In Silico and In Vitro Anticancer Activity of Isolated Novel Marker Compound from Chemically Modified Bioactive Fraction from Curcuma longa (NCCL)

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ABSTRACT

Background: Turmeric (Curcuma longa) is reported to possess wide array of biological activities. Herbal Medicament (HM) is a standardized hexane-soluble fraction of C. longa and is well known for its neuroprotective effect. Objective: In this study, we attempted to synthesize a novel chemically modified bioactive fraction from HM (NCCL) along with isolation and characterization of a novel marker compound (I). Materials and Methods: NCCL was prepared from HM. The chemical structure of the marker compound isolated from NCCL was determined from 1D/2D nuclear magnetic resonance, mass spectroscopy, and Fourier transform infrared. The compound so isolated was subjected to in silico and in vitro screenings to test its inhibitory effect on estrogen receptors. Results: Molecular docking studies revealed that the binding poses of the compound I was energetically favorable. Among NCCL and compound I taken for in vitro studies, NCCL had exhibited good anti-cancer activity over compound I against MCF-7, MDA-MB-231, DU-145, and PC-3 cells. Conclusion: This is the first study about the synthesis of a chemically modified bioactive fraction which used a standardized extract since the preparation of the HM. It may be concluded that NCCL fraction having residual components induce more cell death than compound I alone. Thus, NCCL may be used as a potent therapeutic drug. Key words: Anti-cancer, Curcuma longa, isolation, molecular docking, synthesis

SUMMARY

• In the present paper, a standardized hexane soluble fraction of Curcuma longa (HM) was chemically modified to give a novel bioactive fraction (NCCL). A novel marker compound was isolated from NCCL and was characterized using various spectral techniques. The compound so isolated was investigated for in-silico screenings. NCCL and isolated compound was subjected to in-vitro anti-cancer screenings against MCF-7, MDA MB 231 (breast adenocarcinoma) and DU 145 and PC-3 cell lines (androgen independent human prostate cancer cells). The virtual screenings reveals that isolated compound has shown favourable drug like properties. NCCL fraction having residual components induces more cell death in these four cancer cell lines than isolated compound alone.

Abbreviations used: HM: Herbal Medicament; NCCL: Chemically modified HM; FT-IR: Fourier transform-infrared spectroscopy; NMR: Nuclear magnetic resonance spectroscopy; MS: Mass spectroscopy; HPLC: High-performance liquid chromatography; ER: Estrogen receptor; MTT: 3-(4,5 dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide; MIC: Minimum inhibitory concentration; TAM: Tamoxifen KBr: Potassium bromide; DMSO: Dimethyl sulfoxide; ACN: Acetonitrile; PDB: Protein Data Bank; FDA: Photodiode array detector.

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INTRODUCTION

Nature is one of the greatest chemist and plants are almost countless funds of charming chemical constituent. Natural products and their derivatives have been recognized for many years as a source of therapeutic agents. The use of plants for medicines worldwide still immensely exceeds the use of the current synthetic drug.[1] Turmeric is a spice derived from the rhizomes of Curcuma longa, which is a member of the ginger family (Zingiberaceae) which has been conventionally used in ayurvedic medicine and in the preparation of food.[2] C. longa contains various bioactive components such as tannins, phenols, saponins, alkaloids, flavonoids, glycosides, curcuminoids, and several volatile oils including variety of sesquiterpenes possessing neuroprotective, antithrombotic, anti-inflammatory, and anti-cancer activities.[3-6] It has been reported to possess antifungal,[7] antibacterial,[8] insecticidal,[9] mosquitoocidal,[10] antioxidant,[11] and anti-cancer activities.[12]

Our Institute had developed a standardized hexane-soluble extract of C. longa rhizomes known by the name of Herbal Medicament (HM), as an anti-stroke agent.[12] It has shown potential neuroprotective activity in neurovascular disorder, and the product was licensed to Themis Medicare Ltd., Mumbai for further development as an anti-stroke agent.

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Depending on the wide therapeutic array of \textit{C. longa}, its chemically modified fraction comprising compounds such as curcumen, zingiberene, \(\beta\)-bisabolene, \(\beta\)-sesquiphellandrene, curzerene, residual ar-turmerone, and some cyclic derivatives of the carbonyl compounds, for example, 7,7-dimethyl-5-(2-p-tolypropl)-6, 7-dihydro-1,3,4-oxadiazepin-2-amine (reaction product of ar-turmerone with semicarbazide; compound I), herein referred as NCCL was prepared. Since the anti-cancer effects of curcuminoids or turmerones are well explored\cite{13, 14} the effect of NCCL on these parameters, if any has not been evaluated so far.

Therefore, in this study, we describe the synthesis and anti-cancer screening of a novel chemically modified bioactive fraction NCCL from HM. During this study, we identified a novel marker compound (I) whose isolation, structural elucidation, \textit{in silico} and \textit{in vitro} screenings has been reported in this study.

Estrogens play crucial roles in breast and prostate cancer development. Therefore, much effort has been devoted to block estrogen formation\cite{15, 16}. Tamoxifen (TAM) is the most widely used therapy for antagonizing estrogen receptor (ER) function as it binds to the ER receptor and blocks downstream signaling\cite{17}. Keeping this in mind, molecular docking study of the newly isolated compound was carried out. Oral bioavailability is a desirable property of compounds under investigation in the drug discovery process. Lipinski’s rule of five is a simple model to forecast the absorption and intestinal permeability of a compound\cite{18, 19}. The isolated marker compound obeys the Lipinski rule of five along with obeying some additional parameters\cite{20, 21}. It was then evaluated for anti-cancer activity by 3-(4,5 dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) assay\cite{22} against MCF-7, MDA-MB-231 (breast adenocarcinoma), DU-145 and PC-3 cells (androgen independent human prostate cancer cells). The results of this study are also documented herein.

**MATERIALS AND METHODS**

Hexane (95%), thiosemicarbazide (98%) was purchased from Sigma-Aldrich (St. Louis, MO, USA). HM was prepared in the Medicinal and Process Chemistry Division of Central Drug Research Institute, Lucknow, Uttar Pradesh, India. Milli-Q pure water was obtained from MilliQ water system purification (Millipore India Pvt. Ltd., New Delhi, India). high-performance liquid chromatography (HPLC) grade methanol and acetonitrile (ACN) were purchased from Merck (Mumbai, India). IR spectra was recorded using potassium bromide (KBr) discs, on a PerkinElmer Spectrum RX1 infrared (IR) spectrophotometer. \(^1\)H and \(^13\)C nuclear magnetic resonance (NMR) spectra were recorded in dimethyl sulfoxide (DMSO) on Bruker DRX-300 (400 MHz) and JEOL AL300 Fourier transform (FT)-NMR (400 MHz) systems; chemical shift (δ) is reported in ppm using tetramethyl silane as an internal reference. Mass spectra were recorded on Agilent 6520 mass spectrometer. Elemental analysis was performed on Elementar’s Vario EL III micro-analyzer.

**Preparation of hexane soluble fraction (Herbal Medicament)**

Dried and powdered rhizomes of the \textit{C. longa} (25 Kg) was taken in a percolator and kept in freshly distilled hexane (40 L) for 48 h. After this, hexane was drained and filtered. The extraction was carried out two more times and combined filtrate was concentrated in vacuum at 40°C at 300-mile bar. Finally, when around 1 L of concentrate is left, it is dried over anhydrous sodium sulfate, filtered and further concentrated, first at 300 mb and then under high vacuum to completely remove solvent impurities. Yield of the oil: About 330 g (1.3%).

**Preparation of NCCL extract**

NCCL was prepared from hexane soluble fraction of \textit{C. longa} according to our patented method\cite{23}. The hexane soluble fraction of \textit{C. longa} (HM, 5 g) was mixed with semicarbazide hydrochloride (10 g) and ethyl alcohol (100 ml, 99% pure) with a few drops of glacial acetic acid, refluxed for 48 h. Further addition of semicarbazide hydrochloride (10 g) at an interval of 5–6 h was continued till the total amount of semicarbazide hydrochloride came up to 50 g. The completion of reaction was monitored by thin layer chromatography (TLC) and finally by HPLC for the confirmation of complete removal of carbonyl portion. The reaction mixture was cooled, filtered, residual portion was washed with ethyl alcohol and filtrate was concentrated under vacuum. Further HPLC grade hexane was added to the concentrate and refluxed for 5 h for the extraction of NCCL from ethanol. It was filtered again and concentrated under vacuum. No crystal formation was observed on retaining the NCCL.

**Chromatographic method for fingerprinting of NCCL**

Waters HPLC (Milford, MA, USA) system used was equipped with a binary gradient pump (Model 515, Waters), an autosampler injector (Model 2707, Waters) and a diode array detector (Model 2998, Waters). Waters HPLC interface and Empower software by Waters corporation, USA was utilized for data acquisition. The analytical column RP-18e B Lichrosphere (250 mm × 4 mm, 5 μm, Merck, Germany) at 30°C ± 3°C was used for the study. The mobile phase included ACN water (70:30, v/v) and was degassed before analysis using a Millipore vacuum pump. Column effluent was monitored at 220 nm and 254 nm (photodiode array detection) with a total runtime of 30 min. The flow rate was maintained 1.0 ml/min. and injection volume remained 20 μL.

**Isolation and identification of active marker compound (I) from NCCL**

NCCL (200 mg) was dissolved in ACN (5 ml) and to it a few drops of water were added and injected onto a prep HPLC column C18 (250 mm, 25 mm, 15 μm, Phenomenex). The flow rate was 12 ml/min with isocratic flow of ACN and water (50:50, v/v). The peaks were monitored at 220 and 254 nm and the total run time was 40 min. The fractions were collected according to their ascent in absorbance. The fractions containing compound I was separated and concentrated. This was repeated for 3 times with 200 mg of sample loading. These fractions were further analyzed through analytical HPLC. Its purity was established by TLC and HPLC. Compound I was found to be 99.58% pure. Compound I was characterized by employing different spectral techniques such as FTIR, \(^1\)H, and \(^13\)C NMR, mass.

**Molecular docking studies (in silico)**

Docking was used to predict the binding orientation of drug to their protein target to in-turn predict the affinity and activity of drug which includes docking of ligand to a set of grids describing the target protein. The compound I was docked into the nucleotide-binding pocket of the ERs structure using the program AutoDock 4.0. Interestingly, this method applies a Lamarckian model of genetics, in which environmental adaptations of an individual’s phenotype are reverse transcribed into its genotype and become heritable traits (sic). We have considered Lamarckian genetic algorithm, search method which can handle ligands with more degrees of freedom than the Monte Carlo simulated annealing and traditional genetic algorithm method used in earlier versions of AutoDock, and that the Lamarckian genetic algorithm is the most efficient, reliable,
and successful. AutoDock 4.0, combines energy evaluation through grids of affinity potential employing various search algorithms to find the suitable binding position for a ligand on a given protein.

The structure of ligand molecule (compound I) was analyzed using ChemDraw Ultra 8.0 by CambridgeSoft Corporation (Cambridge Scientific Computing, Inc., USA). 3D, coordinates were prepared using PRODRG server. The protein structure file (Protein Data Bank [PDB]: 210] and 2YJD) taken from PDB (www.rcsb.org/pdb) was edited by removing the heteroatoms, adding C-terminal oxygen. All torsions were allowed to rotate during docking. Steepest descent methods were applied for minimization, using default parameters. While docking, polar hydrogen's were added to ligand using the hydrogen's module in AutoDock tool and thereafter, Kollman united atom partial charges were assigned. Docking to ligand was carried out with standard docking protocol on the basis a population size of 150 randomly placed individuals; a maximum number of 2.5 × 10^7 energy evaluations, a mutation rate of 0.02, a crossover rate of 0.80 and an elitism value of 1. Fifteen independent docking runs were carried out for ligand and results were clustered according to the 1.0 Å rmsd criteria. The grid maps representing the proteins were calculated using auto grid and grid size was set to 60 × 60 × 60 points with grid spacing of 0.375 Å. The active site, i.e., Arg-394 and Glu-305 in the protein interacts with ligand of the substrate and gives increase to the catalytic activity to test ligand that helps in determining the binding pattern of the ligand to the active site of the ER. The docking results were interpreted according to the.pdb file. We have used the coordinates of the minimum energy run created in the.dlg file which was determined using the rmsd table created in the.dlg file itself. The coordinate of the docked protein along with compound I was visualized using UCSF chimera within 6.5 Å region.

Anti-cancer activity (in vitro)

MCF-7, MDA-MB-231, PC-3, and DU-145 cells used in this study were obtained from the American Type Culture Collection, USA and were routinely maintained in Dulbecco's Modified Eagle Medium (Sigma-Aldrich) supplemented with 10% fetal bovine serum (Merck) and 1% antibiotic and antimycotic solution (Merck) at 37°C in a humidified incubator with 5% CO₂. All stock solution of compounds (NCCL and compound I) was prepared in cell culture grade DMSO and stored in −20°C. Compounds were diluted in culture media before use in experiments. Cell viability was assessed using MTT assay, which is based on the reduction of MTT by mitochondrial dehydrogenases of viable cells to form a purple formazan product. In brief, MCF-7, MDA-MB-231, PC-3, and DU-145 cells (5 × 10⁵/well) were plated in 96-well plates. After incubating for overnight, the cells were treated with different concentrations (200, 100, 50, 25, 12.5, and 6.25 µg/ml, in triplicates) of NCCL and compound I for 48 h. Subsequently, 10 µL of MTT (10 mg/ml) was added to each well and incubated for 3 h. The MTT formazan formed by viable cells was dissolved in 100 µL of DMSO and shaken for 10 min. The absorbance was measured at 540 nm on an ELISA reader. Each test was repeated at least 3 times. The concentration of the compound which gives the 50% growth inhibition value corresponds to IC₅₀, calculated using GraphPad Prism 5 by GraphPad Software Inc.

RESULTS

HPLC fingerprinting of blank ACN, HM, and NCCL was done and retention time and % peak area of HM and NCCL at 220 nm and 254 nm is given in Table 1.

| Compound | Structure of 7,7 dimethyl 5 (2 p tolylpropyl) 6, 7 dihydro 1,3,4 oxadiazepin 2 amine (compound I) |
|----------|------------------------------------------------------------------------------------------------|

Table 1: Retention time and percentage peak area of herbal medicament and NCCL at 220 and 254 nm

| HM | NCCL |
|----|------|
| 220 nm | 254 nm | 220 nm | 254 nm |
| Rt | % area | Rt | % area | Rt | % area | Rt | % area |
| 1.967 | 0.04 | 1.233 | 0.02 | 1.659 | 0.21 | 1.722 | 0.17 |
| 2.105 | 0.15 | 1.518 | 0.02 | 2.032 | 0.06 | 2.143 | 1.32 |
| 2.478 | 2.07 | 1.710 | 0.11 | 2.259 | 0.07 | 2.388 | 0.98 |
| 3.374 | 1.99 | 1.990 | 0.14 | 2.513 | 1.39 | 2.570 | 1.68 |
| 3.850 | 3.64 | 2.223 | 0.45 | 2.801 | 0.38 | 2.795 | 2.35 |
| 4.243 | 0.04 | 2.481 | 6.23 | 3.095 | 2.47 | 3.163 | 2.90 |
| 4.601 | 1.67 | 3.168 | 0.30 | 3.365 | 3.27 | 3.719 | 11.44 |
| 5.134 | 0.79 | 3.428 | 0.64 | 3.772 | 2.08 | 4.119 | 1.32 |
| 5.710 | 0.04 | 3.841 | 3.51 | 4.146 | 0.94 | 4.467 | 50.34 |
| 5.941 | 0.04 | 4.245 | 0.54 | 4.468 | 50.25 | 5.091 | 0.99 |
| 6.153 | 0.44 | 4.622 | 1.30 | 5.093 | 13.91 | 5.258 | 1.08 |
| 6.953 | 2.13 | 4.785 | 0.88 | 5.836 | 0.21 | 5.422 | 1.42 |
| 8.796 | 0.29 | 5.159 | 0.91 | 6.570 | 0.76 | 5.840 | 1.51 |
| 9.236 | 0.17 | 6.152 | 2.72 | 7.331 | 1.02 | 6.461 | 2.02 |
| 9.836 | 71.45 | 6.716 | 1.02 | 7.835 | 7.09 | 6.971 | 0.89 |
| 12.030 | 0.07 | 8.870 | 0.80 | 8.983 | 0.06 | 7.840 | 0.74 |
| 12.590 | 3.45 | 9.837 | 50.04 | 9.359 | 0.09 | 8.276 | 0.17 |
| 14.269 | 10.72 | 12.069 | 0.03 | 9.832 | 0.08 | 9.348 | 0.70 |
| 17.681 | 0.25 | 12.708 | 0.80 | 10.305 | 0.78 | 9.827 | 0.23 |
| 18.703 | 0.11 | 14.290 | 12.53 | 10.884 | 0.04 | 10.286 | 1.84 |
| 23.895 | 0.32 | 14.654 | 11.41 | 11.311 | 0.33 | 11.202 | 0.16 |
| 29.328 | 0.12 | 17.873 | 0.10 | 12.042 | 0.57 | 12.728 | 1.30 |
| - | - | 18.720 | 4.23 | 12.639 | 1.74 | 14.969 | 0.02 |
| - | - | 21.826 | 0.06 | 14.164 | 0.10 | 15.441 | 0.32 |
| - | - | 22.473 | 1.15 | 15.428 | 0.28 | 16.042 | 0.25 |
| - | - | 29.177 | 0.07 | 16.001 | 16.17 | 32.35 | 0.10 |
| - | - | - | - | 17.401 | 1.63 | 21.849 | 0.13 |
| - | - | - | - | 21.861 | 0.54 | 26.125 | 2.10 |
| - | - | - | - | 23.798 | 0.14 | 29.146 | 11.40 |
| - | - | - | - | 26.128 | 1.41 | 29.868 | 0.15 |
| - | - | - | - | 29.146 | 7.95 | - | - |

Rt: Retention time; HM: Herbal medicament

Figure 1: Structure of 7,7 dimethyl 5 (2 p tolyl(propyl) 6, 7 dihydro 1,3,4 oxadiazepin 2 amine (compound I)

and concentrated under vacuum. The marker compound, I was characterized as 7,7-dimethyl-5-(2-p-tolylpropyl)-6,7-dihydro-1,3,4-oxadiazepin-2-amine employing different spectral techniques such as FTIR, 1H and 13C NMR, Mass. The purity of compound I was checked by TLC and HPLC and was found >99% pure.

Infrared data

IR (KBr) (cm⁻¹): 3411.7, 3015.2, 1667.2, 1564.5, 1514.7, 1484.0, 1216.1, 1054.7, 759.7, 669.1, 544.7.
Nuclear magnetic resonance data

\[^1\text{H} \text{NMR} (400 \text{ MHz, DMSO-}_d): \delta \ 7.13 \ (d, 2 \text{H}, J = 7.68), \delta \ 7.08 \ (2 \text{H}, J = 7.56), \delta \ 5.93 (5 \text{S}, 2 \text{H}), \delta \ 3.01 (dd, 1 \text{H}, J = 6.90), \delta \ 2.5 (m, 4 \text{H}), \delta \ 2.24 (s, 3 \text{H}), \delta \ 1.31 (s, 3 \text{H}), \delta \ 1.23 (s, 3 \text{H}), \delta \ 1.18 (d, 3 \text{H}, J = 6.72). \]

\[^1\text{C} \text{NMR (400 MHz, DMSO-}_d): \delta \ 155.41, 153.28, 142.96, 135.04, 128.86, 126.69, 61.66, 50.85, 37.98, 36.55, 26.00, 25.78, 22.29, 20.59. \]

Mass data

Mass: M/z 274.5 (M\(^+\)+1), 275.6 (M\(^+\)+2), HRMS m/z 274.1932 (M\(^+\)+1), 275.1956 (M\(^+\)+2).

Elemental analysis data

Anal. Calc. for C\(^\text{71.04}\)H\(^\text{8.77}\)N\(^\text{14.62}\): C 71.04, H 8.77, N 14.62; Found: C 68.179, H 8.517, N 13.247.

The binding poses in the docking studies are energetically favorable. The \textit{in vitro} anticancer activity of NCCL and compound I was checked against four cancer cell lines namely MCF-7, MDA-MB-231, PC-3 and DU-145 at concentrations of 200, 100, 50, 25, 12.5, and 6.25 µg/ml, in triplicates. \(\text{IC}_{50}\) values in µM against various cell lines are given in Table 2.

**DISCUSSION**

NCCL was prepared from HM which is a hexane soluble fraction of \textit{C. longa}. Compound I, 7, 7-dimethyl-5-(2-p-tolylpropyl)-6, 7-dihydro-1,3,4-oxadiazepin-2-amine (C\(^\text{71.04}\)H\(^\text{8.77}\)N\(^\text{14.62}\)), was isolated from NCCL by preparative HPLC. Compound I is characterized using 1D/2D Nuclear magnetic resonance data and infrared spectroscopy (IR), was isolated from \textit{Curcuma longa} (NCCL) by preparative HPLC. Compound I is characterized using 1D/2D Nuclear magnetic resonance data and infrared spectroscopy (IR).

\[\text{IC}_{50}\] values of NCCL and compound I in µM against various cell lines

| Compound | MCF-7 | MDA-MB-231 | PC-3 | DU-145 |
|----------|-------|------------|------|--------|
| NCCL     | 2.093 | 8.57 | 9.24  | 10.78  |
| I        | 17.28 | >20 | 19.82 | >20    |
| TAM      | 5.5   | 11.4 | 10    | 5.5    |
| Epirubicin | 3.7  | 5.6  | 5.7   | 9.9    |

With \textit{in silico} results in hand, it was thought worthwhile to do \textit{in vitro} studies to support the \textit{in silico} studies. The \textit{in vitro} anticancer activity of NCCL and compound I was checked against cancerous (breast and prostate) cells at different concentrations (200, 100, 50, 25, 12.5, and 6.25 µg/ml, in triplicates). The cell lines used in this investigation include MCF-7, MDA-MB-231, PC-3, and DU-145. NCCL significantly inhibited the proliferation of breast cancer cell MCF-7 (\(\text{IC}_{50}=2.1 \mu\text{M}\)), and MDA-MB-231 (\(\text{IC}_{50}=8.5 \mu\text{M}\)) as well as prostate cell lines PC-3 (\(\text{IC}_{50}=9.2 \mu\text{M}\)) and DU-145 (\(\text{IC}_{50}=10.7 \mu\text{M}\)) [Table 2].

**CONCLUSION**

We have reported for the first time, the synthesis of a novel chemically modified bioactive fraction from HM (NCCL) and isolation of a novel marker compound I, 7,7-dimethyl-5-(2-p-tolylpropyl)-6, 7-dihydro-1,3,4-oxadiazepin-2-amine (reaction product of ar-turmerone with semicarbazide). Their inhibitory effects on ERs were evaluated using \textit{in silico} followed by \textit{in vitro} studies. A molecular docking study has revealed that compound I have minimum binding energy with favorable drug-likeness scores and may be considered a good estrogen blocker. The order of NCCL sensitivity was MCF-7 > MDA-MB-231 > PC-3 > DU-145. NCCL fraction having residual components induces more cell death than compound I alone. NCCL turned out to be a potent anticancer agent over compound I against these four cancer cell lines and has emerged as a promising alternative to the marketed TAM.

Further work is in progress in our laboratory to evaluate their \textit{in vivo} anticancer potential and in-depth mechanistic aspects.
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Conflicts of interest
There are no conflicts of interest.

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