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# Fluorescent Detection of Nitrite in Hams Based on Carbon Dots Modify Sensing of Cyanide Ions in Water in Twin Modes

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## ABSTRACT

A novel, price effective, sensitive and environmentally friendly approach employing a physical mixture of carbon dots (CDs) and gold nanoparticles (AuNPs) is according for the detection of cyanide ions (CN-) in water. we tend to determined vital ending of visible radiation of CDs by AuNPs and also the restoration of visible radiation by the removal of AuNPs. A twin mode sensing approach for the estimation of CN- in water is incised out of this observation exploiting each the FRET between CDs and AuNPs and also the well-known gold natural action by CN-. this method doesn't involve any advanced chemistry or any tedious artificial routes and may be used for the sensing of cyanide in binary compound media. the tactic is sensitive up to a degree of one mgL-1 (1 ?g mL-1) of CN- in water and its presence/absence may also be visualized by eye by illuminating the answer exploitation light-weight at 365 nm.

Keywords: Carbon dot; Gold nanoparticle; Visible radiation quenching; Cyanide particle.

#### INTRODUCTION

Through ages, cyanide is that the most feared anions within the surroundings. The moraines and its action on living systems are well documented. CN- causes death of living organisms even at lower concentrations not like serious metal ions that have an effect on health solely at later stage by accumulating within the body. although CN- is very toxicant, it's wide used throughout the globe in varied sectors of trade. Accidental CN-unleash in waste or rivers could lead serious contamination of spring water and even drink [1]. the globe health organization fastened the most acceptable level of cyanide in drink at one.9 nine and also the us Environmental Protection Agency (EPA) has set the most stuff level (MCL) for cyanide in drink at zero.2 mgL.



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the intense toxicity of cyanide in physiological systems, yet because the continued environmental concern caused by its widespread industrial use has diode to the event of various analytical approaches for its detection [2].

Several strategies are according within the literature exploitation fluorimetry, spectrophotometry, activity, titrimetric, chemistry devices and chemodosimetery for the determination of cyanide [3-19]. various studies on AuNP based mostly quantitative analysis sensing of cyanide ions are according [20-23]. Most of those strategies want tedious artificial steps, use of organic solvents, refined instruments and aren't appropriate for on-site cyanide observation.

Semiconductor quantum dots (QDs) became one among the heavily studied materials in various fields because of its high emission quantum yields and size tunable emission profiles. Their serious toxicity even at comparatively low concentrations and restricted binary compound solubility prohibit their applications notably in bio and environmental domains. As compared to the well-established semiconductor QDs, carbon quantum dots (CDs) have attracted tremendous attention due to their fascinating properties, like non-blinking, water solubility, and nontoxicity. AuNPs are accepted as quenchers. This property, in fact, has been utilized in planning novel sensors using fluorophores like semiconductor QDs. visible radiation Resonance Energy Transfer (FRET) between QDs and AuNPs are used for the assay of biomolecules that inhibit the particular interactions between biomolecules. Ending of visible radiation of CDs by serious metal ions are explored and later on used as detection strategies for cations. Recently vital efforts associated with the development of sensors supported CDs are emerged. we tend to determine the ending of visible radiation of CDs by AuNPs.

We reasoned that quenched visible radiation might be revived by the straightforward removal of AuNPs. at the same time analyte induced natural action of Au may also be followed by the reduction in SPR peak of AuNPs. responsibleness associated applications of an analytical approach would be higher and wider if the measure is performed in additional than one mode. gift communication reports for the primary time a one-shot technique exploitation unmodified CDs and AuNPs for the detection and estimation of cyanide ions in water in twin modes by observation visible radiation and optical absorbance.

## EXPERIMENTAL PROCEDURE

#### **Materials**

Chloroauric acid (HAuCl4.3H2O), Trisodium change state, acid anhydrate, PEG organic compound and Rhodamine B were purchased from Sigma-Aldrich, Bangalore, India. cyanide was a present from the pharmacological medicine laboratory, medical specialty Technology Wing, SCTIMST. All alternative chemicals used were of analytical grade from Merck Bharat Ltd, Mumbai, India.

#### **Preparation of Gold Nanoparticles**

AuNPs were synthesized as according by Turkevich et al. Briefly, to a boiling answer of twenty mil of one.0-millimeter HAuCl4, a pair of mils of a tenth answer of trisodium change state dihydrate was another underneath constant stirring. The contents were far away from the new plate once the answer turned red. The cooled contents were unbroken underneath refrigeration till its usage.

#### Synthesis of Carbon Dots (CDs)

Carbon dots were synthesized as according by Goh et al. Briefly, nine mil of glycerin and 600 mg of PEG organic compound (MW ~1800) were taken during a hundred mil three-neck flask degassed with gas. once the temperature was raised to 250°C, 600 mg of acid was another and allowed to react at this temperature for 3hrs. The ensuing product was cooled to temperature and dialyzed against H2O employing an ester qualitative analysis membrane [molecular weight brings to a halt 3500) for two days so as to get rid of any unreacted



reactants. The refined CD was freeze dried and unbroken cold till use.

#### Instrumentation

The UV-Visible absorption spectra of the Au and CD-Au were recorded employing a UV-Visible photometer, Varian, Cary a hundred Bio, Melbourne, Australia and also the visible radiation intensity of the as ready CD and CD-Au were measured employing a Spectrofluorometer, Varian, Cary Eclipse model EL 0507, Melbourne, Australia. Fourier rework Infra-Red (FTIR) spectrum of CD was recorded within the vary 600-4000 cm-1 on a Nicolet 5700 FTIR mass spectrometer, Nicolet INC, Madison, USA employing a Diamond ATR accent. The technique of Dynamic light-weight Scattering (DLS), Malvern Instruments Ltd, UK was used for the determination of the letter potential of the CD, CD-Au and CD-Au- CN. High Resolution Transmission microscopy (HRTEM) was performed during a FEI, TECNAI S Twin research with associate fast voltage of a hundred kV. The sample solutions (CD) were ready by dispersion underneath associate supersonic vibrator. They were then deposited on a formvar coated copper grid and dried during a vacuum at temperature before observation. Transmission microscopy (TEM) pictures were obtained on a Hitachi, H 7650 magnifier, Hitachi, Tokyo, Japan. The sol was deposited onto a 200-mesh copper grid coated with a formvar film and dried long.

#### **Measurement of Fluorescent Quantum Yields of CDs**

The quantum yield (QY) of CDs made up our minds employing a comparative technique. the tactic involves multiple references with identified QYs and provides abundant high accuracy by hard the slope of the road generated by plotting the integrated visible radiation intensity against the absorbance for multiple concentrations of fluorophore. Rhodamine B in water (QY = 31%) was used because the customary to calculate the QY of the CDs. it's according that the answer absorbance at the excitation wavelength for either the sample or the quality mustn't exceed zero.1 to protect against any vital losses because of inner filtration effects. Thus, absorbance values chosen were zero.008, 0.03, 0.05 and 0.1. Emission spectra of the solutions were recorded employing a spectrofluorometer at associate excitation wavelength of 350 nm. Integrated visible radiation intensity against the absorbance and linear match curves we tend tore generated from that we acquire the slope values. Calculation was done supported the subsequent equation, Where m is that the slope of the road, n is that the index of refraction of the solvent, subscript R refers to the quantum yield of the quality. Here for each the quality and also the sample solvent is water; thus n2/nR2 is one.

#### **Detection of Cyanide in Water Samples**

Aqueous solution of cyanide (0.1 M) was ready and diluted one:10 to urge zero.01 M. one mil of CD answer (0.1 mg/ mL) was quenched with gold nanoparticles (AuNPs) (0.1 mg/ mL) and this CD-Au mixture is employed for the detection of CN-. totally different amounts of CN- starting from five mgL-1 (?gmL-1) to one hundred eighty mgL-1 were another to the visible radiation quenched CD-Au and measured each the decrease within the SPR absorption intensity of AuNPs and also the increase in visible radiation emission intensity of the CDs. Dilution issue has been taken into consideration in fixing the intensity of absorption/emission peaks. to substantiate the exactitude and recovery of the probe every set of experiment was distributed in triplicate and similar results inside the most error of 2-3% were obtained.

#### **RESULTS AND DISCUSSION**

We synthesized CDs as according elsewhere. The quantum yield of the CDs made up our minds to be nineteen.3%. FTIR spectrum, delineate, shows characteristic peaks of CDs as according by Goh et al. beside the sturdy C-H stretching vibrations of PEG at 2869 cm-1. The visible radiation of CDs (1 mil, 0.1 mg/mL) is quenched with one mil AuNP (0.1 mg/mL) and also the obtained mixture is called as CD-Au. The interaction of CD with AuNPs doesn't cause any amendment within the optical properties of the AuNPs as evident from



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spectrum. It's additionally clear that the CDs doesn't induce any aggregation of the AuNPs since the SPR most of the AuNPs isn't red shifted within the CD-Au and no further peak is determined on the far side the SPR peak. mass structures would have shown a color amendment or further peak. The decrease in intensity of the SPR absorption of AuNPs is because of the dilution impact caused by the addition of one mil of CD.

The HRTEM micrograph indicates that as ready CD has a mean size of 3-5 nm that agrees with the sooner literature (Figure 1A). the scale of the particles of CD-Au is ~14 nm as shown in TEM micrograph (Figure 1B). It seems that the spherical structures of larger sizes are that of AuNPs. TEM image of CD-Au on interaction with CN- is shown in Figure 1C. Most of the AuNPs [spherical particles in Figure 1B)] are leached away by CNas mirrored by the image. letter potential of CD is -20.9 mV and it's reduced to -3.4 mV for CD-Au. it's reasoned that the reduction is because of the interaction of CD and AuNPs. This feasibleness, in fact, is well mirrored within the increase in letter potential to -36 mV on interaction with CN- that is understood to get rid of Au particles (Table 1).

Sample code	Zeta Potential (mV)
CD	-20.9 ± 4.7
CD-Au	-3.4 ± 4.6
CD-Au-CN	-36.2 ± 6.4

**Table 1:** Change in Zeta potential of CDs when mixed with AuNPs and CN ions.

It is the visible radiation spectra of as ready CDs and CD-Au (Figure 1). It is seen that ~62% ending of the visible radiation emission of CDs were stricken by AuNPs. Complete ending of visible radiation can't be expected since the AuNPs are at random distributed within the medium. Photographic image of CD and CD-Au underneath ultraviolet radiation lamp shown within the inset of Figure 1 clearly replicates the ending of CD by AuNPs. The image is predominately red, characteristic color of AuNPs. On adding CN- to CD-Au mixture, AuNPs are quantitatively removed and at the same time visible radiation intensity of the CDs is reinstated so by enabling estimation of CN- ions in binary compound samples in twin modes.



Figure 1: (A) HRTEM image of CDs. (B) TEM images of CD-Au. (C) TEM images of CD-Au-CN.

The photographic pictures underneath ultraviolet radiation lamp delineates A-E show the ending of the visible radiation of CDs by AuNPs and also the pictures show the restoration of the visible radiation on adding CN-. The photographic pictures of CD-Au and CD-Au-CN underneath actinic radiation. It is seen that CN- fully leaches away the AuNPs (the red color is pale in B) more supporting the TEM depicts the rise in intensity of visible radiation emission of CD-Au and also the decrease in absorbance of the SPR of the AuNPs on adding



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CN- visible radiation intensity is revived proportionately with concomitant reduction within the SPR absorbance.

Interestingly the rise in visible light and therefore the reduction in absorbance square measure found to be linear with relevance the concentration of CN- ions. This technique may be accustomed observe CN-concentration as low as one mg/L (1 ?g/mL) in water and its presence/absence may be envisioned by optic by illuminating the answer victimization light-weight at 365 nm.

We found that unremarkably occurring anions (1000 mgL-1) neither increased the visible light nor reduced the SPR intensity as shown in Figure 2 apparently suggesting that common anions didn't have any adverse influence on the detection of CN ions. many authors have rumored such chromofluorogenic detection ways for the sensing of CN ions. However, a twin sensing technique for the CN particle victimization an equivalent probe isn't found however. Yadav and Singh have rumored a twin ion sensing technique for arsenite and cyanide ions.



Figure 2: (A) Fluorescence emission (B) SPR absorption of CD-Au on interaction with anions.

We used this technique for checking cyanide contamination in water samples collected from our field and conjointly outside the field. None of the samples showed any detectable amount of CN- but water spiked with one hundred mgL-1 showed the restoration of visible light intensity and therefore the corresponding decrease within the SPR absorbance of the CD-Au as delineate.

Earlier reports have shown that visible light of the many fluorophores as well as conjugated polymers may be quenched by the addition of trace amounts of AuNPs. These studies indicate that the mechanism of extinguishing is predominately by energy transfer (FRET). We've got not investigated the potential mechanism of extinguishing of visible light of CDs by AuNPs, we have a tendency to reasoned that FRET is operative here considering the very fact that there's vital spectral overlap of the visible light of CDs and therefore the absorption spectra of AuNPs.

## CONCLUSION

In conclusion, a twin sensing approach for the estimation of CNin water is rumored for the primary time exploiting each the FRET between CDs and AuNPs and therefore the well-known gold natural process by CN-the strategy doesn't involve any complicated chemistry or any tedious artificial routes and might be used for the sensing of cyanide presence in binary compound media. the strategy is sensitive up to a degree of one mg/L (1 ?gmL-1) of CN- in water and thus may be used for the preliminary checking of cyanide contamination in water. the strategy appears to own wider prospects since the analysis may be performed in twin modes.



### REFERENCES

1. Christison T, Rohrer J (2007) Direct determination of free cyanide in drinking water by ion chromatography with pulsed amperometric detection. J Chromatogr A 1155: 31-39.

2. Lee H, Chung Y, Ahn K (2008) Selective fluorescence sensing of cyanide with an o-(carboxamido) trifluoroacetophenone fused with a cyano-1,2-diphenylethylene fluorophore. Tetrahedron Letters49: 5544-5547.

3. Badugua R, Lakowicza J, Geddes J (2005) Cyanide-sensitive fluorescent probes. Dyes and Pigments64: 49-55.

4. Badugua R, Lakowicza J, Geddes C (2004) Excitation and emission wavelength ratiometric cyanidesensitive probes for physiological sensing. Analytical Biochemistry 327: 82-90.

5. Badugua R, Lakowicza J, Geddes C (2005) Enhanced fluorescence cyanide detection at physiologically lethal levels: Reduced ICT-Based signal transduction. J Am Chem Soc 127: 3635-3641.

6. Lee K , Huh J, Kim T, Do Y, Lee M (2008) Electron-beam-induced transition aluminas from aluminum trihydroxide. Scriptamat 59: 1022-1025.

7. Gimeno N, Li X, Durrant J, Vilar R (2008) Cyanide sensing with organic dyes: Studies in solution and on nanostructured Al2O3 Surfaces. Chem. Eur J 14: 3006-3012.

8. Legako J, White B, Harmon H (2003) Rapid reagent-less detection of competitive inhibitors of butyrylcholinesterase. Sensors and Actuators B: Chemical 91: 138-142.

9. Gee H, Lee C, Jeong Y, Woo-Dong J (2011) Highly sensitive and selective cyanide detection via Cu2+ complex ligand exchange. Chem. Commun 47: 11963-11965.

10. Chung S, Nam S , Lim J, Park S, Yoon J (2009) A highly selective cyanide sensing in water via fluorescence change and its application to in vivo imaging. Chem Commun (Camb) 28: 2866-2868.

11. Ajayakumar M, Mukhopadhyay P, Yadav S, Ghosh S (2010) Single-electron transfer driven cyanide sensing: A new multimodal approach. Org Lett 12: 2646-2649.

12. Kim H , Lee H, Lee J, Choi D, Jung J, et al. (2011) Bisindole anchored mesoporous silica nanoparticles for cyanide sensing in aqueous media. Chem Commun 47: 10918-10920.

13. Mashraqui S, Betkar R, Chandiramani M, Estarellas C, Frontera A (2011) Design of a dual sensing highly selective cyanide chemodosimeter based on pyridinium ring chemistry. New J Chem 35: 57-60.

14. Senapati D, Dasary S, Singh A, Senapati T, Yu H, et al. (2011) A label •based sers assay for direct cyanide detection at the parts •trillion level. Chem Eur J17: 8445-8451.

15. Liu C, Tseng W (2011) Colorimetric assay for cyanide and cyanogenic glycoside using polysorbate 40-stabilized gold nanoparticles. Chem Commun 47: 2550-2552.

16. Kim M, Kim S, Jang H, Yi S, Seo S, et al. (2010) A gold nanoparticle-based colorimetric sensing ensemble for the colorimetric detection of cyanide ions in aqueous solution. Tetrahedron Letters 51: 4712-4716.

17. Zhang Z, Zhang J, Qu C, Pan D, Chen Z, et al. (2012) Label free colorimetric sensing of thiocyanate based



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on inducing aggregation of Tween 20-stabilized gold nanoparticles. Analyst 137: 2682-2686.

18. Zhou L, Lin Y, Huang Z, Ren J, Qu X (2012) Carbon nanodots as fluorescence probe for rapid, sensitive, and label-free detection of Hg2+ and biothiols in complex matrices. Chem Commun 48: 1147-1149.