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Quercetin spectrofluorometric quantification in aqueous media using different surfactants as fluorescence promoters†

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Quercetin spectrofluorometric quantification was carried out in aqueous media (pH 7) using micelles of surfactants, namely: CTAB, SDS and TX100 as promoters of quercetin fluorescence, having a critical micelle concentration, CMC, of: 0.94 ± 0.03 , 7.7 ± 0.6 and 0.18 ± 0.3 mM, respectively, measured through UV-Vis spectrophotometry. The thermodynamic binding constant, K , for the quercetin-surfactants' micelles supramolecular complex was estimated to be $(\log(K/M^{-1})) = 2.87 \pm 0.02$, 2.78 ± 0.04 and 2.80 ± 0.04 , respectively, from fluorescence spectrophotometry. With the aid of these parameters it was possible to construct species distribution diagrams and choose the experimental conditions where quercetin can be quantified in aqueous media from fluorescence measurements. The best lowest limit of detection (0.244 ± 0.092) μM was achieved with CTAB micelles while the best sensitivity (2.919 ± 0.054) M^{-1} corresponded to SDS.

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1 Introduction

Quercetin, 3,3',4',5,7-pentahydroxyflavone, see Fig. S1,† is commonly found in fruits and vegetables, although in particular, in apples and onions it occurs at highest levels.^{1,2} Due mainly to its anti-oxidant capacity,³ quercetin exhibits also other very important properties, namely: anti-inflammatory,⁴ anti-atherogenic and anti-carcinogenic.⁵ Furthermore, as caffeine, quercetin is also neuroactive and has GRAS (Generally Recognized As Safe) status. No side-effects have yet been noted in doses of a few grams a day in either humans or animals. In order to quantify quercetin several methods have been reported, namely: spectrophotometry in ethanolic media,^{1,6} using the coloured complex formed on reaction of quercetin with aluminium chloride in aqueous media,⁷ high performance liquid chromatography,^{1,6} electrochemical,^{2,8,9} photo-electrochemical.¹⁰ Flavonols, like the quercetin, containing an -OH group at position 5 (C-5), comprise a special class of non-fluorescent molecules in aqueous solution; since the intermolecular hydrogen bond between the polar groups of solute and H₂O exceeds various intramolecular hydrogen bonds, it displays

a large dihedral angle that makes it very difficult for ESICT (excited state intramolecular charge transfer) to occur.

Thus, the formation of a distorted excited state will become very active in water and no quercetin fluorescence emission is observed.¹¹ It has been shown that under particular conditions, namely: when quercetin binds to intracellular proteins¹² and in methanol media,¹³ the quercetin exhibits strong fluorescence, which was the property herein used for its simple, affordable, quantification. Notwithstanding, the spectrofluorometric quantification of quercetin in aqueous media has not been reported yet. In this work, it is shown that different surfactants namely: cetyl trimethylammonium bromide, CTAB,¹⁴ sodium dodecyl sulphate, SDS,¹⁵ and Triton X-100 (TX-100)¹⁶ can be used as promoters for quercetin fluorescence in aqueous media, allowing its spectrophotometric quantification, in this regard, Dwivedi *et al.*,¹⁷ have shown that the fluorescence properties of an amphiphilic perylene diimide, a bimolecular analogue of *l*-DOPA, can be modulated by means of host-guest interactions with CTAB. It is important to mention that the methods reported so far for quercetin quantification require expensive instrumentation *i.e.* HPTLC^{18,19} or sensing materials namely: multiwall carbon nanotubes² or carbon nanoparticles.²⁰

But most important, all of them need sufficiently long periods of time for quercetin determination, however as has been shown,³ the stability of quercetin and other flavonoids namely: morin, kaempferol, and myricetin are pH and time dependent thus, if these flavonoids are not stabilized during the measuring times their concentration may change. As will be shown in this work the use of surfactants overcomes this

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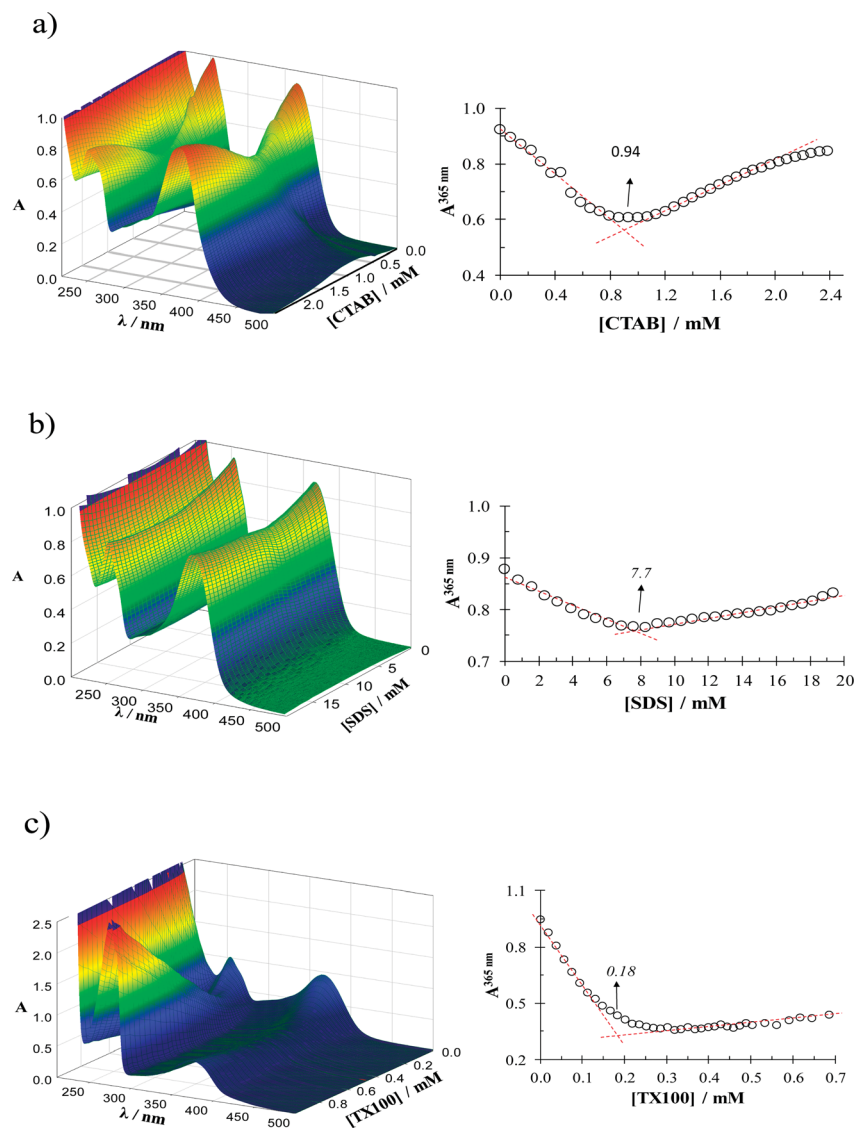


Fig. 1 UV-Vis spectra recorded in the system quercetin–H₂O–surfactant recorded as a function of the concentration of the different surfactants considered: (a) CTAB, (b) SDS and (c) TX-100. In all cases the temperature was $(28.0 \pm 0.1) ^\circ\text{C}$, the pH 7 ± 0.2 and the quercetin concentration was 5×10^{-5} M. The absorbance recorded at 365 nm as a function of the surfactant variation is also shown in each case.

difficulty since they confer enough stability to quercetin during its spectrofluorometric quantification time.

2 Experimental

2.1 Reagents and instruments

All the substances used were reactive grade. Quercetin, CTAB, SDS and Triton X-100 from Sigma-Aldrich, methanol to dissolve the quercetin was J.T. Baker brand, 18 M Ω cm resistivity deionized water from a Millipore Milli-Q equipment was used as solvent for all experiments. To dissolve the reagents, an Ultrasonic (8891 Cole-Parmer, set sonic min). All the experiments were conducted at $28.0 \pm 0.1 ^\circ\text{C}$ aided by a Grant W14 recirculator.

The solutions were protected from light incidence as much as possible. Concentrated solutions of SDS, CTAB and Triton X-100

Table 1 Comparison of the CMC values calculated in this work, from UV-Vis spectrophotometric measurements, with those reported in the literature for the surfactants considered in this work

CMC/mM			
CTAB	SDS	TX-100	
0.94 ± 0.03	7.7 ± 0.6	0.18 ± 0.3	This work
0.92	8.1	^a	21
0.92	8.3	^a	22
0.98	8.2	^a	23

^a CMC reported by the supplier is 0.2–0.9 mM. The uncertainty reported was estimated from at least 5 repetitions.

were prepared in water and a quercetin concentrated one in methanol. Aliquots were taken from each of these solutions to be added to the different systems to obtain the desired





Fig. 2 (a) Fluorescence emission spectra recorded in a 6×10^{-5} M quercetin aqueous solution containing different CTAB concentrations, within the 0 to 2.45 M range, using an excitation wavelength of 350 nm at $(28.0 \pm 0.1)^\circ\text{C}$ and the pH 7 ± 0.2 . (b) Plot $[I/I_0 - 1]^{-1}$ vs. $1/[\text{CTAB}] - \text{CMC}$ obtained from the fluorescence intensity, I , in (a) and CMC in Table 1.

concentrations. The pH of the solutions was measured with a Mettler Toledo pH-meter, with an error of ± 0.01 per pH unit.

2.2 UV-Vis spectrophotometry

Spectrophotometric measurements were carried out with a UV-Vis Lambda 20 spectrophotometer using quartz cells, one-centimetre optical path length, and two transparent faces and the WinLab software. For estimation of the surfactants critical micelle concentration, CMC, three systems were prepared with a volume of 5 mL of water and 5×10^{-5} M quercetin concentration each. To each of the systems, aliquots of the stock solution of one of the surfactants, were added and the respective absorption spectrum was measured after every addition.

2.3 Fluorescence spectrophotometry

Fluorescence measurements were performed with a Perkin Elmer LS50 Luminiscence Spectrometer using quartz cells, one-centimetre optical path length, with the four transparent faces and the WinLab software.

These measurements were used for both: evaluation of the binding constant of the supramolecular complex formed between quercetin and surfactants micelles and for quercetin quantification. Three systems were prepared with a volume of 5 mL of water and a 6×10^{-5} M quercetin concentration. To each of the systems, aliquots of the mother solution of one of the surfactants were added and the respective emission fluorescence spectrum was measured after every addition. For quercetin quantification, calibration plots were obtained by

Table 2 Binding constants between quercetin and micelles of different surfactants^a

Surfactants	Intercept	10^5 slope/M	R^2	$\log(K/M^{-1})$
CTAB	0.1376 ± 0.0061	18.73 ± 0.35	0.9975	2.87 ± 0.02
SDS	0.2823 ± 0.0087	47.08 ± 2.86	0.9819	2.78 ± 0.04
TX-100	0.1262 ± 0.0112	19.79 ± 0.55	0.9947	2.80 ± 0.04

^a The uncertainty reported was estimated from at least 5 repetitions.

adding quercetin (within the 0 to 7.5 mM range) to an aqueous solution having a fixed surfactant concentration (higher than the respective CMC value) and the respective emission fluorescence spectrum was recorded at 350 nm excitation wavelength, for all experiments.

3 Results and discussion

3.1 CMC calculations

Fig. 1 shows a family of UV-Vis spectra recorded in the system quercetin–H₂O–surfactant for a fixed quercetin concentration and different surfactant concentrations. From absorbance variation at 365 nm it is possible to estimate the CMC for each case, see Table 1.^{14–16}

3.2 Quercetin stability

In the absence of surfactant micelles, see Fig. S2,† quercetin is notably unstable, the absorbance at 365 nm decreases linearly with time, referred to its value at $t = 0$, however when the surfactant concentration is present higher than the respective CMC value quercetin becomes stable in time.

3.3 Thermodynamic binding constant for the quercetin–surfactants micelles supramolecular complex

Fig. 2a shows a family of fluorescence emission spectra of quercetin in aqueous media with different CTAB concentrations.

From this figure it becomes plain that in the absence of CTAB molecules, quercetin depicts no fluorescence. Notwithstanding, as the CTAB concentration in the system is increased, a well-defined



Fig. 3 Distribution diagram of the quercetin, Q, species as a function of $p\text{CTAB} = -\log[\text{CTAB}]$.

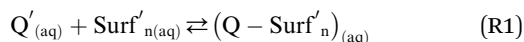




Fig. 4 Fluorescence emission spectra, using an excitation wavelength of 350 nm, recorded in (a) 3.768 ± 0.009 mM CTAB (b) 36.893 ± 0.015 mM SDS and (c) 2.827 ± 0.012 mM TX-100 aqueous solutions added with different quercetin concentrations at $(28.0 \pm 0.1)^\circ\text{C}$ and the pH 7 ± 0.2 . The respective calibration plots are also shown at the right of each spectra family. In these plots the fluorescence intensity of the corresponding surfactant aqueous solution without quercetin was subtracted. The error bars were estimated from at least 5 repetitions.

fluorescence intensity peak is formed at 540 nm, furthermore, quite a similar behaviour was noted when SDS or TX-100 was added, see Fig. S3 in the ESI† of this work.

The supramolecular complex, $Q - \text{Surf}'_n$, formed between quercetin, Q, molecules and the surfactant, Surf, micelles, see Reacc. (1), restricts quercetin mobility and promotes in this way its fluorescence.



where Q' and Surf' represent generalised species. As can be noted from the distribution diagram shown in Fig. S4,† constructed with the $\text{p}K_a$ values reported by Álvarez-Diduk *et al.*,³ at pH 7 the quercetin predominant species correspond to a mono-anionic, Q^- , form while for the surfactant micelles, cations (CTB^+), anions (DS^-) or neutral molecules (TX100) would form depending on the surfactant nature.

According with Guardia *et al.*,²⁴ from fluorescence emission spectra it is possible to determine the binding constants, K , between micelles and chemical systems using eqn (1).

$$\left[\frac{I}{I_0} - 1 \right]^{-1} = \left[\frac{I_M}{I_0} - 1 \right]^{-1} \left[1 + \frac{1}{\gamma K ([\text{surfactant}] - \text{CMC})} \right] \quad (1)$$

where I is the fluorescence intensity of the system at each concentration of the surfactant, I_0 is the fluorescence of the system in the absence of surfactant, and I_M is the maximum fluorescence obtained; γ is the quotient of the molar extinction coefficient, at the excitation wavelength, in the presence (ϵ_p) and in the absence (ϵ_0) of the surfactant.

When the extinction coefficient does not vary in the micellar media $\gamma = 1$. The regression between $[I/I_0 - 1]^{-1}$ and $[[\text{surfactant}] - \text{CMC}]$, gives a straight line and the value of the binding constant between the system and the micelles can be obtained from the ratio of the intercept and the slope of this line, whenever the micelles do not modify the extinction coefficient at the excitation wavelength. From Fig. 2b it can be noted, in the case when CTAB was the surfactant added, that the model associated to eqn (1) fits quite well the experimental data (similar behaviour has been noted in the case of SDS and TX100), see Fig. S3.†



Table 3 Analytical features depicted for the spectrofluorometric quantification of quercetin in aqueous media using different surfactants at fixed concentrations, obtained from the calibration plots shown in Fig. 4^a

Surfactant	[Surfactant]/mM	LOD/ μM	LOQ/ μM	Sensitivity/ μM^{-1}
CTAB	3.768 \pm 0.009	0.244 \pm 0.092	0.81 \pm 0.09	0.718 \pm 0.006
SDS	36.893 \pm 0.015	0.48 \pm 0.19	1.60 \pm 0.18	2.919 \pm 0.054
TX-100	2.827 \pm 0.012	0.56 \pm 0.22	1.88 \pm 0.21	1.346 \pm 0.030

^a The uncertainty reported was estimated from at least 5 repetitions.

Table 2 summarizes the values of binding constant for the quercetin and micelles of the different surfactants considered.

It is important to note that recently, from similar spectroscopic measurements, Szymczyk and Taraba,²⁵ found that $\log(K/M^{-1}) = 3.16$ for the interaction between Triton X-114 and Quercetin (2×10^{-5} M at 20 °C) while Liu and Guo²⁶ reported 3.45 for quercetin (5×10^{-5} M at 25 °C) with Triton X-100 and Singh *et al.*²⁷, 3.25 for sodium bis(2-ethylhexyl)sulfosuccinate and quercetin (3×10^{-5} M at 25 °C).

With the respective K values reported in Table 2 it is possible to construct the species distribution diagram for quercetin and the complex formed in aqueous solution as a function of $p\text{Surfactant} = -\log[\text{surfactant}]$.²⁸ Fig. 3 depicts such diagram in the case of CTAB; similar diagrams, for SDS and TX100 can be found in Fig. S5.†

3.4 Quercetin quantification

Once it was shown that the interaction between quercetin and surfactant promotes the fluorescence of the former in aqueous media, due to the supramolecular interaction with micelles of the amphiphilic molecules, the spectrofluorometric quantification of quercetin in aqueous media was carried out.

In order to do this, an aqueous solution of surfactants with a fixed concentration, 4 times higher than the respective CMC,

was used to ensure the presence of micelles in each case and that the complex quercetin-surfactant predominates in the system, see Fig. 3 and S5.† Fig. 4 shows the fluorescence emission spectra recorded in aqueous surfactant solutions, at a fixed concentration, added with different quercetin concentrations.

It is possible to note in all cases the formation of a fluorescence peak, of which the intensity varies linearly with the quercetin concentration. From these calibration plots, it was possible to calculate the analytical features namely: the limits of detection (LOD) and quantification (LOQ) and the sensitivity in each case, see Table 3.

Table 4 depicts a comparison of the analytical method developed in this work with others reported, particularly regarding quercetin quantification, from which it results plain that the analytical performance of our methodology is plainly comparable or even better than that of others more sophisticated ones.

3.5 Interferences

In order to broaden the usefulness and applicability of the spectrofluorometric quantification of quercetin, experiments were run on other molecules that could cause competition with quercetin in the micelles, namely: morin, rutin, catechin and

Table 4 Comparison of the analytical performance of different methodologies towards quercetin quantification^a

Media	Methodology	10 ⁸ LOD/M	10 ⁸ LOQ/M	10 ⁶ sensitivity/M ⁻¹	Ref.
NH ₃ -NH ₄ Cl buffer solution (0.10 M, pH 9.47)	Fluorescence ^b	9.88	N.R.	9.39	20
Phosphate buffer (0.1 M pH 7.40)	Adsorptive stripping with square wave voltammetry ^c	8.9	N.R.	2.5	2
	Adsorptive stripping with square wave voltammetry ^d	0.75	N.R.	8.7	
	Amperometry ^e	170	N.R.	150	
	Amperometry ^c	140	N.R.	760	
Methanol	Amperometry ^d	20	N.R.	330	
	HPTLC	4700	9500	—	18
	HPTLC	590	1480	—	19
Water (pH 7)	Fluorescence, in the presence of CTAB micelles	24.4	81	0.72	This work
	Fluorescence, in the presence of SDS micelles	47.9	160	2.92	
	Fluorescence, in the presence of TX100 micelles	56.4	188	1.35	

^a N.R. Not reported. ^b Using carbon nanoparticles as fluorescence probe. ^c Using GCE/PEI-CNT. ^d GCE/PAA-CNT. ^e GCE, where: GCE (glassy carbon electrodes), CNT (multiwall carbon nanotubes), PEI (polyethylenimine) and PAA (poly(acrylic acid)).



epicatechin, see Fig. S6,† and thus may provoke interference. It was found that regardless of the nature of the surfactant, when adding morin to an aqueous solution with fixed concentrations of quercetin and surfactant, see Fig. S7,† the fluorescence intensity increases linearly with the morin concentration, contrary, when rutin was added, instead of morin, the fluorescence intensity decreases linearly with its concentration, see Fig. S8,† however, when catechin, see Fig. S9,† or epicatechin, see Fig. S10,† were added the fluorescence intensity remains practically constant. From these results, it can be concluded that while morin is a major interferent, rutin slightly interferes and catechin and epicatechin do not. Furthermore, it was also found that quercetin can be quantified spectrofluorometrically in the presence of a higher concentration of ascorbic acid (see Fig. S11†).

4 Conclusions

It has been shown that micelles of surfactants (anionic, cationic or neutral) in neutral aqueous media interact with quercetin forming stable supramolecular complexes that promote both: time stability and quercetin fluorescence, which allow its quantification by means of spectrofluorometry even in the presence of similar molecules like catechin and epicatechin or higher ascorbic acid concentrations.

Conflicts of interest

There are no conflicts of interest to declare.

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