

# Accepted Manuscript

## Original article

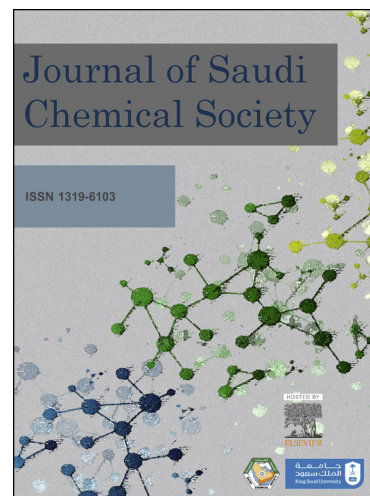
Synthesis, spectral characterization, theoretical, antimicrobial, DNA interaction and *in vitro* anticancer studies of Cu(II) and Zn(II) complexes with pyrimidine-morpholine based Schiff base ligand

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**Synthesis, spectral characterization, theoretical, antimicrobial, DNA interaction and *in vitro* anticancer studies of Cu(II) and Zn(II) complexes with pyrimidine-morpholine based Schiff base ligand**

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**Abstract** Novel Cu(II) (**1**) and Zn(II) (**2**) complexes with 4-(1-(4-morpholinophenyl)ethylideneamino)pyrimidine-5-carbonitrile (**L**) have been synthesized and characterized by various spectroscopic and analytical techniques. DFT (density functional theory) studies result confirms that, LMCT mechanism have been done between **L** and M(II) ions. The antimicrobial studies indicate that the ligand **L** and complexes **1** & **2** exhibit higher activity against the *E. coli* bacteria and *C. albicans* fungi. The groove binding mode of ligand **L** and complexes **1** & **2** with CT-DNA have been confirmed by electronic absorption, competitive binding, viscometric and cyclic voltammetric studies. The electronic absorption titration of ligand **L** and complexes **1** & **2** with DNA have been carried out in different buffer solutions (pH 4.0, 7.0 & 10.0). The  $K_b$  values of ligand **L** and complexes **1** & **2** are higher in acidic buffer at pH 4.0 ( $K_b = 2.42 \times 10^5$ , **L**;  $2.8 \times 10^5$ , **1**;  $2.65 \times 10^5$ , **2**) and the results revealed that, the interaction of synthesized compounds with DNA were higher in the acidic medium than basic and neutral medium. Furthermore, CT-DNA cleavage studies of ligand **L** and complexes **1** & **2** have been studied. The *in vitro* anticancer activities results

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