

# A New Flavonoid From the Rhizomes of *Cyperusstoloniferus*

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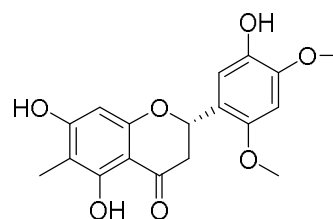
## INTRODUCTION

*Cyperusstoloniferus* Retz. (Cyperaceae) widely grows in Asian countries like India and China. It has been used for treatment of menstrual disorders, dysmenorrhoea, stomachache, and inflammation.<sup>1</sup> The rhizomes of *Cyperus* are also used in the treatment of diabetes, anticancer. Extracts, phytochemical constituents and their derivatives are also used as antirheumatic, excitant, febrifuge and tonic.<sup>2-28</sup> The rhizomes of *C. stoloniferus* were collected in botanical garden, Bhopal University, India in September, 2014 and was identified (voucher specimen No.CSR-113) by the taxonomist of the Department of Botany, Bhopal University. Collected plants, after cutting into small pieces, were dried and pulverized into a coarse powder and stored into an air-tight container.

Although *C. stoloniferus* has shown similar pharmacological effects with *C. rotundus* in India and Vietnamese folk medicine, there are very few reports on the chemical composition and biological activity of this plant. Only three studies on the essential oil composition and the analgesic activity of *C.stoloniferus* were reported so far.

## RESULTS AND DISCUSSION

The air-dried and powdered materials (2.5 kg) were extracted with methanol (6 L) in a sonic bath for 24 hours. The combined extracts were concentrated under a vacuum to obtain a crude residue (210 g), which was then resuspended in water (1 L), and successively extracted by chloroform and ethyl acetate (each 1 L-3 times). The organic layers were concentrated to give 80.1 g and 47.7 g of chloroform and ethyl acetate residues, respectively. The chloroform residue was chromatographed on a silica gel column eluted by a gradient of 1–100% acetone in hexane to afford five fractions F1-F5. Fraction 1 was yielded the following flavonoid.



## Data of the flavonoid

The compound was isolated as yellow semi solid: <sup>1</sup>H NMR (500 MHz, MeOH-d<sub>4</sub>): δ 5.62 (1H), 2.70 (1H), 2.95 (1H), 5.97 (1H, s, H-8), 6.69 (1H), 7.00 (1H), 1.97 (3H), 3.83 (3H), 3.90 (3H); <sup>13</sup>C NMR (125 MHz, MeOH-d<sub>4</sub>): δ 75.2, 43.4, 198.0, 162.6, 105.3, 166.1, 95.2, 162.6, 103.0, 120.5, 151.2, 98.7, 149.5, 141.4, 114.5, 61.9, 56.9, 56.6; ESIMS m/z: 347.13.

## CONCLUSION

Although several flavanones and stilbenes have been previously isolated from *Cyperus* species, the present study is the first report for the isolation of a new flavanone from the *C. stoloniferus* rhizomes. In addition, the C-methyl-flavonoid skeleton was found for the first time from the Cyperaceae family. Piceatannol has been isolated in low yield from other *Cyperus* plants, however this compound was found as a main component (approx. 0.18% dry weight) in the rhizomes of *C. stoloniferus*. Thus compound 1 might be considered as an important chemotaxonomic marker of *C. stoloniferus*.

## REFERENCES

1. Vu VD and Mai TT. Vietnamese J Pharmacol. 1994;223:16.
2. Reddy PP, Tiwari AK, Rao RR, Madhusudhana K, Rao VRS, Ali AZ, Babu KS and Rao JM. New Labdane Diterpenes as Intestinal  $\alpha$ -glucosidase Inhibitor from Antihyperglycemic Extract of *Hedychium spicatum* (Ham. Ex Smith)

- Rhizomes. *Bioorg Med Chem Lett.* 2009;19(9):2562-2565.
- Reddy PP, Rao RR, Rekha KS, Babu K, Shashidhar J, Shashikiran G, Vijaya Lakshmi V and Rao JM. Two New Cytotoxic Diterpenes from the Rhizomes of *Hedychiumspicatum*. *Bioorg Med Chem Lett.* 2009;19(1):192-195.
  - Reddy PP, Rao RR, Shashidhar J, Sastry BS, Rao JM and Babu KS. Phytochemical Investigation of LabdaneDiterpenes from the Rhizomes of *Hedychiumspicatum* and Their Cytotoxic Activity. *Bioorg Med Chem Lett.* 2009;19(21):6078-6081.
  - Reddy PP, Lavekar AG, Babu KS, Rao RR, Shashidhar J, Shashikiran G and Rao JM. Synthesis, Cytotoxic Activity and Structure-Activity Relationships of Hedychenone Analogues. *Bioorg Med Chem Lett.* 2010;20(8):2525-2528.
  - Fajemiroye JO, Galdino PM, Florentino IF, Da Rocha FF, Ghedini PC, Polepally PR, Zjawiony JK and Costa EA. Plurality of Anxiety and Depression Alteration Mechanism by Oleanolic Acid. *J Psychopharmacol.* 2014;98:923-934.
  - Polepally PR, White K, Vardy E, Roth BL, Ferreira D and Zjawiony JK. Kappa-Opioid Receptor-Selective Dicarboxylic Ester-Derived Salvinorin A Ligands. *Bioorg Med Chem Lett.* 2013;23: 2860-2862.
  - Polepally PR, Setola V, Vardy E, Roth BL and Zjawiony JK. New Michael Acceptor-Type of Salvinorin A Ligands to Kappa-Opioid Receptor. *Planta Medica.* 2013;79(05):P41.
  - Polepally PR, White K, Roth BL and Zjawiony JK. Convenient Synthesis and In Vitro Pharmacological Activity of Thioesters of Salvinorin B. *Planta Medica.* 2013;79(05):P43.
  - Polepally PR, Roth BL, White K and Zjawiony JK. Synthesis and Biological Evaluation of New Salvinorin B-Sulfonate Ester Ligands to Opioid Receptors. *Planta Medica.* 2013;79(05):P44.
  - Polepally PR, Roth BL, White K, Ferriera D and Zjawiony JK. Synthesis and In Vitro Biological Evaluation of New Dicarboxylic Ester-Type Salvinorin A Analogs. *Planta Medica.* 2013;79(05): P42.
  - Polepally PR, Huben K, Vardy E, Setola V, Roth BL, Mosier PD and Zjawiony, JK. Michael Acceptor Approach to the Design of New Salvinorin A-Based High Affinity Ligands for the Kappa-Opioid Receptor. *European Journal of Medicinal Chemistry.* 2014;85:818-829.
  - Polepally PR, White K, Roth BL and Zjawiony JK. Synthesis and In Vitro Pharmacological Activity of C-2 Modified New Salvinorin A Analogues. *Planta Medica.* 2013;79(05):P45.
  - Polepally PR, Setola V, Vardy E, Roth BL, Mosier PD and Zjawiony JK. New Salvinorin A-Derived Ligands to Opioid Receptors. *Planta Medica.* 2012;78:PI238.
  - Polepally PR, Setola V, Vardy E, Roth BL, and Zjawiony, JK. Michael Acceptor Approach to the Design of New Salvinorin A-Based High Affinity Ligands for the Kappa-Opioid Receptor. *Planta Med.* 2013;79(05):P45.
  - Polepally PR, White K, Vardy E, Roth BL, Ferreira D and Zjawiony JK. Kappa-Opioid Receptor-Selective Dicarboxylic Ester-Derived Salvinorin A Ligands. *Bioorg Med Chem Lett.* 2013;23: 2860-2862.
  - Polepally PR, White KL, Roth BL and Zjawiony JK. Design, synthesis and pharmacological activity of new C(2)-modified salvinorin A analogues. *Planta Medica.* 2014;80(10):PF8.
  - Rao RR, Tiwari AK, Reddy PP, Babu KS, Ali AZ, Madhusudana K and Rao JM. New Furanoflavonoids, Intestinal  $\alpha$ -glucosidase Inhibitory and Free-Radical (DPPH) Scavenging, Activity from Antihyperglycemic Root Extract of *Derris indica*. *Bioorg Med Chem.* 2009;17(14): 5170-5175.
  - Rao RR, Tiwari AK, Reddy PP, Babu KS, Suresh G, Ali A Z, Madhusudana K, Agawane S B, Badrinaraya, P, Narahari GS and Rao JM. Synthesis of Antihyperglycemic,  $\alpha$ -glucosidase Inhibitory, and DPPH Free Radical Scavenging Furanochalcones. *Med Chem Res.* 2012;21(6): 760-774.
  - Rao RR, Chaturvedi V, Babu KS, Reddy PP, Rao VRS, Sreekanth P, Sreedhar S and Rao JM. Synthesis and Anticancer Effects of Pongamol Derivatives on Mitogen Signaling and Cell Cycle Kinases. *Med Chem Res.* 2012;21:634-641.

21. Raju BC, Pradeep DVS, Reddy PP and Rao JM. CBr<sub>4</sub> Catalyzed Synthesis of Aryl-14H-dibenzo [a,j] Xanthenes Under Solvent-Free Conditions. *Lett in Org Chem.* 2008;5(6):450-454.
22. Reddy PP, Raju BC and Rao JM. A Facile One-Pot Friedlander Synthesis of Quinoline Derivatives. *J Chem Res.* 2008;12(12):679-682.
23. Sałaga M, Polepally PR, Zakrzewski PK, Cygankiewicz A, Sobczak M, Kordek R, Zjawiony JK, Krajewska WM and Fichna J. Novel orally available salvinorin A analog PR-38 protects against experimental colitis and reduces abdominal pain in mice by interaction with opioid and cannabinoid receptors. *Biochemical pharmacology.* 2014;92(4):618- 626.
24. Suresh G, Reddy PP, Babu KS, Shaik TB and Kalivendi SV. Two New Cytotoxic Labdane Diterpenes from the Rhizomes of *Hedychium coronarium*. *Bioorg Med Chem Lett.* 2010; 20(24):7544-7548.
25. Sałaga M, Polepally PR, Sobczak M, Grzywacz D, Sibaev A, Storr M, Dorego JC, Zjawiony JK and Fichna J. Novel orally available salvinorin A analog PR-38 inhibits gastrointestinal motility and reduces abdominal pain in mouse models mimicking irritable bowel syndrome. *J Pharmaceutical Experimental Therapeutics.* 2014;350(1):69-78.
26. White KL, Scopton AP, Rives ML, Bikulatov RV, Polepally PR, Brown PJ, Kenakin T, Javitch JA, Zjawiony JK and Roth BL. Identification of Novel Functionally Selective  $\kappa$ -Opioid Receptor Scaffolds. *Mol Pharmacol.* 2014;85:83-90.
27. White KL, Robinson JE, Zhu H, DiBerto JF, Polepally PR, Zjawiony JK, Nichols DE, Malanga CJ and Roth BL. The G-Protein-Biased  $\kappa$ -Opioid Receptor Agonist RB-64 Is Analgesic with a Unique Spectrum of Activities In Vivo. *Journal of Pharmacology and Experimental Therapeutics.* 2015; 352(1):98-109.
28. Zjawiony JK, Polepally PR, Roth BL, Setola V and Vardy E. Design and Synthesis of Natural-Product Based Ligands with High Affinity to the Kappa-Opioid Receptor. *Planta Medica.* 2011;77(12):SL4.