, 2006, 3252; (h) M. C. Foti, C. Daquino and C. Geraci, *J. Org. Chem.*, 2004, **69**, 2309; (i) M. C. Foti, L. R. Barlay and K. U. Ingold, *J. Am. Chem. Soc.*, 2002, **124**, 12881.
- (a) T. Waki, S. Kobayashi, K. Matsumoto, T. Ozawa, T. Kamada and I. Nakanishi, *Chem. Commun.*, 2013, **49**, 9842; (b) H. P. Indo, I. Nakanishi, K. Ohkubo, H. Yen, M. Nyui, S. Manda, K. Matsumoto, K. Fukuhara, K. Anzai, N. Ikota, H. Matsui, Y. Minamiyama, A. Nakajima, H. Ichikawa, S. Fukuzumi, T. Ozawa, C. Mukai and



- H. J. Majima, *RSC Adv.*, 2013, **3**, 4535; (c) T. Kawashima, S. Manda, Y. Uto, K. Ohkubo, H. Hori, K. Matsumoto, K. Fukuhara, N. Ikota, S. Fukuzumi, T. Ozawa, K. Anzai and I. Nakanishi, *Bull. Chem. Soc. Jpn.*, 2012, **85**, 877; (d) T. Waki, I. Nakanishi, K. Matsumoto, J. Kitajima, T. Chikuma and S. Kobayashi, *Chem. Pharm. Bull.*, 2012, **60**, 37; (e) S. Kobayashi, T. Waki, I. Nakanishi, K. Matsumoto and K. Anzai, *Chem. Pharm. Bull.*, 2010, **58**, 1442; (f) I. Nakanishi, T. Kawashima, K. Ohkubo, H. Kanazawa, K. Inami, M. Mochizuki, K. Fukuhara, H. Okuda, T. Ozawa, S. Itoh, S. Fukuzumi and N. Ikota, *Org. Biomol. Chem.*, 2005, **3**, 626.
- 6 S. B. Kedare and R. P. Singh, *J. Food Sci. Technol.*, 2011, **48**, 412 and references cited therein.
- 7 M. B. Arnao, *Trends Food Sci. Technol.*, 2000, **11**, 419.
- 8 D.-W. Li, F.-F. Tian, Y.-S. Ge, X.-L. Ding, J.-H. Li, Z.-Q. Xu, M.-F. Zhang, X.-L. Han, R. Li, F.-L. Jiang and Y. Liu, *Chem. Commun.*, 2011, **47**, 10713.
- 9 O. Friaa and D. Brault, *Org. Biomol. Chem.*, 2006, **4**, 2417.
- 10 G. Crini, *Chem. Rev.*, 2014, **114**, 10940 and references cited therein.
- 11 I. Nakanishi, S. Fukuzumi, T. Konishi, K. Ohkubo, M. Fujitsuka, O. Ito and N. Miyata, *J. Phys. Chem. B*, 2002, **106**, 2372.
- 12 (a) C. Creutz, *Inorg. Chem.*, 1981, **20**, 4449; (b) N. H. Williams and J. K. Yandell, *Aust. J. Chem.*, 1982, **35**, 1133.
- 13 Clear isosbestic points such as those in Fig. 3a were not observed due to relatively poor solubility of Trolox in phosphate buffer as compared to AsCH₂.
- 14 For 2:1 and 3:1 ethanol–water mixtures, the *k* values were reported to be 8.6×10^3 and $7.0 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$, respectively. See ref. 9.
- 15 (a) M. Spulber and S. Schlick, *J. Phys. Chem. A*, 2010, **114**, 6217; (b) D. Bardelang, L. Charles, J.-P. Finet, L. Jicsinszky, H. Karoui, S. R. A. Marque, V. Monnier, A. Rockenbauer, R. Rosas and P. Tordo, *Chem.–Eur. J.*, 2007, **13**, 9344; (c) H. Karoui and P. Tordo, *Tetrahedron Lett.*, 2004, **45**, 1043; (d) H. Karoui, A. Rockenbauer, S. Pietri and P. Tordo, *Chem. Commun.*, 2002, 3030; (e) C. Ebel, K. U. Ingold, J. Michon and A. Rassat, *Tetrahedron Lett.*, 1985, **26**, 741.

