



*Citation for published version:*

Haque, A, Al Balushi, RA, Al-Busaidi, IJ, Ilmi, R, Al Rasbi, N, Jayapal, M, Khan, MS & Raithby, PR 2019, 'Synthesis, optical spectroscopy, structural, and DFT studies on dimeric iodo-bridged Copper(I) complexes', *Journal of Organometallic Chemistry*, vol. 892, pp. 75-82. <https://doi.org/10.1016/j.jorganchem.2019.04.017>

*DOI:*

[10.1016/j.jorganchem.2019.04.017](https://doi.org/10.1016/j.jorganchem.2019.04.017)

*Publication date:*

2019

*Document Version*

Peer reviewed version

[Link to publication](#)

*Publisher Rights*

CC BY-NC-ND

**University of Bath**

**Alternative formats**

If you require this document in an alternative format, please contact:  
[openaccess@bath.ac.uk](mailto:openaccess@bath.ac.uk)

**General rights**

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

**Take down policy**

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

# Synthesis, Optical Spectroscopy, Structural, and DFT Studies on Dimeric Iodo-bridged Copper(I) complexes

Ashanul Haque,<sup>a,b\*</sup> Rayya A. Al Balushi,<sup>c</sup> Idris Juma Al-Busaidi,<sup>b</sup> Rashid Ilmi<sup>b</sup>, Nawal Al Rasbi,<sup>b</sup> Maharaja Jayapal,<sup>b</sup> Muhammad. S. Khan<sup>b,\*</sup> Paul R. Raithby<sup>d\*</sup>

<sup>a</sup> Department of Chemistry, College of Science, University of Hail, Ha'il 81451, Kingdom of Saudi Arabia.

E-mail: [a.haque@uoh.edu.sa](mailto:a.haque@uoh.edu.sa)

<sup>b</sup> Department of Chemistry, Sultan Qaboos University, P.O. Box 36, Al-Khod 123, Sultanate of Oman.

E-mail: [msk@squ.edu.om](mailto:msk@squ.edu.om)

<sup>c</sup> Department of Basic Sciences, College of Applied and Health Sciences, A'Sharqiyah University, P.O. Box 42, Ibra 400, Sultanate of Oman.

<sup>d</sup> Department of Chemistry, University of Bath, Claverton Down, Bath BA2 7AY, UK.

E-mail: [p.r.raithby@bath.ac.uk](mailto:p.r.raithby@bath.ac.uk)

## Abstract:

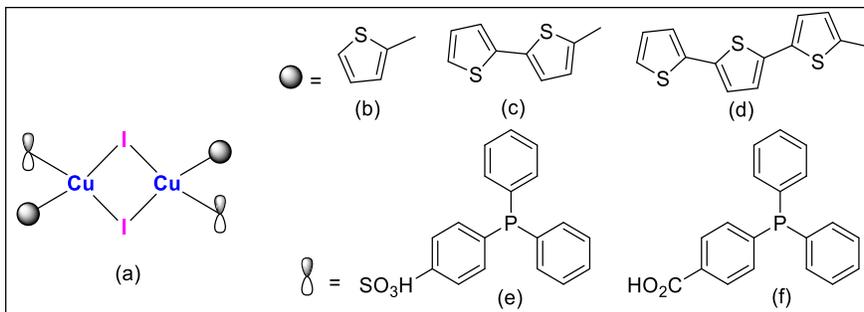
Three new iodo-bridged copper(I) complexes of the type  $[\text{CuI}(\text{PPh}_3)\text{L}]_2$ , where  $\text{L} = \text{Ar}-\text{C}\equiv\text{C}-\text{C}_5\text{H}_4\text{N}$ ,  $\text{Ar} =$  phenyl (**C**<sub>1</sub>), biphenyl (**C**<sub>2</sub>) and fluorenyl (**C**<sub>3</sub>) have been synthesized *via* coordination-driven self-assembly processes. Two of Cu(I) complexes, **C**<sub>2</sub> and **C**<sub>3</sub>, have been characterized by single-crystal X-ray diffraction studies. The complexes have two molecules of the P-donor ligand and two molecules of the N-donor ligand in *trans* configurations, supporting the central  $\text{Cu}_2\text{I}_2$  unit. Absorption properties of the complexes have been investigated. Extensive DFT calculation has been carried out to delineate the influence of aromatic spacers on the optical properties and the nature of excited states. The ease of synthesis of these Cu(I) dimers and the wide range of ethynylpyridine supporting ligands that can be incorporated highlights the potential for these materials to form polymers by linking through the ethynylpyridine ligands.

**Keywords:** Copper(I) complexes; Ethynylpyridine; DFT studies; Self-assembly; X-ray crystal structures

## 1. Introduction

Transition metal complexes have attracted significant attention due to their diverse photophysical properties and applications.[1-5] The last four decades has witnessed a great upsurge in the design and development of  $d^{10}$  ion-based complexes such as those of copper(I) bearing homo- and heteroleptic ligands.[6, 7] Among luminescent transition metal complexes, copper(I)-based systems offer easy synthetic routes, a diverse range of structural arrangements, bright luminescence encompassing a broad range of the visible and UV spectrum, high quantum yields, etc. It has been demonstrated that neutral copper(I) complexes bearing N- and P-coordinating ligands surrounding a halogenido-bridged  $\text{Cu}_2\text{X}_2$  ( $\text{X} = \text{halogen}$ ) core are good candidates for optoelectronic applications.[8] By simple variation of the coordination sphere (i.e. the coordinating and bridging ligands) around the metal, both absorption and emission can be manipulated. For instance, it has been demonstrated that metal to ligand charge transfer (MLCT) or halide to ligand charge transfer (XLCT) emission bands can be significantly tuned by varying the  $\pi^*$  level of ligands.[8] Because of these features, several new copper(I) complexes have been synthesized and tested as components for photovoltaics (PVs), organic-light emitting diodes (OLEDs), sensors, etc.[7, 9] The low toxicity and the pseudo-tetrahedral geometry of the complexes has also prompted researchers to assess their biological activity.[3]

Within the class of  $\text{Cu}_2\text{X}_2$  complexes, synthetic manipulation can readily be achieved by altering the nature of the supporting ligands, usually with donor N- or P-bridgehead atoms.[10-12] The pyridine nitrogen donor atom has a strong tendency to interact with a variety of metals and, as a result, a number of complexes bearing simple to functionalized pyridine (or higher pyridines) have been reported.[3] Not only does the electron-deficient pyridine endow structural rigidity, but a judicious choice of functionalization and macrocyclization through pyridine core can also impart helicity/chirality to the resulting complexes.[13] Bearing these features in mind, we have investigated structural and photophysical behavior of a number of acetylide-functionalized pyridinyl copper(I) complexes. Both dimeric and tetrameric copper(I) complexes bearing ethynylpyridinyl-based coordinating ligands have been investigated.[5, 14, 15] For example, we demonstrated that copper(I) complexes bearing pyridine ligands connected to 4-ethynyl ferrocene show intriguing electrochemical properties. In such complexes, the ferrocene units act as an electron reservoir/sink and was detrimental to the emission properties.[15] Similarly, we recently demonstrated that dye sensitized solar cells (DSSC) based on  $\text{Cu}_2\text{X}_2$  bridged complexes with heteroarylethynyl attached to 4-position of pyridine (**Chart 1**) are highly emissive and show up to 2% light to energy conversion efficiency.[5] The performance of the dye was found to depend significantly on two factors: (i) nature of the aryl group attached to pyridine *via*  $\pi$ -linker; and (ii) presence/absence of an anchoring group associated with the phosphine ligands.



**Chart 1:** Recently reported acetylide-functionalized pyridine-based Cu(I) complexes for PV applications.

To examine the effect of replacing heterocyclic spacers by carbocyclic spacers (such as phenyl, biphenyl and fluorene), we now report the synthesis, the structures and photophysical and DFT studies on three new iodo-bridged copper(I) complexes supported by ethylpyridine-based ligands. The complexes were prepared *via* self-assembly in the solution phase under mild conditions.

## 2. Experimental

### 2.1 General

All chemicals and reagents, except where stated otherwise, were obtained commercially and used without further purification. Unless otherwise stated, syntheses were performed under a dry argon atmosphere. Solvents were pre-dried and distilled before use by standard procedures.[16] NMR spectra were recorded on a Bruker AM-400 spectrometer in  $\text{CDCl}_3$ .  $^1\text{H-NMR}$  spectra were referenced to solvent resonances and  $^{31}\text{P}\{^1\text{H}\}$  NMR spectra were referenced to external phosphoric acid. IR spectra were recorded directly on the sample as attenuated total reflectance (ATR) on Diamond using Cary 630 FT-IR spectrometer. UV-vis spectra were recorded with an Agilent Cary 5000 UV-visible spectrophotometer using a quartz cuvette with a 1 cm path length. Column chromatography was performed on either Kieselgel 60 (230 – 400 mesh) or celite.

### 2.2 General protocol for the synthesis

#### 2.2.1 Synthesis of ligands ( $\text{L}_1\text{-L}_3$ ) and complexes ( $\text{C}_1\text{-C}_3$ )

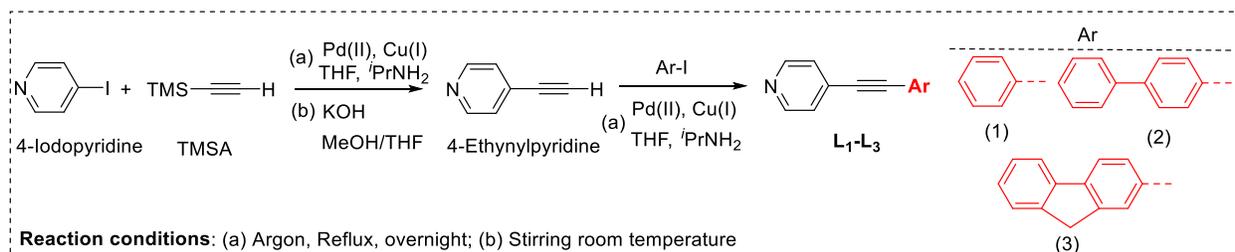
The intermediates and final products were obtained *via* established cross-coupling reactions.[5, 15, 17] Briefly, 4-trimethylsilyl ethynylpyridine was obtained by reacting an equimolar quantity of 4-iodopyridine (2.0 mmol) with trimethylsilyl ethyne (TMSE, 2.0 mmol) in THF/ $i\text{PrNH}_2$  using Pd(II)/Cu(I) catalysts. (**Scheme 1**) The crude product obtained was purified by silica column chromatography using hexane/dichloromethane eluant systems. This was followed by a protodesilylation reaction under basic condition (aq. KOH in MeOH/THF) and the 4-ethynylpyridine was separated by silica column chromatography using pure dichloromethane as eluant. In the next phase of the reaction, respective aryl halides (2.0 mmol) and 4-ethynylpyridine (2.0 mmol) were coupled *via* a Sonogashira cross-coupling reaction. Final products were purified using an alumina column and obtained as tan to light brown solid in quantitative yields.

*Phenylethynylpyridine (L<sub>1</sub>):* IR ( $\nu_{\max}$ )  $\text{cm}^{-1}$ : 2199 ( $\text{-C}\equiv\text{C-}$ ), 1598 (C-N),  $^1\text{H-NMR}$  (700 MHz,  $\text{CDCl}_3$ ):  $\delta$ (ppm) 8.45 (d, 2H,  $J = 6.4$ , H-py), 7.98 (d, 2H,  $J = 5.8$ , H-py), 7.35 (d, 2H,  $J = 6.2$ , H-Ph), 6.70-6.68 (m, 3H, H-Ph). ESI-MS:  $m/z$  179.9 (M<sup>+</sup>).  $\text{C}_{13}\text{H}_9\text{N}$  Anal. Calc.: C, 87.12; H 5.06; N, 7.82%. Found: C, 86.97; H, 4.94; N, 7.77%.

*Biphenylethynylpyridine (L<sub>2</sub>):* IR ( $\nu_{\max}$ )  $\text{cm}^{-1}$ : 2208 ( $\text{-C}\equiv\text{C-}$ ), 1605 (C-N),  $^1\text{H-NMR}$  (700 MHz,  $\text{CDCl}_3$ ):  $\delta$ (ppm) 8.28 (d, 2H,  $J = 5.95$  Hz, H-py), 7.68 (d, 2H,  $J = 4.55$  Hz, H-py), 7.62-7.60 (m, 2H, H-biph), 7.46 (dd, 4H,  $J = 5.53, 13.51$  Hz, H-biph), 7.38-7.35 (m, 3H, H-biph). ESI-MS:  $m/z$  256.1 (M<sup>+</sup>).  $\text{C}_{19}\text{H}_{13}\text{N}$  Anal. Calc.: C, 89.38; H, 5.13; N, 5.49%. Found: C, 8.98; H, 4.83; N, 5.02%.

*Fluorenylethynylpyridine (L<sub>3</sub>):* IR ( $\nu_{\max}$ )  $\text{cm}^{-1}$ : 2205 ( $\text{-C}\equiv\text{C-}$ ), 1608 (C-N),  $^1\text{H-NMR}$  (700 MHz,  $\text{CDCl}_3$ ):  $\delta$ (ppm) 8.61 (d, 2H,  $J = 5.74$  Hz, H-py), 8.27 (d, 2H,  $J = 5.95$  Hz, H-py), 7.81-7.79 (m, 2H, H-fluorene), 7.78 (s, 1H, H-fluorene), 7.73 (m, 2H, H-fluorene), 7.59-7.57 (m, 2H, H-fluorene), 3.93 (s, 2H). ESI-MS:  $m/z$  268.1 (M<sup>+</sup>).  $\text{C}_{20}\text{H}_{13}\text{N}$  Anal. Calc.: C, 89.86; H, 4.90; N, 5.24%. Found: C, 88.26; H, 4.49; N, 4.94%.

### Scheme 1: Synthesis of 4-arylethynylpyridine ligands (L<sub>1</sub>-L<sub>3</sub>).



### 2.2.2 Synthesis of Cu(I) complexes (C<sub>1</sub>-C<sub>3</sub>)

Cu(I) complexes were synthesized following the protocol reported in literature.[5, 15, 17] Complexes **C<sub>1</sub>-C<sub>3</sub>** were obtained by stirring a mixture of the appropriate arylethynylpyridine ligands (1.0 mmol), CuI (1.0 mmol) and triphenylphosphine (1.0 mmol) in an equimolar mixture in degassed dichloromethane. (**Scheme 2**) After stirring for 24 h at room temperature, the mixture was filtered, and the filtrate was concentrated under reduced pressure to produce neutral dimeric complexes **C<sub>1</sub>-C<sub>3</sub>**. All complexes are stable to light and air and were fully characterized. Crystals suitable for X-ray analysis were obtained by slow diffusion of dichloromethane (DCM) in hexane.

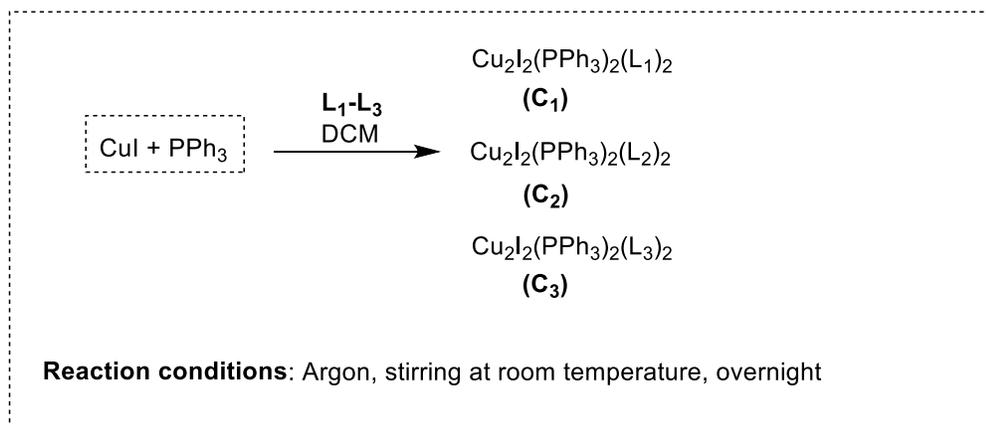
Complex  $[\text{L}_1(\text{CuI})_2(\text{PPh}_3)_2]$  (**C<sub>1</sub>**): IR: 2206  $\text{cm}^{-1}$   $\nu(\text{C}\equiv\text{C})$ , 1488  $\text{cm}^{-1}$   $\nu(\text{C}=\text{N})$ .  $^1\text{H-NMR}$  (700 MHz,  $\text{CDCl}_3$ ):  $\delta$ (ppm) 8.70 (d,  $J = 5.98$  Hz, 4H, H-py), 7.91 (d,  $J = 6.3$  Hz, 8 H, H-py and  $\text{PPh}_3$ ), 7.71-7.42 (m, 36 H, H-Ph and  $\text{PPh}_3$ ).  $^{31}\text{P-NMR}$  (121.53 MHz,  $\text{CDCl}_3$ ):  $\delta = 29.18$  (s,  $\text{PPh}_3$ ) ppm. ESI-MS:  $m/z$  1261.9 (M<sup>+</sup>).  $\text{C}_{62}\text{H}_{48}\text{Cu}_2\text{I}_2\text{N}_2\text{P}_2$  Anal. Calc.: C, 58.92; H, 3.83; N, 2.22%. Found: C, 59.55; H, 4.01; N, 2.17%

Complex  $[\text{L}_2(\text{CuI})_2(\text{PPh}_3)_2]$  (**C<sub>2</sub>**): IR: 2210  $\text{cm}^{-1}$   $\nu(\text{C}\equiv\text{C})$ , 1489  $\text{cm}^{-1}$   $\nu(\text{C}=\text{N})$ .  $^1\text{H-NMR}$  (700 MHz,  $\text{CDCl}_3$ ):  $\delta$ (ppm) 8.65 (d,  $J = 4.48$  Hz, 4 H, H-py), 7.62 (d,  $J = 10.5$  Hz, 12 H, H-py and  $\text{PPh}_3$ ), 7.55-7.34 (m, 40 H,

H-biph and PPh<sub>3</sub>). <sup>31</sup>P NMR (121.53 MHz, CDCl<sub>3</sub>): δ = 29.22 (s, PPh<sub>3</sub>) ppm. ESI-MS: m/z 1413.7 (M<sup>+</sup>). C<sub>74</sub>H<sub>56</sub>Cu<sub>2</sub>I<sub>2</sub>N<sub>2</sub>P<sub>2</sub>: Anal. Calc.: C, 62.76; H, 3.99; N, 1.98%. Found: C, 64.88; H, 4.11; N, 2.01%.

Complex [L<sub>3</sub>(CuI)<sub>2</sub>(PPh<sub>3</sub>)<sub>2</sub>] (**C**<sub>3</sub>): IR: 2208 cm<sup>-1</sup> ν(C≡C), 1493 cm<sup>-1</sup> ν(C=N). <sup>1</sup>H-NMR (700 MHz, CDCl<sub>3</sub>): δ(ppm) 8.64 (m, 4 H, H-py), 7.81–7.79 (m, 4 H, H-py), 7.74–7.57 (m, 30 H, H-flourene and PPh<sub>3</sub>), 7.52–7.34 (m, 14 H, H-flourene and PPh<sub>3</sub>), 3.94 (s, 2H). <sup>31</sup>P NMR (121.53 MHz, CDCl<sub>3</sub>): δ = 29.31 (s, PPh<sub>3</sub>) ppm. ESI-MS: m/z 1440.2 (M<sup>+</sup>). C<sub>76</sub>H<sub>56</sub>Cu<sub>2</sub>I<sub>2</sub>N<sub>2</sub>P<sub>2</sub> Anal. Calc.: C, 63.38; H, 3.92; N, 1.95%. Found: C, 64.88; H, 4.19; N, 2.01 %.

### Scheme 2: Synthesis of Cu(I) complexes (**C**<sub>1</sub>–**C**<sub>3</sub>).



## 2.3 X-ray Crystal Structure Determination

Single-crystal X-ray diffraction experiments were performed at 150(2) K on an Oxford Diffraction Gemini A Ultra CCD diffractometer monochromatic Mo-K $\alpha$  radiation ( $\lambda = 0.71073 \text{ \AA}$ ). The sample temperature was controlled using an Oxford Diffraction Cryojet apparatus; CrysAlis Pro was used for the collection of frames of data, indexing reflections and determining lattice parameters.[18] Structures were solved by direct methods using SIR-92[19] and refined by full-matrix least-squares, on  $F^2$ , using SHELXL2014[20] within the OLEX2 suite of programs.[21] A multi-scan absorption correction was applied in all cases. The hydrogen atoms were generated geometrically and refined with common isotropic thermal parameters. Crystallographic data for the complexes is presented in **Table 1**. In the case of **C**<sub>3</sub>, the fluorene rings showed significant disorder that could not be modelled effectively. As a last resort the unit was assigned isotropic displacement parameters and the carbon atoms were assigned full occupancies. While the refinement was stable with this model several significant residual electron density peaks in the vicinity of the ligand remained. The cif files for the two structures have been deposited with the CCDC: *CCDC 1898577-1898578*

## 2.4 Computational studies

### 2.5 Computational details

All DFT calculations were performed using the Gaussian 09[22] package with the Lee–Yang–Parr correlation function (B3LYP).[23] For copper and iodide, the Hay and Wadt basis-set LANL2DZ pseudopotential.[24, 25] was used. The 6-311+G(d).[26] basis-set was employed for N and P, while the

6-31G(d,p)[27] basis-set was operated for the atomic orbital of C, H to calculate all the optimized structures. TD-DFT calculations were performed using the same functional and the polarizable continuum model, CPCM with dichloromethane ( $\epsilon = 8.93$ ) as the surrounding solvent considering bulk solvent effects.[28] TD-DFT-calculated electronic spectra were depicted by using GaussSum,[29] All the molecular orbitals and spin-density diagrams were plotted using the Chemcraft Visualization program.[30]

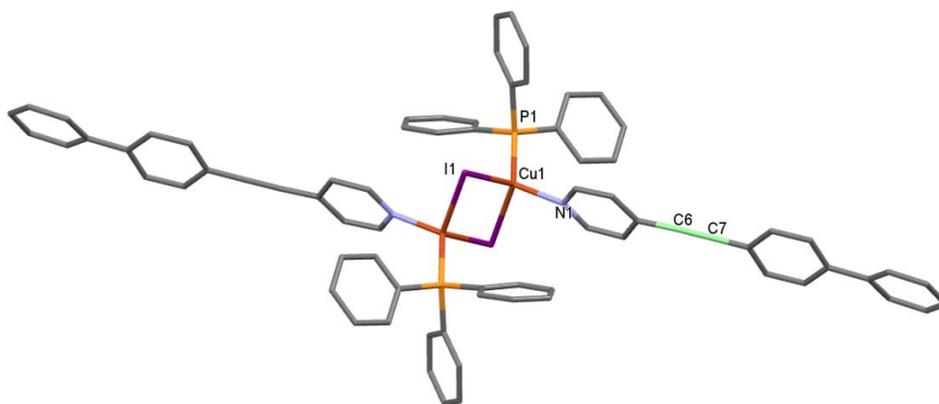
### 3. Result & Discussion

#### 3.1 Synthesis and Spectroscopic Characterization

The ligands **L**<sub>1</sub>–**L**<sub>3</sub> and their corresponding complexes **C**<sub>1</sub>–**C**<sub>3</sub> have been synthesized by adapting previously reported procedure.[5] Briefly, arylhalides (4-iodophenyl, 2-bromobiphenyl or 2-iodofluorenyl) were cross-coupled with TMSA using Sonogashira coupling reaction followed by protodesilylation giving the terminal alkynes (**Scheme 1**). The deprotected ligands obtained were then coupled with 4-iodopyridine to produce ligands **L**<sub>1</sub>–**L**<sub>3</sub> as yellow-brown powders in 60-78% yield. Analytical data (IR, NMR and MS) confirmed the formulation of the ligands. For example, TMSA protection of 4-iodopyridine was confirmed by the observed  $\text{—C}\equiv\text{C—}$  stretching frequency ( $\sim 2165 \text{ cm}^{-1}$ ) while deprotection was confirmed by the presence of the  $\text{—C}\equiv\text{H}$  stretching frequency ( $\sim 3223 \text{ cm}^{-1}$ ) in addition to the  $\text{—C}\equiv\text{C—}$  band. Moreover, the  $\nu_{(\text{C}\equiv\text{C})}$  value of 4-ethynylpyridine was found to be lower than those of the 4-(trimethylsilylethynyl)pyridine due to the fact that terminal alkynes ( $\text{HC}\equiv\text{C—R}$ ) have lower  $\nu_{\text{C}\equiv\text{C}}$  stretching frequencies than their protected counterparts  $\text{RC}\equiv\text{CR}$ .[31] The complexes **C**<sub>1</sub>–**C**<sub>3</sub> were obtained by swift solution phase self-assembled coordination. The reaction of the ligand (**L**<sub>1</sub>–**L**<sub>3</sub>), with copper iodide and triphenyl phosphine (1:1:1 ratio), in dry dichloromethane, under an argon atmosphere produced the complexes **C**<sub>1</sub>–**C**<sub>3</sub>. In addition to the standard analytical techniques, single crystal X-ray diffraction studies provided the full three-dimensional molecular and crystal structures of the three complexes.

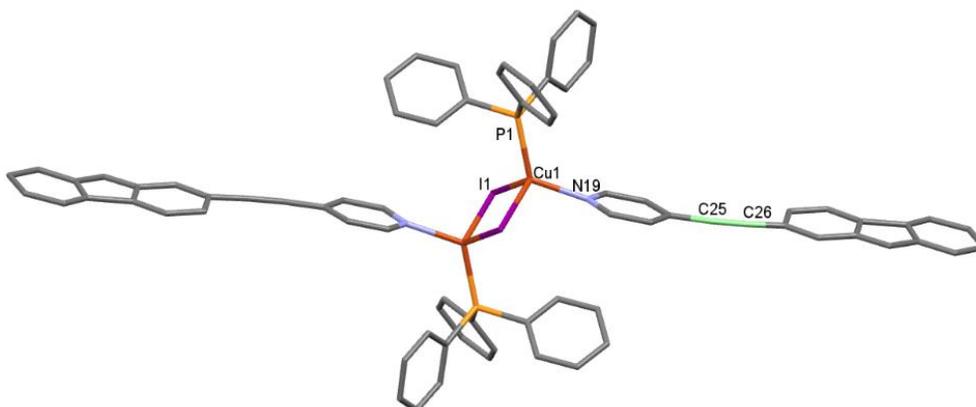
#### 3.2 Structural characterization

The crystal structures of the complexes: **C**<sub>2</sub> and **C**<sub>3</sub> were determined and are shown in **Figures 1** and **2**. Crystal data is summarized in **Table 1** and bond parameters are listed in **Tables ST1-ST4** (*supplementary information*). Single crystals of the complexes were obtained by slow diffusion of hexane into a dichloromethane solution of each complex.



**Figure 1.** The crystal structure of the dimeric complex **C<sub>2</sub>**. Hydrogen atoms are omitted for clarity.

The crystal structures of the dimeric complexes **C<sub>2</sub>** and **C<sub>3</sub>** have a similar planar, parallelepiped  $\text{Cu}_2\text{I}_2$  core and a *trans* arrangement of the two phosphine and the two **L** ligands. In each case the center of the  $\text{Cu}_2\text{I}_2$  core sits on a crystallographic center of symmetry, so that the crystallographic asymmetric unit contains half of a dimeric molecule. Each Cu(I) center is in a tetrahedral coordination environment consisting of two bridging iodo ligands, one  $\text{PPh}_3$  and the bridgehead nitrogen atom of **L<sub>2</sub>** and **L<sub>3</sub>**. The unique bond distance for Cu-I, Cu-P and Cu-N are 2.6487(4), 2.2412(6) and 2.063(2) Å, respectively, in **C<sub>2</sub>** and are 2.651(8), 2.235(2) and 2.059(6) Å, respectively, in **C<sub>3</sub>**. These bond distances are similar to the equivalent bond lengths in a family of previously reported  $[(\text{Py})(\text{Ph}_3\text{P})\text{CuX}]_2$  adducts.[32] The angles at the copper center in the  $\text{Cu}_2\text{I}_2$  core are obtuse, e.g. 111.45(1)° and 108.13(3)° in **C<sub>2</sub>** and **C<sub>3</sub>**, respectively. While the angle at the iodine center is acute, e.g. 68.54(1)° and 71.83(3)° in **C<sub>2</sub>** and **C<sub>3</sub>**, respectively. The Cu...Cu separations in the dimers: **C<sub>2</sub>** (3.0196(5) Å) and **C<sub>3</sub>** (3.1394(5) Å), are too long for a formal Cu-Cu bonding interaction; however, it remains similar to reported analogues.



**Figure 2.** The crystal structure of the dimeric complex **C<sub>3</sub>**. Hydrogen atoms are omitted for clarity.

**Table 1.** Crystallographic data for **C<sub>2</sub>-C<sub>3</sub>**

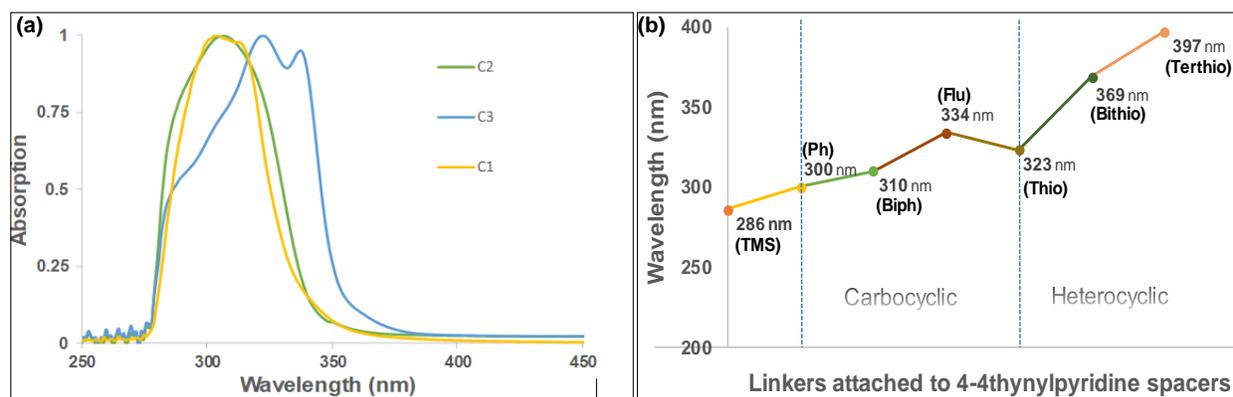
	<b>C<sub>2</sub></b>	<b>C<sub>3</sub></b>
<b>Empirical formula</b>	$\text{C}_{74}\text{H}_{56}\text{Cu}_2\text{I}_2\text{N}_2\text{P}_2$	$\text{C}_{76}\text{H}_{56}\text{Cu}_2\text{I}_2\text{N}_2\text{P}_2$

<b>Formula weight</b>	1416.02	1440.04
<b>T(K)</b>	150(2)	150(2)
<b><math>\lambda</math> (Å)</b>	0.71073	0.71073
<b>Crystal system</b>	Monoclinic	Monoclinic
<b>Space group</b>	<i>P2<sub>1</sub>/n</i>	<i>P2<sub>1</sub>/n</i>
<b>a</b>	9.43370(10)	9.3125(2)
<b>b</b>	22.5428(3)	22.1452(4)
<b>c</b>	14.2331(2)	15.2590(3)
<b><math>\alpha</math></b>	90°	90
<b><math>\beta</math></b>	98.562(1)	101.061(2)
<b><math>\gamma</math></b>	90	90
<b>V (Å<sup>3</sup>)</b>	2993.11(7)	3088.36(11)
<b>Z</b>	2	2
<b><math>\rho_{\text{calc.}}</math> Mg m<sup>-3</sup></b>	1.571	1.549
<b><math>\mu</math> (Mo-K<math>\alpha</math>) (mm<sup>-1</sup>)</b>	1.842	1.787
<b>F(000)</b>	1416	1440
<b>Crystal size (mm)</b>	0.60 x 0.40 x 0.40	0.2 x 0.2 x 0.04
<b><math>\theta</math> range (deg)</b>	2.834 to 31.192	2.971 to 29.689
<b>Reflections collected</b>	79295	40941
<b>Independent reflections [R(int)]</b>	9128 [0.0375]	7906 [0.0385]
<b>Completeness to <math>\theta = 25.242^\circ</math></b>	99.90%	99.80%
<b>Max. and min. transmission</b>	1.00 and 0.730	1.000 and 0.9209
<b>Data / restraints / parameters</b>	9128 / 0 / 370	7906 / 0 / 302
<b>Goodness-of-fit on F<sup>2</sup></b>	1.133	1.034
<b>Final R<sub>1</sub>, wR<sub>2</sub> indices [I&gt;2<math>\sigma</math>(I)]</b>	0.0305, 0.0584	0.0719, 0.1870
<b>R indices (all data)</b>	0.0454, 0.0647	0.0927, 0.2013
<b>Largest diff. peak, hole (e.Å<sup>-3</sup>)</b>	1.101, -0.516	3.528, -1.267

### 3.3 Optical Spectroscopy

The electronic spectra of complexes **C**<sub>1</sub>–**C**<sub>3</sub> were measured in dichloromethane at room temperature (**Figure 3a**). The onset of absorption for all ethylpyridinyl-based ligands and their corresponding Cu complexes were red-shifted compared to ligand precursor 4-(trimethylsilylethynyl)pyridine ( $\lambda_{\text{max}} = 286$  nm). The complex incorporating the fluorenylethynylpyridine ligand (**C**<sub>3</sub>) showed red-shifted band compared to phenyl and biphenylethynylpyridine Cu(I) complexes (**C**<sub>1</sub> and **C**<sub>2</sub>). Complex **C**<sub>1</sub> showed a strong absorption band at 300 nm ( $\lambda_{\text{max}}$ ) whereas complexes **C**<sub>2</sub> and **C**<sub>3</sub> showed absorptions peaking at 310 and 334 nm, respectively. This can be ascribed to the increasing  $\pi$ -conjugation length leading to a lower energy for the complex and absorption at longer wavelengths. The absorption peaks in the UV spectrum ( $\lambda_{\text{max}} < 400$  nm) can be attributed to the  $\pi \rightarrow \pi^*$  transition associated with the arylethynylpyridine group. **Figure 3b** shows the effect of attaching different aromatic spacers to 4-ethynylpyridine ligands on the absorption maxima. As expected, the  $\lambda_{\text{max}}$  values move to the red ( $\geq 10$  nm) upon the incorporation of heterocyclic spacers.[5] Comparison of these data with our previous results [5] clearly indicate that the presence of a heavy atom

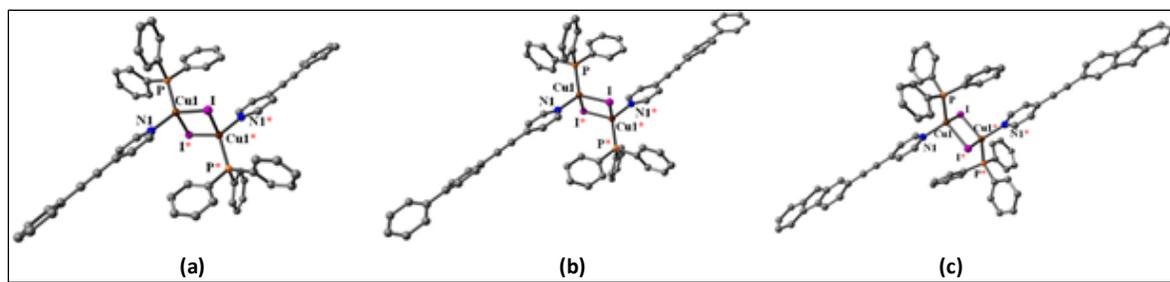
such as sulfur play important role in shifting the  $\lambda_{\text{max}}$  values in dimeric iodo-bridged Cu(I) complexes. As mentioned before, Cu(I) complexes are well known for their emissive properties in both the solid and solution states.[8] In the past, we have demonstrated that the emission properties of ethynylpyridine ligands-based Cu(I) complexes in DCM solution is mainly governed by the electron withdrawing/donating nature of the spacer group. Moreover, there is only little impact of the alkynyl unit (especially for mononynes) on the emission wavelength.[5] Therefore, a trend similar to absorption is expected on the emission properties. We are currently investigating the solid-state as well as solution luminescence, electrochemical, thermal and photovoltaic properties of carbocyclic and heterocyclic spacer incorporated ethynylpyridine ligands-based Cu(I) complexes. The results will be communicated separately.



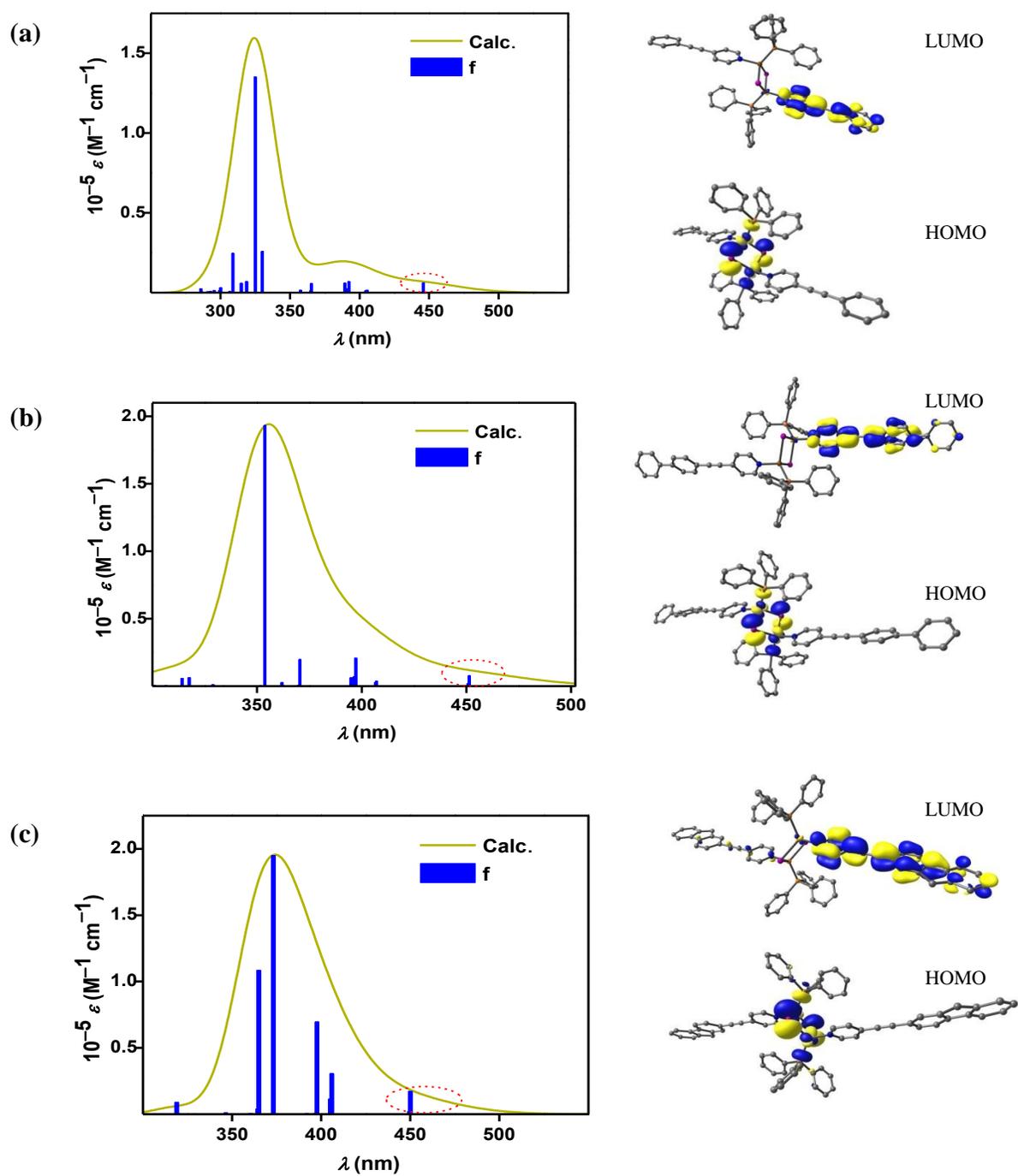
**Figure 3.** (a) Normalized UV-vis spectra of complexes **C**<sub>1</sub>–**C**<sub>3</sub> in dichloromethane, and (b) the effect of different carbocyclic and heterocyclic aromatic spacers (Ar = Phenyl, biphenyl, fluorene, thiophene, bithiophene and terthiophene) attached to 4-ethynylpyridine ligands in dimeric complexes of type [L(CuI)<sub>2</sub>(PPh<sub>3</sub>)<sub>2</sub>] (where L = Ar–C≡C–C<sub>6</sub>H<sub>4</sub>N) on the absorption maxima. Data for the heterocyclic spacers taken from the ref.[5]

#### 4. DFT Calculations

The structures of complexes **C**<sub>1</sub>–**C**<sub>3</sub>, bearing three different carbocyclic groups (with varying conjugation lengths) attached to the ethynylpyridine ligands were confirmed by DFT calculations employing the B3LYP level of theory (see computational details in the experimental section). Geometry-optimization of complexes **C**<sub>2</sub> and **C**<sub>3</sub> was performed using the X-ray crystallographic coordinates and the complex **C**<sub>1</sub> model was optimized using the same hybrid level of theory. The minima were verified *via* computed vibration frequencies. The DFT calculated bond-lengths and bond-angles of the optimized structures are (**Figure 4** and **Table ST5**, *supplementary file*) in good agreement with those obtained experimentally (**Table ST1-4**, *supplementary file*). The time-dependent DFT (TD-DFT) calculations were executed on these optimized geometries to analyze the photophysical properties (**Figure 5**). Other frontier molecular orbitals involved in electronic transitions in complexes **C**<sub>1</sub>–**C**<sub>3</sub>, as obtained by TD-DFT calculation are shown in **Figure SF1-SF3** (*supplementary information*).

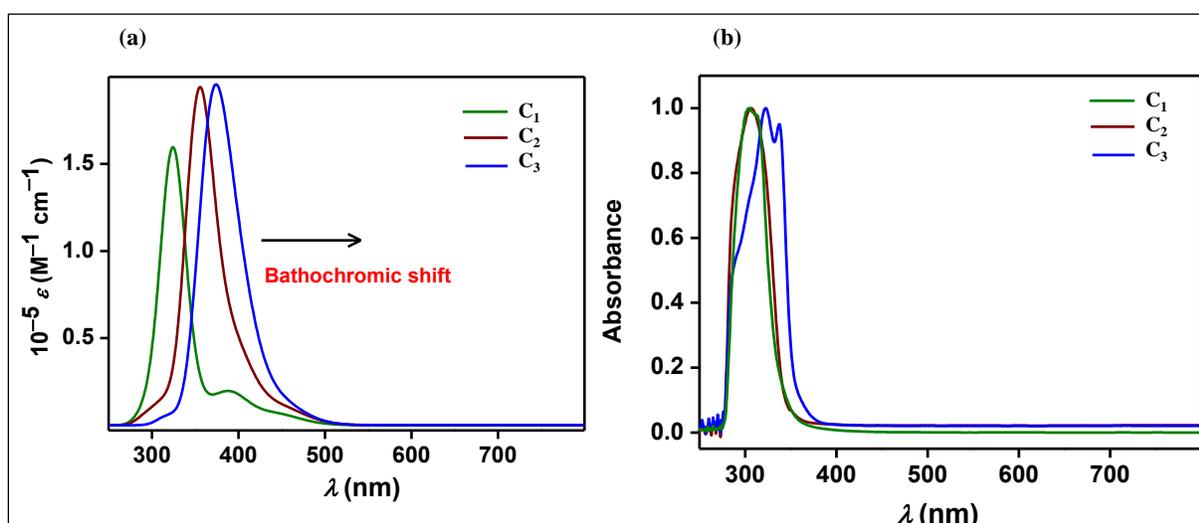


**Figure 4.** DFT optimized structures of (a)  $C_1$ , (b)  $C_2$ , and (c)  $C_3$ .



**Figure 5.** TD-DFT-Calculated electronic spectra for the Cu(I) complexes (a) C<sub>1</sub>, (b) C<sub>2</sub>, and (c) C<sub>3</sub> using the B3LYP level of theory. Each plot shows the simulated absorption profile (green), with the bars marked by blue lines representing oscillator strength. Isosurface plots of the highest-occupied and lowest-unoccupied molecular orbitals (HOMOs/LUMOs) for each system are shown to the right of the simulated spectra.

To obtain deeper insight of the electronic transitions for all three complexes, time-dependent (TD-DFT) calculations have been performed. Here we semi-quantitatively explain the effect of extended conjugation embedded within alkynyl group attached to the pyridyl ligands. The calculated and experimental UV-vis spectra are depicted in **Figure 6**. To analyze the spectroscopic activity of these complexes, we have mainly considered the transition in each case with maximum oscillator strengths,  $f$ , among all the transitions with excitation wavelengths in the 300–400 nm region. From TD-DFT assignments it was revealed that all transitions in this region are largely due to charge-transfer (CT) mainly from the iodide ligands towards the pyridyl rings within the terminally conjugated alkynyl groups. Along with this, there is a substantial contribution of MLCT involving the metal  $\rightarrow$  substituted pyridyl rings. **Tables 2** and **ST6-ST7** (*supplementary information*) lists all excitation energies, wavelengths, oscillator strengths, and an assignment of each state in terms of contributions from individual orbital transitions.



**Figure 6.** TD-DFT simulated (a) and absorption (b) electronic spectra of complexes **C<sub>1</sub>–C<sub>3</sub>** in DCM

**Table 2.** TD-DFT calculated electronic transitions of **C<sub>1</sub>** using B3LYP level of theory.

Excitation energy (eV)	$\lambda$ (nm)	$f$	Transition	Character
2.780	445	0.074	$\alpha/\beta$ -H[~58%L] $\rightarrow$ $\alpha/\beta$ -L[~80%L](92%)	Charge transfer (CT) mainly from iodide towards the pyridyl rings within the terminally conjugated alkynyl groups. A significant contribution of MLCT is also observed involving metal $\rightarrow$ substituted pyridyl rings.
3.756	330	0.310	$\alpha/\beta$ -H-3[~57%M] $\rightarrow$ $\alpha/\beta$ -L+1[~89%L](86%) $\alpha/\beta$ -H-5[~63%L] $\rightarrow$ $\alpha/\beta$ -L+1[~89%L](36%)	Combination of MLCT involving metal $\rightarrow$ substituted pyridyl rings with subsequent CT from iodide towards pyridyl rings within the terminally conjugated alkynyl groups Charge transfer (CT) mainly from iodide towards the pyridyl rings within the terminally conjugated alkynyl groups. An appreciable contribution of MLCT is also observed here involving metal $\rightarrow$ substituted pyridyl rings.
3.8144	325	1.62	$\alpha/\beta$ -H-8[~50%L] $\rightarrow$	Combination of MLCT involving metal $\rightarrow$ substituted pyridyl rings with subsequent CT

			$\alpha/\beta$ -L+1[~89%L](28%) $\alpha/\beta$ -H-7[~53%M]→ $\alpha/\beta$ -L [~80%L](66%)	from iodide towards the pyridyl rings within three terminally conjugated alkynyl groups. MLCT involving Metal → substituted pyridyl rings with terminally conjugated alkynyl groups and CT in substituted pyridyl rings with terminally conjugated alkynyl groups.
4.013	308	0.295	$\alpha/\beta$ -H-10[~77%M]→ $\alpha/\beta$ -L[~80%L](34%)  $\alpha/\beta$ -H-8[~50%L]→ $\alpha$ -L+1[~890%L](46%)	MLCT involving metal → substituted pyridyl rings within the terminally conjugated alkynyl groups. Combination of MLCT involving metal → substituted pyridyl rings with subsequent CT from iodide towards the pyridyl rings within the terminally conjugated alkynyl groups.

The calculated HOMO–LUMO gaps associated with low energy transitions (~ 440–450 nm) are also given in the **Table 3** though they are not clearly visible experimentally due to the broadening of absorption bands. In complex **C<sub>1</sub>**, the brightest transition at nearly 300 nm (experimentally) could be assigned as HOMO–5 to LUMO+1, HOMO–7 to LUMO, and HOMO–8 to LUMO+1 excitations, respectively, with oscillator strengths,  $f = 1.62$ . We also found a considerably weaker transition between the HOMO and LUMO orbitals at 445 nm ( $f = 0.074$ ). A gradual red shift (**Table 3**) in the absorption edge (with maximum  $f$ ) of complex **C<sub>2</sub>** (310 nm) and complex **C<sub>3</sub>** (334 nm) has been observed experimentally, as we move forward from complex **C<sub>1</sub>** towards complex **C<sub>3</sub>**. This can be explained on the basis of their structural configurations. Due to the incorporation of the more delocalized functionalities into the alkynyl group attached to the pyridyl ligands in Cu(I) complexes, in going from **C<sub>1</sub>** to **C<sub>3</sub>** a noticeable bathochromic shift is observed in the spectra. Here, the computational modeling also semi-qualitatively represents the key inferences extracted from the optical spectroscopy. It summarized the importance of extended delocalization of the ligands on the frontier orbitals of complex **C<sub>1</sub>** and its analogues leading to red shift across the metal centers and between the aromatic ligands.

**Table 3.** The calculated HOMO-LUMO gaps associated with low energy transitions, the experimental and calculated  $\lambda$  values.

Complex	$\Delta(\text{HOMO} - \text{LUMO})/\text{kcal/mol}$	$\lambda$ (nm)	
		Expt.	DFT calculated
<b>C<sub>1</sub></b>	64.7838	300	324
<b>C<sub>2</sub></b>	74.0262	310	353
<b>C<sub>3</sub></b>	74.1203	334	375

## 5. Conclusion

We have synthesized a range of [CuI(PPh<sub>3</sub>)L]<sub>2</sub> dimeric complexes, with a Cu<sub>2</sub>L<sub>2</sub> core, where L is a series of ethynyl-functionalized pyridine ligands (Ar-C≡C-C<sub>5</sub>H<sub>4</sub>N), to include the aryl groups phenyl (**C<sub>1</sub>**), biphenyl (**C<sub>2</sub>**) and fluorenyl (**C<sub>3</sub>**) using established coordination-driven self-assembly synthetic strategies.

Crystallographic studies confirm the expected *trans* arrangement of the L ligands across the planar parallelepiped Cu<sub>2</sub>I<sub>2</sub> core. The UV/visible absorption spectra of the three ethylpyridinyl-based ligands and their corresponding Cu complexes were red-shifted compared to ligand precursor 4-(trimethylsilyl ethynyl)pyridine. In the complexes the magnitude of the red shift increases in the order **C**<sub>1</sub> < **C**<sub>2</sub> < **C**<sub>3</sub> which can be ascribed to the increasing  $\pi$ -conjugation length of the arylethynylpyridine ligand. DFT calculations confirm that the bands in the absorption spectra are a mixture of charge transfer from the iodine ligands to the functionalized pyridine groups and MLCT from the copper centers also to the functionalized pyridine ligands.

This study, coupled with our previous work, [5] indicates that these dinuclear copper (I) systems are amenable to coordination with a wide range of functionalized Ar-C $\equiv$ C-C<sub>5</sub>H<sub>4</sub>N ligands, and opens up the possibility of synthesizing one-dimensional polymeric systems *via* further manipulation of the Ar groups to form functionalized linkers through covalent bond formation.

### Acknowledgements

AH acknowledges the Department of Chemistry, University of Hail, Kingdom of Saudi Arabia for providing infrastructure support during the preparation of this manuscript. MSK thanks His Majesty's Trust Fund for Strategic Research (Grant No. SR/SQU/SCI/CHEM/16/02) for funding. PRR is grateful to the Engineering and Physical Sciences Research Council (UK) for continued support (EP/K004956/1).

### References

- [1] I. Ali, W.A. Wani, A. Khan, A. Haque, A. Ahmad, K. Saleem, N. Manzoor, Synthesis and synergistic antifungal activities of a pyrazoline based ligand and its copper (II) and nickel (II) complexes with conventional antifungals, *Microbial pathogenesis* 53(2) (2012) 66-73.
- [2] K. Saleem, W.A. Wani, A. Haque, M.N. Lone, M.F. Hsieh, M.A. Jairajpuri, I. Ali, Synthesis, DNA binding, hemolysis assays and anticancer studies of copper(II), nickel(II) and iron(III) complexes of a pyrazoline-based ligand, *Future Medicinal Chemistry* 5(2) (2013) 135-146.
- [3] A. Haque, R. Ilmi, I.J. Al-Busaidi, M.S. Khan, Coordination chemistry and application of mono- and oligopyridine-based macrocycles, *Coord. Chem. Rev.* 350 (2017) 320-339.
- [4] A. Haque, R.A. Al-Balushi, I.J. Al-Busaidi, M.S. Khan, P.R. Raithby, Rise of Conjugated Poly-ynes and Poly(Metalla-ynes): From Design Through Synthesis to Structure-Property Relationships and Applications, *Chem. Rev.* 118(18) (2018) 8474-8597 and references therein.
- [5] M. Jayapal, A. Haque, I.J. Al-Busaidi, N. Al-Rasbi, M.K. Al-Suti, M.S. Khan, R. Al-Balushi, S.M. Islam, C. Xin, W. Wu, W.Y. Wong, F. Marken, P.R. Raithby, Dicopper(I) Complexes Incorporating Acetylide-Functionalized Pyridinyl-Based Ligands: Synthesis, Structural, and Photovoltaic Studies, *Inorg. Chem.* 57(19) (2018) 12113-12124.
- [6] R. Peng, M. Li, D. Li, Copper(I) halides: A versatile family in coordination chemistry and crystal engineering, *Coord. Chem. Rev.* 254(1) (2010) 1-18.
- [7] R. Ilmi, I. Juma Al-busaidi, A. Haque, M.S. Khan, Recent progress in coordination chemistry, photo-physical properties, and applications of pyridine-based Cu (I) complexes, *J. Coord. Chem.* 71(19) (2018) 3045-3076.
- [8] K. Tsuge, Y. Chishina, H. Hashiguchi, Y. Sasaki, M. Kato, S. Ishizaka, N. Kitamura, Luminescent copper (I) complexes with halogenido-bridged dimeric core, *Coord. Chem. Rev.* 306 (2016) 636-651.
- [9] Y. Liu, S.-C. Yiu, C.-L. Ho, W.-Y. Wong, Recent advances in copper complexes for electrical/light energy conversion, *Coord. Chem. Rev.* (2018).
- [10] L. Qi, Q. Li, X. Hong, L. Liu, X.-X. Zhong, Q. Chen, F.-B. Li, Q. Liu, H.-M. Qin, W.-Y. Wong, Synthesis, characterization and luminescent properties of three-coordinate copper (I) halide complexes containing 2-(diphenylphosphino) biphenyl, *J. Coord. Chem.* 69(24) (2016) 3692-3702.

- [11] B.-L. Chen, L. Liu, X.-X. Zhong, A.M. Asiri, K.A. Alamry, G.-H. Li, F.-B. Li, N.-Y. Zhu, W.-Y. Wong, H.-M. Qin, Synthesis, characterization and luminescent properties of copper (I) halide complexes containing biphenyl bidentate phosphine ligand, *J. Coord. Chem.* 70(23) (2017) 3907-3919.
- [12] X. Hong, B. Wang, L. Liu, X.-X. Zhong, F.-B. Li, L. Wang, W.-Y. Wong, H.-M. Qin, Y.H. Lo, Highly efficient blue–green neutral dinuclear copper (I) halide complexes containing bidentate phosphine ligands, *J. Lumin.* 180 (2016) 64-72.
- [13] K.F. Sultana, S.Y. Lee, J.-E. Lee, J. Seo, S.S. Lee, Mono-and dinuclear silver (I) complexes of benzene-and pyridine-bearing 20-membered thioxaza macrocycles, *Inorg. Chem. Commun.* 10(12) (2007) 1496-1500.
- [14] H.H. Shah, R.A. Al-Balushi, M.K. Al-Suti, M.S. Khan, C.H. Woodall, A.L. Sudlow, P.R. Raithby, G. Kociok-Kohn, K.C. Molloy, F. Marken, New Multi-Ferrocenyl- and Multi-Ferricenyl- Materials via Coordination-Driven Self-Assembly and via Charge-Driven Electro-Crystallization, *Inorg. Chem.* 52(20) (2013) 12012-12022.
- [15] H.H. Shah, R.A. Al-Balushi, M.K. Al-Suti, M.S. Khan, F. Marken, A.L. Sudlow, G. Kociok-Kohn, C.H. Woodall, P.R. Raithby, K.C. Molloy, New di-ferrocenyl-ethynylpyridinyl triphenylphosphine copper halide complexes and related di-ferricenyl electro-crystallized materials, *Dalton Transactions* 43(25) (2014) 9497-9507.
- [16] D.B.G. Williams, M. Lawton, Drying of organic solvents: quantitative evaluation of the efficiency of several desiccants, *The Journal of organic chemistry* 75(24) (2010) 8351-8354.
- [17] H.H. Shah, R.A. Al-Balushi, M.K. Al-Suti, M.S. Khan, C.H. Woodall, K.C. Molloy, P.R. Raithby, T.P. Robinson, S.E. Dale, F. Marken, Long-Range Intramolecular Electronic Communication in Bis (ferrocenylethynyl) Complexes Incorporating Conjugated Heterocyclic Spacers: Synthesis, Crystallography, and Electrochemistry, *Inorg. Chem.* 52(9) (2013) 4898-4908.
- [18] CrysAlis Software package, Oxford Diffraction Ltd, (2011) Oxford, UK, 2011.
- [19] A. Altomare, G. Casciarano, C. Giacovazzo, A. Guagliardi, M. Burla, G.t. Polidori, M. Camalli, SIRPOW. 92—a program for automatic solution of crystal structures by direct methods optimized for powder data, *J. Appl. Crystallogr.* 27(3) (1994) 435-436.
- [20] G.M. Sheldrick, Crystal structure refinement with SHELXL, *Acta Crystallographica Section C: Structural Chemistry* 71(1) (2015) 3-8.
- [21] O.V. Dolomanov, L.J. Bourhis, R.J. Gildea, J.A. Howard, H. Puschmann, OLEX2: a complete structure solution, refinement and analysis program, *J. Appl. Crystallogr.* 42(2) (2009) 339-341.
- [22] M. Frisch, G. Trucks, H. Schlegel, G. Scuseria, M. Robb, J. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. Petersson, 09, Revision D. 01, Gaussian, Inc., Wallingford, CT (2009).
- [23] A. Becke, AD Becke, *J. Chem. Phys.* 98, 1372 (1993), *J. Chem. Phys.* 98 (1993) 1372.
- [24] L. Chiang, A. Kochem, O. Jarjays, T.J. Dunn, H. Vezin, M. Sakaguchi, T. Ogura, M. Orio, Y. Shimazaki, F. Thomas, Radical Localization in a Series of Symmetric NiII Complexes with Oxidized Salen Ligands, *Chemistry—A European Journal* 18(44) (2012) 14117-14127.
- [25] P.J. Hay, W.R. Wadt, Ab initio effective core potentials for molecular calculations. Potentials for the transition metal atoms Sc to Hg, *The Journal of chemical physics* 82(1) (1985) 270-283.
- [26] a. Petersson, A. Bennett, T.G. Tensfeldt, M.A. Al-Laham, W.A. Shirley, J. Mantzaris, A complete basis set model chemistry. I. The total energies of closed-shell atoms and hydrides of the first-row elements, *The Journal of chemical physics* 89(4) (1988) 2193-2218.
- [27] G. Petersson, M.A. Al-Laham, A complete basis set model chemistry. II. Open-shell systems and the total energies of the first-row atoms, *The Journal of chemical physics* 94(9) (1991) 6081-6090.
- [28] S. Miertuš, E. Scrocco, J. Tomasi, Electrostatic interaction of a solute with a continuum. A direct utilizaion of AB initio molecular potentials for the prevision of solvent effects, *Chem. Phys.* 55(1) (1981) 117-129.
- [29] N.M. O'boyle, A.L. Tenderholt, K.M. Langner, Cclib: a library for package-independent computational chemistry algorithms, *J. Comput. Chem.* 29(5) (2008) 839-845.
- [30] G. Zhurko, D. Zhurko, Chemcraft - graphical software for visualization of quantum chemistry computations. <https://www.chemcraftprog.com>.
- [31] J. Lambert, H. Shurvell, D. Lightner, R. Cooks, *Introduction to Organic Spectroscopy* Macmillan, New York (1987) 351.
- [32] L. Engelhardt, P. Healy, J. Kildea, A. White, Lewis-Base Adducts of Group 11 Metal (I) Compounds. XLII.[Cu4X4L4]Cubane'Clusters, L= Hindered N-Base Ligand: Synthesis and Structure of the L= 2-(Diphenylmethyl) pyridine Complexes (X= Cl, Br, I), *Aust. J. Chem.* 42(1) (1989) 107-113.