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AIE active cyanostilbenes for live-cell imaging of lipid droplets†

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Lipid droplets (LDs), owing to their involvement in various physiological processes and diseases such as fatty liver disease, diabetes, and cancer, have attracted considerable attention. Live cell imaging of lipid droplets allows researchers to visualize and track the movement of these organelles within living cells in real time. In this study, we synthesized trifluoromethyl-substituted cyanostilbene fluorophores containing naphthalene or julolidine groups to visualize lipid droplets in live cells and investigate the changes in the size and number of LDs upon oleic acid stimulation. The fluorophores with donor (D) and acceptor (A) substituents show strong solvatochromism due to intramolecular charge transfer (ICT). In addition, these fluorophores show enhanced aggregation-induced emission (AIE) in water, and the lipophilic trifluoromethyl substitution aids the formation of stable organogels. The morphology and size of the aggregates obtained were characterized through scanning electron microscopy and dynamic light scattering experiments. Rheology and differential scanning calorimetry were used to further characterize the organogels. Our findings suggest that these D- π -A-substituted cyanostilbenes are valuable probes for understanding live-cell LD dynamics.

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Introduction

Cyanostilbene derivatives that show characteristic aggregation-induced emission have been investigated extensively as functional materials for various applications in sensing, displays, and bioimaging.^{1–7} The restrictions on molecular rotation imposed by the presence of the nitrile group on the double bond and additional substituents can yield significant changes in either the emission intensity or wavelength. Considering the uniqueness of the scaffold, many cyanostilbene derivatives with several donor or acceptor substituents have been developed. One substituent that has attracted attention is the trifluoromethyl (CF₃) group. Trifluoromethyl groups are important in various areas of chemistry and biology and are used as bioisosteres in medicinal chemistry to create potent substrates with improved bioavailability and specificity.⁸ These groups are known for their strong electron-withdrawing properties, high polarity, high dipole moment and high electric permittivity, and can therefore affect the steric and electronic properties, making them useful in drug design and organic synthesis.⁹ Considering the unique properties that fluorine substitution can offer, Park and co-workers have developed organogels with CF₃ substitution in place of traditional

gelator moieties such as hydrocarbon chain, glycol or amide units, along with their aggregation induced emission properties.^{10–13} Our previous work has also explored the utility of the CF₃ group for the formation of various low molecular weight organogels using various aromatic scaffolds such as pyrene, anthracene and triphenylamine.^{14–17} In addition, trifluoromethyl groups can increase the lipophilicity of molecules, which can improve their ability to penetrate cell membranes and target certain proteins. Thus, we intended (i) to utilize the unique aggregation-induced emission behaviour of cyanostilbene for its utility in biological imaging and (ii) demonstrate the formation of stable organogels. Previously, several derivatives of α -cyanostilbenes have been utilized for cellular imaging. For example, a folic acid conjugate of an unsymmetrical cyanostilbene was utilized for its internalization in a cancer cell line,¹⁸ and α -cyanostilbenes bearing alkylammonium pendants were used for hydroxyl radical bioimaging.¹⁹ By incorporating a mitochondrial-targeting pyridinium group in the cyanostilbene scaffold, mitochondrial uptake and imaging were demonstrated.^{20–22} Cellular internalization was also demonstrated by pyrene- or triphenylamine-containing cyanostilbenes with distribution throughout the cytoplasmic region,²³ monitoring the morphological changes of the plasma membrane,²⁴ and binding to the mitochondria in zebrafish eggs.²⁵ A pyridine-containing acrylonitrile derivative with coumarin substitution was also shown for the imaging the lipid droplets.²⁶ Recently, Niu and co-workers synthesized various acrylonitrile derivatives to image LDs.^{24,27} As demonstrated by their results, the lipophilic properties of the CF₃-containing cyanostilbenes can

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be well-suited to the imaging of lipid droplets. LDs are small, spherical droplets of fats and other lipids that are found in cells, and they are composed of a core of neutral lipids, such as triacylglycerols and cholesteryl esters and surrounded by a phospholipid monolayer. LDs play an important role in energy storage and metabolism and are also involved in the regulation of gene expression and cell signaling.²⁸ Therefore, various research groups have explored the imaging of lipid droplets and underlying phenomena using small-molecule fluorescent probes.^{26,29–35} Utilizing the lipophilic properties of the trifluoromethyl group, and through the incorporation of strong electron-donating groups such as dimethylamino naphthalene or julolidine, we investigated the photophysical properties, organogel formation and utility of cyanostilbene fluorophores for the imaging of lipid-rich organelles. We detail our results below.

Materials and methods

All the necessary chemicals required for the synthesis of the probes were purchased from Aldrich, BLD Pharm, TCI chemicals, Avra synthesis, or SD Fine-chem. The synthesized samples were characterized using ¹H and ¹³C NMR (Bruker Advance III-500 MHz) in chloroform-D or dimethyl sulphoxide (D₆) solvents. Absorption studies were carried out using a Shimadzu UV-Vis 3600i instrument, and steady-state and fluorescence studies were carried out using a Horiba Jobin Yvon Fluorolog-3 spectrofluorimeter. The sample concentrations used for the absorption and fluorescence studies were of the order of 10 μM. The excitation wavelength for recording the emission spectra was typically kept at the absorption maxima of the samples under investigation. Mass spectral data were recorded using a Waters Synapt G2S ESI Q-ToF mass spectrometer. DLS measurements were performed using a Malvern Zetasizer Ultra instrument. The samples were prepared in water and filtered through a micro-filter; air bubbles were also removed before measurements. The organogel was lyophilized and vacuum dried before SEM measurements. Drop cast (10 μM of the sample) SEM images of the aggregate were obtained using a silicon wafer. SEM measurements were performed using an advanced analytical FE-SEM instrument (JEOL JSM-7900F). The rheology measurements were carried out using an Anton Paar (MCR 702) rheometer. TGA-DLS thermal analysis was performed using a Jupiter thermal analyser.

Cell culture and live-cell imaging

The COS-7 cells (monkey kidney fibroblast; ATCC) were grown in cell media containing DMEM (Dulbecco's Modified Eagle Medium; Gibco) supplemented with 10% FBS (fetal bovine serum; Gibco). The cells were maintained in a humidified environment at 37 °C with 5% CO₂. For live-cell imaging, a cell density of 1.5×10^5 cells was seeded on a 35 mm glass-bottom dish 24 h before the dye treatment. The colocalization experiment was performed with the addition of 2 μM of the dye (N-1, N-2, J-1, and J-2, respectively) in FluoroBrite™ DMEM, followed by the addition of 50 nM of Nile red to stain the lipid droplets of

the COS-7 cells, with subsequent incubation for a period of 10 minutes. The cells were then imaged using a 63× oil immersion objective lens (1.40 numerical aperture) of a Leica laser scanning confocal microscope (TCS SP8) in a live cell chamber at 37 °C with 97% humidity and 5% CO₂. The Nile red dye was excited using a 561 nm laser, and the emission signal was collected using a HyD detector. For the synthesized dyes, excitation was *via* a 488 nm laser, and again the HyD detector was used to the record emission signal.

LD accumulation experiments

COS-7 cells were seeded on a 35 mm glass-bottom dish for 24 h, and then treated with 100 μM oleic acid for 4 h. After incubation, the medium was removed, and the cells were washed three times with PBS. The LD accumulation experiment was performed with the addition of J-2 (2 μM) in FluoroBrite™ DMEM. For the analysis of changes in the number and size of the lipid droplets upon oleic acid treatment, 15 cells with and without oleic acid treatment were considered for these calculations.

Cytotoxicity measurements

The MTT assay was used for checking the cytotoxicity level of the dyes. A cell population of 5×10^3 cells per well was seeded in a 96-well plate in Dulbecco's modified Eagle medium (DMEM) complemented with 10% fetal bovine serum (FBS) with 5% CO₂ in an incubator at 37 °C for 24 h. The cells were then treated with different concentrations of each synthesized dye, *i.e.*, 0.25, 0.5, 1, 2 and 4 μM. The dye was incubated with the cells for 4 h. After incubation, the medium was removed from the wells, and 100 μl of MTT solution (0.5 mg mL⁻¹) was added to each well and again incubated for 4 h, leading to the formation of formazan crystals due to the reduction of MTT by viable cells. Using a multichannel pipette, the MTT solution was then removed from the corners of the well without disturbing the purple-colored formazan crystals. DMSO (100 μL) was added to the wells to dissolve the formazan crystals. The plate was shaken for 10 min at RT to dissolve the crystals completely. The absorbance values corresponding to different concentrations were then recorded at 570 nm using a PerkinElmer multimode plate reader. Triplets of each concentration of the probe were prepared for MTT assay. The experiment was repeated twice, and the mean ± standard deviation of the percentage cell viability was plotted against the different concentrations. The percentage cell viability of the dye was calculated with respect to the negative control (cells in DMEM and 10% FBS medium).

Data analysis

ImageJ software (nih.gov) was used to analyze and quantify the live-cell images. For the calculation of the Pearson correlation coefficient (PCC) in the colocalization experiment, the coloc2 plugin of ImageJ was used. Origin 9.1 software was used to plot the data for the cell viability assay.

Results and discussion

Synthesis

The series of LD-CF₃ probes containing trifluoromethyl substituents using the Knoevenagel condensation reaction between the corresponding aldehydes and trifluoromethyl phenylacetonitrile derivatives (Fig. 1). The dimethylamino naphthalene carboxaldehyde was synthesized as per earlier literature.³⁶ The synthetic procedures with characterization and spectral details are provided in the ESI.†

Absorption and emission

The compounds were characterized by their strong electron-donating amino groups (dimethylamine for **N1** and **N2**, and julolidine for **J1** and **J2**) and electron-accepting nitrile and trifluoromethyl groups, which promote strong charge transfer. In addition, the nitrile group on the double bond is expected to aid in the unique aggregation induced emission (AIE) behaviour by increasing the rigidity of the molecule and reducing the degree of intramolecular rotation.² Compound **N1** absorbs at 423 nm in dioxane and **N2** with an additional CF₃ group absorbs slightly higher at 430 nm. Similarly, compound **J1** with one CF₃ group shows an absorption maximum at 434 nm, and **J2** with two CF₃ groups yields a moderate bathochromic shift. These bathochromic absorption shifts are attributed to the electron accepting nature of the CF₃ group. The fused ring nature of the julolidine substituents (**J1** and **J2**) also plays a role in electron donation to yield the bathochromic shifts in absorption. Compound **J2** shows the highest absorption maximum, which is attributed to the increased planarity. Despite the presence of extended conjugation in **N1** and **N2**, the julolidine-containing compounds **J1** and **J2** show a higher absorption, indicating their greater electron-donating nature. The order of absorption wavelength maxima for all the compounds is **N1** < **N2** < **J1** < **J2** (Table 1 and Fig. 2). In water, all the compounds show a

broad absorption spectrum due to their weaker solubility and propensity to form aggregates. The solvent polarity has a weaker effect on the absorption maxima (Fig. 2).

However, the emission of the compounds show a strong solvatochromic shift with the solvent polarity. With a change in the solvent polarity from dioxane to DMSO, **N1** shows a bathochromic shift in emission of 65 nm, **N2** of 63 nm, **J1** of 39 nm, and **J2** of 35 nm. The additional CF₃ moiety shows no major influence on the emission maxima. The naphthalene-substituted compounds **N1** and **N2** show a greater solvatochromic variation than the julolidine derivatives and also exhibit an intense color demarcation (Fig. 2 and Fig. S2, ESI†). The spectroscopic purity of all the compounds is shown by the perfect overlap between the absorption and excitation spectra (Fig. S1, ESI†). As can be seen in Table 1, all the compounds show significant emission wavelength shifts in water attributed to the aggregation induced emission^{2,37} coupled with solvent polarity effects. The emission spectra were also recorded in a dioxane–water binary mixture to learn about the emission behavior through the systematic variation of the solvent polarity. Dioxane is a good solvent in which molecules are readily soluble, whereas water is a poor solvent.

As the percentage of water in dioxane was increased, we noted prominent changes in both the emission wavelength and emission intensity for all the compounds at a water fraction of > 60% (Fig. 3 and Fig. S4, ESI†). Along with the wavelength shifts, a concomitant intensity enhancement was also noted: **N1** and **J1** yield an intensity enhancement of ~6–8-fold, and **N2** and **J2** (with the additional CF₃ substituent) show an intensity enhancement of ~16 fold and ~20 fold, respectively, with distinct colors seen with different water fractions (inset of Fig. 3B and Fig. S4A–C, ESI†). The enhanced emission is attributed to the formation of the aggregates, and the aggregate morphology was studied *via* drop-casting of the aqueous solution on a Si wafer (Fig. 4A, B and Fig. S5a, ESI†). Irregular, spheroid, and rectangular shaped aggregates were obtained for

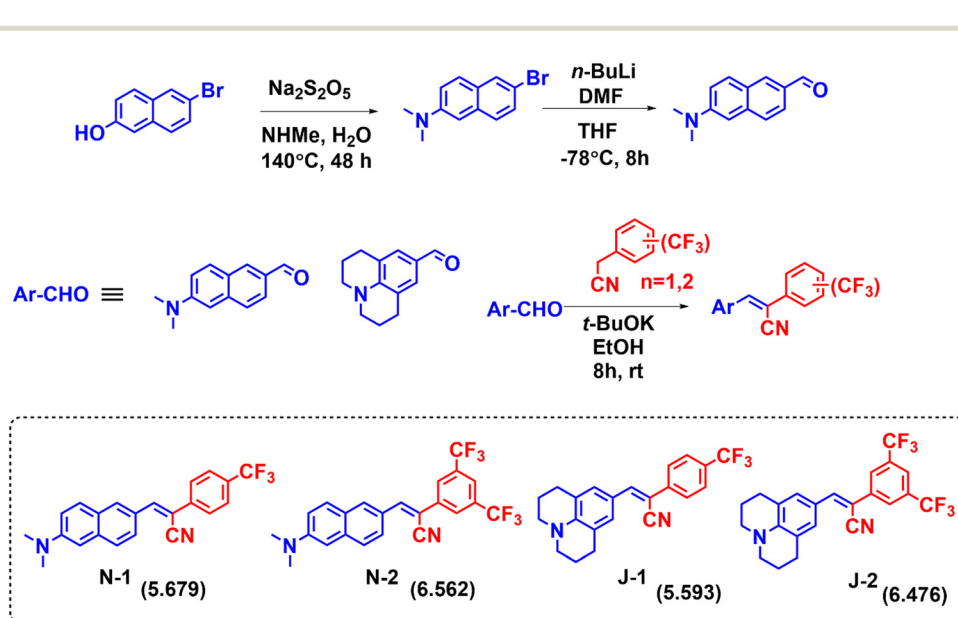


Fig. 1 Synthetic scheme for preparation of the LD-CF₃ probes. (The *clogP* values are given in parentheses.)

Table 1 Absorption and emission properties of all the molecules in different solvents

Solvent	N-1			N-2			J-1			J-2		
	λ_{abs} (nm)	λ_{em} (nm)	ϵ^*	λ_{abs} (nm)	λ_{em} (nm)	ϵ^*	λ_{abs} (nm)	λ_{em} (nm)	ϵ^*	λ_{abs} (nm)	λ_{em} (nm)	ϵ^*
Dioxane	423	528	6.1	430	546	5.3	434	493	5.3	441	508	5.5
THF	425	556	6.0	434	577	5.2	438	551	5.1	448	523	4.9
CH ₃ CN	413	579	5.2	422	595	4.4	435	526	4.0	444	536	4.3
DMF	419	587	4.6	429	602	4.2	441	529	4.1	451	540	4.2
DMSO	420	593	4.9	427	609	2.8	445	532	3.7	452	543	3.6
Ethanol	411	564	5.0	422	587	4.3	428	510	3.8	446	524	4.4
Methanol	411	568	5.2	420	591	4.2	430	513	3.8	444	526	4.2
Water	413	591	3.6	426	614	3.2	428	566	2.4	441	593	3.0
Solid state (λ_{em})	540			648			544			588		

$\epsilon^* = \text{value} \times 10^4 \text{ mol}^{-1} \text{ cm}^{-1}$.

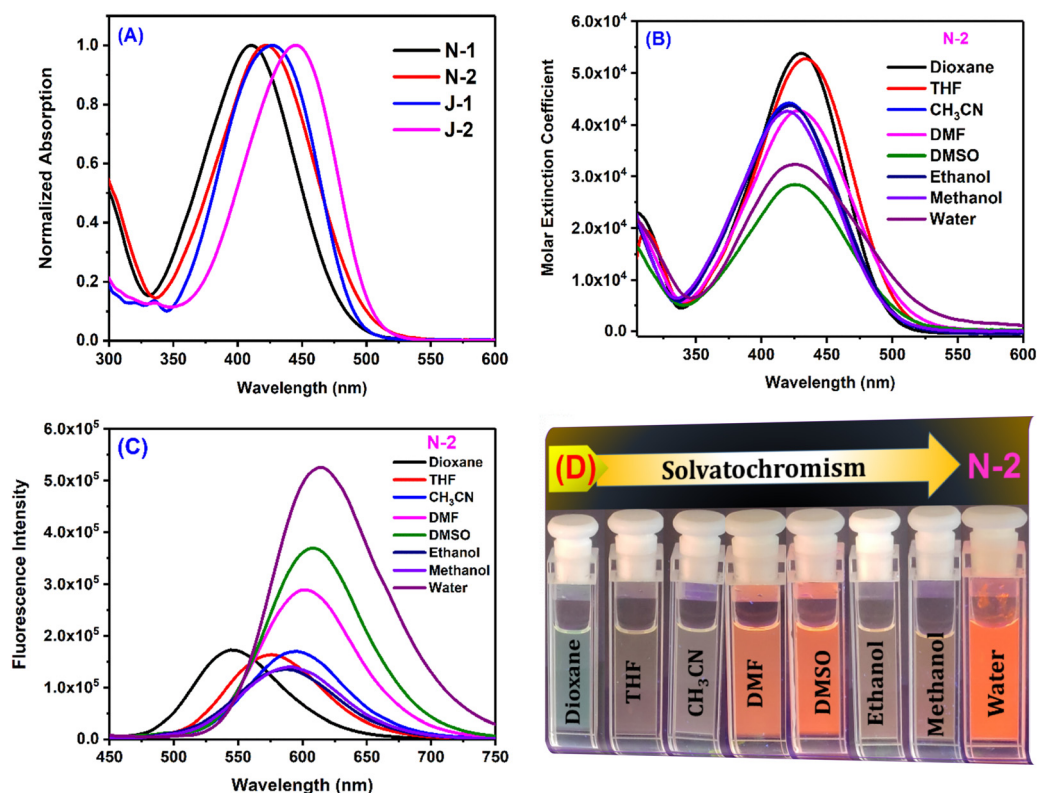


Fig. 2 (A) Absorption of all compounds in ethanol; (B) Absorption of **N2** in all solvents; (C) Emission of **N2** in all solvents; (D) Pictures of solutions for the emission of **N2** in different solvents under UV lamp irradiation of 365 nm. [Concentration = 10 μM]. (Data for the other compounds are given in Fig. S3, ESI[†])

all the compounds in aqueous medium (Fig. 4 and Fig. S5a, ESI[†]). The aggregate particle size was measuring using dynamic light scattering (DLS), which showed a mean hydrodynamic diameter of 397 nm for **N1**, 171.8 nm for **N2**, 152.4 nm for **J1** and 143.1 nm for **J2** (Fig. 4C, D and Fig. S5b, ESI[†]).

Low molecular weight organogels

Small molecule organogels are three-dimensional networks of self-assembled molecules that form a gel-like structure in organic solvents. Organogels based on small molecules have unique properties, such as high stability, ease of preparation, and tunable properties, making them useful for a variety of

applications such as drug delivery, tissue engineering, imaging, tracking and sensing.^{38–40} It has been well established that CF₃ groups and long-chain alkyl groups help to form organogels with higher stability through various non-covalent interactions, packing and bonding interactions.^{10,41} We obtained the formation of a stable organogel with compounds bearing two CF₃ groups (**N2** and **J2**) in ethanol upon heating, followed by sonication and cooling. The organogels were found to be stable for 24 h and converted to the sol state upon heating at 100 °C. The melting and thermal degradation temperatures for the xerogels were obtained *via* differential scanning calorimetry and were found to be 190–200 °C for both **J2** and **N2** (Fig. S7, ESI[†]), indicating their good

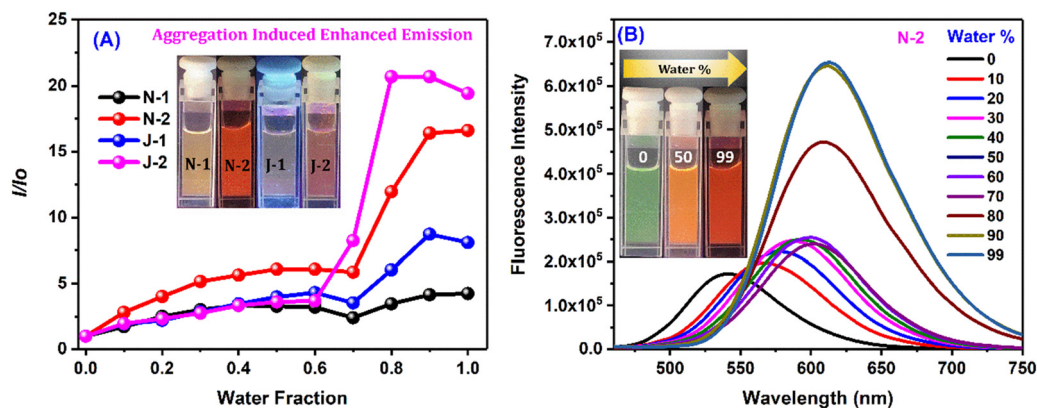


Fig. 3 (A) I/I_0 for all the derivatives corresponding to λ_{\max} of water vs. water fraction; and (B) emission spectra of **N2** in the dioxane/water binary solvent system. [Concentration = 10 μM .]

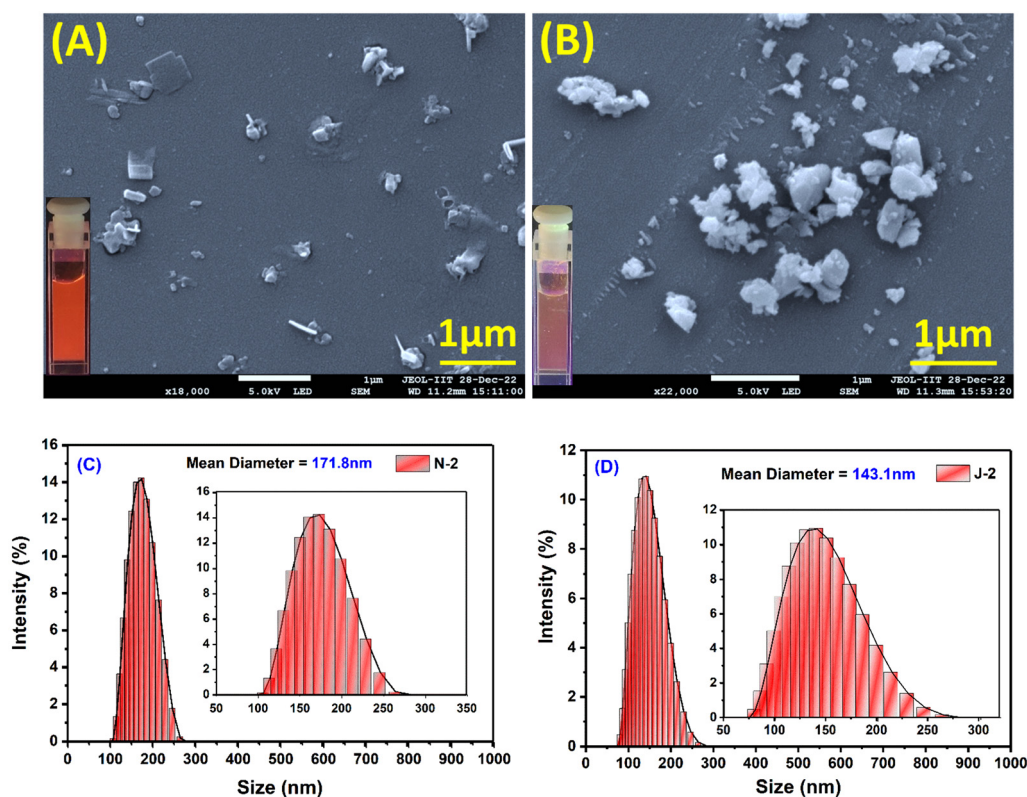


Fig. 4 Scanning electron microscope (SEM) images of 10 μM drop-cast solutions of (A) **N2** and (B) **J2**. The samples were coated with gold for 60 seconds. [The FESEM images of drop-cast solutions of **N1** and **J1** are shown in Fig. S5, ESI.†] (C and D) MA-DLS spectrum depicting the size/mean hydrodynamic diameter of the aggregates of **N2** and **J2**. [DLS spectra of **N1** and **J2** are shown in Fig. S4, ESI.†] [For each sample, a concentration of 10 μM was used.]

stability at higher temperatures. The emission of the organogels shows a red shift compared with their solution state, with **J2** having a higher shift (+69 nm) than **N2** (+46 nm), with the order of solution state < AIE < gel state for **N-2** and solution state < gel phase \leq AIE phase for **J2** (Fig. 5A and B). The viscoelastic properties of the organogels were also investigated using a rheometer. In the rheology experiments, G' and G'' refer to the storage and loss moduli, respectively. The storage modulus (G') represents the

elastic component of a material's response to an applied stress, while the loss modulus (G'') represents the viscous component. The ratio of G'' to G' is known as the loss tangent ($\tan \delta$) and can be used to determine the degree of viscoelasticity of a material. The observed storage modulus for **J2** and **N2** was dominant over their respective viscosity modulus or loss modulus. This indicates that the material can return to its original shape more easily when the applied stress is removed. The moduli are recorded with respect to

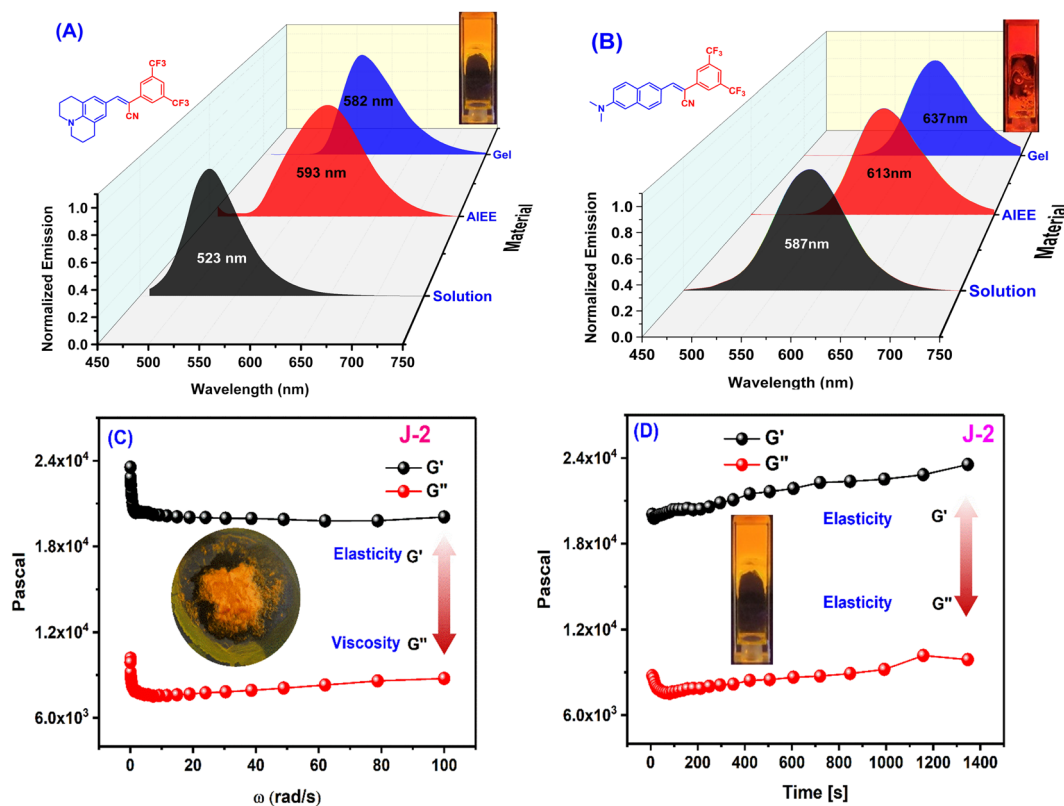


Fig. 5 Emission spectra of (A) **J2** and (B) **N2** in ethanol solution (10 μM), water solution (10 μM , AIE) and the gel state. (C and D) Frequency sweep (C) and time sweep (D) measurements for organogel **J2**. (Insets show gel pictures of **J2** kept in the rheometer after applying pressure; a photo of gel in the cuvette. The frequency sweep and time sweep measurements for organogel **N2** are given in Fig. S6, ESI†)

the angular frequency ω (rad s^{-1}) as well as time (s), and the storage modulus for both was found to be dominant when monitored till $\omega = 100 \text{ rad s}^{-1}$ and $t = 1400 \text{ s}$, which indicates greater gel stability with respect to ω and time (Fig. 5C, D and Fig. S6, ESI†). After a time sweep of 800 s, **N2** was found to lose its elasticity (Fig. S6B, ESI†) although **J2** exhibited an increased elasticity owing to the additional stabilization offered due to the presence of the fused ring system of julolidine. The SEM images of xerogels of the compounds show fibrous and network-like structures of varying thickness, indicating the gel-formation (Fig. 6).

Live cell imaging of lipid droplets

With the strong solvatochromic behavior of the compounds, we extended the utility of the molecules for the imaging of lipid droplets. The synthesized compounds exhibit $c\log P$ values ranging from 5.6 to 6.5, which is a measure of the compound's lipophilicity, *i.e.*, its ability to dissolve in lipids. In the context of lipid droplet imaging, $c\log P$ can be a useful descriptor for predicting a compound's ability to localize to lipid droplets. Compounds with $c\log P$ values greater than ~ 3 are considered to be lipophilic and tend to localize to lipid droplets.⁴² Therefore, given their favorable $c\log P$ values, we investigated the potential of these fluorescent probes (**N1**, **N2**, **J1**, and **J2**) for the labelling of lipid droplets (LDs) in COS-7 cells. We found that when these cells were incubated with the molecules, they quickly entered the cell membrane and produced a distinct,

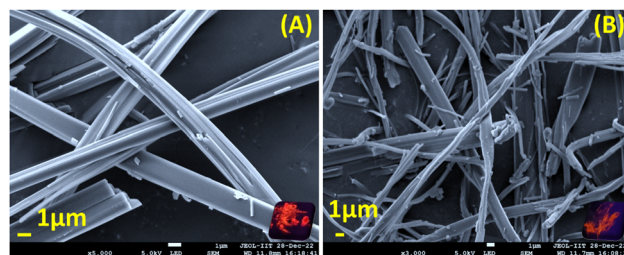


Fig. 6 Scanning electron microscope images of xerogels of (A) **N-2** and (B) **J-2** (where the insets show pictures of xerogels under UV lamp irradiation of 365 nm). The xerogels were obtained via lyophilization and vacuum drying. The samples were coated with gold for 60 seconds.

bright, and uniform punctate like structures throughout the cytoplasm. The images of the cells both in the bright field and the confocal microscope are given in Fig. 7 and Fig. S8 (ESI†) and show structures that resemble lipid droplets. To confirm that these structures were indeed lipid droplets, we also stained the cells with Nile red,⁴³ a known marker for lipid droplets, and found a significant overlap between the two labelling methods, indicating greater selectivity for the lipid droplets.

Among the molecules investigated, **J2** shows the highest PCC value indicating strong selectivity compared with the other fluorophores. We also examined the cytotoxicity *via* MTT assay of the probes at different concentrations for 4 h, and we did not find any adverse cellular behavior (Fig. S9, ESI†).

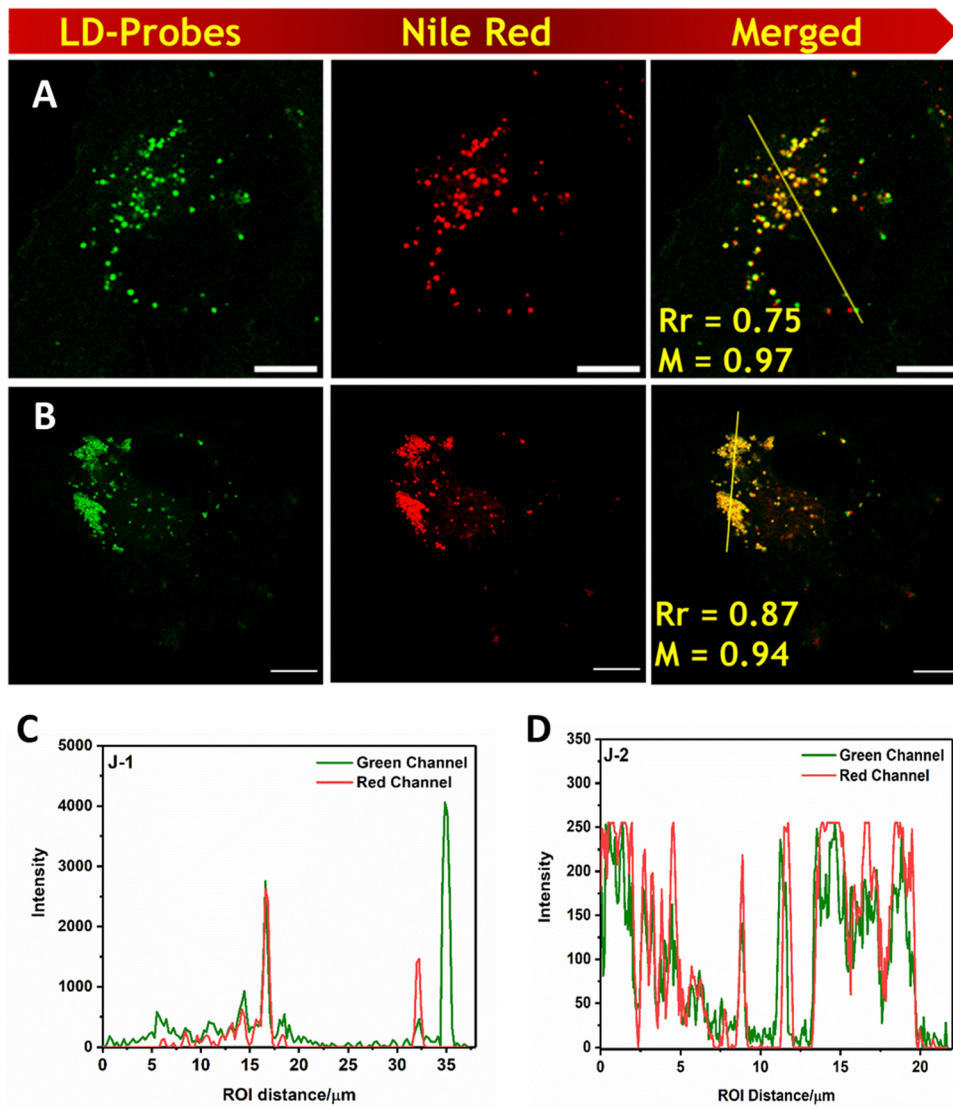


Fig. 7 A and B are cells stained with **J1** and **J2** respectively **J1** and **J2** in COS-7 cells (Column-1). Nile red was used as a lipid droplet marker and merged images are shown in column-3. Scale bar = 10 μm . C and D Intensity profile overlaps of for **J1** and **J2** with Nile red respectively. Bright-field images are shown in the ESI.† ($\lambda_{\text{ex}} = 488 \text{ nm}$, $\lambda_{\text{em}} = 505\text{--}560 \text{ nm}$ for all the molecules; and $\lambda_{\text{ex}} = 561 \text{ nm}$, $\lambda_{\text{em}} = 570\text{--}679 \text{ nm}$ for Nile red.)

LD accumulation

Oleic acid (OA) stimulates the formation of lipid droplets in cells by activating certain signalling pathways, such as the FPAR pathway.⁴⁴ To investigate lipid droplet accumulation, cells were treated with oleic acid (100 μM) for 4 h in cell culture media. Visualization of the cells revealed a significant increase in the number and size of lipid droplets. Fig. 8 clearly illustrates the increase in the number of lipid droplets as well as their size after the addition of OA. The mean diameter (size) of the lipid droplets increased from 261.93 nm to 581 nm upon treatment with oleic acid, and the number of lipid droplets per cell increased from $n = 119$ to $n = 251$. These results indicate that the probes can effectively label and visualize lipid droplets, even at low concentrations (2 μM) and with shorter incubation times (15 minutes).

Conclusions

In summary, we have designed and synthesized four trifluoromethyl-substituted cyanostilbene fluorophores containing naphthalene or julolidine donor groups for effective visualization of lipid droplets in COS-7 cells. These molecules have proven to be good candidates for visualizing changes in lipid droplet number and size upon stimulation with oleic acid, due to their efficient colocalizing ability for lipid droplets. The dynamic visualization of these changes using these potential probes opens opportunities in areas such as detecting accumulated fats in tissues, fatty liver syndrome, and other lipid droplet-associated diseases. Furthermore, these fluorophores, which bear both donor and acceptor substituents, exhibit strong solvatochromism and excellent aggregation-induced emission in aqueous media. Additionally, we demonstrated

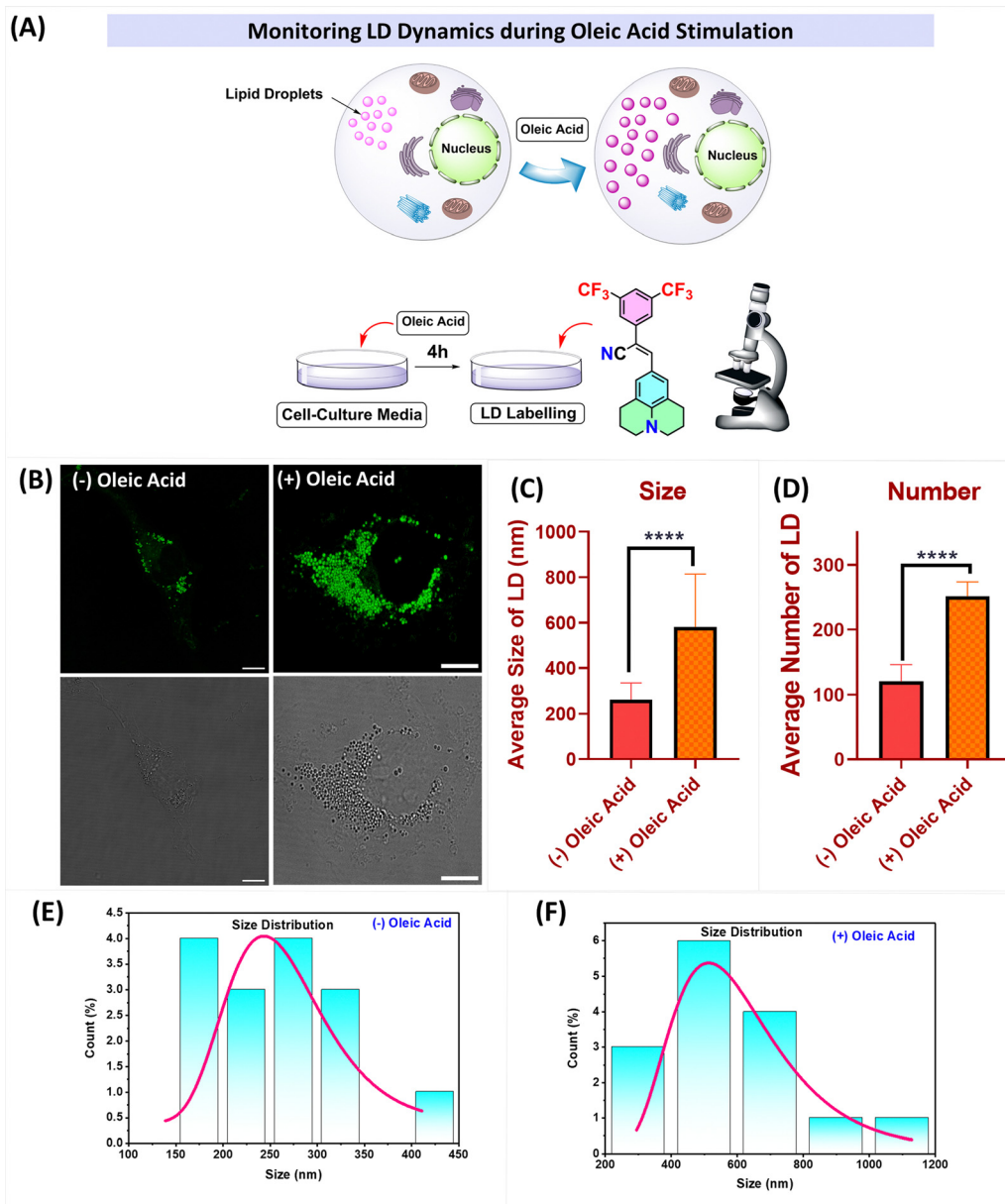


Fig. 8 (A) Pictorial representation of the changes observed after oleic acid stimulation; (B) (left) CLSM images of cells incubated with **J-2** (2 μ M), and (right) the images of oleic acid-treated COS-7 cells followed by incubation with **J-2** (2 μ M). (C and D) Plots representing the changes in LD dynamics with and without oleic acid stimulation: (C) size of LDs, and (D) number of LDs. (E and F) Size distribution histograms showing the sizes of the lipid droplets without (left) and with (right) oleic acid stimulation.

the formation of organogels stabilized by intermolecular interactions using fluorophores bearing two trifluoromethyl substituents.

Author contributions

RD performed the overall investigation and methodology, including the synthesis, fluorescence studies, organogelation studies, analysis of the cellular images and developing the first draft of the manuscript. DS and DR performed cell biology experiments, including colocalization of the probes and cell

viability assay. VS and SK supervised, planned, and conceived the idea and were involved in manuscript editing and writing.

Conflicts of interest

There are no conflicts to declare.

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