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SYNTHESIS OF ISOTHIAZOLOPYRIMIDINE DERIVATIVES
AS POTENTIAL ANTI-INFLAMMATORY AGENTS

Mr. Athiwat Hutchaleelaha

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By Mr. Athiwat Hutchaleelaha

Department Pharmaceutical Chemistry

Thesis Advisor Assistant Prof. Chamnan Patarapanich, Ph.D.
 Associate Prof. Boonardt Saisorn, M.S. in Pharm.



Accepted by the Graduate School Chulalongkorn University in
Partial Fulfilment of the Requirements for the Master's degree.

..... *Thavorn Vajrabhaya* Dean of Graduate School
(Professor Thavorn Vajrabhaya, Ph.D)

Thesis Committee

..... *Sunibhond Pummangura* Chairman

(Associate Prof. Sunibhond Pummangura, Ph.D.)

..... *Boonardt Saisorn* Member

(Associate Prof. Boonardt Saisorn, M.S. in Pharm)

..... *Wallee Vanichseni* .. Member

(Assistant Prof. Wallee Vanichseni, M.Phil.)

..... *Rapepol Bavovada* Member

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..... *Chamnan Patarapanich* Member

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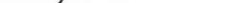
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ลายมือชื่อนิสิต *กานต์ พูล*

ถ่ายทอดความรู้ทางวิชาการที่เปรียบเสมือน...

ลายมือชื่ออาจารย์ที่ปรึกษาawan 



พิมพ์ด้วยน้ำหมึกคั่งชื่อ ทำขานั้นพนธ์ภัยในกรอบสี่เหลี่ยม เนื้อที่อยู่ในเมืองเดียว

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This investigation was to study the synthesis method of 5-substituted-3-(substituted)amino-6-hydroxy [or (substituted) amino] isothiazolo [3,4-d] pyrimidine-4-one which were expected to possess antiinflammatory activity.

The target products were synthesized via 4 steps, ie: 1) synthesis of isothiazole ring by the reaction of ethylcyanoacetate, sodium ethoxide and methyl or phenylisothiocyanate, then cyclisation with cold chloramine solution. 2) Synthesis of isothiazolo [3,4-d] pyrimidine derivatives by the reaction of isothiazole with methyl or phenylisothiocyanate under reflux condition. 3) Methylation of isothiazolo [3,4-d] pyrimidine derivatives with dimethyl sulphate under alkali condition to afford four 5-substituted-6-methylthio-3-(substituted) aminoisothiazolo [3,4-d] pyrimidine-4-ones and three 7-methyl-5-substituted-3-(substituted) aminoisothiazolo [3,4-d] pyrimidine-4-one-6-thiones. 4) Hydrolysis of methylsulphide intermediates with 6N hydrochloric acid under reflux condition to obtain four 5-substituted-3-(substituted) amino-6-hydroxyisothiazolo [3,4-d] pyrimidine-4-ones. When treating 5-methyl-6-methylthio-3-methylamino-isothiazolo [3,4-d] pyrimidine-4-one with piperidine, the expected 5-methyl-3-methylamino-6-(1'-piperidino) isothiazolo [3,4-d] pyrimidine-4-one was obtained.

ศูนย์วิทยทรัพยากร
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ลายมือชื่อนิสิต O. M.

ลายมือชื่ออาจารย์ที่ปรึกษา ...

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