Antichlamydial Sterol from the Red Sea Sponge *Callyspongia* aff. *implexa*

Authors

Usama Ramadan Abdelmohsen^{1,5}, Cheng Cheng¹, Anastasija Reimer², Vera Kozjak-Pavlovic², Amany K. Ibrahim³, Thomas Rudel², Ute Hentschel¹, RuAngelie Edrada-Ebel⁴, Safwat A. Ahmed³

Affiliations

The affiliations are listed at the end of the article

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Correspondence

Safwat A. Ahmed
Department of Pharmacognosy
Faculty of Pharmacy
Suez Canal University
Ring Road Street
Ismailia 41522
Egypt
Phone: + 20 1092 63 83 87
Fax: + 20 643 23 07 41
safwat_aa@yahoo.com

Correspondence

RuAngelie Edrada-Ebel
Strathclyde Institute of
Pharmacy and Biomedical
Sciences
University of Strathclyde
The John Arbuthnott Building
161 Cathedral Street
Glasgow G40 NR
UK
Phone: + 44 141 548 59 68
Fax: + 44 141 552 25 62
ruangelie.edrada-ebel@strath.ac.uk

Abstract

Marine sponges are rich sources of natural products exhibiting diverse biological activities. Bioactivity-guided fractionation of the Red Sea sponge *Callyspongia* aff. *implexa* led to the isolation of two new compounds, 26,27-bisnorcholest-5,16-dien-23-yn-3 β ,7 α -diol, gelliusterol E (1) and C₂₇-polyacetylene, callimplexen A (2), in addition to the known compound β -sitosterol (3). The structures of the isolated compounds were determined by 1D- and 2D-NMR techniques as well as high-

resolution tandem mass spectrometry and by comparison to the literature. The three compounds (1–3) were tested against *Chlamydia trachomatis*, an obligate intracellular gram-negative bacterium, which is the leading cause of ocular and genital infections worldwide. Only gelliusterol E (1) inhibited the formation and growth of chlamydial inclusions in a dose-dependent manner with an IC₅₀ value of 2.3 μ M.

Supporting information available online at <http://www.thieme-connet.de/products>

Introduction

The lack of physical defences in marine sponges makes them susceptible to marine predators such as fish, turtles, and invertebrates. Thus, it is frequently speculated that sponges have developed a wide range of defensive compounds to deter predators [1,2]. They also use their defensive chemicals to keep the offspring of small plants and animals (fouling organisms) from settling on to their outer surfaces [3,4]. These sessile animals are prolific producers of a huge diversity of secondary metabolites that have been discovered over the past years [3,5–8]. The genus *Callyspongia* belongs to the order Haplosclerida, family Callyspongiidae. Members of this genus have provided polyacetylenic compounds including polyacetylenic alcohols [9], sulphate [10], hydrocarbons [11], and amides with diverse activities including antifouling and cytotoxic activities [12]. Moreover, steroidal anti-inflammatory compounds have also been reported from this genus [13].

The gram-negative obligate intracellular human pathogen *Chlamydia trachomatis* is one of the main causative agents of sexually transmitted diseases and infections of the upper inner eyelid (trachoma) [14,15]. The ocular disease is caused

by *C. trachomatis* serovar A to C and, if untreated, may result in blindness; it is thus a major problem in developing countries with poor health care [16,17]. *C. trachomatis* serovars D to K are responsible for infection of the genital tract, which may lead to pelvic inflammatory disease and infertility, whereas serovars L1, L2, and L3 cause lymphogranuloma venereum, a sexually transmitted infection of the lymph nodes [18,19]. *Chlamydia* features a biphasic developmental cycle that occurs in a specialized vacuole, the so-called inclusion. Upon infection, *Chlamydia* suppresses host cell apoptosis to escape the immune response [20]. The infection is initiated by the elementary bodies (EBs), an infectious and metabolically inert chlamydial form. Within the inclusion, EBs differentiate into a replicative form, named reticulate bodies (RBs). At the end of the developmental cycle, a differentiation of the RBs back to the EBs occurs, thus enabling the new progeny to be released and to initiate a new infectious cycle. RBs are also able to convert to a persistent non-replicative state, which leads to a long-term relationship with the host cell [21]. Reinfection and persistence are the reason for prolonged therapy of chlamydial infections, in which different antibiotics are used, including tetracyclines, azithromycin, or erythromycin. An incompetent administra-

tion of antimicrobial therapy is known to induce persistence of *Chlamydia* [22]. Additionally, a vaccine is not available against human *Chlamydia*, which makes it an interesting model organism in search for novel classes of antimicrobial agents. There are few examples of natural products that were found to inhibit chlamydial infections. One is betulin derivatives, of which 32 were examined *in vitro* against *Chlamydia pneumonia*. Only betulin dioxime was highly active with an MIC value of 1 μ M against strain *C. pneumonia* CWL-029 [23]. The novel macrocyclic lactones, saccharocarbins A and B, were isolated from the actinomycete *Saccharothrix aerocolonigenes* subsp. *antibiotica* SCC 1886 and were active against *C. trachomatis* serotype H at an MIC value of 0.5 μ g/mL [24].

In the present study, we aimed to identify new antichlamydial natural products from Red Sea sponges. We herein report the isolation and structure elucidation of two new metabolites, 26,27-bisnorcholest-5,16-dien-23-yn-3 β ,7 α -diol, gelliusterol E (**1**) and C₂₇-polyacetylene, callimplexen A (**2**), along with the previously known inactive metabolite β -sitosterol (**3**), for the first time from *Callyspongia* aff. *implexa*. The metabolites were tested for inhibitory effects on the growth of *C. trachomatis*, and gelliusterol E was found to affect the formation and growth of chlamydial inclusions in a dose-dependent manner. To our knowledge, this is the first report of natural products from this sponge species.

Results and Discussion

In our continuing search for new antichlamydial natural products from marine sponges, the methanol-dichloromethane (1:1) extract of the lyophilized Red Sea sponge *C. aff. implexa* showed activity against *C. trachomatis*. A bioactivity-guided assay was performed to further isolate and characterize the antichlamydial compounds. The crude extract was then fractionated by gel filtration on Diaion HP20 eluting with increasing concentrations of methanol in water. Finally, purification of the bioactive fraction was then accomplished on normal phase HPLC using a dichloromethane (DCM) and ethyl acetate (EtOAc) mixture to afford three compounds, **1–3**.

Compound **1** was isolated as white powder with the molecular formula of C₂₅H₃₆O₂, corresponding to eight degrees of unsaturation. ¹H and ¹³C NMR data for **1** (Table 1) were almost identical to those for gelliusterol A (**4**), which was previously isolated from the marine sponge *Gellius* sp. [25]. The two main differences between the ¹³C NMR chemical shifts for **1** and **4** were observed at C-16 (δ_C 114.2) and C-17 (δ_C 147.0), indicating the presence of a trisubstituted double bond instead of methine and methylene carbons at C-17 and C-16. The absolute configuration at C-3 is assumed to be S, the same as gelliusterol A (**4**) [25]. The relative stereochemistry at other centers was assigned on the basis of a 2D NOESY experiment. The NOESY spectrum showed correlations of H-3 with H-9 and H-14 but no apparent correlations with H-7, H-8, H₃-18, and H₃-19 were observed, indicating that H-3, H-9, and H-14 were on the same side of the molecule in the α -orientation while H-7, H-8, H₃-18, and H₃-19 were in the opposite direction in the β -orientation. Similar NMR shifts of C-20 with gelliusterol A and other sterols show relative configurations at C-20 [25]. The trivial name gelliusterol E was assigned to the new sterol **1**: 26,27-bisnorcholest-5,16-dien-23-yn-3 β ,7 α -diol.

Compound **2** was isolated as a yellow oil with the predicted formula of C₂₇H₃₈ (nine degrees of unsaturation). The ¹H NMR spectrum of **2** (Table 2) showed resonances for one terminal methyl

Table 1 ¹H (400 MHz) and ¹³C (100 MHz) NMR data of compound **1** in CDCl₃.

Position	¹ H	COSY	¹³ C
1	1.11, m	2	37.0
	1.87, m		
2	1.49, m	1	31.5
	1.84, m		
3	3.58, m	2, 4	72.0
4	2.34, m	3	41.4
5			147.7
6	5.61, d	7	124.5
7	3.85, br s	6, 8	65.9
8	1.49, m	7	39.1
9	0.91, t		44.0
10			38.0
11	1.49, m		21.6
12	1.11, m		40.9
	2.00, m		
13			45.6
14	1.08, m	15	47.2
15	2.05, m	16	25.0
16	5.15, d	15	114.2
17			147.0
18	0.68, d		12.0
19	1.00, s		18.0
20	1.5, m	21, 22	35.8
21	1.01, s		19.0
22	2.33, d	20	26.2
23			78.9
24			77.7
25	1.69, m		3.3

at δ_H 0.87 and one terminal acetylene at δ_H 3.01 (s, H-1). The IR spectrum also established the presence of terminal acetylenic C–H and the C $\equiv$ C bands [26]. Two olefinic protons for a Z-double bond at C-3/C-4 ($J^{3,4}$ 10.8 Hz), together with signals for nine methylenes, accounted for the remaining protons in the molecule. Multiplets at δ_H 2.15 and 2.33 were assigned to alkyl protons next to the olefinic group. The ¹³C NMR spectrum of **2**, combined with HMQC experiment results (Table 2), showed resonances for eight acetylenic carbons [δ_C 81.7, 80.4, 80.4 (overlapped C)], two *sp*² carbons [δ_C 145.2 and 108.8], and sixteen methylene carbons. Interpretation of the ¹H-¹H COSY and HMBC spectra led us to the assembly of the C-1/C-7 unit. The quaternary acetylenic carbons were established from the HMBC experiment as correlations between H-1/C-2 (²*J*_{CH}), H-3/C-1 (³*J*_{CH}), H-4/C-2 (³*J*_{CH}), and H-1/C-2 (²*J*_{CH}), determining the location of the terminal acetylene and olefinic carbons. Further support for the C-1/C-11 unit came from comparing the chemical shifts of **2** with aikupikanyne B isolated from the Red Sea sponge *Callyspongia* sp. whose ¹³C and ¹H NMR spectra are very similar [10]. Significant fragmentation ion peaks (*m/z* 131, 145, 173, 211, 277, and 333) in the LRMS confirmed the position of the isolated triple bond and supported the structure of **2** (Fig. 1). Also, a combination of COSY and HMBC experiments and comparison with the spectra of callipentayne, which was isolated from the Japanese sponge *Callyspongia* sp. [27], enabled us to establish the structure of **2**. The name callimplexen A was assigned to the new C₂₇-polyacetylene. More functionalized C₂₇-polyacetylenes have been isolated before from a Caribbean sponge, *Petrosia* sp. [28]. The molecular ion peak of the third compound (**3**) was afforded by GC-EIMS as *m/z* 414.34 with the predicted formula of

Table 2 ^1H (400 MHz) and ^{13}C (100 MHz) NMR data of compound **2** in CDCl_3 .

Position	^1H	^{13}C
1	3.01, s	81.7
2		80.4
3	5.47, d	108.8
4	5.97, m	145.2
5	2.15, m	32.0
6	1.54, m	19.3
7	2.33, m	32.0
8		80.4
9		80.4
10	2.33, m	32.0
11	1.58, m	19.3
12	1.58, m	19.3
13	2.33, m	32.0
14		80.4
15		80.4
16	2.36, m	32.0
17	1.54, m	19.3
18	1.54, m	19.3
19	2.36, m	32.0
20		80.4
21		80.4
22	2.33, m	32.0
23	1.52, m	32.0
24–26	1.25, m	32.0
27	0.87, m	14.8

$\text{C}_{29}\text{H}_{50}\text{O}$ and identified as sitosterol from the online NIST11 database. By comparing the ^1H and ^{13}C spectra with those of β -sitosterol ($\geq 97\%$, Sigma-Aldrich), the compound was confirmed as β -sitosterol.

Compounds **1–3** were tested for their antichlamydial activity using epithelial HeLa cells with *C. trachomatis* as a model system. Only gelliusterol E had an effect on *Chlamydia* infection, while at the same time, it did not have an adverse effect on host cells (● Fig. 2a,b). Gelliusterol E inhibited the formation and growth of chlamydial inclusions in a dose-dependent manner. HeLa cells were infected in the presence of 1.6, 8.0, and 40.0 μM of gelliusterol E. The higher the concentration of the compound, the smaller the inclusion size (● Fig. 2b). However, there was no fragmentation of inclusions as had been observed with other antichlamydial agents (data not shown). At the highest concentration of 40 μM , no inclusions could be observed at all, analogous to the effect of the tetracycline (Tet) control (● Fig. 2a). The IC_{50} value, defined as the concentration at which the size of chlamydial inclusion relative to the cell surface was reduced by 50%, was $2.34 \pm 0.22 \mu\text{M}$ (● Fig. 2c).

The inhibitory effect of gelliusterol E was even greater when we addressed the ability of *C. trachomatis* to create viable progeny under the treatment with gelliusterol E (● Fig. 3a,b). An EB progeny assay was performed in order to analyze the effect of the compound on the developmental cycle of *Chlamydia*. HeLa cells were infected with *C. trachomatis* in the presence of the DMSO and Tet controls and in the presence of 1.6 μM , 8 μM , and 40 μM gelliusterol E. For the analysis of the EB primary infection, the infected cells were harvested after 24 h. A parallel set of samples was infected for 48 h, with the EB progeny being used to infect new cells without any treatment, and these cells were harvested after 24 h. A Western blot was performed with the cell lysates using an anti- β -tubulin antibody as a loading control. For determi-

Fig. 1 Structures of isolated compounds **1–3** and gelliusterol A (**4**).

nation of the chlamydial content, an antibody was applied against the chlamydial protein OmpA. As suggested by our previous results, the compound affected the primary infection in a dose-dependent manner – the higher the concentration, the less the chlamydial protein OmpA was detected. However, gelliusterol E also affects the progeny, which is less infectious. At a concentration of 40 μM , practically no viable progeny was formed. Thus, the compound not only inhibits the formation but also affects the developmental cycle of *Chlamydia*. *C. trachomatis* is known to acquire lipids necessary for its growth, including cholesterol, from the host cell [29]. Due to the structural similarity of gelliusterol E to cholesterol, we speculate that the substance could be in some way inhibiting the lipid acquisition of the chlamydial inclusion and therefore inhibiting its growth. Recent reports, however, suggest that *C. trachomatis* can successfully grow in cells defective in cholesterol biosynthesis [30], but it does not exclude the possibility that gelliusterol E is being recruited to the inclusion, similarly to cholesterol, where it could affect the stability and growth of the inclusion membrane. Future experiments will be performed to explore the exact mechanism of action.

In conclusion, two new metabolites, 26,27-bisnorcholest-5,16-dien-23-yn-3 β ,7 α -diol, gelliusterol E (**1**) and C_{27} -polyacetylene, callimplexen A (**2**), along with the previously known metabolite β -sitosterol (**3**) were isolated and identified from the Red Sea sponge *Callyspongia* aff. *implexa*. An important finding from this study is the potent antichlamydial activity of gelliusterol E against *C. trachomatis*. This finding highlights the potential of marine sponges for novel bioactivities for biomedical applications.

Fig. 2 Gelliusterol E inhibited *Chlamydia* infection in a dose-dependent manner. **a** HeLa cells were infected with *C. trachomatis* expressing GFP at a MOI 1 for 24 h in the presence of DMSO and 40 μ M gelliusterol E. The chlamydial inclusions appear in the green channel (arrow), cell nuclei were stained with DAPI (blue channel), and the actin cytoskeleton with Phalloidin647 (red channel). Scale bars are 25 μ m. **b** The graph represents mean values $\pm$ SD of the surface of inclusions relative to the cell surface from six independent repetitions of the experiment. **c** IC_{50} of gelliusterol E. (Color figure available online only.)

Fig. 3 Gelliusterol E affects the primary and progeny chlamydia infection in a dose-dependent manner. **a** Western blot resulting from the EB progeny assay showing the EB primary (Prim) and progeny (Prog) infection of *C. trachomatis* in the presence of the DMSO and tetracycline (Tet) controls and depicted gelliusterol E concentrations. **b** The signals from Western blot as described in **a** were quantified using Image J. The graph represents mean values $\pm$ SD from three independent experiments.

Materials and Methods

Biological material, collection, and identification

C. aff. implexa, Topsent 1892, was collected using Scuba at depths of 15–20 m from the Red Sea (Safaga, Egypt). The sponge material was frozen immediately and kept frozen at -20°C until processed. The sponge was identified by Dr. R. W. M. van Soest (Faculty of Science, Zoological Museum, Amsterdam). A voucher specimen was deposited in the herbarium section of the Pharmacognosy Department (Faculty of Pharmacy, Suez Canal University, Ismailia, Egypt) under registration number SAA-12 and at the Zoological Museum of the University of Amsterdam under registration No. ZMAPOR. 19758.

Extraction and isolation of compounds from *Callyspongia aff. implexa* extract

After freeze-drying, the sponge material (0.7 kg) was extracted with methanol-dichloromethane (1 : 1) (4 L $\times$ 4) at room temperature to yield 65 g of crude extract. The extract was fractionated by gel filtration on a Diaion HP20 open column (100 cm $\times$ 10 cm; the volume of the column was 7 L) using a gradient of H_2O (A) and MeOH (B) (0–100% B). Similar fractions were grouped together, then concentrated under reduced pressure, and monitored by TLC using silica gel G₆₀ F₂₅₄ plates to give three fractions, F1 (23 g), F2 (16 g) and F3 (20 g). Fraction 2 showed a potential inhibitory effect against *C. trachomatis* and was subjected to a flash silica gel column (I.D. $\times$ L 5 cm $\times$ 25 cm) with acetone and MeOH to yield fractions F4–15. Only fraction 4 (650 mg) showed a potential inhibitory effect against *C. trachomatis*. Further separation was performed on a semipreparative normal-phase HPLC column [5 μ m, 250 $\times$ 10 mm (i.d.); Luna, Phenomenex, USA] using a DCM

and EtOAc solvent mixture (70% DCM:EtOAc to 100% EtOAc over 40 min at a flow rate of 5 mL/min), giving three compounds, **1** (11 mg, Rt = 14), **2** (5 mg, Rt = 9) and **3** (38 mg, Rt = 17).

Gelliusterol E (1), purity 95%: white powder, amorphous powder; IR (film) $\nu_{\max}$ 2245, 3313 cm^{-1} ; ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz), **Table 1**; HREIMS m/z 367.2638 [$\text{M} - \text{H}$] $^-$ (calcd. for $\text{C}_{25}\text{H}_{35}\text{O}_2$, 367.2637).

Callimplexen A (2), purity 96%: yellow oil; IR (film) $\nu_{\max}$ 2230, 3298 cm^{-1} ; $\lambda_{\max}$ (log ϵ): 205, 255; ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz), **Table 2**; HREIMS m/z 361.2859 [$\text{M} - \text{H}$] $^-$ (calcd. for $\text{C}_{27}\text{H}_{37}$, 361.2895). Combustion analysis showed 89.41% carbon and 10.59% hydrogen.

Cell culture

HeLa cells were cultured in RPMI 1640 (Gibco) media supplemented with 10% (v/v) h.i. FBS (Biocrom) at 37°C under 5% CO_2 . *C. trachomatis* C2 (L2/434/BU) green fluorescent protein (GFP)-expressing strain was generated essentially as described in Wang et al. [31].

Antibodies

β -Tubulin antibody was purchased from Santa Cruz Biotechnology (Santa Cruz), and the OmpA antibody was raised in rabbits against the cytosolic domain of OmpA carrying a His-tag. Phalloidin647 was obtained from Thermo Scientific and DAPI (4',6-diamidino-2-phenylindole) from Sigma.

Bioactivity assay

The assay was performed in black bottom 96-well plates (Greiner). 1.2×10^4 HeLa cells were seeded per well. Compounds, solubilized in DMSO, were added to final concentrations of 1.6 μM , 8 μM , and 40 μM . To the same amount of DMSO and 11.25 μM tetracycline (Sigma Aldrich, purity 99%) negative and positive controls were added, respectively. After 1 h incubation, cells were infected with *C. trachomatis* at MOI 1 for 24 h at 35°C in media supplemented with 5% h.i. FBS in the presence of the compounds. The cells were then washed with PBS and fixed with 4% PFA. After washing, cell nuclei were stained with DAPI (4',6-diamidino-2-phenylindole, 0.2 ng/mL) and the actin cytoskeleton with Phalloidin647. The analysis was performed with the Operetta high content imaging system (Perkin Elmer) at six pictures per well and 20 $\times$ magnification. The ratio between HeLa cell surface and the *C. trachomatis* inclusion area was calculated by the system. 11×10^3 Cells have been analyzed, on average. The calculated percentage of inclusion area of the bioactivity assay was used to determine the IC_{50} . A logarithmic trend line was used to calculate the agent concentration at 50% inclusion surface.

Elementary body progeny assay

4×10^5 HeLa cells were seeded in two 12-well plates (plate A and B), which were treated in the same manner. Treatment of the cells with the compounds and infection with bacteria were performed as described in the bioactivity assay. Plate A (primary infection) was harvested after 24 h using Laemmli buffer. Forty-eight h after infection, to allow *C. trachomatis* to finalize the developmental cycle, host cells of plate B were lysed with glass beads. The progeny infectious particles were released, and 2 μL of the suspension was used to infect fresh HeLa monolayers in a 12-well plate (plate C, progeny infection). After 24 h of infection, plate C was harvested in the same manner as the plate A. Cell lysates were analyzed by Western blot using antibodies against β -tubulin and the chlamydial protein OmpA. The measured OmpA intensities were

normalized with the β -tubulin intensities, where the normalization factor was calculated by dividing the β -tubulin intensities with the intensity of the β -tubulin DMSO control.

Supporting information

1D and 2D NMR spectral data of **1** and **2** are available as Supporting Information.

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Conflict of Interest

▼ The authors declare no conflict of interest.

Affiliations

- ¹ Department of Botany II, Julius-von-Sachs Institute for Biological Sciences, University of Würzburg, Würzburg, Germany
- ² Department of Microbiology, Biocenter, University of Würzburg, Würzburg, Germany
- ³ Department of Pharmacognosy, Faculty of Pharmacy, Suez Canal University, Ismailia, Egypt
- ⁴ Strathclyde Institute of Pharmacy and Biomedical Sciences, University of Strathclyde, Glasgow, UK
- ⁵ Permanent address: Department of Pharmacognosy, Faculty of Pharmacy, Minia University, Minia, Egypt

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