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Chemical Constituents of *Rumex abyssinicus* Roots and Evaluation of Its Antimicrobial Activities

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Abstract: Medicinal plants are usually screened for phytochemicals that may lead to further isolation, purification, and characterization of their active principles. The roots of *Rumex abyssinicus* are used traditionally in Ethiopia for treating different diseases such as hepatitis, hemorrhoids, gonorrhea, typhus, and wounds. Thus, this study was conducted to investigate the phytochemicals and *in vitro* antimicrobial activity of the roots of *R. abyssinicus* extracts. The roots were collected, chopped, and air-dried under shade in the laboratory. Then, the dried plant material was subjected to sequential extraction using the maceration technique by soaking 1 kg of the sample in petroleum ether, chloroform, acetone, and methanol. Meanwhile, the filtrates were concentrated under reduced pressure using a rotary evaporator. Afterward, the crude extracts were dried and screened for *in vitro* antimicrobial activity tests using four bacterial and two fungal strains. Based on its activity, the acetone extract was used for column chromatographic isolation. Herein, two known anthraquinones were isolated and characterized: chrysophanol and emodin. The microbial activity of both extracts and isolated compounds were evaluated against medically important strains by the agar disc diffusion method. The concentration of the extracts and isolated compounds was 100 mg/mL tested against *S. aureus* ATCC25903, *K. pneumoniae* NCTC13368, *E. coli* ATCC 25722, *P. aeruginosa* DSMZ 1117 and *C. albicans* and *S. cerevisiae*. The zone of growth inhibition of extracts and isolated compounds were compared with standard drugs like Gentamycin and Mancozeb. The results showed that the isolated compounds (**1** and **2**) have higher bacterial growth inhibition than the extracts. However, compound **2** was not active against *S. cerevisiae* strain.

Keywords: *Rumex abyssinicus*; chrysophanol; emodin

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INTRODUCTION

Medicinal plants (MPs) are the abundant bioresources of medical substances of traditional systems of medicine, modern medicines, nutraceuticals, food supplements, folk medicines, pharmaceutical intermediates, and chemical entities for synthetic drugs (1). Since time immemorial, different parts of medicinal herbs have been used to cure specific ailments (2). They are recognized for their ability to produce a wealth

of secondary metabolites, and mankind has used many species for centuries to treat a variety of diseases (1). Many of the natural products (NPs) have been shown to present interesting biological and pharmacological activities and serve as the starting point in the development of modern medicine (3).

A number of modern drugs have been derived from natural sources, many based on their traditional medicinal claims. The plant products used in the

treatment of diseases could be obtained from barks, leaves, flowers, roots, fruits, or seed parts of the plants (4). The use of biological resources for various therapies has been known in many different parts of the world, especially in remote regions where traditional medicines (TMs) provide an alternative to the modern health care system (5). Many people in developing countries still use TM due to its low cost and availability (6). Over 80% of people in developing countries rely on TM for primary health care needs, of which a major proportion corresponds to plant extracts (6,7). The phytochemicals isolated from these plants have shown *in vitro* and *in vivo* biological activities (8).

Ethiopia is the home of many plant species that are commonly used in many disease treatments by local healers (9). Some of the factors attributed to these facts are accessibility, affordability as compared to modern drugs, socio-cultural background, and their effectiveness against several health problems (1). For example, *R. abyssinicus* is a widely spread medicinal plant in the highlands of tropical Africa and is a common weed of cultivated lands or disturbed grounds ranging from North Africa to Ethiopia at altitudes between 1200 and 3300 m (10). It is a perennial herb, which grows up to 3 m tall, with a thick, fleshy rhizome. The plant is endemic to Ethiopia and it is referred to as *mekmeko* in the Amharic language (11,12).

The genus *Rumex* belongs to the family *Polygonaceae*, which consists of about 200 species. Most of the species of *Polygonaceae* contain a large number of bioactive compounds such as anthraquinones, flavonoids, terpenes, naphthalenes, stilbenoids, and steroids, flavonoid glycosides, leucoanthocyanidins, and phenolic acids (11). Traditionally, people use different parts of *Rumex* species to treat several health problems such as infections, diarrhea, constipation, mild diabetes, oedema, jaundice, skin, liver, and gallbladder disorders and inflammation, and as an antihypertensive, diuretic and analgesic preparation (9,13). The decoction of the leaves or root powder of *R. abyssinicus* is taken as a vermifuge. It is also used to treat malaria, gonorrhea, poisoning, hepatitis, constipation, sciatic neuralgia, hypertension, migraine, rheumatism, breast cancer, stomach distention, earache, hepatic diseases, hemorrhoids, typhus, rabies, and wound (14). The roots of *R. abyssinicus* are reported to possess antibacterial activity against *Streptococcus pyogenes* (15). Its antifungal activities were also tested against several fungal pathogens (*Aspergillus fumigatus*, *Aspergillus flavus*, *Aspergillus versicolor*, *Aspergillus niger*, *Blastoschizomyces capitatus*, *Fusarium oxysporum*, *Fusarium moniliforme*, *Fusarium semitectum*, *Pythium sp.*, *Rhizopus sp.*,

Sporotrichum sp., and *Thermomyces ssp.*) and results showed its promising activity (16).

For this reason, medicinal plants are considered a chemical factory as it contains a multitude of chemical compounds which are suggested to be natural bioactive like alkaloids, glycosides, saponins, resins, oleoresins, sesquiterpene, lactones, anthraquinones, and essential oils and fixed oils (17). The emergence of antibiotic-resistant microbial strains (including bacterial and protozoan parasites) and the increasing failure of available chemotherapeutics (18) urge the search for microbiologically active MPs. Hence, the bioactive phytochemical constituents present in the plant play a significant role in the development of medicine and drug discovery. Therefore, herein it was aimed to carry out phytochemicals isolation and evaluation of their *in vitro* antimicrobial activities of extracts from the roots of *R. abyssinicus*.

EXPERIMENTAL SECTION

Chemicals

Chemicals used for this study were petroleum ether, chloroform, acetone, ethyl acetate, silica gel 60-120 mm mesh size. Dimethyl sulfoxide (DMSO), Mueller Hinton agar, and nutrient broth were used for the antimicrobial test. All chemicals and reagents used were of analytical grade.

Apparatus and Instruments

Apparatuses such as rotary evaporator (Heidolph, Germany, laboratory 4000), round bottom flasks, measuring cylinder, filter paper (cotton swab), weighing balance (model NWT100001X) oven (N50C GENLAB WIDNES, England) were used. Analytical Thin Layer Chromatography (TLC) was performed on precoated silica gel 60 F₂₅₄ plates. Incubator (Gene lab incubator) and Hood (CLB-201-04, vertical laminar cabinet) were used for the antimicrobial activity studies. Glass column chromatography (500 mm, B-34/35) and UV chamber (254 and 365 nm) were used for isolation of compounds. The NMR spectra were recorded on an Avance 600 MHz spectrometer (Bruker, Billerica, MA, USA) using the DMSO solvent.

Sample Collection and Preparation

The roots of *R. abyssinicus* were collected from Gomma District, Jimma Zone, Oromiya Regional State, Ethiopia, in July 2019. Then, it was roughly washed with tap water, chopped into smaller pieces, and air-dried under a shade at room temperature in the laboratory. The air-dried sample was homogenized into a suitable size to increase the rate of solvent penetration into the cells by using mechanical grinding. Finally, it was subjected to organic solvent extraction.

Extraction and Isolation

The air-dried root sample of *R. abyssinicus* (1 kg) was sequentially extracted by soaking in petroleum ether, chloroform, acetone, and methanol (2 x 2.5 L) each using a maceration technique at room temperature and was kept for 48 h. The extracts were filtered with filter paper, plugged in cotton, and concentrated using a rotary evaporator under reduced pressure; the yellowish crude extracts were kept in a desiccator until dryness for further experimental purposes. Then, the crude products weighing 5.45 g (0.55%), 8.50 g (0.85%), 14.50 (1.45%), and 14 g (1.40%), respectively were obtained. Meanwhile, the four extracts were applied for *in vitro* antimicrobial activity test and TLC analysis to continue the isolation of pure compounds, with the product having promising bioactivity against the selected bacterial and fungal strains.

Isolation of pure compounds was continued using the crude products demonstrating promising activity against the strains and good TLC profile. Among the four crude extracts, acetone extract has shown the highest zone of growth inhibition against bacterial and fungal strains. Thus, based on its bioactivity and TLC profile, it was subjected

to column chromatographic separation. Petroleum ether/ethyl acetate combination showed better separation of the compounds on TLC tests. Initially, the column chromatography was packed with silica gel (60-120 mesh size) just by activating in an oven at 100 °C for 2 h. Then, the fine powder of the sample acetone extract (14.5 g) was loaded to the column chromatography.

The elution was started with petroleum ether and increased the polarity using 1 up to 100% ethyl acetate. Meanwhile, a total of 148 fractions were collected in a 20 mL small beaker for each. The first 30 fractions were merged after TLC analysis. Using TLC and R_f value results, fractions with similar R_f values were combined. Fractions 52-65 were combined and washed repeatedly with n-hexane to afford compound **1** (0.40 g). The elution was continued by increasing the polarity of the solvent. The second pure compound **2** was obtained from fractions 66-81. It showed one spot and some impurities which were later purified by continuing washing it with n-hexane. Then, a crystalline orange color product of compound-**2** (0.38 g) was obtained. The isolated compounds were characterized by using $^1\text{H-NMR}$, $^{13}\text{C-NMR}$, and 2D NMR spectroscopy techniques.

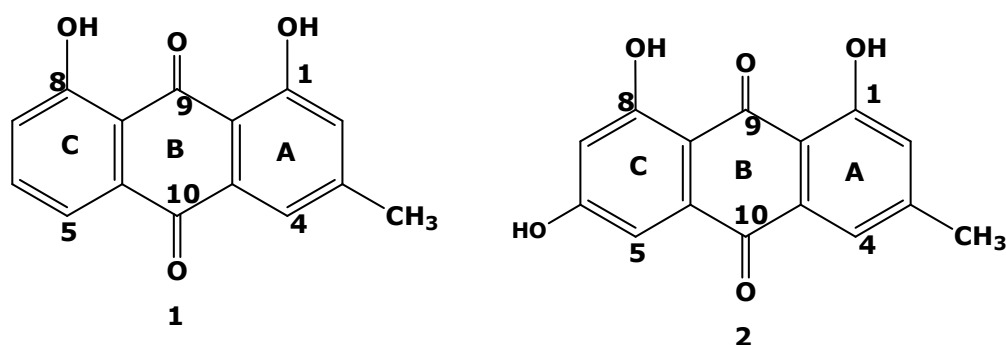


Figure 1. The proposed structure of compounds **1** and **2**.

Antimicrobial Activity Test

Test strains and preparation of test samples:

Microorganisms used for evaluation of *in vitro* antimicrobial activities were *S. aureus* ATCC25903, *K. pneumoniae* NCTC13368, *E. coli* ATCC 25722, *P. aeruginosa* DSMZ 1117, *C. albicans*, and *S. cerevisiae*. These strains were obtained from Jimma University Microbiology Research Laboratory, Biology Department. The *in vitro* antibacterial and antifungal activities of the crude extracts and isolated compounds were determined against four bacteria and two fungal strains mentioned above using the agar disc diffusion method and was carried out following a standard procedure as used earlier by Abdissa *et al.* (19). Then, bacterial and fungal stock cultures were incubated for 24 h at 37 °C on nutrient agar and potato dextrose agar (PDA) media following refrigeration storage at 4 °C. The bacterial strains were grown in Mueller-Hinton agar (MHA) plates at

37 °C (the bacteria were grown in the nutrient broth at 37 °C and maintained on nutrient agar slants at 4 °C), while the fungal strains were grown in PDA media at 28 °C. The stock cultures were maintained at 4 °C. The bacterial and fungal stock cultures were incubated for 24 h at 37 °C on nutrient agar and PDA medium.

Active cultures for this experiment were prepared by transferring a loop full of bacterial cells from the stock cultures to test tubes of Mueller-Hinton broth (MHB) that incubated without agitation for 24 h at 37 °C. A cell suspension of each organism was freshly prepared by transferring isolated colonies selected from a 24 h agar plate into a broth and the cell density was adjusted to turbidity equivalence of 0.5 McFarland turbidity standards (1×10^8 CFU/mL) in sterile saline solution. The bacterial and fungal cultures were then streaked onto MHA plate with a sterile cotton swab to obtain

a uniform thick lawn of growth. Meanwhile, the sterile paper discs (6 mm diameter, Whatman No.3) were separately soaked in pre-prepared crude extracts and pure compounds, whose stock solutions were prepared at a concentration of 100 mg/mL by dissolving 0.1 g (100 mg) of the crude extracts and pure compounds were separately dissolved in DMSO before aseptically placed on an already inoculated MHA plate. It was then allowed to diffuse for 5 min at ambient temperature and then incubated at 37 °C for 24 h. Gentamycin and Mancozeb (10 µg) were used as a standard positive control against bacterial and fungal strains, respectively, DMSO was used as a negative control. Finally, the zone of growth inhibition was reported by measuring diameter using a transparent ruler after 24 h of incubation.

RESULTS AND DISCUSSION

Characterization and Antimicrobial activities

Compound-**1** was isolated as a yellowish crystalline solid obtained from the fractions 52-65. Its R_f value was 0.65 using 70% petroleum ether in ethyl acetate as an eluent. The ^1H -NMR spectrum is given in supporting information (**SI1**), which showed that the presence of five aromatic protons at δ_{H} 7.22, 7.39, 7.55, 7.71, and 7.81 ppm (Table 1). Besides, hydroxyl proton (-OH) groups were observed at δ_{H} 11.93. And ^1H -NMR spectroscopic data of compound-**1** showed a singlet proton signal at δ_{H} 2.44 ppm representing three protons attached to C-3 of the aromatic ring A. The two broad singlet peaks observed at δ_{H} 7.22, 1H and 7.55, 1H represent the protons attached to C-2 and C-4 of the aromatic ring, respectively. The aromatic protons that are attached to C-5 and C-7 are a doublet of a doublet at δ_{H} 7.72, *dd*, $J=12$,

1H, and 7.39, *dd*, $J=6$, 1H, respectively. The *pseudo*-triplet proton was observed at δ_{H} 7.82, *t*, $J=6,12$ (m,1H). These three mutually coupled aromatic protons at C-5, C-6, and C-7 are in the ABX spin system. Moreover, there are two chelated protons at the *peri*-position (C-1 and C-8) of the aromatic ring, a highly deshielded proton at δ_{H} 11.93 ppm and downfield shifted carbonyl signals at δ_{C} 192.0 and 181.0 ppm (Table 1). Thus, the presence of a 1,8-dihydroxyanthraquinone moiety clearly indicates that as it is the chrysophanol (19). And its molecular formula is $\text{C}_{15}\text{H}_{10}\text{O}_4$ (12).

Its ^{13}C -NMR spectrum (**SI2**) showed 15 signals that represented 15 carbons including one methyl group at δ 22.0, five aromatic carbons at δ_{C} : C-2 (124.6), C-4 (120.0), C-5 (119.8), C-6 (137.8), and C-7 (124.9) as evidenced from HSQC data (**SI3**). Besides, the spectra showed two-hydroxyl carbons at δ_{C} 161.8 and 162.0, which represent carbons at *peri*-position and two aromatic carbonyl groups were also observed at δ_{C} C-9 (192.0) and C-10 (181.0). Therefore, based on the combined spectroscopic data (COSY (**SI4**), HSQC, HMBC (**SI5**), and ^1H - ^{13}C), the proposed structure was suggested chrysophanol (Compound **1**, 0.40 g). Based on this research finding and literature review reports (19,20), Compound-**1** was identified as 1,8-dihydroxy-3-methyl-9,10-anthraquinone (chrysophanol) (Table 2, Figure 1). It was reported from the roots of *R. abyssinicus*, where pure compounds were isolated using n-hexane, ethyl acetate, and methanol (20). Other studies reports also indicated that chrysophanol is present in *Polygonaceae*, *Rhamnaceae*, *Fabaceae*, *Liliaceae*, *Asphodelaceae*, *Buphorbiaceae*, *Meliaceae*, *Podocarpaceae*, *Picramniaceae*, and *Hemerocallidaceae* plant families (21–23).

Table 1. ^1H (600 MHz) and ^{13}C (150 MHz) NMR Spectral Data of Compounds **1** and **2** in DMSO- d_6 .

Compound- 1			Compound- 2	
position	$\delta^1\text{H}$ (m, J in Hz)	$\delta^{13}\text{C}$	$\delta^1\text{H}$ (m, J in Hz)	$\delta^{13}\text{C}$
1	11.93, s	161.8	11.96, s	161.9
2	7.22, s	124.6	7.11, <i>b</i> , s	124.5
3	-	149.6	-	149.7
4	7.55, s	120.0	7.42, s	121.0
5	7.72 (<i>dd</i> , $J=12$)	119.8	7.06, s	109.2
6	7.82 (<i>t</i> , $J=6,12$)	137.8	-	164.9
7	7.39 (<i>dd</i> , $J=6$)	124.9	6.55, s	108.3
8	-	162.0.	12.04, s	166.0
9	-	192.0	-	190.1
10	-	181.0	-	181.7
11	-	133.8	-	135.5
12	-	116.3	-	109.3
13	-	114.2	-	113.7
14	-	133.5	-	133.2
C-Me	2.44, s	22.1	H-Me 2.50, s	21.9

Compound-**2** (0.38 g) was isolated as an orange crystalline solid with R_f value 0.58 using 70% petroleum ether in ethyl acetate. The ^1H -NMR spectrum of the compound (**SI6**) shows the presence of four aromatic protons at δ_{H} 7.42, 7.11, 7.06, and 6.55 ppm. It has also revealed the presence of one set of methyl protons at δ_{H} 2.50 and two-hydroxyl protons at δ_{H} 11.96 and 12.04 ppm.

Its ^{13}C -NMR spectrum data shows (**SI7**) the presence of 15 carbon signals (Table 1). Among these carbons, there are four CH aromatic ring protons at δ_{C} : C-2 (124.5), C-4 (121.0), C-5 (109.2) and C-7 (108.3) ppm, two-carbonyl carbons that attached to benzene ring at δ_{C} C-9 (190.1) and C-10 (181.7) ppm, three oxygenated carbons of the aromatic ring at δ_{C} : C-1 (161.9), C-6 (164.9), C-8 (166.0) and five quaternary carbons indicated at C-3 (149.7), C-11 (135.5), C-12 (109.3), C-13 (113.7) and C-14 (133.2) ppm. The ^{13}C NMR spectrum also showed the presence of one methyl carbon at δ_{C} of 21.9 and one methyl-substituted carbon observed at δ_{C} 149.7(C-3). Furthermore, COSY (**SI8**), HMBC (**SI9**), and HSQC (**SI10**) spectrum correlation data have confirmed that the proposed compound structure was named as 3-methyl-1,6,8-trihydroxyanthraquinone (Emodin)(23–25) (Table 2, Figure 1). Emodin is found in *Rhamnus* and *Rumex* of higher plant species, and it is a metabolite of some *Penicillium* species (12,24,26).

From Table 2 above (compound **1**), the COSY correlation spectrum data show that there are three pairs of couplings between partner protons of 3J value observed at 7.72 (H-5) with H-6 (7.82), and H-6 (7.82) with H-7 (7.39) including the weak *meta*-coupled protons of 4J value of H-5 coupling with H-7. Similarly, a triplet proton for H-6, and a doublet of doublet for its partners. The other important heteronuclear multiple bonds correlation spectrum is HSQC, which correlates the carbon-proton of a single bond (1J). For instance, the correlation of H-2 (7.22) with C-2 (124.6), H-4 (7.55) with C-4 (120.0), H-5 (7.72) with C-5

(119.8), and H-6 (7.82) with C-6 (137.8). On the other hand, the HMBC spectrum shows the correlation of protons with carbon in two to four bond distances. For example, it is observed that at proton δ_{H} : H-2 (7.22) correlated with C-11 (133.8), C-4 (120.0), and the correlation with C-1 (161.8). The correlation at δ_{H} 4 (7.55) showed a long-range correlation with C-10 (181.0), C-2 (124.6), and C-13 (114.2). Based on the combined spectroscopic data, the structure of compound **1** was proposed as Chrysophanol in Figure 1.

From Table 2 above (compound **2**), HMBC spectrum showed the correlations of protons H-2 (7.11) with C-1 (161.9), C-4 (121.0) C-11 (135.5) and C-Me (21.9) and H-4 (7.42) to C-1 (161.9), C-4 (121.0) C-13 (113.7) C-9 (190.1) and C-Me (21.9). These correlations give us the information of the partial aromatic structure of compound-**2** indicated as ring A. Correspondingly, the correlations of protons H-5 (7.06 ppm) to C-6 (164.9), C-9 (190.1), C-12 (109.3) and H-7 (6.55) to C-5 (109.2), C-8 (166.0) give us the presence of another partial aromatic cyclic structure of ring C.

The correlation of H-7 with C-8 and C-6 indicates the presence of two alcohols in this *meta* position of the aromatic cycle. The correlations of H-5 to the carbonyl signal C-10 and of H-2 (7.11 ppm) to carbonyl signal C-9 confirm the presence of anthraquinone. Moreover, the correlation of H-5 to C-6 (an oxygenated quaternary carbon) indicates the presence of an alcohol group (Figure 2). And the COSY correlation spectrum (^1H - ^1H) helps to indicate the partner protons coupling each other like the two *meta*- coupling protons H-2 with H-4 (ring A) and H-5 with H-7 (ring C). Equally, the single bond coupling of the HSQC correlation spectrum (^1H - ^{13}C) plays a vital role in suggesting the structure of the compound. The HSQC correlation of the aromatic ring A of H-2 to C-2 and H-4 to C-4 equivalently for ring B, H-5 correlated to C-5 and H-7 to C-7, which helps to propose the skeleton of the compound. Based on the combined NMR spectrum, compound **2** was proposed as Emodin.

Antibacterial and Antifungal Activity Evaluation

The *in vitro* antibacterial and antifungal activities of crude extracts and isolated compounds (**1** and **2**) were carried out using four human pathogenic bacterial strains: *P. aeruginosa*, *K. pneumoniae*, *S. aureus*, and *E. coli* and two fungal strains: *C. albicans* and *S. cerevisiae* by using agar disc diffusion method. The bioassay test was conducted using the petroleum ether, chloroform, acetone, and methanol extracts of *R. abyssinicus* roots. Its crude extracts have demonstrated a good yield of the antimicrobial zone of growth inhibition (**SI11**), but petroleum ether extract is not active against *S. aureus* and *K. pneumoniae*. The previous research report finding shows that the n-hexane extract of leaves, shoots, and roots of *Rumex dentatus* did not show any activity against several bacterial strains tested and the methanol extract has no prominent inhibition against fungal strains (27). Therefore, based on its promising antimicrobial activity in demonstrating a better zone of growth inhibition, good TLC profile, and a high percentage yield, the acetone extract was selected for the isolation of pure compounds. It has demonstrated zone of growth inhibition against *S. aureus* 21.0 ± 1.4 , *K. pneumoniae* 15.5 ± 0.4 , *P. aeruginosa* 22.5 ± 0.7 , and *E. coli* 18.0 ± 0.4 mm and the positive standard (Gentamicin) has 34.5 ± 0.7 , 30.0 ± 0.7 , 30.0 ± 0.1 , and 32.0 ± 0.1 mm, respectively (Table 3). The previous report also indicated that *R. abyssinicus* is used for wound healing purposes in Ethiopia (12).

The pure compound-**1** (Chrysophanol) and compound-**2** (Emodin) have also exhibited a diameter of better zone of growth inhibition against *S. aureus* (23.0 ± 0.1 mm) and *K. pneumoniae* (22.5 ± 0.4 mm) than *P. aeruginosa* (18.0 ± 0.2 mm) and *E. coli* (18.0 ± 0.4) in mm and the positive standard (Gentamicin) has 26.0 ± 0.3 mm zone of growth inhibition (**SI11**). Compound-**2** has also shown a better zone of growth inhibition against *K. pneumoniae* (20.0 ± 0.7) and *S. aureus* (20.0 ± 0.2) than *P. aeruginosa* (19.0 ± 0.3) and *E. coli* (17.0 ± 0.7) in mm. The results indicated that compounds **1** and **2** have shown better

antibacterial activity against *K. pneumoniae* and *S. aureus* strains and their activity was close to the positive standard (Gentamicin). These anthraquinones themselves have little therapeutic activity and need to be in the form of water-soluble glycosides to exert their action (26). Similarly, the antifungal activity of compounds **1** and **2** (**SI12**), was active over *C. albicans* with a zone of inhibition 20.0 ± 0.4 and 22.0 ± 0.7 mm, respectively which is almost equal to the zone of inhibition exhibited by the positive standard (Mancozeb). Conversely, both compounds **1** and **2** were not active on the *S. cerevisiae* fungal strains (Table 3). Thus far, it was reported that chrysophanol (**1**) and emodin (**2**) are antimicrobial and wound healing active constituents isolated from the rhizomes of *R. abyssinicus* and other plant species (12,28).

CONCLUSION

The main objective of this study was to carry out a phytochemical constituent investigation and *in vitro* antimicrobial activity screening of the root extracts of *R. abyssinicus*. Based on its promising bioactivity and good TLC profile, the acetone crude extract was subjected to column chromatographic isolation packed with silica gel using petroleum ether. As a result, from the root extracts of *R. abyssinicus*, two compounds, Chrysophanol (Compound **1**) and Emodin (Compound **2**) anthraquinones were isolated and evaluated against four bacterial and two fungal strains. It was screened against *S. aureus*, *P. aeruginosa*, *E. coli*, *K. pneumoniae*, *C. albicans*, and *S. cerevisiae* using Gentamycin and Mancozeb as positive standards, and DMSO as negative standard. The antibacterial activity of isolated compound **1** showed higher activity against *S. aureus* 22.5 ± 0.4 mm, *P. aeruginosa* 18.0 ± 0.2 mm, *E. coli* 18.0 ± 0.4 mm, *K. pneumoniae* 23.0 ± 0.1 mm than **2**; while antifungal activity **2** is higher than *C. albicans* 22 ± 0.7 mm than **1** for *C. albicans* 20 ± 0.4 mm. On the contrary, compound **2** was not active against the *S. cerevisiae* strain. And the structures of these compounds were elucidated using 1D and 2D-NMR spectroscopic techniques.

Table 2. COSY, HSQC, and HMBC Correlation Data of Compounds **1** and **2** in DM

	Position δC	Compound 1 δH	HSQC (1J)	COSY ($^3J, ^4J$)	HMBC (3J)
2	124.6	H-2 (7.22)	C-C-2 \longleftrightarrow H-2	-	H-2 \rightarrow C-4, C-
4	120.0	H-4 (7.55)	C-4 \longleftrightarrow H-4	-	H-4 \rightarrow C-2, C-10
5	119.8	H-5 (7.72)	C-5 \longleftrightarrow H-5	H-5 \longleftrightarrow H-6	H-5 \rightarrow C-10, C-7
6	137.8	H-6 (7.82)	C-6 \longleftrightarrow H-6	H-5 \longleftrightarrow H-7	H-6 \rightarrow C14, C-8
7	124.9	H-7 (7.39)	C-7 \longleftrightarrow H-7	H-6 \longleftrightarrow H-7	H-7 \rightarrow C-5, C-13
Me	22.1	H-Me (2.44)	-	-	H-Me \rightarrow C-2, C-4
Compound-2					
2	124.5	H-2 (7.11)	H-2 \longleftrightarrow C-2	-	H-2 \rightarrow C-4, C-13
4	121.0	H-4 (7.42)	H-4 \longleftrightarrow C-4	-	H-4 \rightarrow C-2, C-10,
5	109.2	H-5 (7.06)	H-5 \longleftrightarrow C-5	-	H-5 \rightarrow C-7, C-10,
7	108.3	H-7 (6.55)	H-7 \longleftrightarrow C-7	-	H-7 \rightarrow C-5, C-12
8	166.0	H-8 (12.04)	OH-8 \longleftrightarrow C-8	-	-
Me	21.9	H-Me (2.50)	H-Me \longleftrightarrow C-Me	-	H-Me \rightarrow C-2, C-4

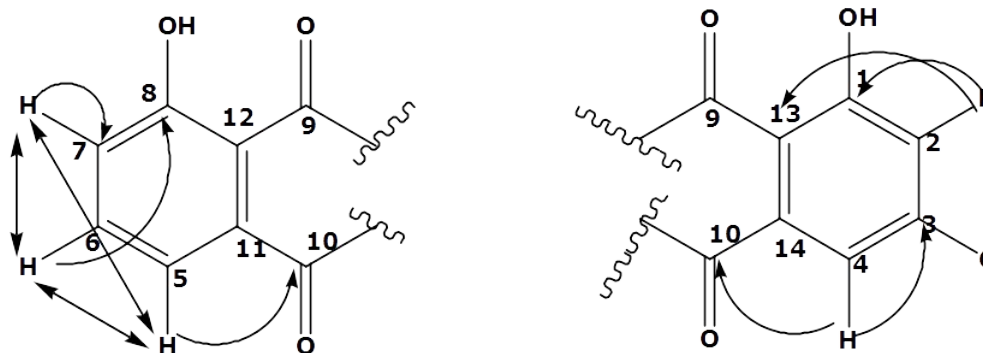

Figure 2. COSY, HSQC, and HMBC correlations of the aromatic cycle A, B, and

Table 3. *In-vitro* Antimicrobial Activities of Extracts and Isolated Compounds from Roots of *R. a*

Bacterial strain	The diameter of zone of inhibition in mm (M±SD)				Gentamycin	DMSO
	PE	CE	AE	ME		
<i>S. aureus</i>	9.5±0.7	17.5±1.4	21.0±1.4	18.0±0.1	34.5±0.7	NI
<i>P. aeruginosa</i>	NA	12.5±0.7	15.5±0.4	13.0±0.1	30.5±0.7	NI
<i>E. coli</i>	10.5±0.7	14.0±0.2	18.0±0.4	13.5±0.7	32.0±0.1	NI
<i>K. pneumoniae</i>	NA	11.5±0.7	22.5±0.7	15.0±0.2	30.0±0.1	NI
Gentamycin	-	-	-	-	-	-
DMSO	-	-	-	-	-	-
Fungal strain						Mancozeb
<i>C. albicans</i>	9.0±1.4	12.0±0.1	17.5±0.7	13.0±3	20.5±.07	NI
<i>S. cerevisiae</i>	7.5±0.7	10.0±0.7	15.0±0.7	15.0±1.4	24.5±0.7	NI
Mancozeb	-	-	-	-	-	-
DMSO	-	-	-	-	-	-

Key :

PE= petroleum ether extract, CE=chloroform extract, AE=acetone extract, ME=methanol extract, C-1 Compound-1, C Not active, DMSO = Dimethyl sulfoxide, M±SD = Mean ± Standard Deviation

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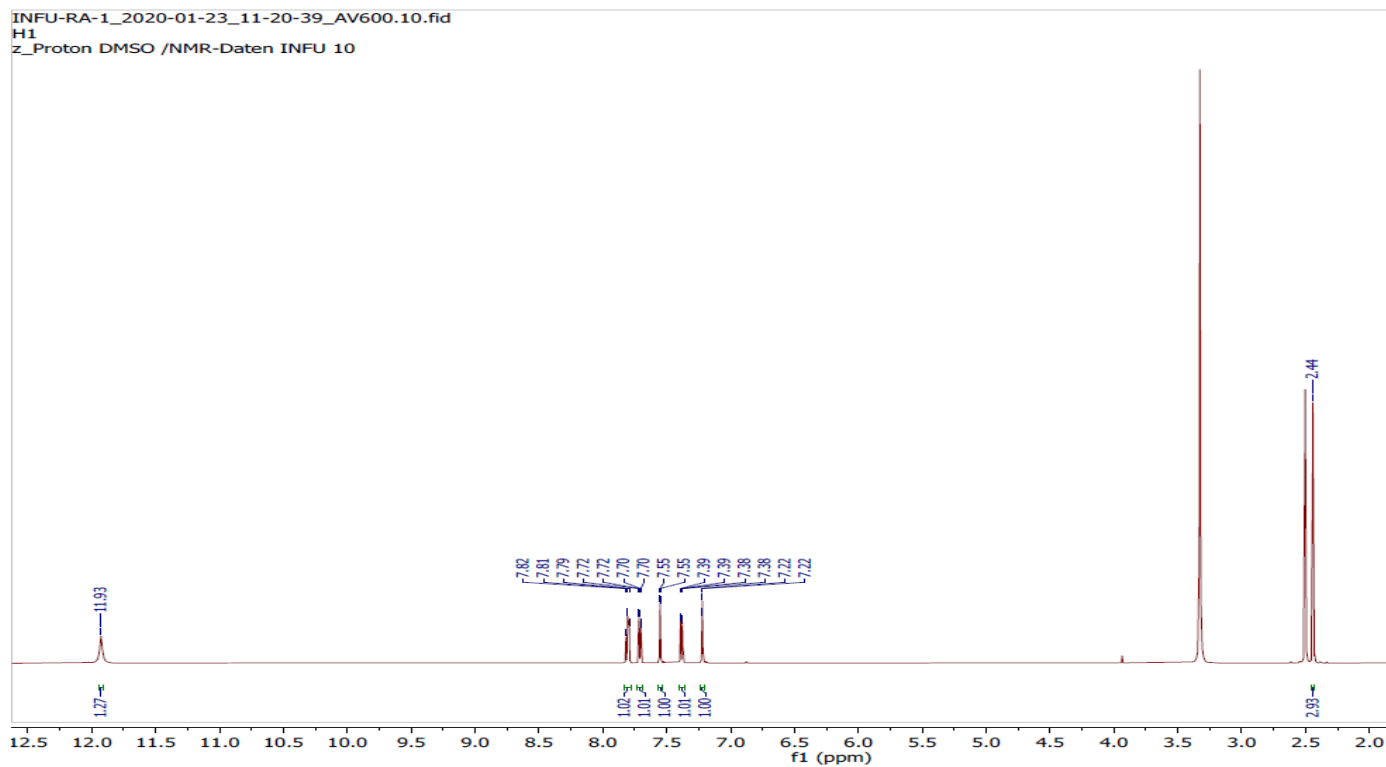
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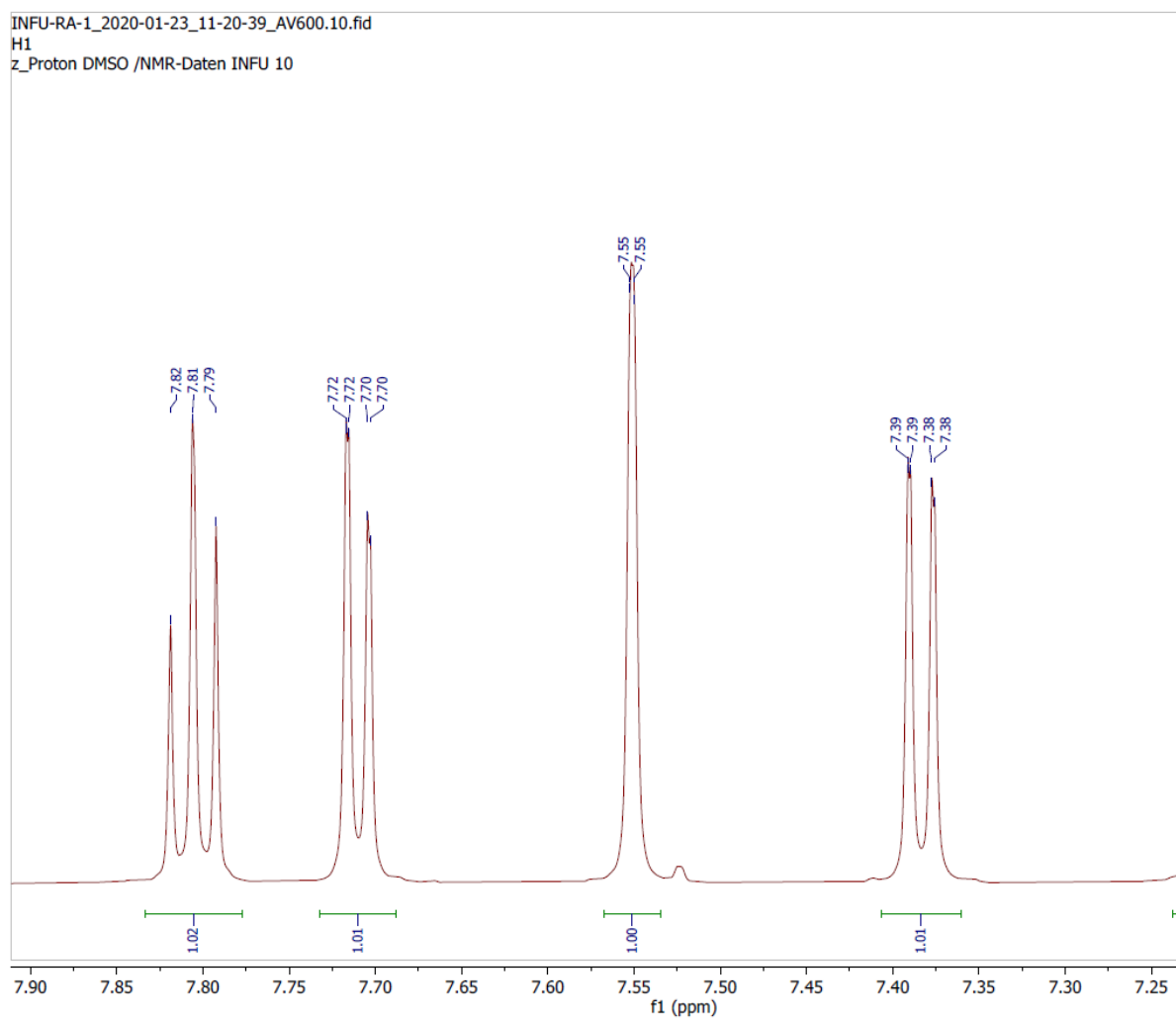
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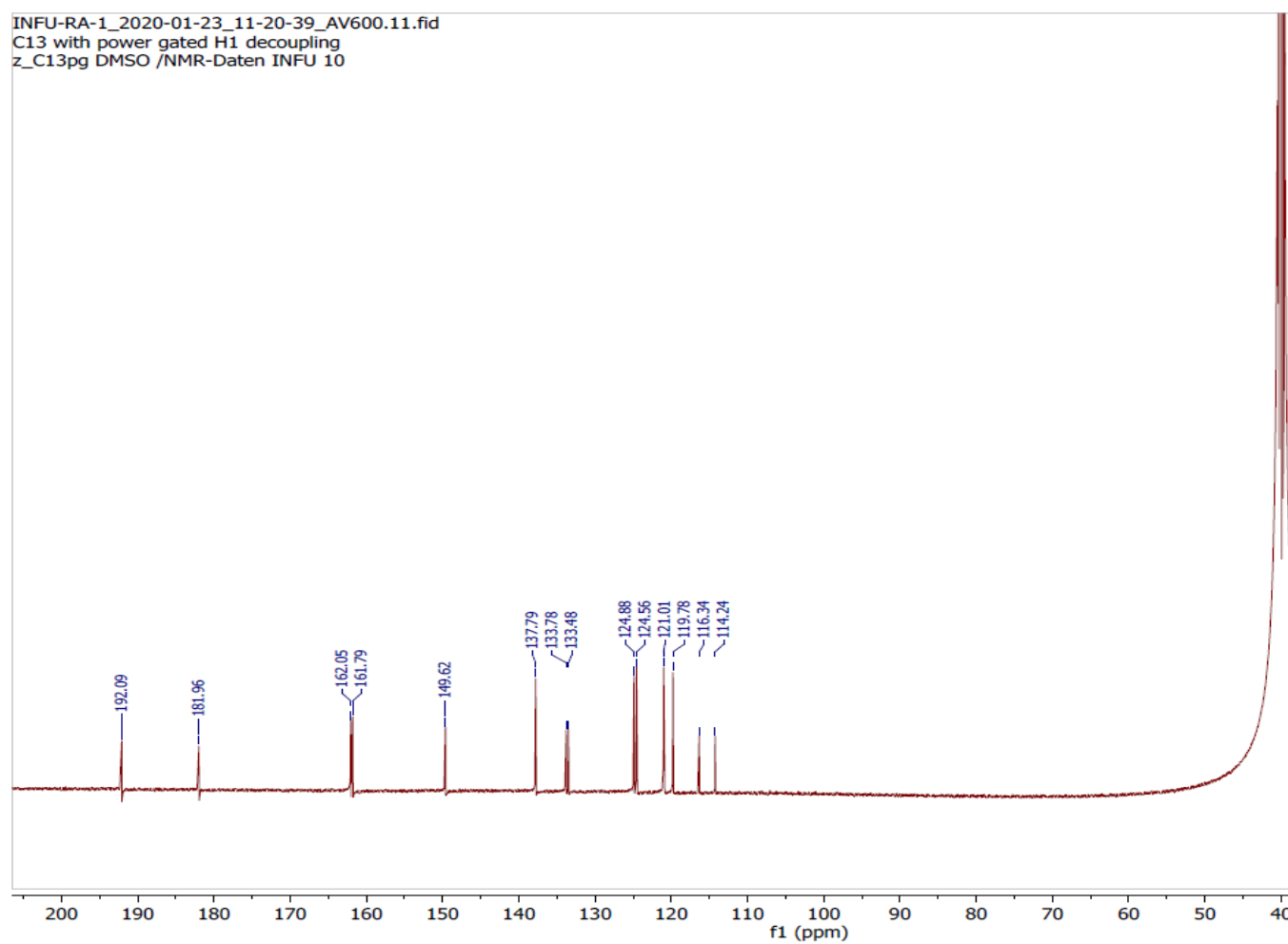
Supporting information
"Chemical Constituents of *Rumex abyssinicus* Roots and Evaluation of Its
Muaz Shifa, Dele Abdissa, Tsegaye Girma Asere*



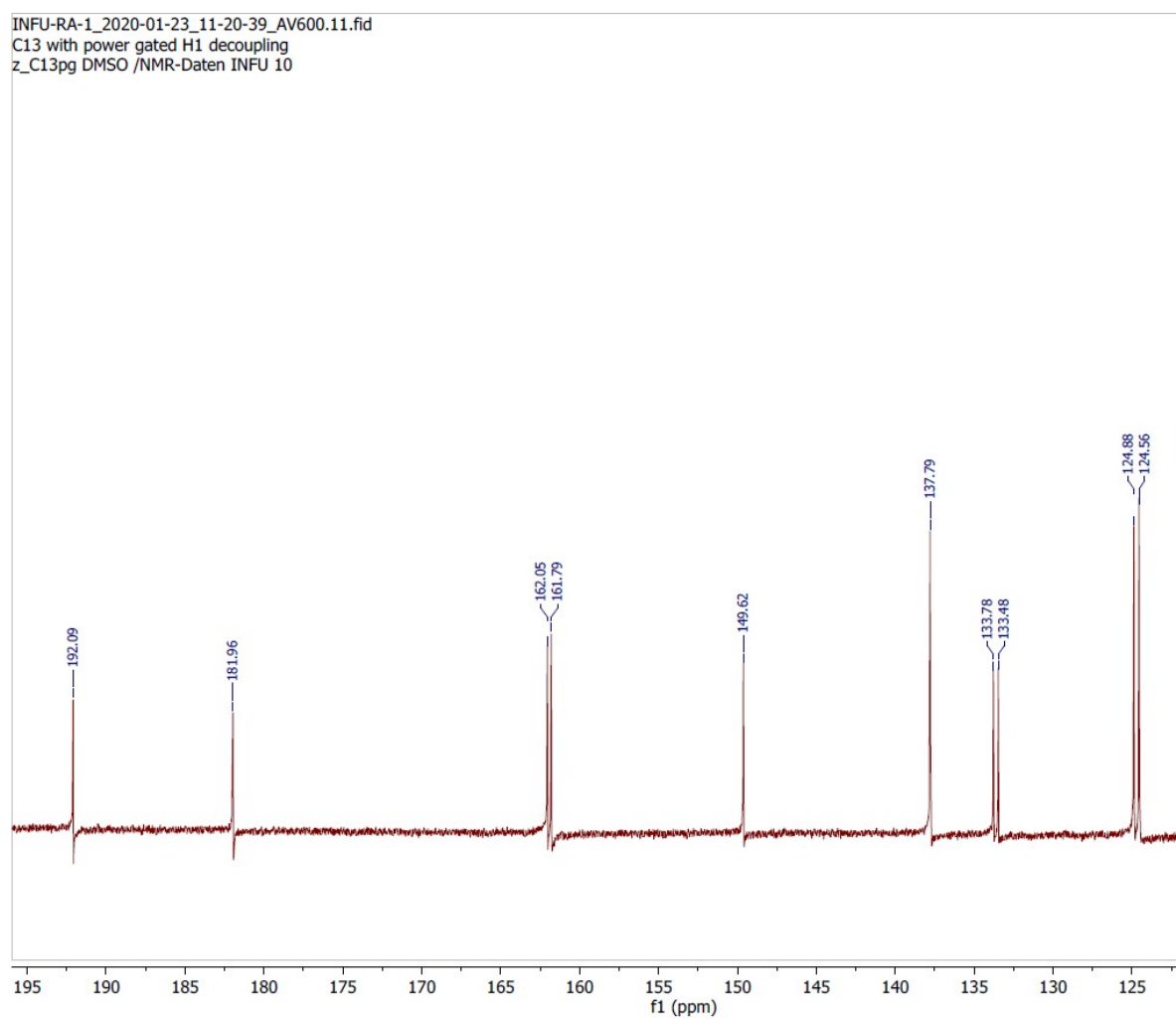
SI1a: ¹H-NMR spectral data of compound-**1**.



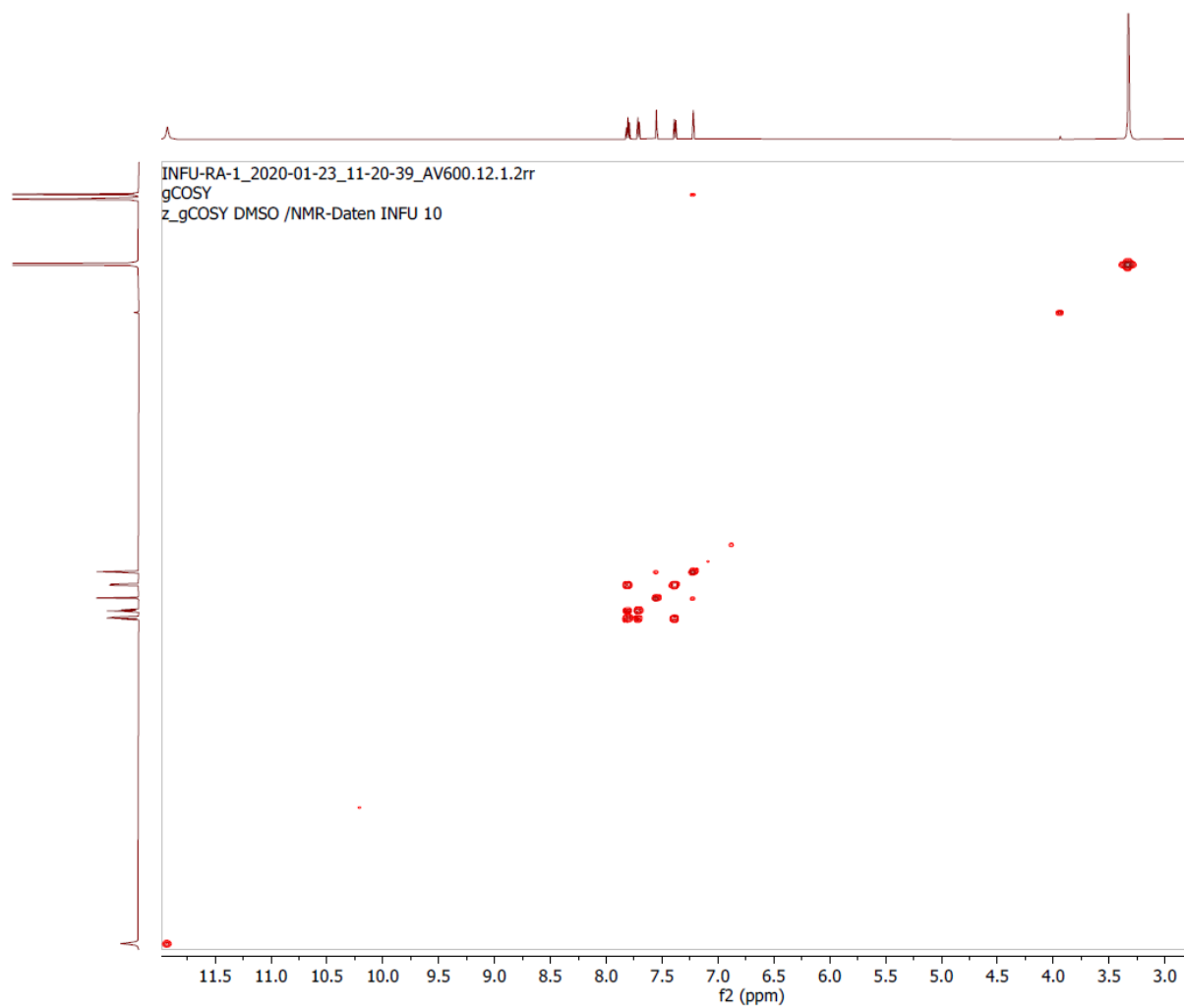
SI1b: Expanded ^1H NMR spectral data of compound-**1** from 7.15-7.90 pp



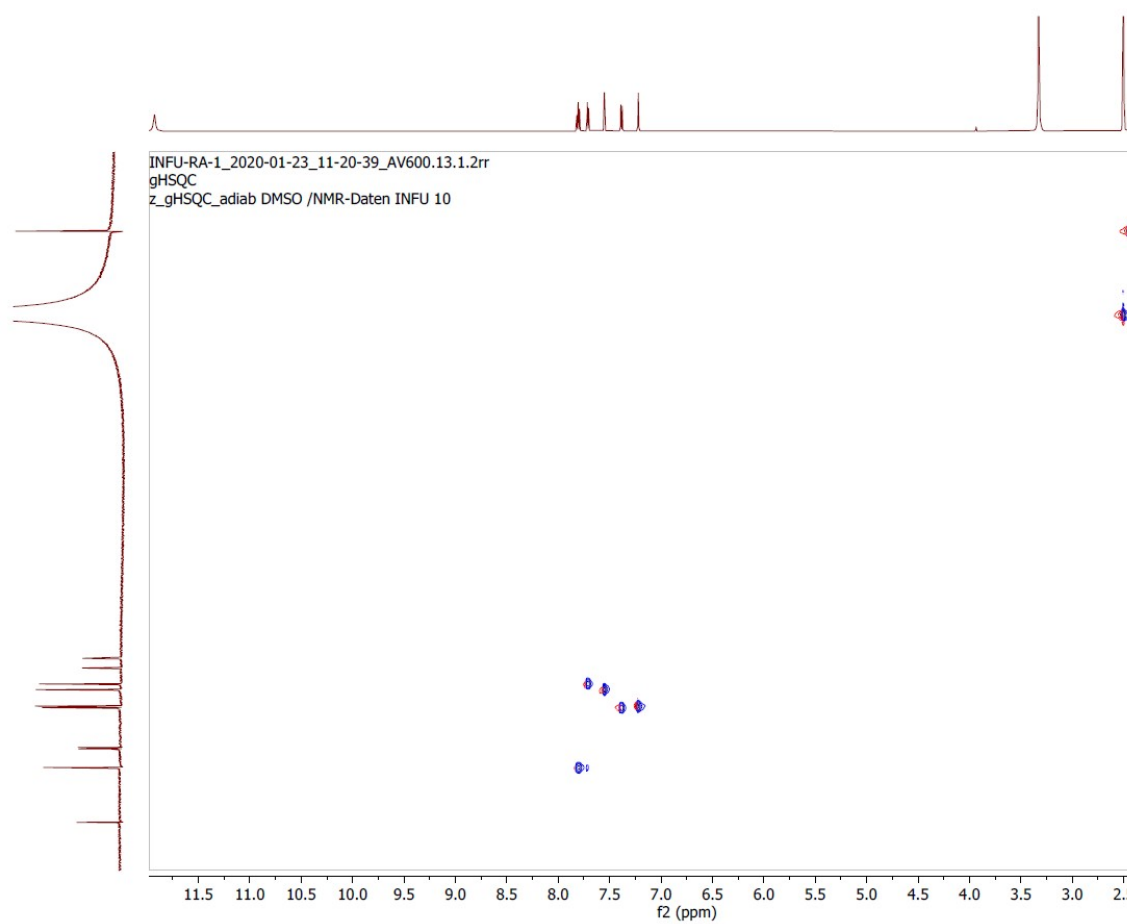
SI2a: ^{13}C NMR spectral data of compound-**1**.



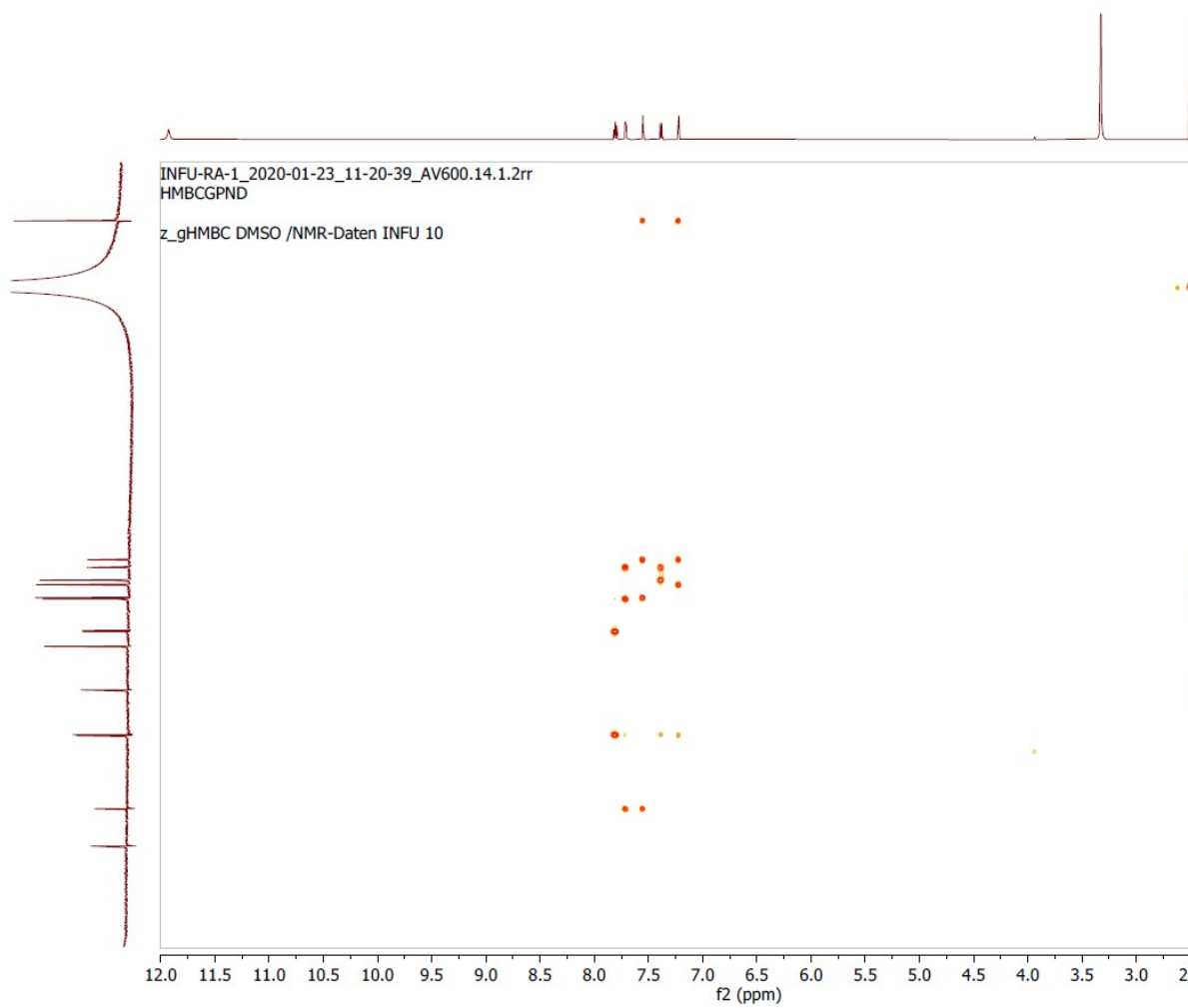
SI2b: Expanded ^{13}C NMR spectral data of compound-**1** from 115-195 pp



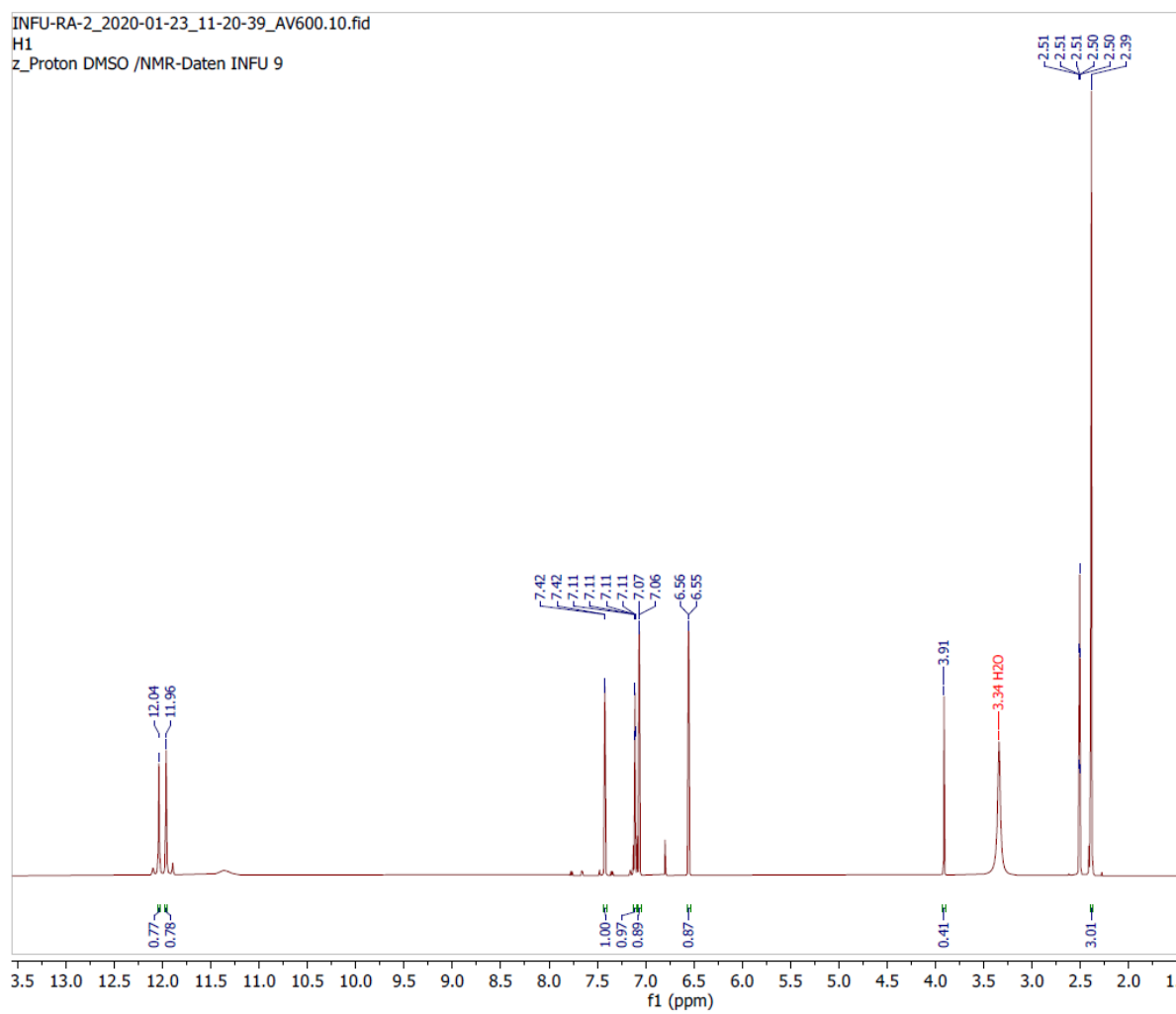
SI3: COSY spectral data of compound-**1**.



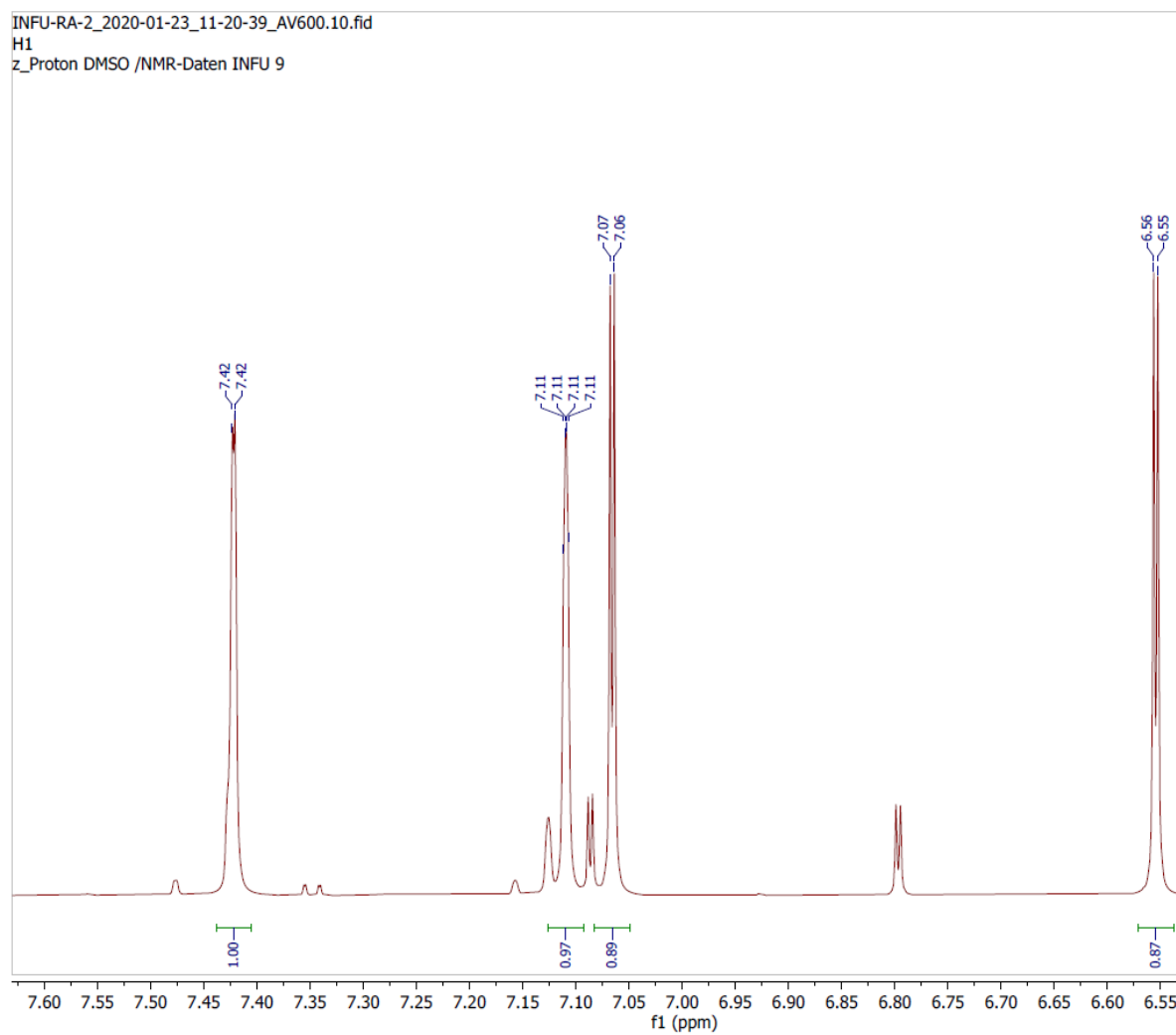
SI4: HSQC spectral data of compound-**1**.



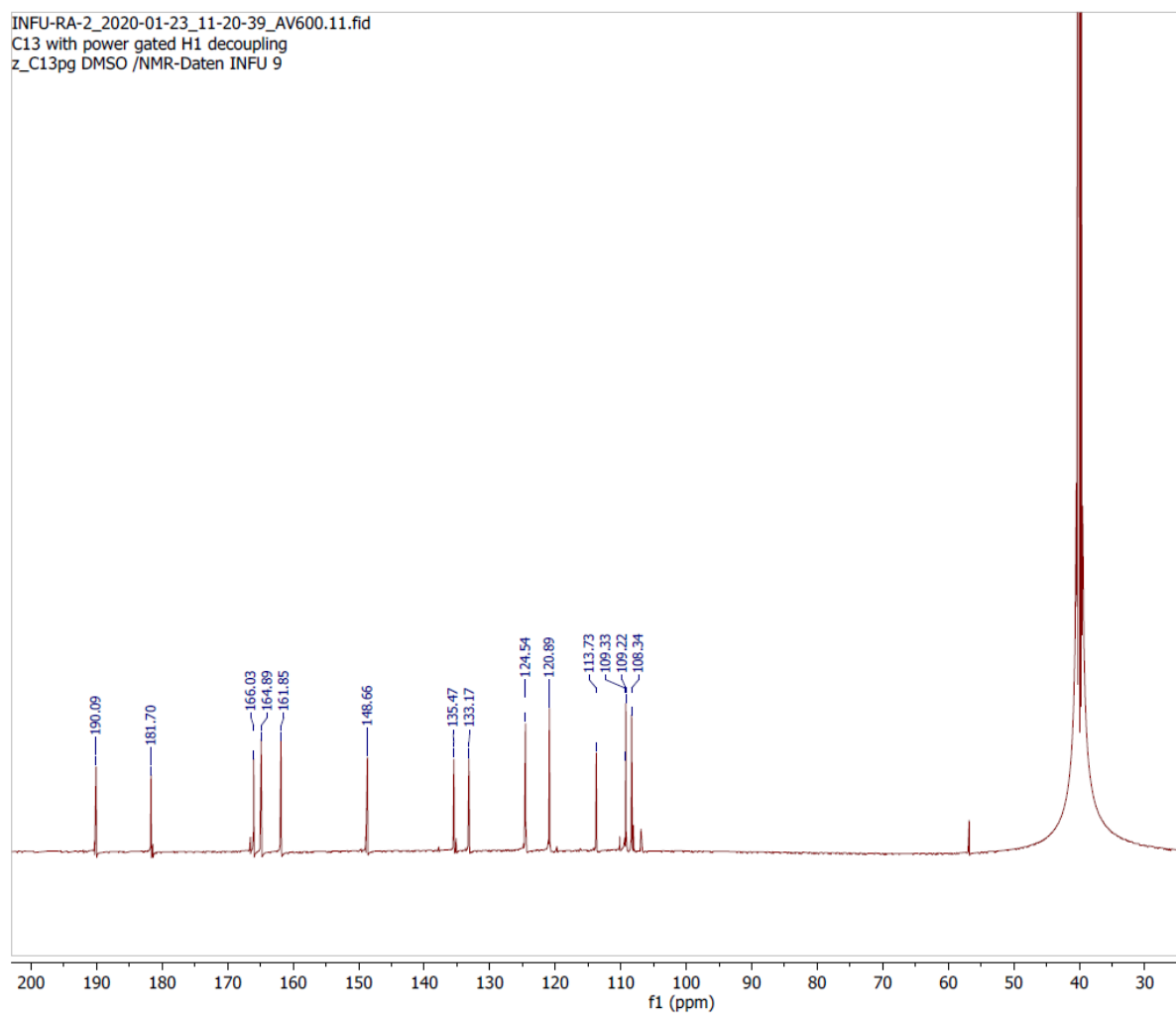
SI5: HMBC spectral data of compound-**1**.



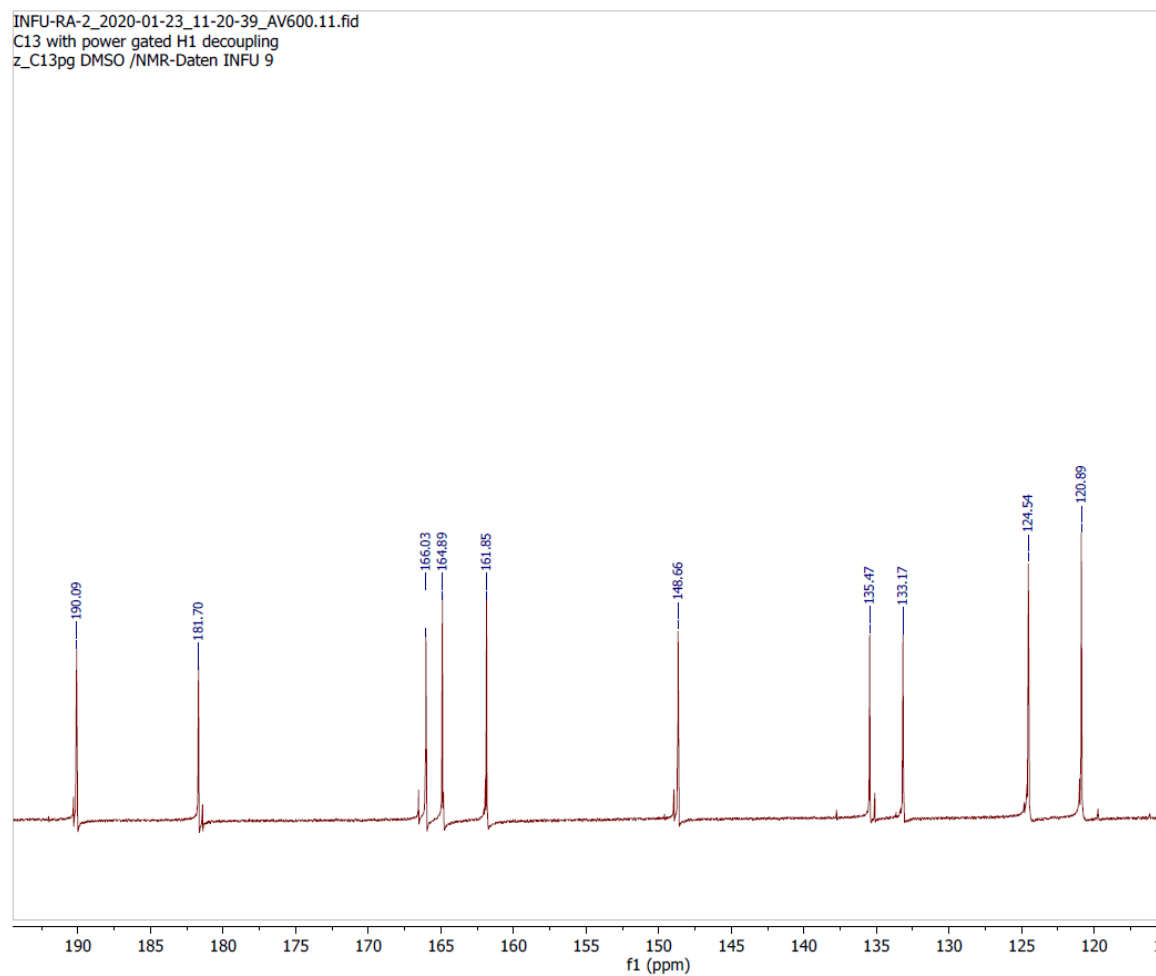
SI6a: ¹HNMR spectral data of compound-**2**.



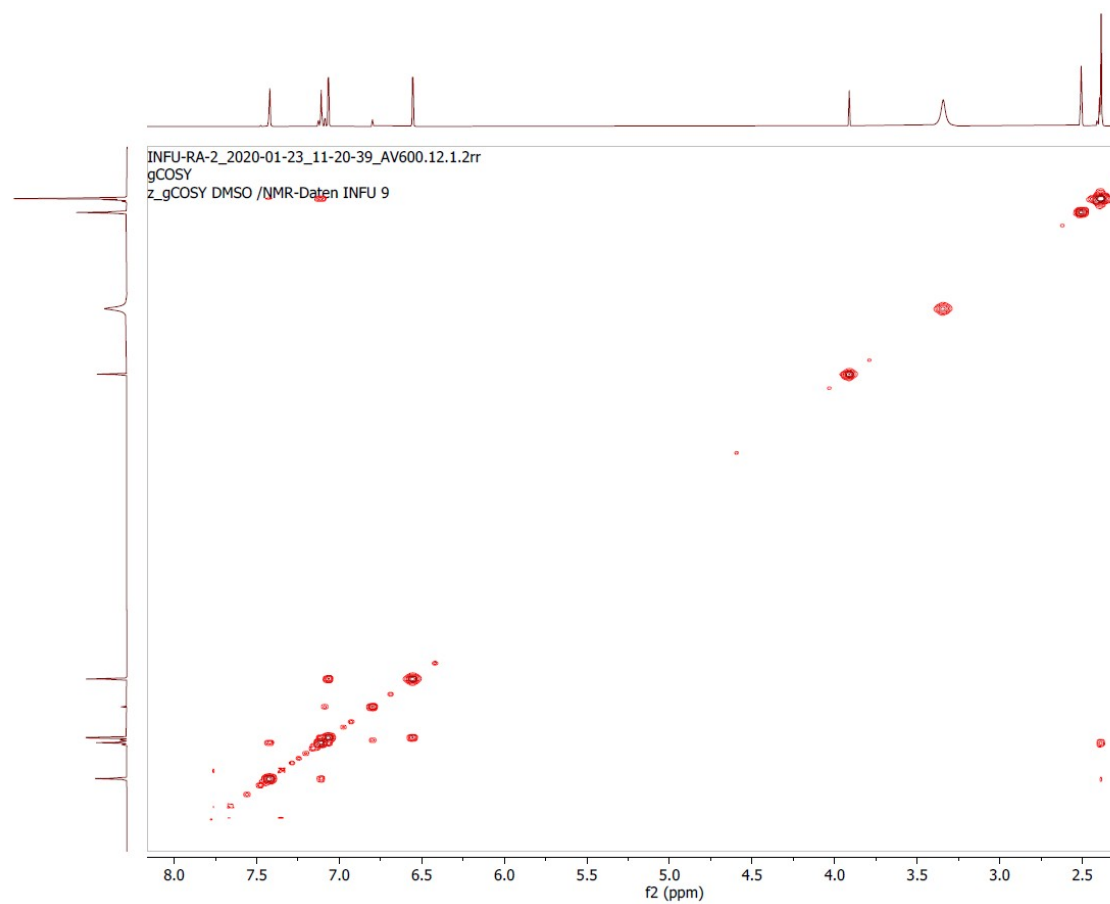
SI6b: Expanded ¹H NMR spectral data of compound-**2** from 6.40–7.60 ppm



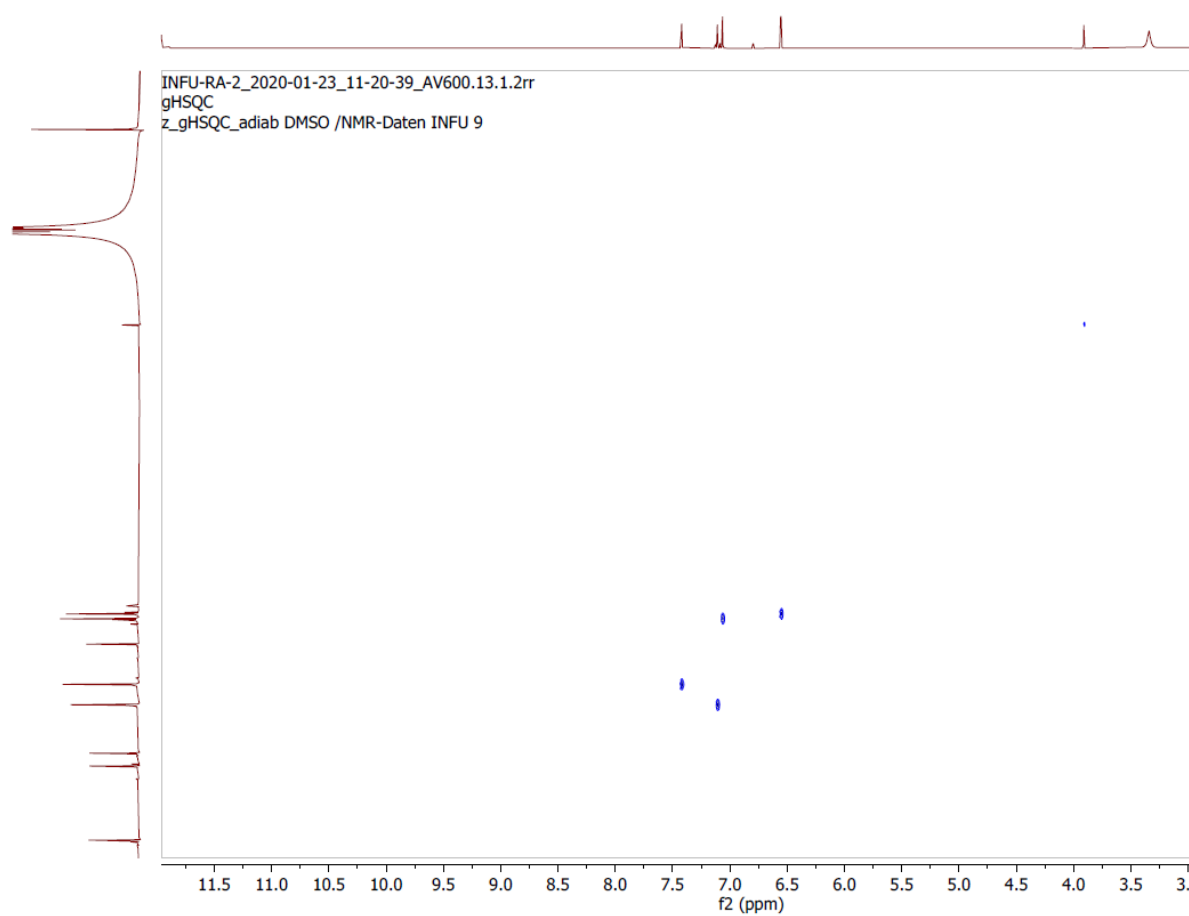
SI7a: ^{13}C NMR spectral data of compound-**2**.



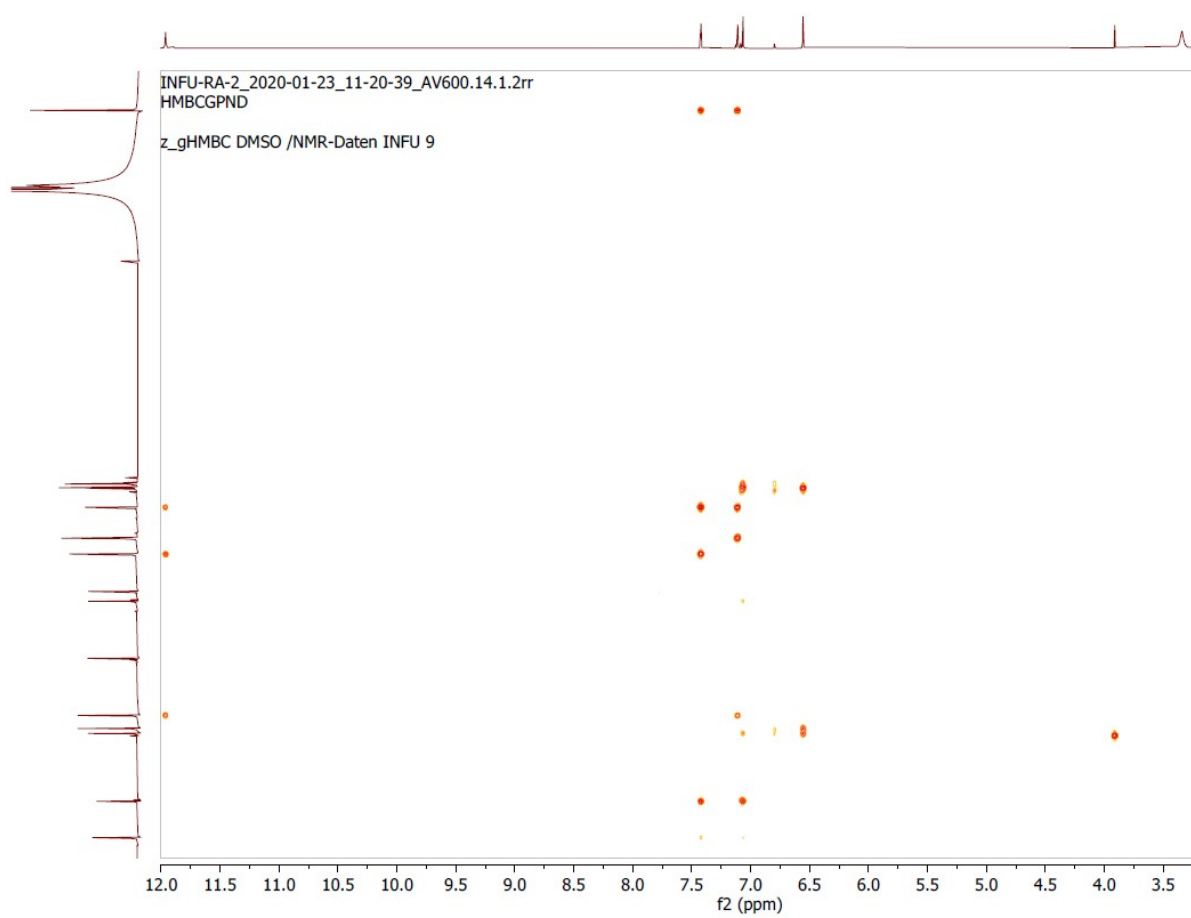
SI7b: Expanded ^{13}C NMR spectral data of compound-**2** from 110-190 pp



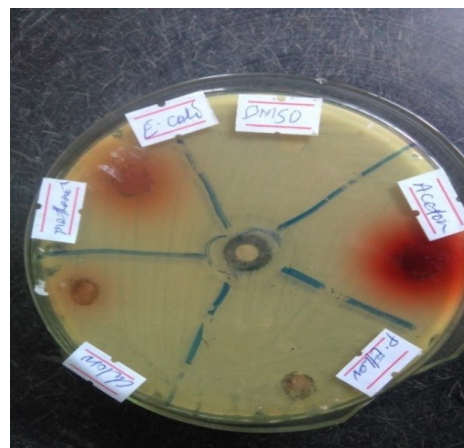
S18: COSY spectral data of compound-**2**.



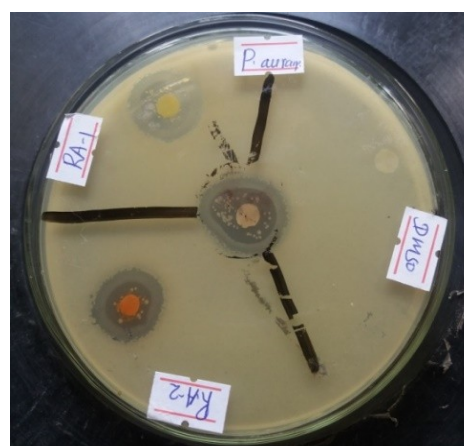
SI9: HSQC spectral data of compound-**2**.



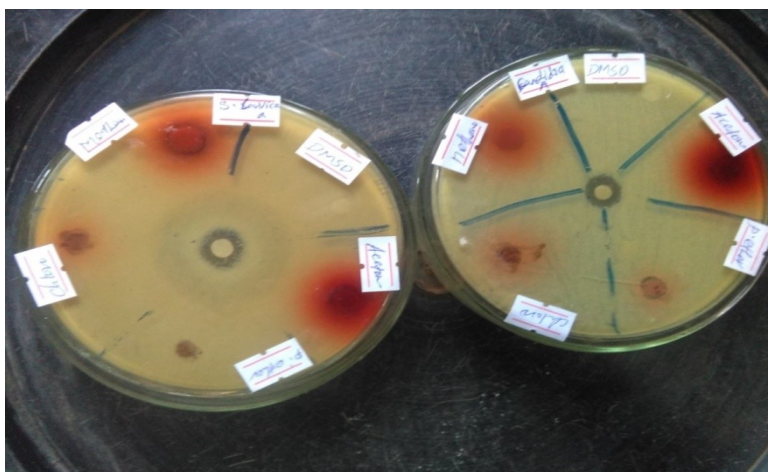
SI10: HMBC spectral data of compound-**2**.



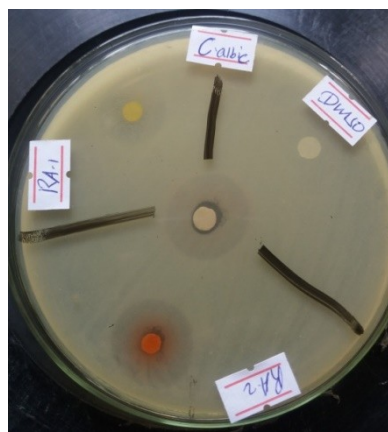
SI11a: Antibacterial activity of crude extract of root of *R. abyssinicus*



SI11b: Antibacterial activity of isolated compounds (**1** and **2**) from the root of *R*.



SI12a: Antifungal activity of crude extract of root of *R. abyssinicus*.



SI12b: Antifungal activity of isolated compounds (**1** and **2**) from the root of *R. a*.