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LIST OF ABBREVIATIONS

 $[\alpha]_D$ specific rotation

% v/v % volume/volume % w/v % weight/volume

% w/w % weight/weight

μg microgram

μL microliter

μM micromolar

AB_q AB quatet

ABTS 2,2'-azinobis-(3-ethylbenzothiazoline-6-sulfonic acid)

Ac acetate (¹³C-NMR)

Ac₂O acetic anhydride

AcOH acetic acid

AcONH₄ ammonium acetate

AIBN 2,2'- azobisisobutyronitrile

AllTMS allyltrimethylsilane

AlMe₃ trimethylaluminium

APT attached proton test

Ar aromatic (¹³C-NMR)

ASAP atmospheric pressure solids analysis probe

BnBr benzyl bromide

BnCl benzyl chloride

BnNH₂ benzyl amine

Boc tert-butyloxycarbonyl
Boc₂O di-tert-butyl dicarbonate

bp boiling point

br broad (¹H-NMR)

brd broad doublet (¹H-NMR)

brs broad singlet (¹H-NMR)

^tBuOH *tert*-butanol

Bu₃SnH tributyltin hydride

c concentration °C degree Celsius

CAM ceric ammonium molybdate solution

cat. catalyst

CbzCl benzyl chloroformate

CDCl₃ deuterated chloroform

CD₃OD deuterated methanol

 $Ce(SO_4)_2$ cerium(IV) sulphate

CF₃CO₂H trifluoroacetic acid

CFU colony forming unit

CHCl₃ chloroform

CH₂Cl₂ dichloromethane

CH₃CN acetonitrile
CH₃NO₂ nitromethane

cm centimeter

¹³C NMR carbon-13 nuclear magnetic resonance

CO₂ carbon dioxide

CrO₃ chromium(VI) oxide

Cu(OAc)₂ copper(II) acetate

d doublet (¹H-NMR)

DCM dichloromethane

dd doublet of doublets (¹H-NMR)

ddd doublet of doublets (¹H-NMR)

DFT density functional theory

DHAP dihydroxyacetone phosphate

dm decimeter

DMAP 4-dimethylaminopyridine

DMF dimethylformamide

D₂O deuterium oxide

DPPH 2,2'-diphenyl-1-picrylhydrazyl

 δ chemical shift

EC₅₀ half maximal effective concentration

ED₅₀ median effective dose

ESI + electrospray ionization (positive ion mode)

et al. and others

Et₃N triethylamine

Et₂O diethyl ether

EtOAc ethyl acetate

EtOH ethanol

Et₃SiH triethylsilane

eV electron volt

exp experimental

g gram

GC-FID gas chromatography-flame ionization detector

GC-MS gas chromatography-mass spectrometry

gHSQC gradient Heteronuclear Single Quantum Correlation

Grubbs' II catalyst benzylidene[1,3-bis(2,4,6-trimethylphenyl)-2-

imidazolidinylidene]dichloro(tricyclohexylphosphine)

ruthenium

h hour

H₂ hydrogen gas

HCl hydrochloric acid

HCOONH₄ ammonium formate

He helium

H₂NCO₂Bn benzyl carbamate

¹H-NMR proton nuclear magnetic resonance

H₂O₂ hydrogen peroxide

HRMS high-resolution mass spectra

H₂SO₄ sulfuric acid

Hz hertz

IC₅₀ half maximal inhibitory concentration

i.d. inner diameter in vacuo in a vacuum

IR infrared spectroscopy

J coupling constants (NMR)

K₂CO₃ potassium carbonate

kg kilogram

KO^tBu potassium tert-butoxide

KOH potassium hydroxide

L liter

lit. literature

m meter

m multiplet (¹H-NMR)

MeOH methanol

Me₂S dimethyl sulfide

MeSO₂Cl methanesulfonyl chloride (mesyl chloride)

mg milligram

MgSO₄ magnesium sulfate

MHz megahertz

MIC minimum inhibitory concentration

mL milliliter

mm millimeter

mM millimolar

mmol millimole

MsCl methanesulfonyl chloride (mesyl chloride)

MS mass spectrometry

MTT 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium

bromide

m/z mass-to-charge ratio

n- normal-

N₂ nitrogen gas

NaBH₄ sodium borohydride

Na₂CO₃ sodium carbonate

NaH sodium hydride

NaHCO₃ sodium bicarbonate

NaIO₄ sodium periodate

NaOH sodium hydroxide

NEt₃ triethylamine

NH₃ ammonia

(NH₄)₆MoO₂₄ ammonium molybdate

NIST national institute of standards and technology

NISTREP NIST replicates correction

nm nanometer

NMO 4-methylmorpholine N-oxide

NOESY Nuclear Overhauser Effect Spectroscopy

o/n over night

OsO₄ osmium tetroxide

Pd/C palladium on carbon

PdCl₂ palladium(II) chloride

Pd(OH)₂/C Palladium hydroxide on carbon

petrol petroleum spirit

PFU plaque forming unit

Ph phenyl

PhMe toluene

Ph₃P⁺CH₃Br⁻ methyltriphenylphosphonium bromide

Ph₃P=CHCOCH₃ acetylmethylenetriphenylphosphorane

p.o. per os (by mouth)

POCl₃ phosphorus oxychloride

ppm parts per million

PPTS pyridinium *p*-toluenesulfonate

p-TsOH *p*-toluenesulfonic acid

Py pyridine

q quatet (¹H-NMR)

RA relative area

RCM ring closing metathesis

 R_f retention factor

RhuA L-rhamnulose-1-phosphate aldolase

RI Kovàts retention indices

ROESY rotating frame overhauser effect spectrometry

rpm revolutions per minute

rt room temperature

s singlet (¹H-NMR)

t triplet (¹H-NMR)

TBAI tetrabutylammonium iodide

TBDMSCl *tert*-butyldimethylsilyl chloride

*t*BuOK Potassium *t*-butoxide

TCDI thionocarbonyl-1,1'-diimidazole

TES-Cl triethylchlorosilane

THF tetrahydrofuran

TiCl₄ titanium tetrachloride

Ti(Oⁱ-Pr)₄ titanium isopropoxide

TLC thin layer chromatography

TMS trimethylsilyl

TMSOTf trimethylsilyl trifluoromethanesulfonate

TrCl trityl chloride

UDP-Galf uridine diphosphogalactofuranose

Yb(OTf)₃ ytterbium(III) trifluoromethanesulfonate

Z Carboxybenzyl group

Zn Zinc

ข้อความแห่งการริเริ่ม

- วิทยานิพนธ์นี้เป็นการศึกษาฤทธิ์ทางชีวภาพ ได้แก่ ฤทธิ์ต้านมะเร็ง ฤทธิ์ยับยั้งวัณโรค ยับยั้งเชื้อ 1) เฮอร์ปีส์ ซิมเพล็กซ์ ไวรัส ไทป์ 1 ฤทธิ์ต้านจุลชีพ และฤทธิ์ต้านอนุมูลอิสระของสารสกัดหยาบ, น้ำมัน และสารที่แยกออกมาได้จากพืชสมุนไพรไทยจำนวน 3 ชนิค ได้แก่ ใบเงิน ขนไก่ทองคำ และผักคืด นอกจากนี้ยังรายงานผลการวิเคราะห์ด้วยเทคนิค จีซี-เอฟไอดี และจีซี-เอ็มเอสของ สารสกัดหยาบและน้ำมันของพืชสมุนไพรทั้ง 3 ชนิดได้ทำการศึกษาหาโครงสร้างทางเคมีของ สารสกัดบริสุทธิ์ทั้ง 6 ชนิดจากพืชสมุนไพรดังกล่าวโดยอาศัยการวิเคราะห์ด้วยเทคนิคทาง สเปกโตรสโคปีและจีซี-เอ็มเอส พร้อมทั้งเปรียบเทียบข้อมูลที่ได้กับข้อมูลที่เคยรายงานมาก่อน โดยการศึกษาในครั้งนี้สามารถแยกสควอถีน และสติคมาสเตอรอลออกมาจากสารสกัดเฮกเซน และแยก 1,4-ใคใกลโคโลอิล-เบนซีนจากสารสกัดเอทิล อะซิเตทของต้นใบเงิน ส่วนน้ำมัน หอมระเหยของต้นขนไก่ทองคำนำมาแยกสกัดและทำให้บริสุทธิ์จนได้องค์ประกอบหลักคือ คูบีนอล และสปาฐลินอล ซึ่งคูบีนอลแสดงฤทธิ์ต้านมะเร็งปอด และสารสกัดคลอโรฟอร์มของ ผักดีดที่แสดงฤทธิ์การยับยั้งเชื้อเฮอร์ปีส์ ซิมเพล็กซ์ ไวรัส ไทป์ 1 นำมาแยกได้สารสโคโพเลติน ในวิทยานิพนธ์นี้เป็นการรายงานครั้งแรกขององค์ประกอบทางเคมี และฤทธิ์ทางชีวภาพของ น้ำมัน และสารสกัดจากใบของต้นใบเงิน, การแยกองค์ประกอบทางเคมี และทดสอบฤทธิ์ทาง ชีวภาพของสารที่สกัดจากน้ำมันของใบสดจากต้นขนไก่ทองคำ และการแยกสารที่มีฤทธิ์ทาง ชีวภาพจากสารสกัดของกิ่งต้นผักดีค จากผลการศึกษาครั้งนี้จะใช้เป็นข้อมูลในการนำพืช สมุนไพรเหล่านี้ไปศึกษาต่อ และการศึกษาค้นคว้ายาใหม่ต่อไป
- 2) วิทยานิพนธ์นี้ได้ทำการพัฒนาวิธีการสังเคราะห์สารจากธรรมชาติ (-)-สตีเวียมีน และอนุพันธ์ ของสตีเวียมีน เพื่อลดขั้นตอนและระยะเวลาในการสังเคราะห์ ลดการใช้สารเคมีในปริมาณมาก และให้ผลิตภัณฑ์ที่มีประสิทธิภาพ และมีปริมาณผลผลิตที่ดี วิธีการสังเคราะห์นี้สามารถใช้เป็น แนวทางในการสังเคราะห์สารโพลีไฮดรอกซีเลเต็ด อินโคลิซิดีน อัลคาลอยด์ชนิดอื่นๆ นอกจากนี้ผลการทดสอบฤทธิ์การยับยั้งไกลโคซิเดสของสารที่สังเคราะห์ได้ในวิทยานิพนธ์นี้ จะเป็นข้อมูลประกอบการพัฒนาโครงสร้างสารโพลีไฮดรอกซีเลเต็ด อินโคลิซิดีน อัลคาลอยด์ เพื่อให้มีฤทธิ์ที่ดีขึ้น

STATEMENT OF ORIGINALITY

- In this research, the biological activities; anticancer, antimycobacterial, anti-1) herpes simplex virus type-1, antimicrobial and antioxidant activities of the crude extracts, the essential oils and the isolated compounds of the three Thai medicinal plants (Graptophyllum pictum (L.) Griff., Gynura divaricata (L.) DC. and Solanum spirale Roxb.) were studied. Furthermore, the GC-FID and GC-MS data of the extracts and the essential oils of these plants were also revealed. The structural elucidations of six bioactive compounds from these plants were analyzed using the spectroscopic techniques and GC-MS, and then the results were compared with the previous reports. Squalene and stigmasterol were isolated from the hexane fraction, while 1,4-diglycoloyl-benzene was isolated from the EtOAc fraction of G. pictum. Cubenol and spathulenol were separated and purified from the essential oil of G. divaricata. Cubenol inhibited the NCI-H187 cell line (small cell lung cancer). The CHCl₃ extract of S. spirale, which showed the anti-herpes simplex virus type-1 activity, was separated and purified to obtain scopoletin as a bioactive compound. This is the first report on the chemical constituents and the biological activities of the essential oil and the extracts of the leaves of G. pictum, the isolation of chemical compounds from the essential oil and its biological activities of the fresh leaves of G. divaricata and the isolation another bioactive compound from the stems of S. spirale were revealed. The information from this study will be useful for further investigation and drug discovery. by Chiang Mai University
- A synthetic method for the preparation of natural (-)-steviamine and its analogues in order to reduce the synthetic procedure, time consuming, the large amounts of the chemical reagents and provide the effective products in good percentage yields were developed. This concise synthesis may be used as a guideline for the synthesis of other polyhydroxylated indolizidine alkaloids. Furthermore, the results of glycosidase inhibitory activities of the synthetic compounds in this study may provide the information for the structural development of the

polyhydroxylated indolizidine alkaloids to increase their glycosidase inhibitory activities.

