Vol. 14/ No.1/2-12/Nov-Jan/2024

ISSN: (print) 2231–3087 / (online) 2230-9632

CODEN: HLEEAI http://heteroletters.org



# **Graphical abstract**

Paper-1

Heterocyclic Letters 14: iss.-1 (2024), 13-16

An efficient copper catalyzed synthesis of 2-Methoxy-3-(1H-1,2,3-triazol-1-yl) methyl)quinoxalines under Click conditions

Shashikala Kethireddy<sup>1</sup>, AsraBanu Syeda<sup>2</sup>, Balaraju Vudari<sup>3</sup>, Ravi Kumar Bommera<sup>4</sup> and Laxminarayana Eppakayala<sup>3\*</sup>

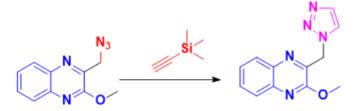
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Paper-2

Heterocyclic Letters 14: iss.-1 (2024), 17-27

Synthesis of 2-bromo-6-(4,5-diphenyl-1-((1-phenyl-1*H*-1,2,3-triazol-4-yl) methyl)-1*H*-imidazol-2-yl) pyridine derivatives: Evaluation of anticancer activity and molecular docking studies

Lakavath Ramdas, Dharmasothu Veeranna Guguloth Ravi Jadhav Ramchander Ramchander

<sup>1</sup>Department of Chemistry, University College of Science, Osmania University, Hyderabad, Telangana-500 007, India. Corresponding author: Jadhav Ramchander

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Paner-

Heterocyclic Letters 14: iss.-1 (2024), 29-42

Synthesis, characterization and biological activity of 3-((4-substituted phenyl-1*h*-1,2,3-triazol-1-yl)methyl)-2*h*-chromen-2-ones

# M. Mahesha, P. Sanjeevaa, G. Manjunatha, G. Bheemarajua, V. Kamala Prasadb and P. Venkata Ramanaa\*

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<sup>b</sup>Denisco Chemicals Pvt. Ltd., Hyderabad-500055, Telangana, India.

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In the present study, a novel series of 3-((4-substituted phenyl-1*H*-1,2,3-triazol-1-yl)methyl)-2*H*-chromen-2-ones were synthesized in good yields using substituted 3-(azidomethyl)-2*H*-chromen-2-one as a precursor. Substituted 3-(azidomethyl)-2*H*-chromen-2-ones were in turn synthesized using substituted 2-oxo-2*H*-chromene-3-carboxylic acid as starting material. All the newly synthesized compounds were characterized by IR, NMR, mass spectra and elemental analyses. The newly synthesized chromen-2-ones were screened for their antibacterial and antifungal activities by disc diffusion method.

$$R = -H, -CH_3, -CI$$
(i) TEA, ECF, THF, - 15°C, 30 min, NaBH<sub>4</sub>, H<sub>2</sub>O, 0°C, 5 min.

(ii) Imidazole, PPh<sub>3</sub>, I<sub>2</sub>, Dry DCM, Over night

(iii) DMF, NaN<sub>3</sub>, rt, 3hr
(iv) CuSO<sub>4</sub>, Sodium ascorbate, rt,

Scheme

Paper-4

Heterocyclic Letters 14: iss.-1 (2024), 43-53

Bi (iii) catalyzed one pot-two component approach: Synthesis of 2-aminoaryl Benzimidazoles

# S N Murthy Boddapati, $^{\rm a*}$ Bhuvaneswari Chalapaka, $^{\rm a}$ Satish V A N $^{\rm a}$ Srinivas Pilli $^{\rm a}$

 $t-BuOH + H_2O(1:1) rt$ 

a Dept. of Chemstry, Sir C. R. Reddy College, Eluru, Andhra Pradesh, India-534007. \*Correspondence:snmurthyboddapati@gmail.com

A simple rapid one-pot two component method was developed for the synthesis of 2-aminoaryl Benzimidazoles from various *o*-phenylenediamine reacts with phenylisothiocyanates using an efficient, inexpensive and readily available Bi(NO3)<sub>2</sub>.5H<sub>2</sub>O Catalyst. The protocol proceeds through Nucleophilic addition and Bi(III) promoted desulfurization and C-N bond formation reactions.

$$R \xrightarrow{\parallel} NH_2 + R_1 - N = C = S \xrightarrow{1 \text{ eq. } Cs_2CO_3} EtOH:H_2O \\ tt, 2h R_1 \xrightarrow{\parallel} NHR_1 \xrightarrow{R} R R \xrightarrow{R} R R \xrightarrow{R} R R \xrightarrow{R}$$

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Paper-

Heterocyclic Letters 14: iss.-1 (2024), 55-63

An approach to rapid one-pot multi-component synthesis of 1,4-dihydropyrano [2,3-c] pyrazole catalyzed by core shell NiO-C<sub>03</sub>O<sub>4</sub>@CuO nanoplates under solvent-free conditions.

S.V. Thakare, a\* A.V. Borhade, b S. L. Sangle c

a Department of Chemistry K.S.K.W. Arts, Commerce and Science College CIDCO Nashik 422008 Maharashtra India

b Department of Chemistry H.P.T. Arts & R.Y.K. Science College Nashik 422 005, Maharashtra India

c Department of Chemistry Shri Swami Shatkopacharyaji Maharaj (S.S.S.M.) Arts, Science and Commerce College, Saikheda Corresponding author Email: <a href="mailto:thakare.savita1@gmail.com">thakare.savita1@gmail.com</a>

Core shell NiO-Co3O4@CuO nanoplates is novel, recoverable and cost-effective nanoscale heterogeneous solid catalysts for one-pot multi-component solvent free synthesis of 1,4-dihydropyrano [2,3-c] pyrazole derivatives.

R= H, 4-F, 4-Cl, 4-OH, 4-Me, 3-N0<sub>2</sub>, 4-OMe

Paper-6

Heterocyclic Letters 14: iss.-1 (2024), 65-72

Synthesis, characterization and antimicrobial activity of novel chalcone derivatives having 1-[4-(benzyloxy)-3 chlorophenyl] ethanone moeity.

Shreekant Prajapati<sup>a</sup>, Sanjay S. Shah<sup>b</sup>, Saddam. S. Sipai<sup>c</sup>

<sup>a</sup>Ph.D. Scholar, Department of Chemistry, H. N. G. U, Patan

<sup>b</sup>Professor, Department of Chemistry, Shri U P Arts, Smt. M. G. Panchal Science & V.L. Shah Commerce College, Pilvai, Gujarat, India

<sup>c</sup>Research Assistant, Department of Chemistry, Shri U P Arts, Smt. M. G. Panchal Science & V.L. Shah Commerce College, Pilvai, Guiarat. India

Corresponding Author Email: <a href="mailto:shreekant12041991@gmail.com">shreekant12041991@gmail.com</a>

The simple and highly effective clasian Schmidt reaction occur between novel ketone and different aldehyde and gives chalcone. Reaction performed at room temperature for 4 hour.

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Paper-7

Heterocyclic Letters 14: iss.-1 (2024), 73-81

Synthesis of coumarin-coupled pyrazole and isoxazole compounds

#### Abhimanyu Pawara\*, Kishor Naktodea, Kishore Purib and Santosh Gaikwadb, c

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- <sup>b</sup> Department of Chemistry, Shri Shivaji College of Arts, Commerce and Science, Akola 444 003 (Maharashtra), India.
- <sup>c</sup> Department of Chemistry, Late Pundalikrao Gawali Arts and Science College, Shirpur (Jain), Dist. Washim 444504, Maharashtra, India.

Corresponding author E-mail: pawarabhay909@gmail.com

In the current study, we synthesized new compounds, namely 3-(substituted-(Furan-2-yl) acryloyl)-2H-chromen-2-one, 3-(5-substituted-(furan-2-yl)-4,5-dihydro-1H-pyrazol-3-yl)-2H-chromen-2-one, 3-(3-(5-substituted-furan-2-yl)-4,5-dihydro-1-(2,4-dinitrophenyl)-1H-pyrazol-5-yl)-2H-chromen-2-one, and 3-(5-(5-substituted-furan-2-yl)-4,5-dihydroisoxazol-3-yl)-2H-chromen-2-one, along with derivatives. Characterization of these novel compounds was achieved using infrared (IR), 1H NMR, 13C-NMR spectroscopy, and elemental analysis. These reactions were conducted in the presence of sodium acetate as a catalyst and ethanol as the solvent.

Paper-8

Heterocyclic Letters 14: iss.-1 (2024), 83-88

An Efficient Synthesis of Bis Indolyl Methane Using PMImBr

# Rakesh P. Chaudharia, Ganesh R. Chaudharia, Hemant A. Mahajanba

<sup>a</sup>Department of Chemistry, Arts and Science College, Bhalod, Taluka-Yawal, Dist-Jalgaon MS, India.

<sup>b</sup>Department of Chemistry, MTES's Smt. G.G. Khadse College, Muktainagar, Taluka- Muktainagar, Dist-Jalgaon MS, India.

A simple and efficient protocol has been developed for the synthesis of a series of bisindolyl methanes using an ionic liquid and a few drops of sulfuric acid. The reaction involves a three-component, one-pot condensation of two molecules of indole and aromatic aldehydes. The environmentally friendly protocol has high yields, quick reaction times, and easy product recovery.

+ R—CHO 
$$\frac{\text{PMImBr/2 drops of H}_2\text{SO}_4}{80\,^{\circ}\text{C}}$$

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Paper-9

Heterocyclic Letters 14: iss.-1 (2024), 89-100

In silico approaches for antidepressant activity of novel pyrazole derivative

Arunkumar Malaviya<sup>a\*</sup>, Zakirhusen Gadhawala<sup>b</sup>, Vishwaskumar Panchal<sup>c</sup>

a,c Department of Chemistry, H.N.G.U., Patan, Gujarat, India

<sup>b</sup>Department of Chemistry, The HNSB LTD Science College, Himatnagar, 383001, Gujarat, India

In this study, we conducted a comprehensive structural analysis of six newly synthesized compounds, denoted as BB1-BB6, all of which featured carbaldehydes with various substituents. Structural confirmation was achieved through a battery of analytical techniques, including IR, 1H-NMR, 13C-NMR, and ESI-MS. Subsequently, we utilized advanced In Silico methodologies to evaluate the antidepressant potential of these compounds, employing molecular docking to assess their interactions with relevant receptors.

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 $R^4$ 
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 $R^4$ 

Paper-10

Heterocyclic Letters 14: iss.-1 (2024), 101-107

Screening of microbes with ability to produce surface-active biomolecules from petroleum contaminated sites.

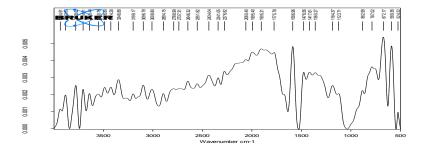
# Malode, U.A.a N.S. Satone and K.A.Kaleb

,\*Department of Microbiology, Bajaj College of Science, Wardha.442001 Maharashtra, India

<sup>b</sup> Department of Biotechnology, Bajaj College of Science, Wardha.442001 Maharashtra, India

\*Corresponding author: ulka9malode@gmail.com

Bacterial strain isolated from petroleum contaminated soil of Wardha region producessurface active biomolecule.



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Paper-1

Heterocyclic Letters 14: iss.-1 (2024), 109-119

Synthesis of N-substituted azepines *via* ionic liquid of Cu(II)Catalyzed oxidative cyclization of Dimethyl 2-(Arylamino) maleate ion with 2,5- dimethoxytetrahydrofuran under Microwave irradiaton

Kommakula Ashok<sup>1\*</sup>, Boda Sakram<sup>2</sup>, B. Manjula<sup>1</sup>, A.Chandrakar<sup>1</sup>

1. Department of Chemistry, Kamala Institute of Technology and Science, Huzurabad, Karimnagar–505468, Telangana, INDIA

2.Department of Chemistry, Osmania University, Hyderabad-500007, Telangana State, India

Email: ashok.kommakula@gmail.com

Paper-12

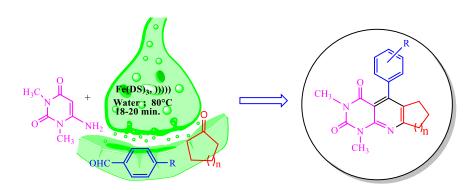
Heterocyclic Letters 14: iss.-1 (2024), 121-129

A clean and efficient synthesis of pyrimido[4,5-b]quinoline and pyrido[2,3-d]pymidine under sonication using fe(ds)<sub>3</sub> as lewis acid-surfactant-combined catalyst

Aakash Singha, SunitaYadavb, and Ruby Singhb,\*

<sup>a</sup>Department of Applied Sciences and Humanities, Dr. Shakuntla Mishra National Rehabilitation University, Lucknow, 226017, Uttar Pradesh, India.

<sup>b</sup>Department of Chemistry, School of Basic Sciences, Jaipur National University, Jaipur, 302017, Rajasthan, India \*E-mail address: drrubychem@yahoo.com



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ISSN: (print) 2231-3087 / (online) 2230-9632

CODEN: HLEEAI http://heteroletters.org



Paper-13

Heterocyclic Letters 14: iss.-1 (2024), 131-141

Synthesis, characteristics and anticancer activity of novel 2-thiohydantoin analogues

# Prashant. A. Gotmare<sup>1\*</sup>, and Sanjay V. Kolhe

\*1Department of Chemistry, Shri Shivaji Arts, Commerce and Science college, Akot, Dist.Akola-444101, (Maharashtra), India

\*E-Mail:prashantpatilchem2763@gmail.com

Aurone (0.01 M) and N-substituted thiourea (0.01 M) were added to a round bottom flask along with 10% KOH and Ethanol as a solvent. A reaction mixture was reflux about 3 hr. The mixture had been cooled, poured into ice cold water and filtered using a suction pump. The final product was recrystallized with Ethanol.

Paper-14

Heterocyclic Letters 14: iss.-1 (2024), 143-152

"MsCl-DMF/DMAc: A Vilsmeier-Haack type reagent for the synthesis of 2-chloro-3-formyl quinolines/ 2-chloro-3-acetyl quinolines"

#### Mahesh P More<sup>a</sup> and Tanuja V Kadre<sup>a\*</sup>.

<sup>a</sup>Department of Chemistry, Dr. A. P. J. Abdul Kalam University, Indore, Madhya Pradesh, India-452016.

\*Correspondence Author Email: tanujavkadre@gmail.com

2-Chloro-3-formyl quinolines and 2-chloro-3-acetyl quinolines were synthesized by reaction of N, N-dimethyl formamide (DMF)-methane sulfonyl chloride or N, N-dimethyl acetamide (DMA) - methane sulfonyl chloride (MsCl) system with corresponding acetanilide respectively. Methane sulfonyl chloride adducts with N, N'-dimethyl formamide (DMF) and N, N'-dimethyl acetamide (DMA) were explored as an alternative to Vilsmeier –Haack (VH) type reagents for effective synthesis of 2-chloro-3-formyl quinolines and 2-chloro-3-acetyl quinolines. Reaction times and temperature reduced significantly using this adduct as compaired with N, N'-dimethyl formamide - phosphoryl chloride adduct or N, N-dimethyl acetamide - phosphoryl chloride adduct. The synthesized compounds were characterized by melting points, <sup>1</sup>H- NMR and mass spectral data.

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CODEN: HLEEAI http://heteroletters.org



#### Paper-15

Heterocyclic Letters 14: iss.-1 (2024), 153-168

Polyborate supported perchloric acid as an efficient solid acid catalyst for one-pot synthesis of 12-aryl-8, 9, 10, 12-tetrahydrobenzo[a]xanthene-11-one derivatives under solvent-free condition

Dilip Aute<sup>a</sup>, Tushar Bendre<sup>a</sup>, Amol Parhad<sup>a</sup>, Vaishali Murade<sup>a</sup>, Gopinath Shirole<sup>b</sup>, Vijay Kadnor<sup>c</sup>, Bhagwat Uphade<sup>a</sup> and Anil Gadhave<sup>\*a</sup>

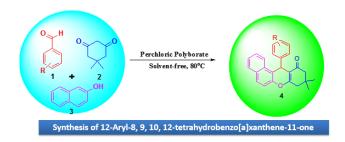
<sup>a</sup>Department of Chemistry and Research Centre, Padmashri Vikhe Patil College, Pravaranagar, Dist.Ahmednagar-413736 (MS), India.

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Paper-16

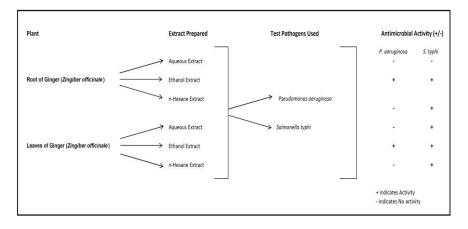
Heterocyclic Letters 14: iss.-1 (2024), 169-174

Comparative study of potential antimicrobial activity of root and leaves of ginger against selected bacteria.

# R.D. Morea, S.D. Chavanb, S.P. Moharirc, M.V. Bankard, N.P. Bhosalec, M.D. Jadhavf

<sup>a\*,b,c</sup>Department of Chemistry, Siddharth Arts, Commerce & Science College, Jafrabad, Jalna-431206(M.S.) India. <sup>d,e</sup>Department of Botany, Siddharth Arts, Commerce & Science College, Jafrabad, Jalna-431206(M.S.) India. <sup>f</sup>Department of Microbiology, Siddharth Arts, Commerce & Science College, Jafrabad, Jalna-431206(M.S.) India.

Antimicrobial activity of root and leave of ginger was studied using aqueous, ethanol and n-Hexane extract against two test pathogens namely *P.aeruginosa* and *S. typhi*.



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ISSN: (print) 2231-3087 / (online) 2230-9632

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Paper-17

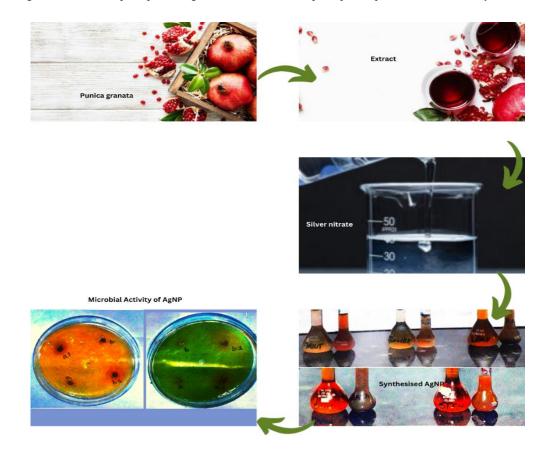
Heterocyclic Letters 14: iss.-1 (2024), 175-180

Phytofabrication of silver nanoparticles using various parts of punica granatum and their antimicrobial actvities

# Angelin Pushpa Ba\*, T. Sahaya Maria Jeyaseelib

- <sup>a</sup>Assistant Professor, Department of Nano science, Sarah Tucker College, (Autonomous) Tirunelveli 627 007, affiliated to Manonmaniam Sundaranar University, Tirunelveli, Tamilnadu, India.
- <sup>b</sup>Assistant Professor, Department of Nanoscience, Sarah Tucker College, (Autonomous) Tirunelveli 627 007, affiliated to Manonmaniam Sundaranar University, Tirunelveli, Tamilnadu, India.
- \*Corresponding author E-mail: angelin0809@gmail.com Mobile Number: 9486164199

In the present work, we described the green synthesis of silver nanoparticles (SNPs) using various parts of *Punica granatum* and their antimicrobial activities. The Bio-reduction of silver ions have been performed by the various parts of extract of *Punica granatum*. The antimicrobial activity of green synthesised SNPs is compared with AgNO<sub>3</sub> and the extract. *Punica granatum* extract showed good antibacterial activity against E.coli. The phytochemical investigation of *Punica Granatum* aqueous extract showed that it is rich in Xanthoproteins, Tannins, Terpenoids, Flavonoids, Phenols, Lacking steroids, Glycosides Coumarins, Fatty acids and Steroids. The biosynthesized SNPs were characterized by UV– visible spectrometry. Finally the phytochemical screening indicates that the plant parts are good source of bioactive principle for pharmaceutical industry.



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CODEN: HLEEAI http://heteroletters.org



Paper-18

Heterocyclic Letters 14: iss.-1 (2024), 181-201

Synthesis, characterization and application studies of Tio<sub>2</sub>-phosphotungstic acid nanocomposits prepared by blending method

# Praveendaniel M<sup>1</sup>, Rajesh Anantha Selvan P<sup>2</sup>, Paramanantha Swami Doss D<sup>3</sup>

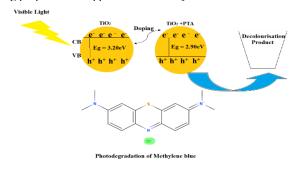
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\*Corresponding author Email: rajesh.chem@stjohnscollege.edu.in

TiO<sub>2</sub>-PTA nanocomposites were synthesized by blending method and characterized by UV-visible, FT-IR, XRD, EDX, and SEM. The results showed that there is a strong interaction between TiO<sub>2</sub> and PTA, with a decreased crystallite size and uniformly distributed dopant. The nanocomposites showed excellent photocatalytic activity, with a degradation efficiency of 99.64%. The electrical conductivities of the nanocomposites increased with increasing temperature, and the dielectric constants were positive. The nanocomposites also had significant antimicrobial activity against Streptococcus pyogenes. These results suggest that the TiO<sub>2</sub>-PTA nanocomposites have promising properties for applications in many different fields.



# **REVIEWS**

Review No.1

Heterocyclic Letters 14: iss.-1 (2024), 203-210

Mini-review on the synthesis of oxazole derivatives using non-conventional methods.

Figueroa-Valverde Lauro<sup>1\*</sup>, Rosas-Nexticapa Marcela<sup>2</sup>, Alvarez-Ramirez Magdalena<sup>2</sup>, Ortega-Cervantes Catalina<sup>2</sup>, Melgarejo-Gutierrez, Montserrat<sup>3</sup> Mateu-Armand<sup>2</sup>, López-Ramos Maria<sup>1</sup>, Mijangos-Sánchez Juliette<sup>1</sup>.

Laboratory of Pharmaco-Chemistry, Faculty of Chemical Biological Sciences, University Autonomous of Campeche, Av. Agustín Melgar s/n, Col Buenavista C.P. 24039 Campeche, Camp., México.

$$\begin{array}{c} R_2 \\ R_1 \\ \hline \end{array} \qquad \begin{array}{c} R_2 \\ R_3 \\ \hline \end{array} \qquad \begin{array}{c} \blacksquare \quad \text{Non-Conventional Methods} \\ \blacksquare \quad \text{Higher yield} \\ \blacksquare \quad \text{Higher selectivity} \\ \hline \end{array} \qquad \begin{array}{c} R_1 \\ \hline \end{array} \qquad \begin{array}{c} R_2 \\ \hline \end{array}$$

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Review No.2

Unlocking the Biological Potency of 2H-Chromene: A Comprehensive Review

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"R. R. Mehta college of science & C. L. Parikh college of commerce, Palanpur, Gujarat, India.

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Anti-diabetic activity

Antiproliferative activity

Anticancer activity

Anti-convulsant and anti-anxiety activity