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SYNTHESIS AND CHARACTERIZATION OF VANADIUM(IV)-FLAVONOID COMPLEX AND ITS RADICAL SCAVENGING PROPERTIES TOWARDS SUPEROXIDE FREE RADICAL

ABSTRACT

In this project, VO(3OH-6M-Fl)₂ complex synthesized using vanadium(IV) acetylacetonate $(VO(acac)_2)$ and hydroxy-6-methylflavone (3OH-6M-Fl). The complex stoichiometry and stability were checked using FTIR and UV-vis spectroscopies along with elemental (C, H) analysis. V=O peak around 990 cm⁻¹ in IR spectrum along with shifted ligand-based peaks confirm the coordination of V(IV) with the ligand in a bidentate fashion. In UV-Vis spectrum peaks around 400-450 nm were observed that correspond to the ligand to metal charge transfer (LMCT) transitions. The radical scavenging ability of newly synthesized complex along with vanadium complexes other investigated using DPPH (2,2-diphenyl-1picrylhydrazyl). The synthesized complex exhibits strong radical scavenging activity compared to VO(acac)₂ and BHA (butyl hydroxyanisole), with IC_{50} value of (105, 95 and 96) mM, respectively.

OBJECTIVES

- To synthesis, characterization and to study the stability of vanadium (IV)flavonoid complexes using FTIR and UV-Vis.
- To investigate the anti-oxidant and radical scavenging ability of the complex.

LITERATURE REVIEW

Flavonoids are considered as good chelating ligands especially the ones that are having hydroxyl group in one or more of their rings can interact with metals. Some recent research suggested that the presence of OH group in the position 3 or 5 on the ring can have a better chelating effect than being in ortho position on the ring (P.G. Pietta, 2000). DPPH assay is a standard radicle assay that is stable in solution and possess a purple colour at 517nm absorbance in ethanol. The main role of DPPH is to accept a hydrogen atom (H) donated by the scavenger complexes or the antioxodant molecule, resulting in the reduction of DPPH to DPPH₂, the purple colour change to yellow and the absorbance decreases as a result (M.S. Blois, 1958).

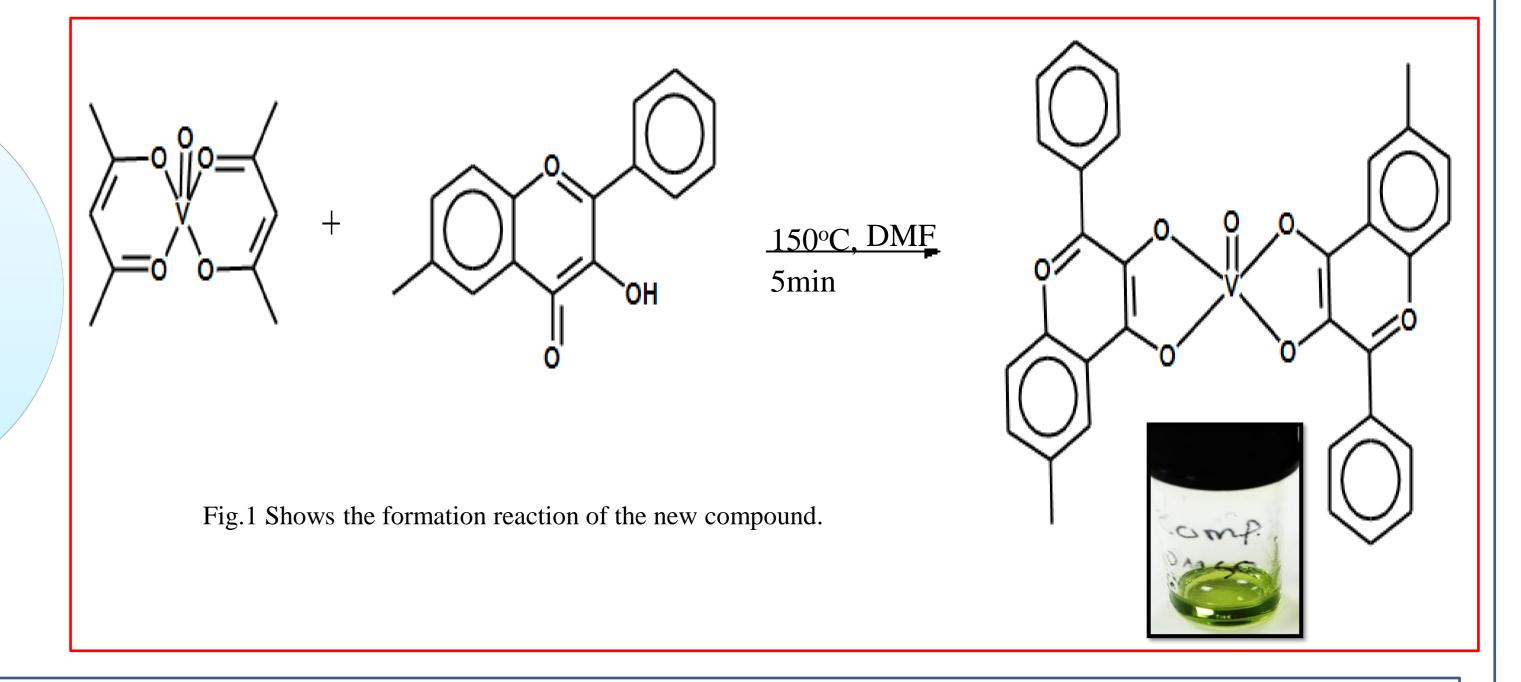
METHODOLOGY

Step one was the synthesis:

VO(acac)2 6-methyl flavanol

DMF (solvent)

Microwav e initiator (150 °C, 5 min)



RESULTS and DISCUSSION

IR Spectroscopy

The IR test was used to assign the peaks of the functional groups, to investigate the formation of product.

Fig.2 shows the FTIR spectrum of the starting materials and the newly synthesised compound in region of 1700-500 cm⁻¹.

Wave number cm

Functional group	Frequency cm ⁻¹
VO(acac) ₂	
V=O	980 - 996
C=O	1518 -1555
C=C	2000-2500
С-Н	2928-3027
Ligand	
C=O	1545-1570
C=C	2354-2486
С-Н	2600-3019
О-Н	3332-3581
Compound	
V-O	790-810
V=O	980 - 996
C=O	1450-1575
C=C	2354-2578

2922-3076

Table.1 shows the functional group of VO(acac)2, Ligand and

UV-Vis Analysis

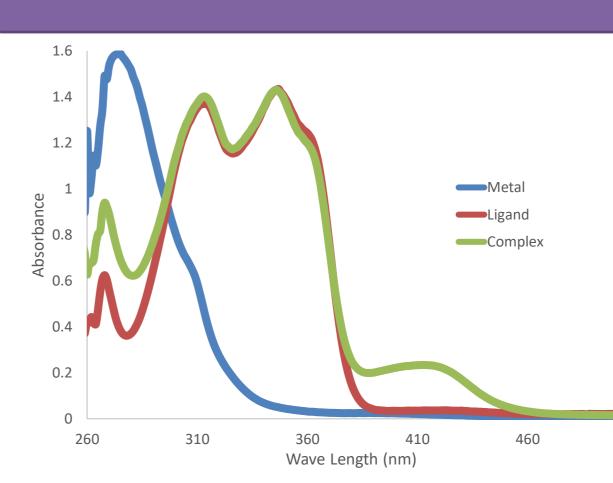
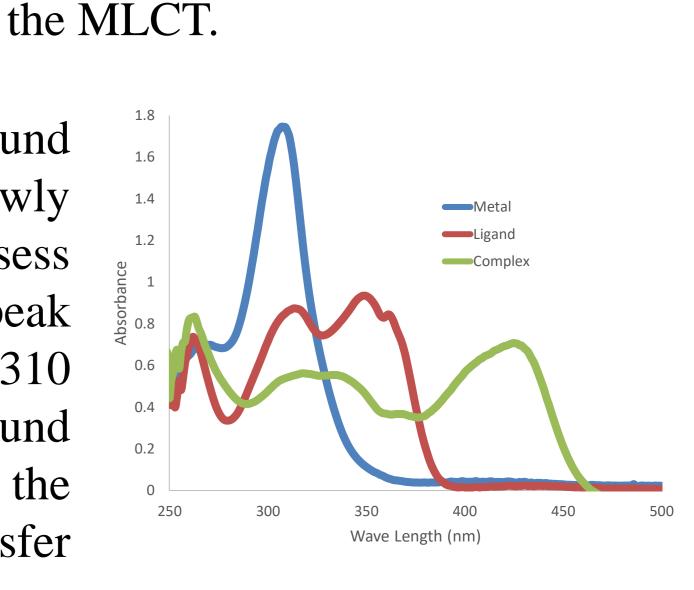


Fig.5 Shows the UV-vis spectrum of VO(acac)₂, ligand and the synthesized complex in DMF.

VO(acac)₂ peaks are shown around 270nm, while the ligand and newly synthesized complex possess approximately the peak same positions and intensity around 310 and 350 nm, except the peak around 410-430 nm which correspond to the metal-ligand charge transfer (MLCT).



In DMSO the VO(acac), peaks

appears around 310nm, as for the

ligand and complex, they retain the

same peaks position but differs in

the absorbance intensity, with an

extra peak shown in the complex

spectrum around 430nm, indicating

Fig.6 Shows the UV-vis spectrum of VO(acac)₂, ligand and the synthesized complex in DMSO.

Radical scavenging

C-H

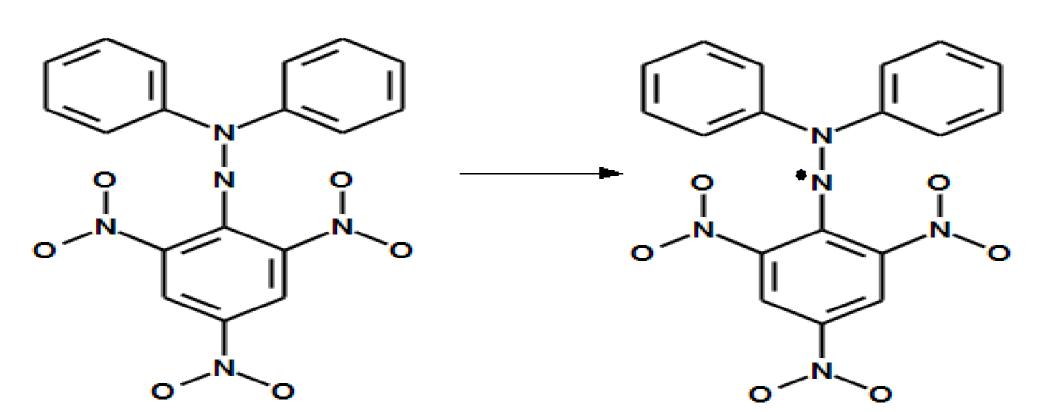
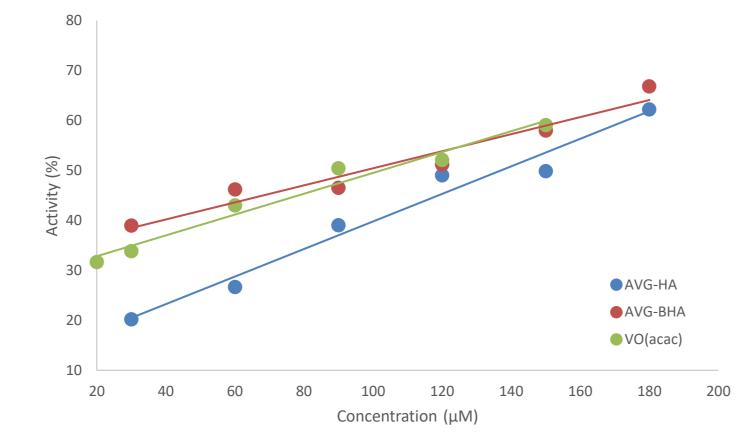


Fig.3 The DPPH complex before and after the reduction by anti-oxidant compounds



The IC₅₀ of the complex, VO(acac)₂ and BHA were found to be (105, 95 and 96) μ M, respectively.

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Fig.4 Shows the DPPH free radical scavenging activity of standard BHA, $Vo(acac)_2$ and the Complex.

CONCLUSION

In FTIR spectrum the main functional groups especially V=O peak around 900-1000cm⁻¹ is observed. Disappearance of –OH peak also indicated deprotonation of the –OH group and its coordination with metal centre. Based on the UV-vis spectra and comparison with literature, it might be suggested that the complex geometry could be a square pyramidal.

Radical scavenging studies show that metal complexes as well as metal salts possess scavenging/inhibition potential against superoxide radicals such as DPPH. The activity of these types of complexes is influenced by its nature and the substituted groups position on the ligands.

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