

# Cocculus hirsutus: Molecular Docking to Identify Suitable Targets for Hepatocellular Carcinoma by *In silico* Technique

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## ABSTRACT

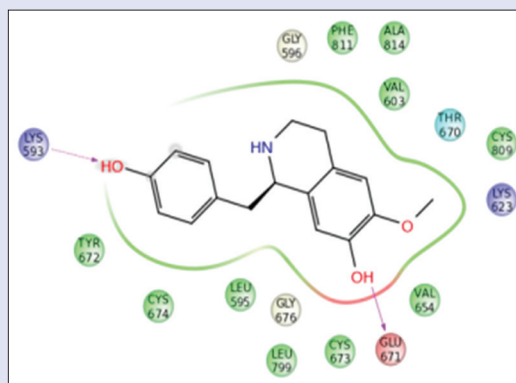
**Background:** Protein–ligand interaction plays a major role in identification of the possible mechanism by which a ligand can bind with the target and exerts the pharmacological action. **Objective:** The aim is to identify the best candidate of *Cocculus hirsutus* which binds with the hepatocellular carcinoma (HCC) targets by docking studies. **Materials and Methods:** The reported phytoconstituents such as coclaurine, hirsutine, cohirsine, cohirsinine, lirioreosinol, cohirsitinine, haiderine, jantinine, isotrilobine, shaheenine, jantine, and coccoline present in the plant, *C. hirsutus* were docked with the HCC targets such as Aurora kinase, c-Kit, fibroblast growth factor, nuclear factor kappa B (NF-κB), B-cell lymphoma-extra large, and vascular endothelial growth factor (VEGF) using *in silico* technique with the software Grid-Based Ligand Docking with Energies. **Results:** Haiderine, shaheenine, and coclaurine had good interaction with Aurora kinase with the glide score and glide energy of – 7.632, –7.620, –7.464; and – 56.536, –55.203, –52.822, respectively. Coclaurine, lirioreosinol, and haiderine possess good binding with c-Kit with the glide score and glide energy of – 8.572, –6.640, –6.478; and – 56.527, –57.138, –20,522, respectively. Lirioreosinol, hirsutine, and coclaurine exhibit good binding with c-Kit with the glide score and glide energy of – 5.702, –5.694, –5.678; and – 48.666, –35.778, –41,673, respectively. Similarly, coclaurine, haiderine, and hirsutine had good interaction with NF-κB. Haiderine, jantinine, and coclaurine had good binding with VEGF receptors (VEGFR) and coclaurine, lirioreosinol, and haiderine exhibit good bonding with VEGFR. **Conclusion:** Coclaurine, haiderine, and lirioreosinol exhibited good hydrogen bonding interactions and binding energy with the select targets. Hence, these compounds have to be taken up for experimental work against hepatocellular carcinoma.

**Key words:** *Cocculus hirsutus*, hepatocellular carcinoma, molecular docking, phytoconstituents

## SUMMARY

- Compounds of interest showed good interaction and binding with the

selected targets. Hence these compounds has to be explored further to study their anticancer potentials.



**Abbreviations used:** HCC: Hepatocellular Carcinoma, Bcl-xL: B-cell lymphoma-extra large, FGF: Fibroblast Growth Factor, VEGF: Vascular Endothelial Growth Factor, DLA: Dalton's Lymphoma Ascites.

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## INTRODUCTION

Hepatocellular carcinoma (HCC), the third common cancer, accounts for more than 626,000 new cases per year worldwide.<sup>[1]</sup> This study aimed to perform docking of ligands, which were reported in the plant *Cocculus hirsutus* with the macromolecular targets which involves in various apoptosis cell signaling pathways.<sup>[2]</sup> Our laboratory findings indicate that this plant is found to have *in vitro* cytotoxic activity in MCF-7, Hep-2, and Hela cancer cell line<sup>[3]</sup> and *in vivo* anticancer activity against DLA cells in mice.<sup>[4]</sup> To find out the best candidate of the plant constituents, the reported phytoconstituents of *C. hirsutus* were docked with the various targets responsible for different signaling pathways such as Aurora kinase, proto-oncogene c-Kit (c-Kit), B-cell lymphoma-extra large (Bcl-xL), nuclear factor kappa B (NF-κB), fibroblast growth factor (FGF), and vascular endothelial growth factor (VEGF).

Plants are rich in secondary metabolites such as alkaloids, glycosides, tannins, terpenoids, flavanoids, etc. *C. hirsutus* (L.) Diels is a climbing shrub belongs to the family *Menispermaceae* is rich in alkaloid, flavanoids, and phenolic compounds. *C. hirsutus* is widely used in the indigenous

system of medicine for curing various ailments due to its different medicinal properties. The plants are reported to have antioxidant, cytotoxic,<sup>[5]</sup> hepatoprotective,<sup>[6]</sup> anticancer, and hypotensive.<sup>[7]</sup> The reported phytoconstituents of *Cocculus hirsutus* are coclaurine, coccoline, cohirsine, cohirsinine, cohirsitinine, haiderine, hirsutine, isotrilobine, Jantine, jantinine, lirioreosinol, and shaheenine.<sup>[8,9]</sup> Attempt has been made to dock these plant constituents with the HCC targets such as Aurora kinase, c-Kit, FGF, NF-κB, Bcl-xL, and VEGF using *in silico* technique with the software Grid-Based Ligand Docking with Energies (GLIDE).

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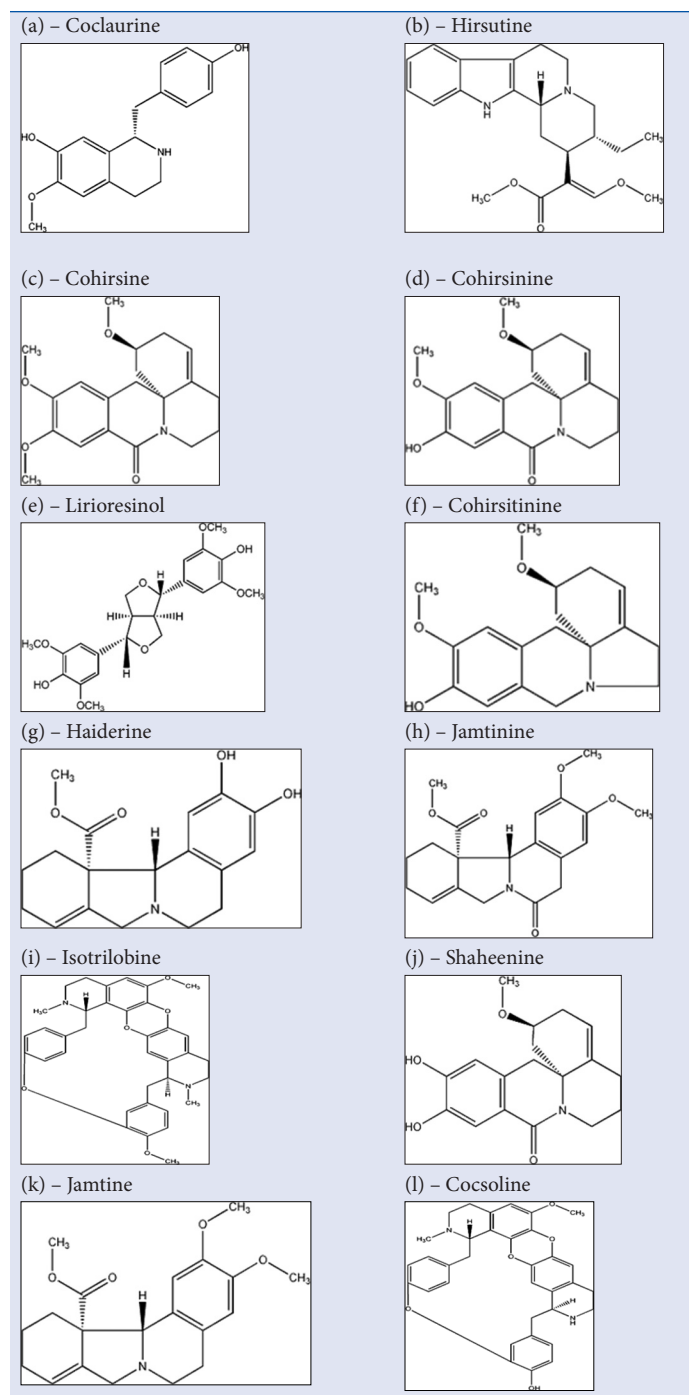
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## MATERIALS AND METHODS

Previously reported phytoconstituents of *C. hirsutus* [Table 1] were downloaded from the database and subjected to docking studies. Maestro build panel was used for building the ligands. LigPrep is a utility of Schrodinger software (USA) that generates three-dimensional structures from two-dimensional representation. The active protein site with which the ligand will be docked was prepared using the wizard, protein preparation of Schrodinger software. Ligand and protein were docked using the software GLIDE. The prepared ligands were docked against such as Aurora kinase, c-Kit, FGF, NF-kB, Bcl-xL, and VEGF cancer targets.

**Table 1:** Compounds of *Cocculus hirsutus* taken for docking studies



$GS_{\text{score}} = b \times \text{Coul} + a \times \text{vdW} + \text{Metal} + \text{Lipo} + \text{RotB} + \text{Hbond} + \text{BuryP} + \text{Site}$   
Where Coul is Coulomb's energy, vdW is Van der Waal's energy, Metal is metal-binding term, Lipo is lipophilic contact term, RotB is penalty for freezing rotatable bonds, Hbond is hydrogen-bonding term, BuryP is penalty for buried polar groups, site is polar interactions at the active site, and the coefficients of vdW and Coul are  $a = 0.065$  and  $b = 0.130$ , respectively. All docking computations were carried out using Linux is an operating system manufactured by Linus Torvalds.

## RESULTS

The results of the best candidate with good binding efficiency of the phytoconstituent with good G-score, glide energy, and amino acid interactions along with the bond distance are summarized in Table 2.

## DISCUSSION

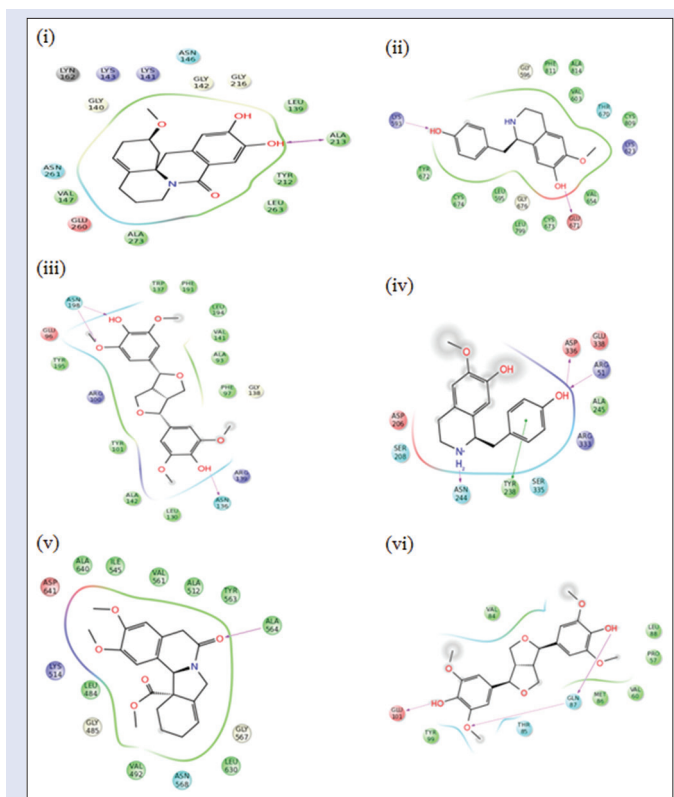
From the results, it is observed that out of 12 compounds of *C. hirsutus*, coclaurine, haiderine, and lirioresinol are found to have good glide score and glide energy toward Aurora kinase, c-Kit, FGF, NF-kB, Bcl-xL, and VEGF.

Aurora kinase which is crucial for cell cycle control is overexpressed in tumor cells.<sup>[10]</sup> Inhibiting Aurora kinase in such increased levels will control

**Table 2:** Phytoconstituents having good interactions with cancer targets

Name of the cancer targets	Phytoconstituents of <i>Cissampelos pareira</i>	Glide score	Glide energy	Interactions	Bond distance (D...A) Å		
Aurora kinase	Haiderine	-7.632	-56.536	Ala (213)	2.042		
				Leu (139)	1.883		
	Shaheenine	-7.620	-55.203	Ala (213)	1.941		
	Coclaurine	-7.464	-52.822	Leu (139)	1.640		
				Ala (213)	2.475		
C-kit	Coclaurine	-8.572	-56.527	Pro (214)	1.865		
				Lys (593)	1.908		
				Glu (671)	1.773		
Bcl-xL	Lirioresinol	-6.640	-57.138	Arg (815)	1.976		
				Lys (593)	2.374		
	Haiderine	-6.478	-20.522	Cys (673)	1.740		
				Asn (136)	1.957		
NF-κB	Lirioresinol	-5.702	-48.666	Asn (198)	2.211		
				Asn (198)	2.464		
				Asn (198)	1.935		
	Haiderine	-5.694	-35.778	Asn (198)	1.935		
				Coclaurine	-5.678	-41.673	-
				Coclaurine	-4.199	-32.578	Asp (336)
FGF	Haiderine			Asn (244)	2.446		
				Asp (206)	1.662		
				Asp (206)	1.820		
	Coclaurine	-3.993	-27.769	Tyr (238)	2.078		
				Gln (50)	1.950		
				Arg (51)	2.010		
				Ala (564)	2.185		
VEGFR	Haiderine	-6.498	-54.607	Ala (564)	1.926		
				Ala (564)	2.264		
	Coclaurine	-6.469	-52.314	Ala (564)	2.264		
				Glu (562)	1.854		
				Asp (641)	1.582		
VEGFR	Coclaurine	-3.775	-24.443	Thr (85)	1.762		
				Glu (101)	1.894		
	Lirioresinol	-3.714	-32.780	Gln (87)	1.897		
				Leu (88)	1.899		
				Gln (87)	2.069		
VEGFR	Haiderine	-3.580	-23.317	Gln (87)	2.069		
				Glu (101)	1.596		

FGF: Fibroblast growth factor; Bcl-xL: B-cell lymphoma-extra large; NF-κB: Nuclear factor kappa B; VEGFR: Vascular endothelial growth factor receptor



**Figure 1:** (i). Docking of Shaheenine with Arorakinase, (ii) Docking of Coclaurine with C- kit, (iii) Docking of Lirioresinol with Bcl-xL, (iv) Docking of Coclaurine with NF-κB, (v) Docking of Jaminine with FGF, (vi) Docking of Lirioresinol with VEGF

the rapid cell growth. c-Kit receptor is present in hematopoietic cells and other tissue cells. c-Kit signaling plays a vital role in regulation of the red blood cell production, lymphocyte proliferation, etc. Downstream signal transduction regulates cell growth, proliferation, and differentiation.<sup>[11]</sup> Overexpression of Bcl-xL relates with resistance to chemotherapy and radiation therapy in multiple cancer.<sup>[12]</sup> NF-κB, i.e. NF-κB is a transcription factor, which will play a vital role in carcinogenesis. Therefore, inhibition of NF-κB activation plays a tumor suppressor role in Liver.<sup>[13]</sup> FGF target is the FGF, which will increase in solid, and metastatic tumors and gives drug resistance.<sup>[14]</sup> Thus, inhibitions of FGF will results in the entry of chemical compounds into the target cell and thereby cell toxicity and retardation of cell growth.<sup>[15]</sup> VEGF-receptors (VEGFRs) are useful of the angiogenesis (development of blood vessels from pre-existing vasculature) and vasculogenesis (development of the circulatory system). Overexpression of VEGF will lead to the abnormal growth of tumor cells.<sup>[16]</sup> Therefore, an inhibitor of VEGFR will control the growth of new cells binding with the above targets and inhibiting their activity will show a significant tumor control in liver tissues. Thus, this study suggests that these phytoconstituents, especially the best candidates such as coclaurine,

haiderine, and lirioresinol, will exhibit its activity by effective binding with the targets as explained by GLIDE software [Figure 1].

## CONCLUSION

From this study, it is clear that out of 12 compounds docked with various targets of HCC, coclaurine, haiderine, and lirioresinol have found to have good interaction and binding efficiency with all targets. Therefore, these best candidates for HCC have to be taken up for further research.

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## Conflicts of interest

There are no conflicts of interest.

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**Dr. B. Samuel Thavamani** is working as associate professor in the department of Pharmacognosy, PSG College of Pharmacy, Coimbatore. He has been working in the area of phytochemical studies of plants especially of anticancer property. To his credit he has national and international publication in this area.