Geranylated Coumarins From Thai Medicinal Plant *Mammea siamensis* With Testosterone 5α-Reductase Inhibitory Activity

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Geranylated coumarin constituents, kayeassamin I (1) and mammeasins E (2) and F (3) were newly isolated from the methanol extract of the flowers of *Mammea siamensis* (Calophyllaceae) originating in Thailand, along with five known isolates, such as mammea E/BC (23), deacetylmmammea E/AA cyclo D (31), deacetylmmammea E/BB cyclo D (32), mammea A/AA cyclo F (34), and mammea A/AC cyclo F (35). These compounds (1–3) were obtained as an inseparable mixture (ca. 1:1 ratio) of the 3’R and 3’S forms, respectively. Among the isolated coumarins from the extract, mammeasins E (2, 22.6 µM), A (4, 19.0 µM), and B (5, 24.0 µM), kayeassamins E (9, 33.8 µM), F (10, 15.9 µM), and G (11, 17.7 µM), surangin C (13, 5.9 µM), and mammeas A/AA (17, 19.5 µM), E/BB (22, 16.8 µM), and A/AA cyclo F (34, 23.6 µM), were found to inhibit testosterone 5α-reductase.

Keywords: *Mammea siamensis*, mammeasin, 5α-reductase inhibitor, geranylated coumarin, calophyllaceae

INTRODUCTION

The Calophyllaceae plant *Mammea siamensis* (Miq.) T. Anders. is a small evergreen tree distributed in Thailand (locally called “Sarapi” or “Saraphi”), Laos, Cambodia, Vietnam, and Myanmar. The flowers of this plant have traditionally been used as a heart tonic, fever-lowering, and enhancement of appetite in Thailand (locally called “Sarapi” or “Saraphi”), Laos, Cambodia, Vietnam, and Myanmar. The flowers of this plant have traditionally been used as a heart tonic, fever-lowering, and enhancement of appetite in Thailand (locally called “Sarapi” or “Saraphi”), Laos, Cambodia, Vietnam, and Myanmar. Here, we conducted the isolation and structural verification of 1–3, as well as examined the testosterone 5α-reductase...
inhibitory activity of its coumarin constituents (1–35), including five new isolates, such as mammea E/BC (23), daecetylmaimmea E/AA cyclo D (31), daecetylmaimmea E/BB cyclo D (32), mammea A/AA cyclo F (34), and mammea A/AC cyclo F (35).

**MATERIALS AND METHODS**

**General Experimental Procedures**

The following instruments were used to obtain physical data: a SEPA-300 digital polarimeter (Horiba Ltd., Kyoto, Japan, l = 5 cm) for specific rotations; an UV-1600 spectrometer (Shimadzu Co., Kyoto, Japan) to record UV spectra; a FTIR-8100 spectrometer (Shimadzu Co.) to measure IR spectra; a JNM-ECA800 (800 MHz), JNM-ECA700 (700 MHz), JNM-ECA500 (500 MHz), and JNM-ECS400 and JNM-AL400 (400 MHz) spectrometers (JEOL Ltd., Tokyo, Japan) to determine 1H NMR spectra; JNM-ECA800 (200 MHz), JNM-ECA700 (175 MHz), JNM-ECA500 (125 MHz), and JNM-ECS400 and JNM-AL400 (100 MHz) spectrometers (JEOL Ltd.) to record 13C NMR spectra in CDCl3 at room temperature (25°C) with tetramethysilane as an internal standard; an Exactive Plus Orbitrap mass spectrometer (Thermo Fisher Scientific Inc., Waltham, MA, USA) to measure ESIMS and HRESIMS; an HPLC detector, SPD-10Avp UV-Vis (Shimadzu Co.); and Cosmosil 5C18-MS-II (Nacalai Tesque, Inc., Kyoto, Japan) HPLC columns (4.6 mm i.d. × 250 mm and 20 mm i.d. × 250 mm) for analytical and preparative purposes, respectively.

The following materials and experimental conditions were used for the column chromatography (CC): normal-phase silica gel CC, silica gel 60N (Kanto Chemical Co., Ltd., Tokyo, Japan); for the preparative purposes, respectively.

**Plant Material**

The flowers of *Mammea siamensis* were collected from the Nakhonsithammarat Province, Thailand, in September 2006, a s...
0.0023%) (Mahidol et al., 2007) together with kayeassamins E (9, 28.6 mg, 0.0113%), F (10, 98.7 mg, 0.0390%), and G (11, 43.4 mg, 0.0171%), deacetylmammea E/BC cyclo D (33, 18.6 mg, 0.0073%), and benzoic acid (10.9 mg, 0.0043%).

**Kayeassamin I (1)**
Pale yellow oil; [α]D 25°− 50.4 (c 0.63, CHCl3) ([α]D 25°− 35.52 (c 0.90, CHCl3) (Win et al., 2008a)); 1H and 13C NMR spectroscopic data (see Table 1); Negative-ion ESIMS m/z 439 [M – H]−; HRESIMS m/z 439.2116 (calcd for C26H31O6, 439.2115) (Figures S3–S7).

**Mammeasin E (2)**
Pale yellow oil; [α]D 25°− 58.9 (c 0.12, CHCl3); UV (MeOH) λmax nm (log ε): 223 (4.01), 278 (4.12), 302 (4.12); IR (KBr) νmax cm−1: 1,740, 1,713, 1,613, 1,454, 1,408, 1,284, 1,126, 1,049; 1H and 13C NMR spectroscopic data (see Table 2); Negative-ion ESIMS m/z 453 [M – H]−; HRESIMS m/z 453.2272 (calcd for C27H33O6, 453.2272) (Figures S8–S12).

**Mammeasin F (3)**
Pale yellow oil; [α]D 25°− 42.1 (c 0.45, CHCl3); UV (MeOH) λmax nm (log ε): 224 (3.89), 298 (3.82); IR (KBr) νmax cm−1: 1,732, 1,713, 1,605, 1,454, 1,381, 1,261, 1,126, 1,049; 1H and 13C NMR spectroscopic data (see Table 2); Negative-ion ESIMS m/z 453 [M – H]−; HRESIMS m/z 453.2287 (calcd for C27H33O6, 453.2272) (Figures S13–S17).

**DDQ Oxidation of Kayeassamin A (8) and Surangins C (13) and D (14)**
A solution of kayeassamin A (8, 9.0 mg) in dry-toluene (2.0 mL) was treated with 2,3-dichloro-5,6-dicyano-1,4-benzquione (DDQ, 10.0 mg) and the solution stirred at room temperature (25°C) for 4 h. The aqueous solution was saturated with sodium hydrogen carbonate (NaHCO3) and extracted with EtOAc. The EtOAc extract was washed with brine then dried over anhydrous magnesium sulfate (MgSO4) and filtered. Removal of the solvent under reduced pressure gave a residue, which was purified by HPLC [Cosmosil 5C18-MS-II, MeOH–1% aqueous AcOH (85:15, v/v)] to give kayeassamin I (1, 3.8 mg, 46%). Through the similar procedure, mammeasin E (2, 3.3 mg, 38%) and mammeasin F (3, 2.0 mg, 17%) were obtained from surangins D (14, 9.6 mg) and C (13, 12.7 mg), respectively.

**Assay for Testosterone 5α-Reductase Inhibitory Activity**
The experiment was performed in accordance with previously reported methods (Matsuda et al., 2001; Lee et al., 2012; Koseki et al., 2015) with slight modifications. In brief, the assay was performed in 48-well microplates (Sumitomo Bakelite Co., Ltd., Tokyo, Japan). The reaction solution was pre-incubated with or without a test sample (5 µL/well, dissolved in DMSO), in a potassium phosphate buffer (40 mM, pH 6.5, 490 µL/well) containing substrate (0.35 nmol of testosterone, Tokyo Chemical Industry Co., Ltd., Tokyo, Japan) and NADPH (10 nmol, Oriental Yeast Co., Ltd., Tokyo, Japan) at room temperature (25°C) for 20 min. The enzymatic reaction was initiated by the addition of rat liver S9 fractions (10 µL/well, dissolved in the phosphate buffer, 20.6 µg/well, Oriental Yeast Co., Ltd., Tokyo, Japan, lot no. 109031513) at 37°C for 30 min. After incubation, the reaction mixture was immediately heated in boiling water for 2 min to stop the reaction. Then the reaction solution of each well was transferred to a microtube and extracted with 500 µL of EtOAc. After the microtube was centrifuged (10,000 rpm, 5 min), an aliquot of each EtOAc phase (300 µL) was transferred into another tube. The solvent in the tube was evaporated and the residue was dissolved in 30 µL of acetonitrile containing an internal standard (I.S.) fluindocortisone acetate (20 µg/mL, Sigma-Aldrich, Co., LLC, St. Louis, USA). An aliquot of 2 µL was injected into the HPLC under the following conditions [Instrument: a series LC-20A Prominence HPLC system (Shimadzu Co., Kyoto, Japan); Detection: UV (254 nm); Column: Cosmosil 5C18-MS-II (Nakalai Tesque Inc., Kyoto, Japan, 5 µm particle size, 2.0 mm i.d. × 150 mm); Column temperature: 40°C; Mobile phase: MeOH–H2O (60:40, v/v); Flow rate: 0.2 mL/min; retention time: 13.5 min for testosterone and 8.0 min for I.S. A similar procedure that described above was carried out for the control tubes. The 5α-reductase inhibitory activity was determined from the following equation using the peak area ratios (r = testosterone/I.S.). Experiments were performed in triplicate or quadruple, and IC50 values were determined graphically. The 5α-reductase inhibitor finasteride (Tokyo Chemical Industry Co., Ltd., Tokyo, Japan) was used as a reference compound.

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\text{Inhibition(%) = } \left( \frac{r(T) - r(C)}{r(B) - r(C)} \right) \times 100
\]

Control (C): enzyme (+), test sample (−); Test (T): enzyme (+), test sample (+); Blank (B): enzyme (−), test sample (−).

## Statistics
Values are expressed as mean ± S.E.M. One-way analysis of variance (ANOVA), followed by Dunnett’s test, was used for statistical analysis. Probability (p) values <0.05 were considered significant.

### RESULTS AND DISCUSSION

**Effects of the Methanol Extract From the Flowers of *M. siamensis* on Testosterone 5α-Reductase**
The male sex hormones, androgens, play a crucial role in the development, growth and function of the prostate, and other androgen-sensitive peripheral tissues. In the prostate gland, androgens are involved in benign prostatic hyperplasia and prostate cancer, as well as in skin disorders, such as acne, seborrhea, androgenic alopecia, and hirsutism. Among the androgens, testosterone is the most abundant in serum and secreted primarily by the testicles and ovaries. The enzyme steroid 5α-reductase catalyzes the conversion of testosterone to the most potent natural androgen, 5α-dihydrotestosterone (Yamana et al., 2010; Yao et al., 2011; Azzouni et al., 2012). Therefore, inhibition of testosterone 5α-reductase could be useful for the treatment of the above diseases. To date, three types
of 5α-reductases, chronologically named types 1, 2, and 3 5α-reductases, have been described (Yamana et al., 2010; Azzouini et al., 2012; Titus et al., 2014). A type 2 and 3 5α-reductase inhibitor, finasteride, is currently marketed worldwide as a drug for benign prostatic hyperplasia and is also used in the treatment of hair loss (Heinzl, 1999; Tosti and Piraccini, 2000) and in the prevention of prostate cancer (Coltman et al., 1999). Therefore, 5α-reductase is considered a useful therapeutic target in the treatment and prevention of the above deceases. In particular, many heterocyclic compounds based on oxygen and nitrogen atoms often have good antiproliferative activity against a variety of solid tumor cell lines and are expected to be seeds of new anticancer agents (Sharma et al., 2018; Petel et al., 2019).

During our characterization studies on bioactive constituents from Thai natural medicines (Manse et al., 2017; Morikawa et al., 2018; Tanabe et al., 2018; Kobayashi et al., 2019), a methanol extract of the flowers of M. siamensis was found to inhibit 5α-reductase activity (IC₅₀ = 2.4 μg/mL). In order to investigate new 5α-reductase inhibitors, we conducted a search for the bioactive constituents from the flowers of M. siamensis.

### Isolation

In our previous report we described the isolation of 26 coumarins: mammesins A (4, 0.0293%), B (5, 0.0115%), C (6, 0.0008%), and D (7, 0.0047%), kayeassams A (8, 0.0578%), E (9, 0.0113%), F (10, 0.0390%), and G (11, 0.0171%), surangins B (12, 0.0271%), C (13, 0.0571%), and D (14, 0.0632%), 8-hydroxy-5-methyl-7-(3,7-dimethoxy-octa-2,6-dienyl)-9-(2-methyl-1-oxobutyl)-4,5-dihydropyrano[4,3,2-de]chromen-2-one (15, 0.0015%), 8-hydroxy-5-methyl-7-(3,7-dimethoxy-octa-2,6-dienyl)-9-(3-methyl-1-oxobutyl)-4,5-dihydropyrano[4,3,2-de]chromen-2-one (16, 0.0012%), mammee A/AA (17, 0.0494%), A/AB (18, 0.0048%), A/AC (19, 0.1056%), A/AD (20, 0.0022%), E/BA (21, 0.0045%), E/BB (22, 0.0194%), A/AA cyclo D (24, 0.0305%), A/AB cyclo D (25, 0.0097%), A/AC cyclo D (26, 0.0109%), B/AB cyclo D (27, 0.0016%), B/AC cyclo D (28, 0.0062%), E/BC cyclo D (29, 0.0058%), and deacetylmammae A/BC cyclo D (33, 0.0073%), as described previously (Morikawa et al., 2012; Ninomiya et al., 2016). In the present study, we additionally isolated kayeassamin I (1, 0.0072%) and mammesins E (2, 0.0099%) and F (3, 0.0015%).

### Table 1

| Position | 1α | 1β | 1α | 1β |
|----------|----|----|----|----|
|          | δH | δC | δH | δC |
| 2        | 159.6 | 159.6 | 159.6 | 159.6 |
| 3        | 6.60 (br s) | 6.61 (d, 0.9) | 6.61 (s) | 107.0 |
| 4        | 160.6 | 160.6 | 160.6 | 160.6 |
| 4a       | 101.0 | 101.0 | 101.1 | 101.0 |
| 5        | 155.9 | 155.9 | 155.9 | 155.9 |
| 6        | 105.8 | 106.0 | 105.8 | 105.8 |
| 7        | 162.9 | 162.9 | 162.9 | 162.9 |
| 8        | 104.5 | 104.6 | 104.5 | 104.5 |
| 8a       | 157.2 | 157.2 | 157.3 | 157.3 |
| 1′       | 5.43 (br t, ca. 8) | 71.8 | 5.43 (br t, ca. 8) | 71.7 |
| 2′       | 1.51, 1.95 (both m) | 30.7 | 1.53, 1.96 (both m) | 30.5 |
| 3′       | 1.11 (3H, t, 7.4) | 10.2 | 1.09 (3H, t, 7.4) | 10.1 |
| 4′       | 83.0 | 83.1 | 83.0 | 83.0 |
| 5′       | 5.53 (d, 10.2) | 125.0 | 5.55 (d, 10.2) | 124.8 |
| 6′       | 6.78 (d, 10.2) | 116.5 | 6.79 (d, 10.2) | 116.8 |
| 7′       | 2.09 (2H, m) | 23.0 | 2.09 (2H, m) | 23.2 |
| 8′       | 5.06 (qt, 0.9, 7.1) | 123.1 | 5.06 (qt, 0.9, 7.1) | 123.0 |
| 9′       | 1.64 (3H, d, 0.9) | 25.6 | 1.67 (3H, d, 0.9) | 25.6 |
| 10′      | 1.55 (3H, s) | 17.6 | 1.57 (3H, s) | 17.7 |
| 11′      | 206.4 | 206.4 | 206.4 | 206.4 |
| 2′′      | 3.26 (2H, t, 7.1) | 46.7 | 3.28 (2H, t, 7.1) | 46.7 |
| 3′′      | 1.78 (2H, t, 7.4, 7.1) | 18.0 | 1.78 (2H, t, 7.4, 7.1) | 18.0 |
| 4′′      | 1.04 (3H, t, 7.4) | 13.8 | 1.03 (3H, t, 7.4) | 13.8 |
| 2′′′-OH3 | 1.52 (3H, s) | 27.2 | 1.48 (3H, s) | 27.5 |
| 7-OH     | 14.47 (s) | 14.47 (s) | 14.48 (brs) | 14.48 (brs) |

*Measured by 800 MHz for 1H NMR and 200 MHz for 13C NMR.

*Reported in Win et al. (2008b) by 400 MHz for 1H NMR and 100 MHz for 13C NMR.

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**Geranylated Coumarins From M. siamensis**

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from the methanol extract of *M. siamensis* flowers as shown in Figure 1, together with six coumarins: mammeas E/BC (23, 0.0076%) and E/BD cyclo D (30, 0.0015%), deacetylmammeas E/AA cyclo D (31, 0.0005%) and E/BB cyclo D (32, 0.0023%), and mammeas A/AA cyclo F (34, 0.0010%) and A/AC cyclo F (35, 0.0068%), using normal-phase silica gel and reversed-phase ODS column chromatographic purification steps, and finally by HPLC (Figure 2).

**Structures of Kayeassamin I (1) and Mammeasins E (2) and F (3)**

Compound 1 was obtained as pale yellow oil with a negative optical rotation ([α]D

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= −50.4 in CHCl₃), and its molecular formula was deduced to be C₂₅H₃₂O₄ by high-resolution ESIMS (HRESIMS) measurement. As shown in Figure 3, the HPLC analysis suggested that 1 was obtained as an inseparable mixture (ca. 1:1 ratio). The ¹H and ¹³C NMR spectra spectroscopic properties (Table 1, CDCl₃) of 1, which were assigned with the aid of DEPT, DQF-COSY, HSQC, and HMBC experiments, were in accordance with those of kayeassamin I except for the observation of duplicate signals (1a and 1b) measured by high resolution 800 MHz NMR spectrometer: two primary, a tertiary, and two vinyl methyls [1a: δ = 1.04 (3H, t, J = 7.4 Hz, H-2‴), 1.11 (3H, t, J = 7.4 Hz, H-3‴), 1.52 (3H, s, 2‴-CH₃), 1.55 (3H, s, H-3′), 1.64 (3H, d, J = 0.9 Hz, H-9‴)], [1b: δ = 1.03 (3H, t, J = 7.4 Hz, H-2‴), 1.09 (3H, t, J = 7.4 Hz, H-3‴), 1.48 (3H, s, 2‴-CH₃), 1.57 (3H, s, H-3′), 1.67 (3H, d, J = 0.9 Hz, H-9‴)], five methylenes [1a: δ = 1.51, 1.95 (1H each, both m, H-2′, 1.71, 1.91 (1H each, both m, H-2‴), 1.78 (2H, qt, J = 7.4, 7.1 Hz, H-2‴)], 2.09 (2H, m, H-6″), 3.26 (2H, t, J = 7.1 Hz, H-2‴)], a methine bearing an oxygen function [1a: δ = 3.41, 3.43 (1H, br t, J = ca. 8 Hz, H-1‴), 1b: δ = 3.43 (1H, br t, J = ca. 8 Hz, H-1‴)], four olefinic protons [1a: δ = 5.06 (1H, qt, J = 0.9, 7.1 Hz, H-7‴), 5.53 (1H, d, J = 10.2 Hz, H-3‴), 6.60 (1H, br s, H-3), 6.78 (1H, d, J = 10.2 Hz, H-4‴), 1b: δ = 5.06 (1H, qt, J = 0.9, 7.1 Hz, H-7‴), 6.60 (1H, br s, H-3), 6.78 (1H, d, J = 10.2 Hz, H-4‴), 14.7 (3H, s, 2‴-CH₃), 14.8 (3H, s, H-3′), 15.2 (3H, s, H-3′)].

**Table 2** ¹H and ¹³C NMR spectroscopic data (CDCl₃) for mammeasins E (2) and F (3).

| Position | 2a | 2b | 3a | 3b |
|----------|----|----|----|----|
| δH       | δC | δH | δC | δH |
| 1        | 5.40 (br t, ca. 8) | 71.8 | 5.40 (br t, ca. 8) | 71.7 | 5.43 (m) | 71.8 | 5.43 (m) | 71.8 |
| 2        | 1.53, 1.96 (both m) | 30.7 | 1.53, 1.96 (both m) | 30.6 | 1.52, 1.96 (both m) | 30.7 | 1.52, 1.96 (both m) | 30.6 |
| 3        | 1.12 (3H, t, 7.3) | 10.2 | 1.09 (3H, t, 7.1) | 10.1 | 1.12 (3H, t, 7.1) | 10.2 | 1.10 (3H, t, 7.1) | 10.1 |
| 4        | 83.0 | 83.1 | 83.0 | 83.1 |
| 5        | 5.53 (d, 10.2) | 125.0 | 5.54 (d, 10.2) | 124.9 | 5.54 (d, 10.2) | 124.9 | 5.54 (d, 10.2) | 124.8 |
| 6        | 6.79 (d, 10.2) | 116.5 | 6.78 (d, 10.2) | 116.5 | 6.79 (d, 10.2) | 116.6 | 6.79 (d, 10.2) | 116.6 |
| 7        | 1.71, 1.90 (both m) | 41.6 | 1.71, 1.90 (both m) | 41.8 | 1.71, 1.91 (both m) | 41.7 | 1.71, 1.91 (both m) | 41.9 |
| 8        | 2.08 (2H, m) | 23.0 | 2.08 (2H, m) | 23.2 | 2.09 (2H, m) | 23.0 | 2.09 (2H, m) | 23.3 |
| 9        | 5.06 (qt, 1.0, 1.7) | 123.1 | 5.06 (qt, 1.0, 1.7) | 123.0 | 5.06 (dt, 1.3, 7.1) | 123.1 | 5.06 (qt, 1.3, 7.1) | 123.0 |
| 10       | 1.52 (3H, s) | 17.6 | 1.54 (3H, s) | 17.7 | 1.57 (3H, s) | 17.6 | 1.57 (3H, s) | 17.7 |
| 11       | 206.2 | 206.2 | 210.7 | 210.7 |
| 2        | 3.14 (2H, d, 6.7) | 53.6 | 3.14 (2H, d, 6.7) | 53.6 | 3.89 (m) | 47.0 | 3.89 (m) | 47.0 |
| 3        | 2.27 (m) | 25.6 | 2.27 (m) | 25.5 | 1.25 (3H, d, 6.7) | 16.8 | 1.26 (3H, d, 6.7) | 16.6 |
| 4        | 1.03 (3H, d, 6.6) | 22.6 | 1.03 (3H, d, 6.6) | 22.6 | 1.46, 1.89 (both m) | 27.2 | 1.46, 1.89 (each m) | 27.2 |
| 5        | 1.03 (3H, d, 6.6) | 22.6 | 1.03 (3H, d, 6.6) | 22.6 | 0.98 (3H, t, 7.5) | 11.7 | 0.98 (3H, t, 7.5) | 11.7 |
| 2‴-CH₃   | 1.51 (3H, s) | 27.3 | 1.48 (3H, s) | 27.5 | 1.52 (3H, br s) | 27.3 | 1.47 (3H, br s) | 27.5 |
| 7-CH₃    | 14.51 (s) | 14.51 (s) | 14.44 (s) | 14.44 (s) |

*Measured by 700 MHz for ¹H NMR and 175 MHz for ¹³C NMR.
*Measured by 800 MHz for ¹H NMR and 200 MHz for ¹³C NMR.
Based on this, oxidation of 3-methyl-1-oxobutyl moiety in the 8-position. The absolute configuration of the 1′-position in 1 has been assumed to be S by comparison of the optical rotation with that of similar compounds (Win et al., 2008b). To confirm the stereochemistry, we carried out chemical correlation between 1 and kayeassamin A (8), which has been reported to be in the 1′S form by the modified Mosher’s method (Win et al., 2008a). Thus, oxidation of 8 with 2,3-dichloro-5,6-dicyano-p-benzoquinone (DDQ) gave 1. Consequently, the absolute configuration in the 1′′ position of 1 was confirmed to be S.

Mammeeasin E (2) was also obtained as an inseparable mixture (ca. 1:1 ratio, Figure S1) with a negative optical rotation ([α]$_D^{27}$ = −58.9 in CHCl$_3$). In the negative-ion ESIMS of 2, a quasimolecular ion peak was observed at m/z 453 [M − H]$^−$, and HRESIMS analysis indicated the molecular formula was C$_{27}$H$_{33}$O$_6$. The $^1$H and $^{13}$C NMR spectra (Table 2, CDCl$_3$) of 2 were similar to those of 1, except for the signals due to the 3-methyl-1-oxobutyl moiety in the 8-position [2a: δ 1.03 (6H, d, J = 6.6 Hz, H$_3$-4′′′ and H$_3$-5′′′), 2.27 (1H, m, H-3′′′)], 3.14 (2H, d, J = 6.7 Hz, H$_2$-2′′′); 2b: δ 1.03 (6H, d, J = 6.6 Hz, H$_3$-4′′′ and H$_3$-5′′′), 2.27 (1H, m, H-3′′′)], 3.14 (2H, d, J = 6.7 Hz, H$_2$-2′′′)] instead of the 1-oxobutyl moiety of 1. As shown in Figure S2, the connectivity of the quaternary carbons in 2 were elucidated on the basis of DQF-COSY and HMBC experiments. Thus, the DQF-COSY experiment on 2 indicated the presence of the following partial structures: C-1′-C-3′; C-3′′-C-4′; C-5′′-C-7′′; and C-2′′′-C-5′′′ shown in bold lines. In the HMBC experiment, long-range correlations were observed between the following proton and carbon pairs: H-3 [2a: δ 6.61 (1H, d, J = 0.9 Hz); 2b: δ 6.59 (1H, d, J = 1.0 Hz)] and C-2 (2a: δC 159.6; 2b: δC 159.6), C-4a (2a: δC 101.0; 2b: δC 101.1); the hydrogen-bonded hydroxy proton [2a: δ 14.51 (1H, s); 2b: δ 14.51 (1H, s)] and C-6 (2a: δC 105.8; 2b: δC 106.0), C-7 (2a: δC 163.0; 2b: δC 163.0), C-8 (2a: δC 104.5; 2b: δC 104.6); H-1′ [2a: δ 5.40 (1H, br t, J = ca. 8 Hz); 2b: δ 5.40 (1H, br t, J = ca. 8 Hz)] and C-3 (2a: δC 107.0; 2b: δC 107.1), C-4a; H-3′′′ [2a: δ 5.53 (1H, d, J = 10.2 Hz); 2b: δ 5.54 (1H, d, J = 10.2 Hz)] and C-6, C-2′′′ (2a: δC 83.0; 2b: δC 83.1), 2′′′-CH$_3$ (2a: δC 27.3; 2b: δC 27.5); H-4′′′ [2a: δ 6.79 (1H, d, J = 10.2 Hz); 2b: δ 6.78 (1H, d, J = 10.2 Hz)] and C-5 (2a: δC 156.0; 2b: δC 156.0), C-6; H-2′′′ [2a: δ 1.71, 1.90 (1H each, both m); 2b: δ 1.71, 1.90 (1H each, both m) and C-2′′′, 2′′′-CH$_3$; H-7′′′ [2a: δ 5.06 (1H, qt, J = 1.0, 7.1 Hz); 2b: δ 5.06 (1H, qt, J = 1.0, 7.1 Hz)] and C-9′′′ (2a: δC 25.5; 2b: δC 25.6), C-10′′′ (2a: δC 17.6; 2b: δC 17.7); H-9′′′ [2a: δ 1.64 (3H, d, J = 1.0 Hz); 2b: δ 1.67 (3H, d, J = 1.0 Hz)] and C-7′′′ (2a: δC 123.1; 2b: δC 123.0), C-8′′′ (2a: δC 132.6; 2b: δC 132.5), C-10′′; H-10′′′ [2a: δ 1.52 (3H, s); 2b: δ 1.54 (3H, s)] and C-7′′′-9′′′; and H-2′′′ and C-1′′′ (2a: δC 206.2; 2b: δC 206.2). On the other hand, the molecular formula of mammeeasin F (3) was determined to be the same as that of 2, C$_{27}$H$_{33}$O$_6$, by HRESIMS measurement. The $^1$H and $^{13}$C NMR spectroscopic properties (Table 2, CDCl$_3$) of 3, which were observed to be duplicate signals caused by its inseparable mixture (ca. 1:1 ratio, Figure S1), were quite similar to those of 2 except for the signals due to the 2-methyl-1-oxobutyl moiety in the 8-position [3a: δ 0.98 (3H, t, J = 7.5 Hz, H$_3$-5′′′), 1.26 (3H, d, J = 6.7 Hz, H$_3$-3′′′), 1.46, 1.89 (each 1H, both m, H$_2$-4′′′), 3.89 (1H, m, H$_2$-2′′′); 3b: δ 0.98 (3H, t, J = 7.5 Hz, H$_3$-5′′′), 1.26 (3H, d, J = 6.7 Hz, H$_3$-3′′′), 1.46, 1.89 (each 1H, both m, H$_2$-4′′′), 3.89 (1H, m, H$_2$-2′′′)]. Finally, 2 and 3 were derived by DDQ oxidation of surangins D (4) (Ngo et al., 2010) and C (13) (Verotta et al., 2004; Yagi et al., 2006), respectively. Based on this.
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**FIGURE 2** | Coumarin constituents (4–35) from the flowers of *M. siamensis*.
evidence, the stereostructures of 2 and 3 were determined to be as shown.

**Effects of Coumarin Constituents of the Flowers of M. siamensis on Testosterone 5α-Reductase**

To characterize the active constituents of this plant material, the inhibitory effects of 30 isolates (1–13, 17–20, 22–29, 31–35) against 5α-reductase were examined. As shown in Table 3, mammeeasins E (2, 22.6 µM), A (4, 19.0 µM), and B (5, 24.0 µM), kayeassamins E (9, 33.8 µM), F (10, 15.9 µM), and G (11, 17.7 µM), surangin C (13, 5.9 µM), and mammeeas A/AA (17, 19.5 µM), E/BB (22, 16.8 µM), and A/AA cyclo F (34, 23.6 µM), were found to inhibit testosterone 5α-reductase (Table S1).

**CONCLUSIONS**

The structures of geranylated coumarin constituents, kayeassamin I (1) and mammeeasins E (2) and F (3), newly isolated from the methanol extract of the flowers of M. siamensis, were determined. Of the isolated coumarins, mammeeasins E (2, 22.6 µM), A (4, 19.0 µM), and B (5, 24.0 µM), kayeassamins E (9, 33.8 µM), F (10, 15.9 µM), and G (11, 17.7 µM), surangin C (13, 5.9 µM), and mammeeas A/AA (17, 19.5 µM), E/BB (22, 16.8 µM), and A/AA cyclo F (34, 23.6 µM) were active 5α-reductase inhibitors. Although the intensity of the 5α-reductase inhibitory activity of these coumarins is moderate compared to a positive control having a steroid skeleton finasteride, to the best of our knowledge, there are few reports of the 5α-reductase inhibitors with non-steroidal skeletons (Dörsam and Altheim, 2009; Aggarwal et al., 2010; Chaudhary and Turner, 2010; Wu and Kapoor, 2013). Therefore, these active coumarins may be useful candidates for seed compounds of new non-steroidal 5α-reductase inhibitors. Further studies are required to elucidate the detailed structure activity relationships as well as the mode of action including the enzymatic inhibitory activity of these coumarins.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/Supplementary Material.

**AUTHOR CONTRIBUTIONS**

TM, FL, YM, HS, SS, and KN performed the experiments. TM, OM, and KN conceived and designed the experiments. SC and YP collected and identified the plant material. TM and FL wrote the paper. All authors have approved the final version of the manuscript.
**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2020.00119/full#supplementary-material

These data include HPLC chromatograms of mameesans E (2a, 2b) and F (3a, 3b) (Figure S1), 1H-1H COSY and HMBC correlations of 2 and 3 (Figure S2), 1D and 2D NMR spectra of 1–3 (Figures S3–S17), and inhibitory effects of coumarin constituents (1–35) from *M. siamensis* on testosterone 5α-reductase (Table S1).

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**ACKNOWLEDGMENTS**

The authors gratefully thank the Division of Joint Research Center, Kindai University for the NMR and MS measurements. We would like to thank Editage (www.editage.com) for English language editing.

**FUNDING**

This work was supported in part by the JSPS KAKENHI, Japan [Grant Numbers 18K06726 (TM) and 18K06739 (KN)].
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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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