

# 2-Methyl-4-oxo-*N*-(4-oxo-2-phenyl substituted-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamides—A New Range of Fluorescent Whiteners: Synthesis and Photophysical Characterization

Vikas S. Patil · Vikas S. Padalkar · Nagaiyan Sekar

Received: 14 October 2013 / Accepted: 7 April 2014 / Published online: 23 May 2014  
© Springer Science+Business Media New York 2014

**Abstract** Fluorescent quinazolinones were synthesized from ethyl 2-methyl-4-oxo-3,4-dihydroquinazoline-5-carboxylate intermediate. The photophysical properties of the compounds were evaluated in DMF solvent. The experimental absorption and emission of the compounds were compared with the vertical excitation and emission obtained Density Functional Theory (DFT) and Time Dependent Density Functional Theory (TD-DFT) computation. Application of the fluorescent compounds as a fluorescent brightening agent was tested on polyester fiber. Changes in the electronic transition, energy levels, and orbital diagrams of quinazolin-4(3*H*)-one analogues were investigated using the DFT computations and were correlated with the experimental spectral data. The experimental absorption and emission wavelengths are in good agreement with those predicted using the DFT and TD-DFT.

**Keywords** Quinazoline · DFT · TD-DFT · Absorption · Emission · Photophysical properties

## Introduction

A large number of alkaloids contain quinazolinone nucleus and they are known to have biological activities [1]. In addition to pharmaceutical applications, quinazolinone derivatives

are widely used as coloring materials in dyestuff industry [2, 3]. The azo colorants based on quinazolinone have industrial importance [3, 4]. The substituted quinazolinone are reported as fluorescent compounds for different applications [5–10]. Quinazolinone derivatives are known to be fluorescent brightening agents [2, 3, 8]. Optical whiteners are characterized by a strong absorption below 400 nm with well separated intense emission beyond 400 nm [11]. The optical bleaching agents are chemicals that increase the whiteness when applied on the natural and synthetic textile materials [12]. The surface of textile materials therefore appear whiter to the human eye [13]. Fluorescent brighteners are substances which normally have a system of conjugated double bonds and electron-donating groups to show high fluorescence [14, 15].

Fluorescent brightening agents (FBAs) are used in textile industry to enhance the whiteness and brightness of textile substrates; they also significantly increase the UV-blocking properties of the medium to which they are applied [16, 17]. They are also used in pulp and paper industries for the improvement of brightness [18]. In addition to the above mentioned applications they are also used in pH chemosensing materials [19], chemosensors [20], photo-induced electron transfer sensors [21], light emitting diodes [22], biological staining [23] and polyurethane fluorescent brightener dispersions [24].

Fluorescent whitening agents are derivatives of triazine [25], stilbene [26], pyrazole [27], naphthalimide [28], quinoline [29], iso-quinoline [30], quinoxaline [31], oxadiazole [32], triazole [33], benzoxazole [34], benzotriazole [35] and tinopol-CBX [36] based compounds. The fluorescent quinazolinone are also reported as fluorescent brightening agents for polyester fibers [2, 3]. A fluorescent brightener should present a high quality of whiteness and fastness. However, as the fluorescent brightener is exposed to the sunlight, its whiteness is diminished [3]. This happens because the

V. S. Patil · V. S. Padalkar · N. Sekar (✉)  
Tinctorial Chemistry Group, Institute of Chemical Technology  
(Formerly UDCT), N. P. Marg, Matunga, Mumbai 400  
019, Maharashtra, India  
e-mail: n.sekar@ictmumbai.edu.in

V. S. Patil  
e-mail: vikasudct@gmail.com

V. S. Padalkar  
e-mail: vikaspadalkar@gmail.com

chemical structure of fluorescent brightener is destroyed by the autoxidation from free radical generated from the sunlight [3]. Fluorescent whiteners with enhanced chemical, thermal and photostability are needed today for high performance textile substrates.

Reported quinazolinone derivatives are 2 and 6-substituted in which the electron donor is at the 2-position and the acceptor at the 6-position [2, 3]. A study of the literature reveals that, electron donor substitution at the 2-position along with an acceptor at the 6-position gives red shifted absorption and hence red shifted emission, because such a substitution pattern leads to an effective donor acceptor chromophore [9]. To achieve blue shifted emission it is necessary to reduce the donor strength.

In this paper, attempts have been made to synthesize fluorescent quinazolinone thiazolidines. The synthesized quinazolinone derivatives contain electron donor at the 2-position and an acceptor at the 5-position. The photophysical properties like absorption, emission and quantum yields of the compounds were recorded in the solvent dimethylformamide (DMF). Fluorescent compounds were used for whitening polyester fiber for optical brightening study. The changes in the electronic transition, energy levels, and orbital diagrams of the quinazolinone derivatives were studied by TD-DFT computation. The computational results were correlated with the experimental photophysical properties.

## Methodology

### Materials and Methods

All the commercial reagents and solvents were purchased from S. D. Fine Chemicals Pvt. Ltd. and they were used without purification and all the solvents were of spectroscopic grade. The absorption spectra of the dyes were recorded on a Spectronic Genesys 2 UV-Visible spectrophotometer, and emission spectra were recorded on Varian Cary Eclipse fluorescence spectrophotometer using freshly prepared solutions in solvents of different polarities at the concentration of  $1 \times 10^{-6}$  mol L<sup>-1</sup>. The excitation wavelength used for the fluorescence measurements was absorption maxima of the compounds in respective solvents. The FT-IR spectra were recorded on a Perkin-Elmer Spectrum 100 FT-IR Spectrometer. <sup>1</sup>H NMR spectra were recorded on VXR 400 MHz instrument using TMS as an internal standard. Mass spectra were recorded on Finnigan mass spectrometer.

### Computational Methods

The ground state geometries of the compounds **7a–7e** in their C<sub>1</sub> symmetry were optimized using the tight criteria in the gas

phase using (DFT) [37]. The functional used in this study was B3LYP [38, 39]. The basis set used for all atoms was 6-31G(d) [40–42]. The vibrational frequencies of the optimized structures were computed using the same method to verify that the optimized structures correspond to local minima on the potential energy surface. The vertical excitation energies at the ground-state equilibrium geometries were calculated with TD-DFT [43–45]. The low-lying first singlet excited state (S<sub>1</sub>) of each tautomer was relaxed using the TD-DFT to obtain its minimum energy geometry. The difference between the energies of the optimized geometries in the first singlet excited state and the ground state was used in computing the emissions [46–52]. All electronic structure computations were carried out using the Gaussian 09 program [53].

### Relative Quantum Yield Calculations

The quantum yields of compounds **7a–7e** in DMF were evaluated. Anthracene was used as the standard. Quantum yields were calculated using the comparative method [54, 55]. The absorption and emission characteristics of the standards and for the compounds in polar solvents were measured at different concentrations (1, 2, 3, 4, and 5 ppm level). The emission intensity values were plotted against absorbance values and linear plots were obtained. The gradients were calculated for the compounds in each solvent and for the standards. All the measurements were done by keeping the parameters such as solvent and slit width constant. The relative quantum yields of the synthesized compounds in different solvents were calculated by using the Eq. 1 [54, 55].

$$\phi_x = \Phi_{st} \times \frac{Grad_x}{Grad_{st}} \times \frac{\eta_x^2}{\eta_{st}^2} \quad (1)$$

Where:

$\Phi_x$	Quantum yield of compound
$\Phi_{st}$	Quantum yield of standard sample
$Grad_x$	Gradient of compound
$Grad_{st}$	Gradient of standard sample
$\eta_x$	Refractive index of solvent used for synthesized compound
$\eta_{st}$	Refractive index of solvent used for standard sample

### General Procedure of Dyeing

Dyeing of polyester fabric was carried out using high temperature high pressure method in Rossari Labtech Flexi Dyer dyeing machine at a material to liquor ratio of 1:20. 2 %

Fluorescent compounds were used for the dyeing (calculated on weight of the fabric). All the synthesized fluorescent compounds are having less solubility in water. Initially the compounds were dissolved in 5 mL DMF and diluted with 15 mL buffered solution of pH 5 made by using sodium acetate and acetic acid in water. The mixture was ultrasonicated for 15 min to obtain a fine dispersion. Metamol was used as a dispersant. The polyester fabric was dyed using the above solution and metamol as the dispersing agent. The dye bath temperature was raised at a rate of 3 °C min<sup>-1</sup> to 130 °C, maintained at this temperature for 60 min, and rapidly cooled to room temperature as shown in Fig. 1. The dyed fabrics were rinsed with cold water and allowed to dry in the open air.

## Experimental

### Synthesis Details

#### *Ethyl 2-methyl-4-oxo-3,4-dihydroquinazoline-5-carboxylate 4*

The intermediate **4** was prepared according to the known procedure [56].

#### *2-Methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide 5*

Ethyl-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carboxylate (5 g) on reaction with hydrazine hydrate (15 mL) for 2 h under reflux condition gave 2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5**.

Yield: 89 %, Melting point => 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.20 (s, 3H), 5.26 (s, 2H), 7.72 (d, 1H, *J*=22.00 Hz), 7.79 (d, 1H, *J*=6.7 Hz), 7.82 (dd, 1H, *J*=22.00, 6.7 Hz).

**FT-IR (KBr)**: 3165, 3103, 2981, 2917, 2871, 1665, 1607, 1562, 1478, 1383, 1325, 1261, 1193, 1158, 1103, 1041, 843, 788, 721, 663 cm<sup>-1</sup>.

#### *2-Methyl-4-oxo-N'-[(Z)-phenylmethylidene]-3,4-dihydroquinazoline-5-carbohydrazide 6a*

2-Methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** (1 g, 4.58 mmol) was reacted with benzaldehyde (0.46 g, 4.58 mmol) in methanol in presences of catalytic amount of sulfuric acid at reflux condition for 2 h to give 2-methyl-4-oxo-*N'*-[(*Z*)-phenylmethylidene]-3,4-dihydroquinazoline-5-carbohydrazide **6a**.

Yield: 67 %, Melting point => 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.36 (s, 3H), 2.50 (s, 1H), 3.16 (s, 1H), 7.45 (d, 2H, *J*=7.7 Hz), 7.78 (d, 1H, *J*=8.5 Hz), 7.86 (m, 5H, *J*=8.0, 11.09 Hz), 12.01 (s, 1H)

**FT-IR (KBr)**: 3011, 1678, 1613, 1567, 987, 673 cm<sup>-1</sup>.

**Mass**: *m/z* 307 (M<sup>+</sup>).

#### *2-Methyl-N'-[(Z)-(4-nitrophenyl)methylidene]-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide 6b*

2-Methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** (1 g, 4.58 mmol) was reacted with *p*-nitro benzaldehyde (0.69 g, 4.58 mmol) in methanol in the presences of catalytic amount of sulfuric acid at reflux condition for 6 h to give 2-methyl-*N'*-[(*Z*)-(4-nitrophenyl) methylidene]-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6b**.

Yield: 87 %, Melting point => 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.23 (s, 3H), 7.34 (d, 2H, *J*=7.7 Hz), 7.62 (d, 2H, *J*=7.7 Hz), 7.75 (dd, 2H, *J*=7.7, 9.1 Hz), 8.15 (d, 2H, *J*=13.2 Hz), 8.3 (d, 1H, *J*=13.2 Hz), 11.75 (s, 1H).

**FT-IR (KBr)**: 2965, 2900, 2644, 1691, 1665, 1584, 1478, 1406, 1309, 1264, 908, 740, 711. cm<sup>-1</sup>.

**Mass**: *m/z* 349 (M-2).

#### *N'-(anthracen-9-ylmethylidene)*

#### *-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide 6c*

2-Methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** (1 g, 4.58 mmol) was reacted with anthracene 9-carbaldehyde (0.94 g, 4.58 mmol) in methanol in the presences of catalytic amount of sulfuric acid at reflux condition for 4 h to give *N'*-(anthracen-9-ylmethylidene)-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6c**.

Yield: 83 %, Melting point => 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.24 (s, 3H), 7.34 (d, 1H, *J*=7.7 Hz), 7.59 (d, 1H, *J*=7.0 Hz), 7.62 (dd, 4H, *J*=8.1, 7.9 Hz), 7.76 (d, 4H, *J*=8.1, 9.7 Hz), 7.73 (s, 1H), 9.25 (s, 1H), 11.72 (s, 1H).

**FT-IR (KBr)**: 2991, 1778, 1623, 1637, 1557, 967, 656 cm<sup>-1</sup>.

**Mass**: *m/z* 407 (M<sup>+</sup>).

#### *2-Methyl-N'-[(Z)-(4-N, N-diethyl, 2-hydroxy)methylidene]-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide 6d*

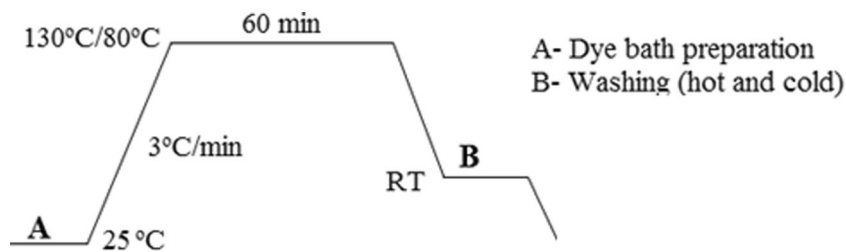
2-Methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** (1 g, 4.58 mmol) was reacted with 4-(diethylamino)-2-hydroxybenzaldehyde (0.88 g, 4.58 mmol) in methanol in the presences of catalytic amount of sulfuric acid at reflux condition for 7 h to give 2-methyl-*N'*-[(*Z*)-(4-N, N-diethyl, 2-hydroxy)methylidene]-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6d**.

Yield: 79 %, Melting point => 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.21 (s, 3H), 2.48 (t, 3H, *J*=7.0 Hz), 3.35 (d, 2H, *J*=7.0 Hz), 7.30 (d, 1H, *J*=8.1 Hz), 7.60 (d, 1H, *J*=7.7 Hz), 7.74 (dd, 1H, *J*=9.8, 7.7 Hz), 8.81 (m, 3H, *J*=9.8, 6.9 Hz), 12.21 (s, 1H), 10.39 (s, 1H),

**FT-IR (KBr)**: 3027, 1781, 1622, 1645, 1534, 2511, 978, 656 cm<sup>-1</sup>.

**Fig. 1** Dyeing profile of polyester used



**Mass:**  $m/z$  395 ( $M^+$ ).

*N'*-[(*Z*)-(1-*Tert*-butyl-4-chloro-1*H*-pyrazol-5-yl)methylidene]-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6e**

2-Methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** (1 g, 4.58 mmol) was reacted with 2-*tert*-butyl-5-chloro-1*H*-imidazole-4-carbaldehyde (0.85 g, 4.58 mmol) in methanol in the presences of catalytic amount of sulfuric acid at reflux condition for 4 h to give *N'*-[(*Z*)-(1-*tert*-butyl-4-chloro-1*H*-pyrazol-5-yl)methylidene]-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6e**.

Yield: 85 %, Melting point => 300 °C.

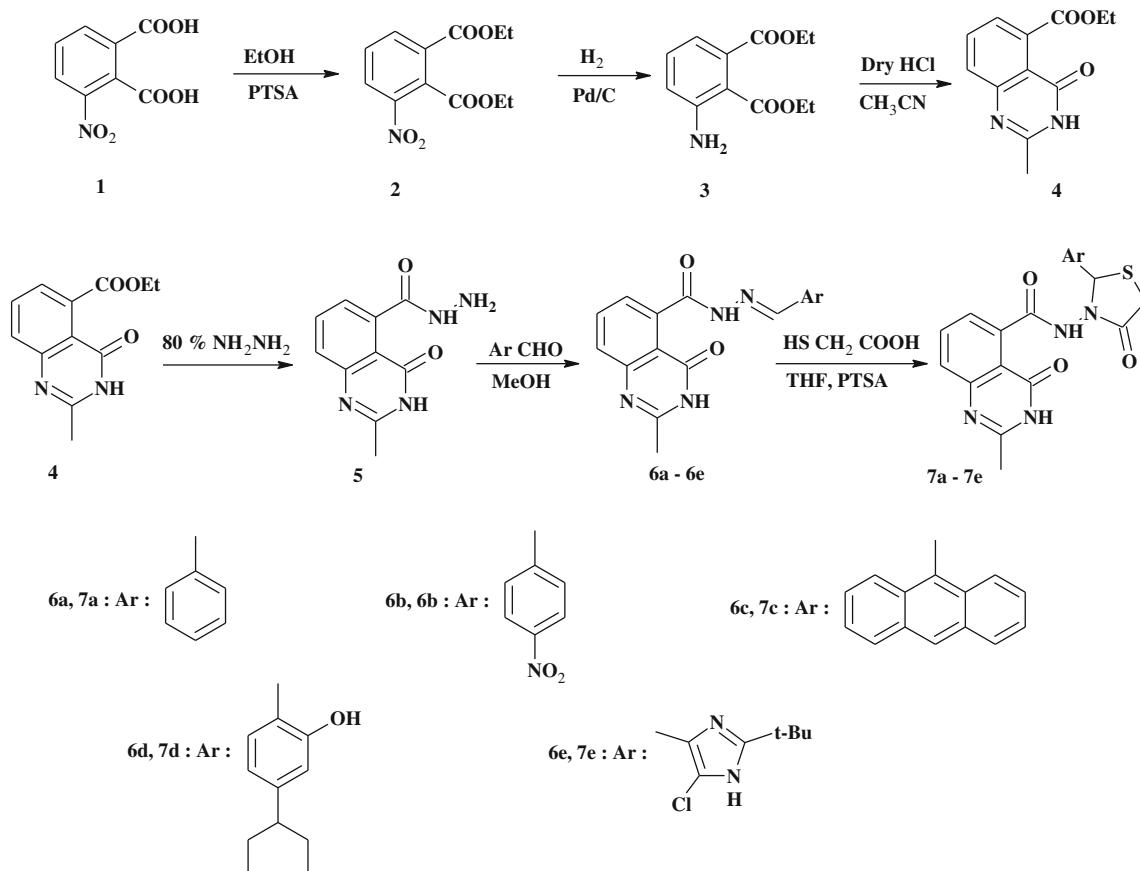
<sup>1</sup>H-NMR ( $CD_3)_2SO$ :  $\delta$  2.5 (s, 3H), 2.65 (s, 9H), 7.30 (d, 1H,  $J=8.1$  Hz), 7.60 (d, 1H,  $J=7.7$  Hz), 7.74 (dd, 1H,  $J=8.1, 7.7$  Hz), 11.72 (s, 1H).

**FT-IR (KBr):** 3014, 1771, 1643, 1655, 1521, 1456, 1378, 1218, 978, 698, 651  $cm^{-1}$ .

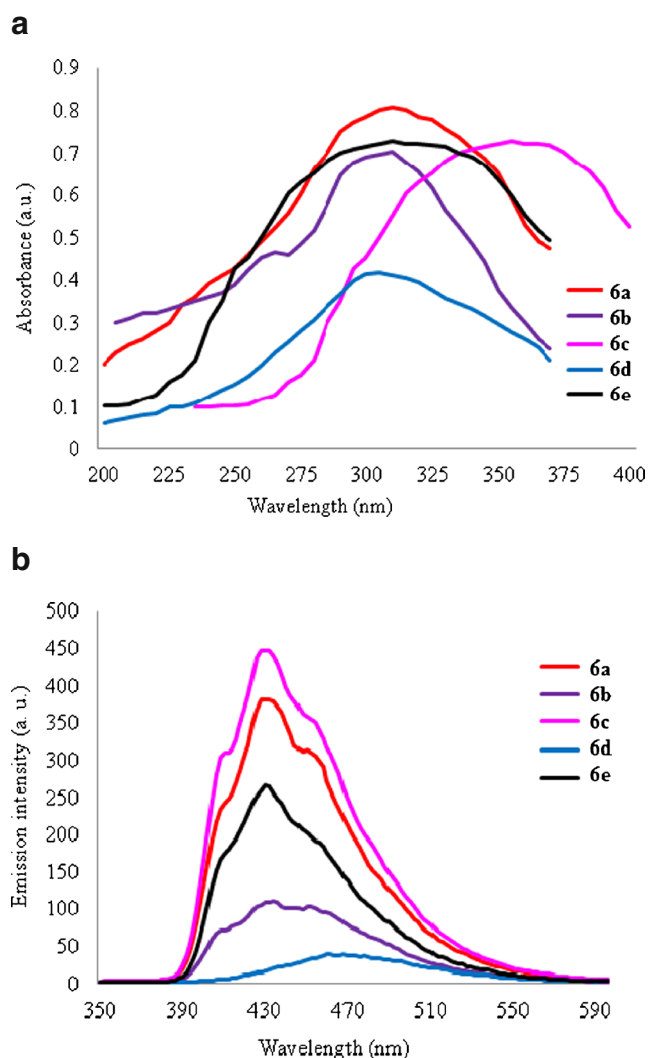
**Mass:**  $m/z$  388 ( $M^+$ ).

2-Methyl-4-oxo-*N*-(4-oxo-2-phenyl-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide **7a**

2-Methyl-4-oxo-*N'*-[(*Z*)-phenylmethylidene]-3,4-dihydroquinazoline-5-carbohydrazide **6a** (1.42 mmol) was reacted with thioglyconic acid (1.42 mmol) in tetrahydrofuran (THF) at reflux condition for 6 h to give 2-methyl-4-oxo-*N*-(4-



**Scheme 1** Synthesis of 2-methyl-4-oxo-*N*-(4-oxo-2-phenyl substituted-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamides



**Fig. 2** Absorption and emission spectra of compounds **6a–6e**. **a** Absorption spectra of compounds **6a–6e**. **b** Emission spectra of compounds **6a–6e**

oxo-2-phenyl-1,3-thiazolidin-3-yl]-3,4-dihydroquinazoline-5-carboxamide **7a**.

Yield: 69 %, Melting point => 300 °C.

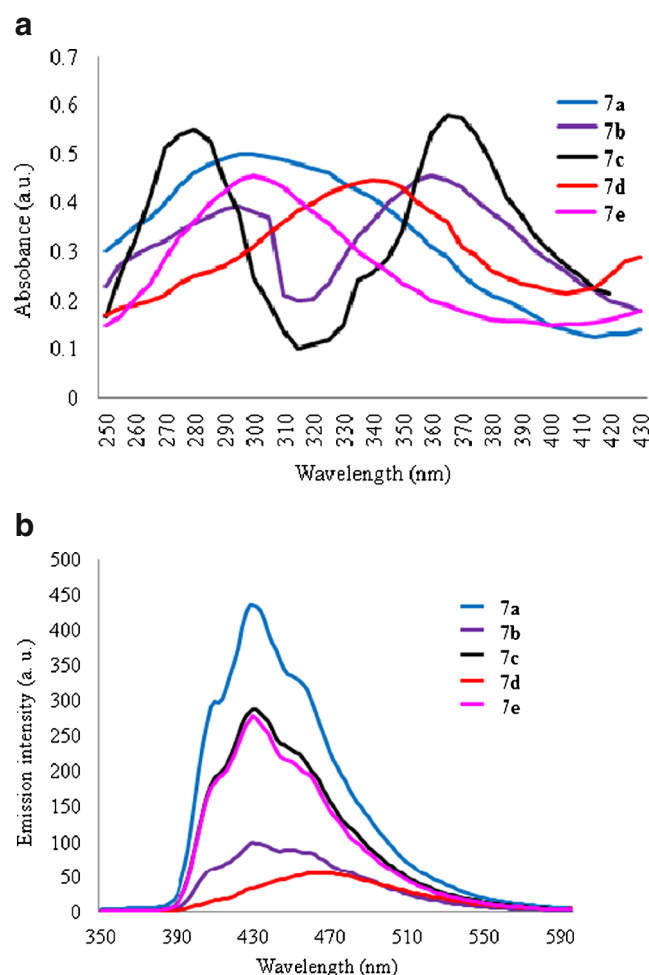
**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.25 (s, 3H), 2.5 (s, 2H), 3.45 (s, 1H), 7.33 (d, 1H, *J*=7.7 Hz), 7.63 (d, 1H, *J*=7.7 Hz), 7.74 (dd, 1H, *J*=8.1, 7.3 Hz), 7.80–7.88 (m, 3H, *J*=8, 8.1, 7.7 Hz), 8.15 (d, 1H, *J*=8.1 Hz), 8.36 (d, 1H, *J*=8 Hz), 9.71 (s, 1H), 11.78 (s, 1H)

**FT-IR (KBr)**: 2978, 1789, 1768, 1628, 1667, 1581, 1458, 1335, 1267, 1211, 970, 689, 651 cm<sup>-1</sup>.

**Mass**: *m/z* 388 (M<sup>+</sup>).

*2-Methyl-N-[2-(4-nitrophenyl)-4-oxo-1,3-thiazolidin-3-yl]-4-oxo-3,4-dihydroquinazoline-5-carboxamide 7b*

*2-Methyl-N'-[(Z)-(4-nitrophenyl)methylidene]-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide 6b* (1.42 mmol) was reacted with thioglycolic acid (1.42 mmol) in THF at reflux



**Fig. 3** Absorption and emission spectra of compounds **7a–7e**. **a** Absorption spectra of compounds **7a–7e** in DMF. **b** Emission spectra of compounds **7a–7e** in DMF

condition for 12 h to give 2-methyl-*N*-[2-(4-nitrophenyl)-4-oxo-1,3-thiazolidin-3-yl]-4-oxo-3,4-dihydroquinazoline-5-carboxamide **7b**.

Yield: 61 %, Melting point => 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**: δ 2.28 (s, 1H), 2.50 (s, 1H), 3.65 (s, 1H), 7.37 (d, 1H, *J*=7.7 Hz), 7.69 (d, 1H, *J*=7.7 Hz), 7.69 (d, 1H, *J*=7.7 Hz), 7.76 (dd, 1H, *J*=8.1, 7.3 Hz), 8.16 (d, 2H, *J*=8.8 Hz), 8.37 (d, 2H, *J*=8.8 Hz), 9.71 (s, 1H), 11.85 (s, 1H)

**FT-IR (KBr)**: 2981, 2823, 2761, 1629, 1558, 1478, 1419, 1338, 1167, 1089, 1044, 876, 779, 698 cm<sup>-1</sup>.

**Mass**: *m/z* 427 (M<sup>+</sup>).

*2-Methyl-4-oxo-N-(4-oxo-2-anthracene-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide 7c*

*N'*-(Anthracen-9-ylmethylidene)-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6c** (1.42 mmol) was reacted with thioglycolic acid (1.42 mmol) in THF at reflux condition for 14 h to give 2-methyl-4-oxo-*N*-(4-oxo-2-

**Table 1** Observed UV-visible absorption, emission and computed vertical excitation and emission of compounds **7a–7e** in DMF

Comp	$\lambda_{max}^{Exp}(nm)$	TD-DFT vertical excitation		$f$	% D	$\lambda_{Em}^{Exp}(nm)$	$\lambda_{Em}^{DFT}$	% D	<sup>a</sup> Stokes shift $\Delta\lambda$ (cm <sup>-1</sup> )	<sup>b</sup> $\Phi$
		nm	eV							
7a	300	307	4.029	0.120	2	431	345	19	10131	0.002
7b	341	362	3.417	0.003	6	432	426	1	6177	0.017
7c	354	402	3.083	0.010	13	464	440	4	6696	0.021
7d	342	383	3.233	0.0003	11	431	473	9	6037	0.102
7e	312	303	4.083	0.007	2	428	470	9	8686	0.253

<sup>a</sup> Stokes shift in cm<sup>-1</sup><sup>b</sup> Quantum yieldAnalyses were carried out at room temperature (25 °C); experimentally observed  $\lambda_{max}$ 

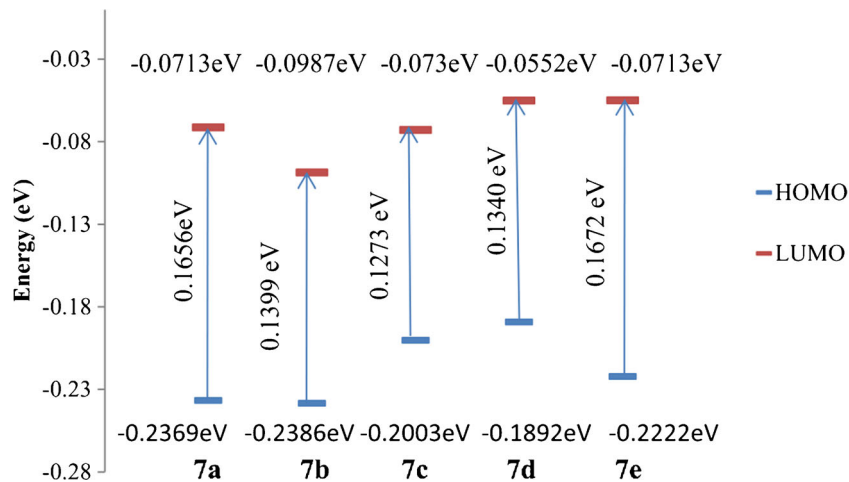
(% D) % Deviation between vertical excitation and experimental absorption and experimental emission and computed (TD-DFT) emission

 $f$ : Oscillator strengthanthracene-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide **7c**.

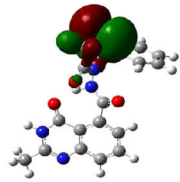
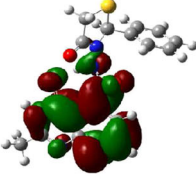
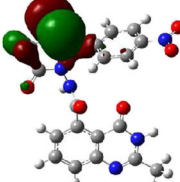
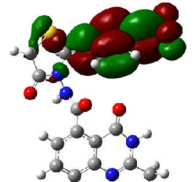
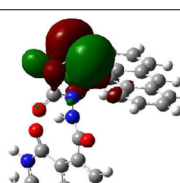
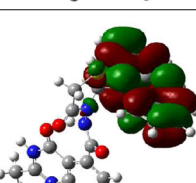
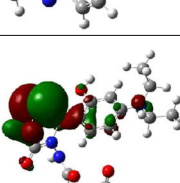
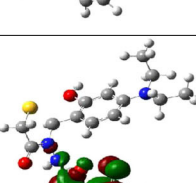
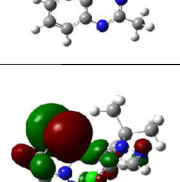
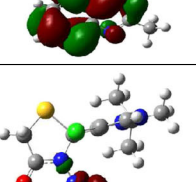
Yield: 71 %, Melting point =&gt; 300 °C.

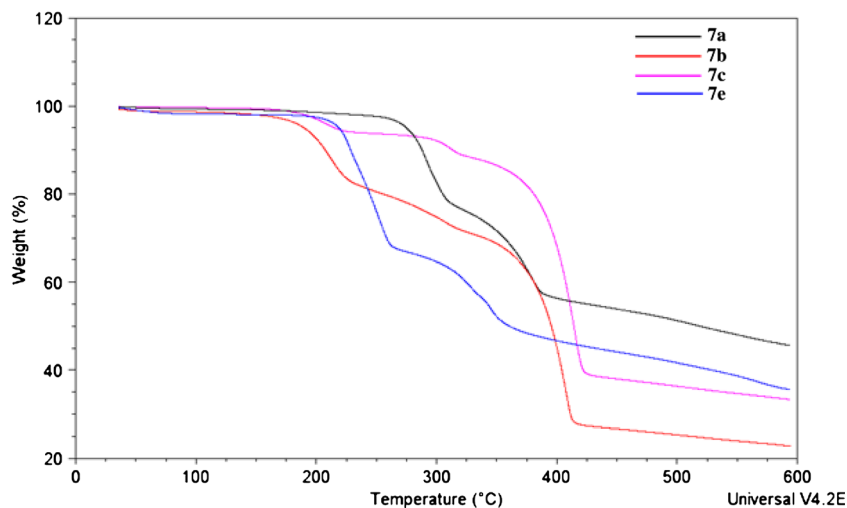
**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**:  $\delta$  2.28 (s, 3H), 2.50 (s, 2H), 3.65 (s, 1H), 7.37 (d, 1H,  $J=7.7$  Hz), 7.67 (d, 1H,  $J=7.7$  Hz), 7.70–7.82 (m, 12H,  $J=18.6, 7.7, 8.0$  Hz), 11.85 (s, 1H).**FT-IR (KBr)**: 3018, 1801, 1757, 1625, 1681, 1580, 1448, 1345, 1255, 1210, 978, 667, 644 cm<sup>-1</sup>.**Mass**:  $m/z$  488 (M<sup>+</sup>).2-Methyl-4-oxo-*N*-(4-oxo-2-(4-(diethylamino)2-phenol-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide **7d**2-Methyl-*N'*-[(*Z*)-(4-*N*, *N*-diethyl, 2-hydroxy)methylidene]-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6d** (1.42 mmol) was reacted with thioglycolic acid (1.42 mmol) in THF at reflux condition for 10 h to give 2-methyl-4-oxo-*N*-(4-oxo-2-(4-(diethylamino)2-phenol-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide **7b**.

Yield: 69 %, Melting point =&gt; 300 °C.

**<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO**:  $\delta$  2.43 (s, 3H), 2.51 (6H, t,  $J=14.3$  Hz), 2.64 (2H, s), 3.36 (1H, s), 3.92 (q, 4H,  $J=14.3$  Hz), 6.92 (s, 1H), 7.63 (d 1H,  $J=7.7$  Hz), 7.72 (d, 1H,  $J=8.1$ ), 7.92 (dd, 1H,  $J=7.7, 8.1$  Hz), 8.03 (d, 1H,  $J=9.3$  Hz), 8.23 (d, 1H,  $J=7.9$  Hz), 8.88 (dd, 1H,  $J=9.3, 7.9$  Hz), 10.39 (s, 1H), 12.25 (s, 1H), 12.51 (s, 1H).**FT-IR (KBr)**: 3028, 1777, 1759, 1631, 1667, 1568, 1440, 1320, 1245, 1264, 957, 678, 621 cm<sup>-1</sup>.**Mass**:  $m/z$  469 (M<sup>+</sup>).2-Methyl-4-oxo-*N*-(4-oxo-2-(1-*tert*-butyl-4-chloro-1*H*-pyrazole-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide **7e***N'*-[(*Z*)-(1-*tert*-butyl-4-chloro-1*H*-pyrazol-5-yl)methylidene]-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **6e****Graph 1** HOMO-LUMO energies of compounds **7a–7e** in different solvents

**Table 2** Frontier molecular orbitals of compounds **7a–7e**

Compounds	HOMO	LUMO
<b>7a</b>		
<b>7b</b>		
<b>7c</b>		
<b>7d</b>		
<b>7e</b>		

**Fig. 4** TGA of compounds **7a**, **7b**, **7c** and **7e**

(1.42 mmol) was reacted with thioglyconic acid (1.42 mmol) in THF at reflux condition for 18 h to give 2-methyl-4-oxo-*N*-(4-oxo-2-1-*tert*-butyl-4-chloro-1*H*-pyrazole-1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide **7e**

Yield: 61 %, Melting point => 300 °C.

<sup>1</sup>H-NMR (CD<sub>3</sub>)<sub>2</sub>SO: δ 2.22 (s, 3H), 2.48 (s, 9H), 3.16 (s, 1H), 3.35 (s, 1H), 7.30 (d, 1H, *J*=8.7 Hz), 7.61 (d, 1H, *J*=8.7 Hz), 7.74 (dd, 1H, *J*=6.1, 8.7 Hz), 11.72 (1H, s)

FT-IR (KBr): 3034, 1759, 1721, 1645, 1621, 1545, 1435, 1341, 1257, 1222, 950, 669, 602 cm<sup>-1</sup>.

Mass: *m/z* 462 (M<sup>+</sup>).

## Result and Discussion

### Chemistry

Synthesis of 2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** was performed by reacting ethyl-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carboxylate with 98 % hydrazine hydrate under reflux condition for 2 h. Initially ethyl-2-methyl-4-oxo-3,4-dihydroquinazoline-5-carboxylate was insoluble in hydrazine hydrate at room temperature and it gets solubilized after heating at reflux condition. The product was separated from the reaction mass as the reaction goes to completion. On completion of the reaction the reaction mixture was cooled, the product was separated by filtration, washed with cold water, and dried.

The intermediate 2-methyl-4-oxo-3,4-dihydroquinazoline-5-carbohydrazide **5** was reacted with different aldehydes in methanol at reflux condition to give the corresponding Schiff base 2-methyl-4-oxo-*N'*-[(*Z*)-phenyl substituted methylidene]-3,4-dihydroquinazoline-5-carbohydrazide (**6a–6e**). In this step the formation of Schiff base requires nearly 2 to 7 h. The reaction was initiated in the presence of catalytic amount of H<sub>2</sub>SO<sub>4</sub>. On completion of the reaction the reaction

mixture was cooled, and neutralized with  $\text{Na}_2\text{CO}_3$  at low temperature to precipitate the products. The products were filtered and dried. The yields of the compounds **6a–6e** range from 67 to 87 %.

2-Methyl-4-oxo-N-(4-oxo-2-phenyl substituted -1,3-thiazolidin-3-yl)-3,4-dihydroquinazoline-5-carboxamide (**7a–7b**) was prepared by reacting the Schiff base 2-methyl-4-oxo-*N'*-[(*Z*)-phenyl substituted methylidene]-3,4-dihydroquinazoline-5-carbohydrazide (**6a–6e**) with thioglycolic acid in THF in the presence of catalytic amount of PTSA. The reaction details are summarized in the experimental procedure and the synthetic strategy is presented in Scheme 1.

### Photophysical Properties

The absorption and emission properties of the compounds **6a–6e** and **7a–7e** were studied in the solvent DMF. All the absorption-emission studies were performed at room temperature using solutions of concentration  $1 \times 10^{-6}$  M. The synthesised compounds are fluorescent in solution under irradiation of UV light. The absorption and emission spectra of the compounds **6a–6e** and **7a–7e** are presented in Figs. 2 and 3 respectively. The effect of electron donor and electron acceptor groups on the absorption and the emission properties of the compounds were studied. All the compounds absorb in the ultraviolet region and emit in the visible region. The absorption and the emission properties of the compounds show that they are well suitable to function as fluorescent brightening agents. In other words they absorb in the ultra-violet region and emit in the visible region. The observed photophysical properties of the compounds are compared with the computational results obtained by DFT and TD-DFT and the results are summarised in Table 1.

The absorption and emission properties of the compounds depend on the quinazolone unit as there is no direct

conjugation between quinazolone and thiazolyl units. This is also supported by the fact that the compound **6a–6e** and **7a–7e** have similar photophysical properties. The compounds **6a–6e** absorb at 300 nm except the compound **6c** which absorbs at 354 nm in DMF. The experimental absorption properties of the compounds are well in agreement with the computed energy difference between HOMO and LUMO Graph 1. In the case of the compound **6c** the energy difference between HOMO and LUMO is less as compared to the other compounds and it shows a red shifted absorption. The analogous behavior is observed for the compounds **7a–7e**. The compounds **6a–6e** emit at 430 nm except the compound **6c** which emits at 470 nm in DMF. The emission properties of the compound **6a–6e** and the compounds **7a–7e** are almost the same in DMF. This clearly indicates that the quinazolone unit is responsible for the absorption as well as the emission properties. The compounds **7b** and **7c** show dual absorption. The short wavelength absorption is in the range 274–294 nm and the long wavelength absorption is at 350 nm. In the case of the emission spectra of the compound **7a–7e** a single intense emission was observed. The quantum yields of the compounds are in range of 0.002 to 0.353. The compound **7e** shows a higher quantum yield as compared to the compounds **7a–7d** in DMF.

### Computational Study

The observed experimental absorption properties of the compounds were compared with the vertical excitation data obtained computationally and they are in good agreement with each other. The maximum difference is observed for the compounds **7c** and **7d**. In the case of the compounds **7a**, **7b** and **7e** the experimental absorption and the vertical excitation are almost the same. In the case of the emission a large difference was observed between the experimental emission and the calculated emission for the compound **7a** (19 %) and a

**Table 3** Color properties of compounds **7a–7e**

	Standard blank polyester	7a	7b	7c	7d	7e
X	48.025	56.532	51.414	57.805	48.921	53.635
Y	50.82	59.421	54.689	61.766	51.823	56.874
Z	54.138	56.284	55.871	61.848	54.815	57.327
L*	76.57	81.522	78.862	82.789	77.175	80.109
a*	−0.396	0.525	−1.111	−1.797	−0.54	−0.696
b*	0.416	6.879	2.687	3.915	0.799	3.451
C*	0.574	6.899	2.908	4.308	0.964	3.52
H*	133.608	85.601	112.491	114.682	124.075	101.434
K/S	0.2515	0.5506	0.3152	0.5626	0.2739	0.4152
Berger whiteness		−299.914	−145.533	−275.681	−16.865	−168.12
Stensbay whiteness		62.46	67.468	65.653	73.158	67.668
Taube whiteness		46.873	59.417	62.094	63.791	58.686



larger difference was observed for the compounds **7d** and **7e**. In the case of the compounds **7c** and **7d** the experimental emission and the calculated emission are very close to each other. A large Stokes shift was observed for the compounds **7c** and **7e**. The % deviation between the experimental photophysical properties and the calculated photophysical properties and oscillator strength obtained by DFT computation are summarised in Table 1.

The most probable electronic transitions occurring in the molecules were understood using the frontier molecular orbitals (HOMO and LUMO) generated using Gaussview 05 program for compounds **7a–7e** in the solvent DMF Table 2. All the compounds may be considered to be consisting of three cores—quinazolinone, thiazolidine and aromatic cores. The electron density for the compounds **7a–7e** is concentrated on the thiazolidine core, but the LUMO energy distribution pattern is not linear. In the case of compounds **7a**, **7d** and **7e** the electron density is spread over quinazolinone unit. The electron distribution is on the aromatic system of the thiazolidine for compounds **7c** and **7b**. The electron distribution patterns of compounds **7a–7e** indicates that the thiazolidine core acts as donor and quinazolinone and aromatic cores of the compound **7c** and **7b** act as acceptor units.

#### Thermogravimetric Analysis

Thermal stability of the compounds **7a–7e** was evaluated by thermogravimetric analysis (TGA). The compounds are thermally stable up to 200 °C. The compounds start decomposing after 200 °C and the decomposition is complete at 600 °C. Compound **7c** is thermally more stable as compared to the compounds **7a**, **7b** and **7e**. The compound **7d** is a semi-solid and so thermal stability of the compound **7d** was not evaluated. The thermogravimetric studies have been carried out in the temperature range 50–600 °C under nitrogen gas at a heating rate of 10 °C min<sup>-1</sup> Fig. 4.

#### Color Assessment

The colorimetric parameters of the whitened polyester fabrics using synthesized fluorescent molecules **7a–7e** were recorded on a reflectance spectrophotometer CE-7000A Gretag-Macbeth. CIE 1976 Color Space method was used to evaluate the color values of the synthesized compounds **7a–7e** on polyester fabrics in terms of L\*, a\* and b\* (Table 3). All the compounds have good affinity towards the polyester fabrics at high temperature and gave whitening with a blue tinge on polyester fabrics. The whiteness index values of the fabrics dyed with compounds **7a–7e** are summarized in Table 3.

#### Conclusion

Fluorescent compounds are synthesised from the intermediate ethyl 2-methyl-4-oxo-3, 4-hydroquinazoline-5-carboxylate. Photophysical properties of the compounds in DMF were evaluated experimentally and the results are compared with the theoretical data. The experimental results are in good agreements with the theoretical results. The % deviation between the experimental absorption and the emission is in the range between 1 and 19 %. The fluorescent compounds show good brightness on polyester fibres and have good thermal stability.

**Acknowledgments** Vikas Patil and Vikas Padalkar are thankful to Institute of Chemical Technology and they have contributed equally.

#### References

- Mhaske S, Argade N (2006) The chemistry of recently isolated naturally occurring quinazolinone alkaloids. *Tetrahedron* 62:9787–9826
- Bhatti HS, Seshadri S (2004) Synthesis and fastness properties of styryl and azo disperse dyes derived from 6-nitro substituted 3-aryl-2-methyl-4(3H)-quinazolinone. *Color Technol* 120:151–155
- Bhatti HS, Seshadri S (2004) Chromophoric potential of the 4(3H)-quinazolinones: a review. *Color Technol* 120:101–107
- Jung H, Kurtz W (1976) Neue pigment farbstoffe, DE2507908. BASF, Germany
- Rodgers JL (1965) 2-Ortho-hydroxy-phenyl-4-(3 h)-quinazolinones J P US3169129
- Golomolzin BV, Shcherak LO, Ya I, Postovskii (1969) Bisquinazolinone fluorescent whitening agents. *Khim Geterotsikl Soedin* 6:1131
- Cevasco AA (1972) Method for emitting fluorescent light using fluorescent quinazolones, DE2209872
- Purnaprajna V, Seshadri S (1973) Studies in the Vilsmeier Haack reaction part XIII: application of Vilsmeier Haack reaction to pyrazolo [5, 1-b] quinazolone system. *Indian J Chem* 15B:109–117
- Masaaki O, Yoshiro T, Katsunobu S, Sadaharu A (1973) Manufacture of basic dyes. *Jpn Kokai* 73(00):723
- Chamberlain TR, McCrae JM (1979) *Proc. 7th Int. Colour Symp., Interlaken, Switzerland*
- Venkataraman K (1971) *Chemistry of synthetic dyes*. 5. Academic Press. Inc, London
- Hunger K (2003) *Industrial dyes*, 3rd edn. Wiley VCH, Weinheim
- Tutak M (2011) Optical whitening of cationised cotton: effect on whiteness and whiteness tint. *Color Technol* 127:340–345
- Hurd RP, Reagan BM (1990) Exhaustion, photostability, and photosensitising properties of triazinylaminostilbene fluorescent brightening agents on cotton. *J Soc Dyes Colour* 106:49–54
- Leaver IH, Milligan B (1984) Fluorescent whitening agents—a survey. *Dyes Pig* 5:109–144
- Reinher D, Luther H, Mertzger G (2003) Use of fluorescent whitening agents. US20030103924A1
- Farrar JM, Jackson AJ (2005) Optical brighteners compositions their production and their use US6890454 B2
- Hussain M, Shamey R, Hinks D, Shafei A, Ali S (2012) Synthesis of novel stilbene-alkoxysilane fluorescent brighteners, and their performance on cotton fiber as fluorescent brightening and ultraviolet absorbing agents. *Dyes Pig* 92:1231–1240

19. Georgiev NI, Bojinov VB, Marinova N (2010) Novel PAMAM light-harvesting antennae based on 1,8-naphthalimide: synthesis, energy transfer, photophysical and pH sensing properties. *Sensors Actuators B* 150:655–666
20. Bojinov VB, Konstantinova TN (2007) Fluorescent 4-(2,2,6,6-tetramethylpiperidin-4-ylamino)-1,8-naphthalimide pH chemosensor based on photoinduced electron transfer. *Sensors Actuators B* 123: 869–876
21. Tian H, Gan J, Chen K, He J, Liang Song Q, Yuan Hou X (2002) Positive and negative fluorescent imaging induced by naphthalimide polymers. *J Mater Chem* 12:1262–1267
22. Zhu W, Hu C, Chen K, Tian H (1998) Luminescent properties of copolymeric dyad compounds containing 1,8-naphthalimide and 1,3,4-oxadiazole. *Synth Met* 96:51–154
23. Harrington BJ (2009) The staining of oocysts of cryptosporidium with the fluorescent brighteners Uvitex 2B and calcofluor white. *Lab Med* 40:219–223
24. Xian HH, Xing YZ, Jia DB (2011) Synthesis and characterization of a novel waterborne stilbene-based polyurethane fluorescent brightener. *Chin Chem Lett* 22(8):997–1000
25. Um S, Lee JK, Kang Y, Baek DJ (2006) The synthesis and properties of triazine-stilbene fluorescent brighteners containing the phenolic antioxidant [II]. *Dyes Pig* 70:84–90
26. Um S, Kang Y, Lee JK (2007) The synthesis and properties of triazine-stilbene fluorescent brighteners containing a monophenolic antioxidant. *Dyes Pig* 75:681–686
27. Wang X, Li W, Zhang XH, Liu DZ, Zhou XQ (2005) A study on the synthesis and photophysical performances of some pyrazole and triazole fluorescent brightening agents. *Dyes Pig* 64:141–146
28. Bojinov V, Grabchev I (2001) A new method for synthesis of 4-allyloxy-1,8-naphthalimide derivatives for use as fluorescent brighteners. *Dyes Pig* 51:57–61
29. Rangnekar DW, Rajadhyaksha DD (1987) Synthesis of phthalimid-3-yl and -4-yl aminoethylenes and pyrroloquinolines and a study of their fluorescence properties. *Dyes Pig* 8:1–10
30. Rangnekar DW, Shenoy GR (1987) Synthesis of 7H-benzo [d] triazolo [5, 1-a] isoquinolin-7-one derivatives and study of their fluorescent properties. *Dyes Pig* 8:291–299
31. Rangnekar DW, Tagdiwala PV (1986) Synthesis of 6-acetamido-2-substituted quinoxaline derivatives and their use as fluorescent whiteners for polyester fibres. *Dyes Pig* 7:445–455
32. Rangnekar DW, Phadke RC (1985) Synthesis of 2-hetaryl-5-phenyl-1,3,4-oxadiazole and bis-1,3,4-oxadiazole derivatives and their use as fluorescent whiteners for polyester fibres. *Dyes Pig* 6:293–302
33. Rangnekar DW, Tagdiwala PV (1986) Synthesis of 2,4-dihydro-6-methyl-4-phenyl 2-(4-substituted phenyl) pyrazolo[3,4-triazole derivatives and their use as fluorescent whitener for polyester fibres. *Dyes Pig* 7:289–298
34. Um S (2007) The synthesis and properties of benzoxazole fluorescent brighteners for application to polyester fibres. *Dyes Pig* 75:185–188
35. Bojinov V, Panova I, Grabchev I (2005) Benzotriazoles: Novel adducts of a 2-(2-hydroxyphenyl)-benzotriazole and a blue emitting benzo[de]isoquinoline-1,3-dione for “one-step” fluorescent brightening and stabilization of polymers. *Polym Degrad Stab* 88:420–427
36. Mishra H, Zhang Y, Geddes CD (2011) Metal enhanced fluorescence of the fluorescent brightening agent Tinopal-CBX near silver island film. *Dyes Pig* 91:225–230
37. Treutler O, Ahlrichs R (1995) Efficient molecular numerical integration schemes. *J Chem Phys* 102:346–354
38. Becke AD (1993) A new mixing of Hartree–Fock and local density-functional theories. *J Chem Phys* 98:1372–1377
39. Lee C, Yang W, Parr RG (1988) Development of the Colle–Salvetti correlation-energy formula into a functional of the electron density. *Phys Rev B* 37:785–789
40. Kim CH, Park J, Seo J, Park SJ, Joo T (2010) Excited State Intramolecular Proton (ESIPT) transfer and charge transfer dynamics of a 2-(2'-hydroxyphenyl)benzoxazole derivative in solution. *J Phys Chem A* 114:5618–5629
41. Santra M, Moon H, Park MH, Lee TW, Kim Y, Ahn KH (2012) Dramatic substituent effects on the photoluminescence of boron complexes of 2-(benzothiazol-2-yl)phenols. *Chem Eur J*. doi:10.1002/chem.201200726
42. Li H, Niu L, Xu X, Zhang S, Gao F (2011) A comprehensive theoretical investigation of intramolecular proton transfer in the excited states for some newly-designed diphenylethylene derivatives bearing 2-(2-hydroxy-phenyl)-benzotriazole part. *J Fluoresc* 21: 1721–1728
43. Hehre WJ, Radom L, Schleyer PVR, Pople J (1986) *Ab Initio molecular orbital theory*. Wiley, New York
44. Bauernschmitt R, Ahlrichs R (1996) *Chem Phys Lett* 256:454–464
45. Furche F, Rappaport D (2005) Density functional theory for excited states: equilibrium structure and electronic spectra. In: Olivucci M (ed) *Computational photochemistry*. Elsevier, Amsterdam, Vol 16, Chapter 3
46. Lakowicz JR (1999) *Principles of fluorescence spectroscopy*, 2nd edn. Kluwer, New York
47. Valeur B (2001) *Molecular fluorescence: principles and applications*. Wiley-VCH Verlag, Weinheim
48. Phatangare KR, Gupta VD, Tathe AB, Padalkar VS, Patil VS, Ramasami P, Sekar N (2013) ESIPT inspired fluorescent 2-(4-benzo[d]oxazol-2-yl)naphtho[1,2-d]oxazol-2-yl)phenol: experimental and DFT based approach to photophysical properties. *Tetrahedron* 69:1767–1777
49. Padalkar VS, Ramasami P, Sekar N (2013) A combined experimental and DFT-TDDFT study of the excited-state intramolecular proton transfer (ESIPT) of 2-(2'-hydroxyphenyl) imidazole derivatives. *J Fluoresc* 23:839–851
50. Gupta VD, Tathe AB, Padalkar VS, Patil VS, Phatangare KR, Umape PG, Ramasami P, Sekar N (2013) TDDFT investigation of the electronic structures and photophysical properties of fluorescent extended styryl push-pull chromophores containing carbazole unit. *J Fluoresc* 23:1121–1138
51. Padalkar VS, Ramasami P, Sekar N (2013) TD-DFT study of excited-state intramolecular proton transfer (ESIPT) of 2-(1,3-benzothiazol-2-yl)-5-(N, N-diethylamino)phenol with benzoxazole and benzimidazole analogues. *J Fluoresc* 18:797–805
52. Patil VS, Padalkar VS, Tathe AB, Sekar N (2013) Synthesis, photophysical and DFT studies of ESIPT inspired novel 2-(2',4'-dihydroxyphenyl) benzimidazole, benzoxazole and benzothiazole. *J Fluoresc* 23:1019–1029
53. Frisch MJ, Trucks GW, Schlegel HB, Scuseria GE, Robb MA, Cheeseman JR, Scalmani G, Barone V, Mennucci B, Petersson GA, Nakatsuji H, Caricato M, Li X, Hratchian HP, Izmaylov AF, Bloino J, Zheng G, Sonnenberg JL, Hada M, Ehara M, Toyota K, Fukuda R, Hasegawa J, Ishida M, Nakajima T, Honda Y, Kitao O, Nakai H, Vreven T, Montgomery JA, Peralta JE, Ogliaro F, Bearpark M, Heyd JJ, Brothers E, Kudin KN, Staroverov VN, Kobayashi R, Normand J, Raghavachari K, Rendell A, Burant JC, Iyengar SS, Tomasi J, Cossi M, Rega N, Millam NJ, Klene M, Knox JE, Cross JB, Bakken V, Adamo C, Jaramillo J, Gomperts R, Stratmann RE, Yazyev O, Austin AJ, Cammi R, Pomelli C, Ochterski JW, Martin RL, Morokuma K, Zakrzewski VG, Voth GA, Salvador P, Dannenberg JJ, Dapprich S, Daniels AD, Farkas O, Foresman JB, Ortiz JV, Cioslowski J, Fox DJ (2010) *Gaussian 09*, revision C.01. Gaussian, Inc., Wallingford
54. Williams AR, Winfield S, Miller JN (1983) Relative fluorescence quantum yields sing a computer controlled luminescence spectrometer. *Analyst* 108:1067–1071
55. Padalkar VS, Patil VS, Sekar N (2011) Synthesis and characterization of novel 2, 2'- bipyrimidine fluorescent derivative for protein binding. *Chem Cent J* 5(72):1–7
56. Patil VS, Padalkar VS, Sekar N (2012) Intrinsic catalytic activity of an acidic ionic liquid as a solvent for quinazoline synthesis. *Catal Sci Tech* 2:1681–1684