Evaluation of the effects of sound exposure and low field electromagnetism on growth and antibiotics susceptibility of some microorganisms

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Abstract

Background: The emergence of antibiotic-resistant microorganisms has been largely associated with drug misuse, drug abuse, and indiscriminate disposal of drugs; however, the interactions between some environmental factors and antibiotic resistance by microorganisms have been understudied. In this study, the effects of sound and electromagnetic field on the growth and antibiotic susceptibility of selected microorganisms to antibiotics were investigated.

Results: Microorganisms used include Gluconobacter oxydans ATCC 19,357, Rhodobacter sphaeroides ATCC 17,023, Citrobacter freundii ATCC 33,128, Yersina pestis ATCC 11,953, Bacillus subtilis 6633, Acetobacter aceti ATCC 15,973, Escherichia coli ATCC 25,922, Pseudomonas aeruginosa ATCC 9027, Streptococcus pyogenes ATCC 19,613, Klebsiella pneumonia ATCC 25,955, Staphylococcus aureus ATCC 25,923 and Serratia marcescens ATCC 14,766. The antibiotics used were: Ciprofloxacin 5 μg, Imipenem 10 μg, Ampicillin 10 μg, Ceftazidime 30 μg and Tetracycline 30 μg for Gram-negative bacteria while Pefloxacin 10 μg, Gentamycin 10 μg, Amplicillin + Cloxacillin 30 μg, Cefuroxime 20 μg, Amoxicillin 30 μg, Ceftriaxone 25 μg, Ciprofloxacin 10 μg, Streptomycin 30 μg, Co-trimoxazole 30 μg, and Erythromycin 10 μg for Gram-positive bacteria, respectively. Acoustic treatment had varying effects on the antibiotics susceptibility profile of all test bacterial culture. Before exposure, P. aeruginosa had the highest zone of inhibition of 34 ± 3.4 mm, while B. subtilis had least inhibition zone of 12 ± 2.8. After exposure to acoustic treatment at 5000 Hz/90 dB at 72 h, C. freundii had highest zone of inhibition of 32 ± 0.7 mm and the least zone of 11 ± 1.4 mm observed in P. aeruginosa. At 1125 Hz/80 dB after 72 h, R. sphaeroides had highest zone of 34 ± 0.7 mm while A. aceti had least zone of inhibition of 10 ± 0.0 mm. Effect of electromagnetic flux treatment of 15 min showed E. coli to be the most inhibited having a growth rate of 0.08 log cfu/mL, antibiotics testing showed G. oxydans to have the highest zone of inhibition of 28 ± 3.5 mm and least zone was observed in B. subtilis having a zone of 13 ± 2.8 mm.

Conclusion: This study showed that environmental factor such as sound and electromagnetic flux (EMF) could interfere with the physiology of bacteria including resistance/susceptibility to antibiotics. However, further investigation will be needed to understand full mechanisms of action of sound and electromagnetic field on bacteria.

Keywords: Audible sound, Electromagnetic flux, Typed strain culture, Music

Background

Several biophysical factors such as sound, electromagnetic radiation and electric current resulting from natural and anthropogenic activities have been reported to affect the growth and metabolism of various life forms...
Microorganisms used in this study were *Gluconobacter oxydans* ATCC 19,357, *Rhodobacter sphaeroides* ATCC 17,023, *Citrobacter freundii* ATCC 33,128, *Yersina pestis* ATCC 11,953, *Bacillus subtilis* ATCC 6633, *Acetobacter acetii* ATCC 15,973, *Escherichia coli* ATCC 25,922, *Pseudomonas aeruginosa* ATCC 9027, *Streptococcus pyogenes* ATCC 19,613, *Klebsiella pneumonia* ATCC 25,955, *Staphylococcus aureus* ATCC 25,923 and *Serratia marcescens* ATCC 14,766. The organisms were purchased from the American Type Culture Collection (ATCC), and they were stored at 4 °C on Nutrient Agar (Oxoid Uk) slants and sub-cultured fortnightly.

### Antibiotics susceptibility test pattern
Antibiotic susceptibility of the organisms was performed by using eighteen to twenty four (18–22) hours old bacterial cultures using different antibiotics discs for both Gram-positive and Gram-negative organisms according to the Kirby-Bauer technique as described by Baker et al. (1980). The antibiotics used were: Ciprofloxacin 5 μg, Imipenem 10 μg, Ampicillin 10 μg, Ceftazidime 30 μg and Tetracycline 30 μg for Gram-negative bacteria while Pefloxacin 10 μg, Gentamycin 10 μg, Ampicillin + Cloxacillin 30 μg, Cefuroxime 20 μg, Amoxicillin 30 μg, Ceftriaxone 25 μg, Ciprofloxacin 10 μg, Streptomycin 30 μg, Co-trimoxazole 30 μg, and Erythromycin 10 μg for Gram-positive bacteria, respectively. The discs were purchased from Oxoid chemicals (UK). The diameters of the inhibition or stimulation zone were measured after 24 h, 48 h and 72 h, respectively.

### Effect of sound of different frequencies on growth and antibiotic sensitivity pattern
Exposure of the different organisms to different acoustic treatment was done in an anechoic chamber constructed locally as described by Guet al. (2010) using polyurethane foam of density 1 g/cm³, wood and cotton fabric material in a cylindrical shape. The dimensions of the cylinder were 15 inches height, 12 inches diameter and had a rectangular orifice of dimension 5.6 inches by 6 inches plate 1. Each of the test organism was inoculated into sterile nutrient broth and incubated for 24 h, 48 h and 72 h, respectively, and thereafter exposed to acoustic treatment (produced using a Bluetooth speaker which was remotely powered and placed directly above the inoculated nutrient broth-containing the test organisms) at different exposure time ranging between 24 and 72 h, frequencies and decibel of 5000 Hz/90 dB and 1125 Hz/80 dB and growth observed on Jenway Spectrophotometer at 540 nm. Sample without sound treatment served as control group. Incubation was done at 37 ± 1 °C and growth was also checked on spectrophotometer after 24–72 h incubation duration (Guet al. 2010). Antibiotic susceptibility pattern was done before and after the acoustic exposure experiment as described earlier.
microorganism, actively growing bacterial cultures were inoculated into sterile nutrient broth and incubated at 37 C for 24 h and these were then exposed to low electromagnetic field as described by Garuba et al. (2020). The tubes of the bacteria to be exposed were placed in the middle of the coil by using supports to get a homogenous and higher magnetic field strength in which the ends of the solenoid were connected to variac fed from the mains (220 V). The magnetic field intensity was measured by means of hand held Gauss/Tesla meter, as the result proved its homogeneity among all the volume and was within ±5% at the ends. Equal volumes of the bacterial culture were exposed for 15 min at their mid-to-late exponential phase (maximum rate of active growth) to the electromagnetic field at an intensity of (5 mT), after exposure the growth rate was measured in a spectrophotometer and compared with the control (Mona et al. 2013).

Statistical analysis
Results obtained in this study were subjected to analysis of variance using ANOVA, and separation of means was carried out by Duncan's multiple range test (Duncan 1955).

Results
Twelve bacteria were subjected to two different acoustic treatments (5000 Hz/90 dB and 1125 Hz/80 dB) for duration of 24–72 h after which growth and antibiotic susceptibility pattern of each of the isolate measured. The results showed that all the bacteria culture at 5000 Hz/90 dB had an increase growth (p < 0.05) as compared with control except E. coli, K. pneumoniae and B. subtilis which decreased at 72 h acoustic exposure (Fig. 1). Similarly, exposure to sound of 1125 Hz/80 dB for 72 h had varied effects on the growth of all the organisms investigated except for S. marcescens and A. aceti which had a decline after 72 h exposure (Fig. 2).

Results of the antibiotics susceptibility pattern after the cultures were exposed to sound treatment at 5000 Hz/90 dB and 1125 Hz/80 dB is presented in Tables 1, 2, 3 and 4. The results revealed that the exposure of S. marcescens to acoustic treatment at 5000 Hz/90 dB showed increased resistance (p < 0.05) to Ciprofloxacin at 24 h (30 mm) and at 72 h (31 mm) and maintained its resistance profile to tetracycline and Ampicillin throughout the exposure duration. P. aeruginosa became susceptible to Ampicillin after 72 h (11 mm) having an initial resistance profile and treatment to Tetracycline showed resistance. G. oxydans which was initially susceptible became resistant to Ceftazidime after 72 h exposure while A. aceti became resistant to Ampicillin after exposure to treatment; B. subtilis showed increased susceptibility to Ampicillin after 24 h (27 mm) at p < 0.05 and then became resistant after 48 and 72 h exposure, respectively, as shown in Tables 1 and 2. Similarly, an acoustic treatment of 1125 Hz/80 dB, R. sphaeroides showed increased susceptibility at 72 h (28 mm) (at p < 0.05) to Ceftazidime and also to Imipenem (34 mm), respectively. A. aceti which had an initial susceptibility profile (16 mm) became resistant to Ampicillin at 48 h and 72 h exposure treatment, exposure of S. aureus to acoustic treatment showed increased resistance to Pefloxacin, Gentamycin, Ampicillin, Amoxacillin, Streptomycin all through the exposure duration, respectively, as shown in Tables 3 and 4.
Exposure of the test organisms to electromagnetic flux of 5 mT for 15 min was carried out and the growth alongside antibiotic susceptibility pattern investigated. The results presented in Tables 5 and 6 and Fig. 3 showed that Y. pestis, K. pneumonia and A. aceti had an increased growth (p < 0.05) rate compared to control while other test bacteria had a decrease in growth compared to the control (Tables 5 and 6).

Electromagnetic exposure was also found to have varying effects on the antibiotic susceptibility pattern of the organisms under investigation. Exposure of G. oxydans showed an increase susceptibility to Tetracycline (18 mm), Y. pestis also showed an increase in susceptibility to Ceftazidime (19 mm) after electromagnetic treatment. B. subtilis became resistant to Ampicillin and S. aureus became resistant to Gentamycin, Ampicillin, Cefuroxime and Amoxacillin, respectively. S. pyogenes was observed to have an increase susceptibility to Pefloxacin (20 mm), Gentamycin (17 mm), Ampicillin (13 mm), Ceftriaxone (19 mm), Ciprofloxacin (23 mm), Streptomycin (23 mm), Cotrimoxazole (24 mm) and Erythromycin (24 mm), respectively, as shown in Table 5 and 6.

Discussion
In natural world, nearly all life forms are surrounded by different kinds of sound and electromagnetic fields from different sources (Dånet 2005,) and there are few reports documenting the interactions between microorganisms, sound and electromagnetic fields (Gu et al. 2016; Movahedi et al. 2019; Robinson et al. 2021). However, the data provided by literature is not adequate in elucidating the physiological response to interactions between biological systems and these physical factors (that either occurs naturally or as a result of anthropogenic activities) (Gu et al. 2010; Said-Salman et al. 2019). Furthermore, the use of physical signals such as sound waves, by the microorganisms for their intercellular communication has not received due attention (Chandra et al. 2018).

In this study, effect of low electromagnetic field and acoustic sound on the growth and antibiotics susceptibility pattern of twelve bacteria were investigated and some organisms were observed to respond to sound stimulation with a positive effect on growth, whereas sound waves had an inhibitory effect on the growth of some other microorganisms. This suggests that the action of sound on microbial growth showed obvious
frequency peculiarities. This peculiarity has been previously reported by Kothari et al. (2018). Similarly, Souvik et al. (2018) investigated the effect of three types of sound frequencies between 20 Hz and 20 kHz on the growth and the antibiotic susceptibility pattern of *E. coli*. The results indicated that *E. coli* growth increased when exposed to sound frequency below 1 kHz (432 Hz) while at frequencies above 1 kHz, a poor growth was recorded. The altered growth rate observed has been suggested to be due to an altered movement of key molecules across the cell membrane of the organism as sound has been reported to increase membrane permeability in some microorganisms (Sarvaiya and Kothari 2015). The increased growth observed could be as a result of an adaptive response of the test bacteria to the sound exposure having an influence on microbial growth.

**Fig. 2** Growth rate of selected Gram-negative (a) and Gram-positive bacteria (b) after exposure to acoustic treatment at 1125 Hz/80 dB compared with unexposed control.
due to increase in membrane permeability while reduced growth observed could be as a result of mechanical stress created by the sound exposure after a long duration which could trigger the stress response of the bacteria hence resulting in the death of the organisms (Shah et al. 2016; Sarvaiya and Kothari, 2017; Kothari et al. 2018). This is because sound waves have been reported to travel through any medium giving rise to sonic vibrations which are thought to be sensed by the test microbial population through mechanosensory receptors (which act as safety valves against osmotic shock, and in higher organisms they participate in sensing touch and sound waves) (Kung et al. 2010; Ward et al. 2014; Shah et al. 2016). The microbial population may then modulate its behaviour in accordance to the magnitude and duration of the sonic vibration by generating appropriate physiological responses (Martinac 2012) such as opening of mechanosensitive ion channels which in turn affects movement of certain key ions across the cell membrane, ultimately resulting in an altered pattern of growth and metabolism (Vitkauskaite and Grubliauskas 2018). Antibiotic susceptibility pattern of the bacteria investigated in this study was also found to vary considerably with some of the becoming more susceptible while other became resistant as previously reported (Sarvaiya and Kothari 2017; Souvik et al. 2018). The altered antibiotic pattern of antibiotic sensitivity by the test organisms has been suggested to result from an altered transport (increased uptake) of the antibiotic used through the cell membrane via the mechanosensitive channel which responds to mechanical stress by

Table 1 Antibiotics susceptibility profile for selected Gram-negative bacteria as read in mean diameter of zones of inhibition (mm) after exposure to acoustic treatment at 5000 Hz/90 dB compared with control

| Bacteria          | CAZ | CIP | TET | IMP | AMP |
|-------------------|-----|-----|-----|-----|-----|
| *S. marcescens* ATCC14756 | 26±1.4* | 28±2.8 | R   | 26±1.4 | R   |
| b) 17±0   | b) 30±0.7 | b) R   | 19±2.1 | b) R   |
| c) 25±1.4 | c) 28±2.1 | c) R   | 24±1.4 | c) R   |
| d) 26±1.4 | d) 31±1.4 | d) R   | 25±0.7 | d) R   |
| *P. aeruginosa* ATCC9027 | 26±1.4 | 34±3.5 | R   | 26±1.4 | R   |
| b) 23±0.7 | b) 29±1.4 | b) R   | 28±2.1 | b) R   |
| c) 20±1.4 | c) 21±0.7 | c) R   | 24±0.7 | c) R   |
| d) 22±0.0 | d) 22±3.5 | d) R   | 29±1.4 | d) 11±1.4 |
| *G. oxydans* ATCC19357 | 27±2.8 | 26±0.7 | R   | 26±1.4 | 23±0.7 |
| b) R     | b) 27±3.5 | b) 17±2.8 | 27±2.8 | 28±0.7 |
| c) 29±1.4 | c) 21±2.1 | c) 12±0.7 | 24±2.8 | 14±2.1 |
| d) 19±49 | d) 21±0.7 | d) R   | 28±0.7 | d) R   |
| *R. sphaeroides* ATCC17023 | 24±0.7 | 24±0.7 | R   | a) R   |
| b) 16±0   | b) 15±0.0 | b) R   | 19±0.7 | b) R   |
| c) R     | c) 20±2.1 | c) 25±0.7 | 21±0.7 | c) R   |
| d) 14±57 | d) 18±2.1 | d) R   | 23±5.7 | d) R   |
| *C. freundii* ATCC33128 | 14±0.7 | 26±0.7 | R   | 29±4.2 | 30±2.1 |
| b) 31±64 | b) 22±5.7 | b) 12±2.8 | 30±0.7 | 21±9.2 |
| c) 16±42 | c) 16±2.1 | c) 15±2.1 | 28±2.1 | c) R   |
| d) 14±2.1 | d) 19±7.1 | d) 20±2.1 | 32±0.7 | d) R   |
| *Y. pestis* ATCC11953 | 13±1.4 | 25±0.0 | R   | 23±1.4 | 15±0.0 |
| b) 13±2.1 | b) 20±0.7 | b) R   | 18±5.7 | b) R   |
| c) 15±0.7 | c) 22±0.7 | c) R   | 24±1.4 | c) 13±0.7 |
| d) 13±14 | d) 21±0.7 | d) R   | 25±0.0 | d) 21±2.1 |
| *K. pneumoniae* ATCC25923 | 26±1.4 | 31±2.1 | R   | 22±2.8 | 24±0.7 |
| b) 26±0 | b) 20±1.4 | b) 16±1.27 | 20±1.4 | 14±2.1 |
| c) 26±1.4 | c) 23±3.5 | c) 17±4.2 | 25±4.9 | 14±6.4 |
| d) 28±5.7 | d) 18±1.4 | d) R   | 31±1.4 | d) R   |
| *A. aceti* ATCC15973 | 19±4.2 | 21±9.9 | R   | 13±4.2 | 17±1.4 |
| b) 21±2.8 | b) 19±1.4 | b) 15±9.2 | 30±24 | b) R   |
| c) 23±28 | c) 19±0 | c) 13±1.4 | 26±0.7 | c) R   |
| d) 27±14 | d) 21±2.8 | d) 17±0.7 | 28±0.0 | d) R   |
| *E. coli* ATCC25922 | 29±1.4 | 22±1.4 | R   | 19±0   | 33±4.2 |
| b) 20±1.4 | b) 21±0.7 | b) R   | 30±0.7 | b) R   |
| c) R     | c) 18±3.5 | c) 14±7.8 | 30±2.1 | c) R   |
| d) 15±14 | d) 24±0.7 | d) R   | 29±1.4 | d) R   |

a) Control b) 24 h c) 48 h d) 72 h R—Resistance CAZ—Ceftazidime CIP—Ciprofloxacin
TET—Tetracycline IMP—Imipenem AMP—Ampicillin *Data are mean of replicate ± Standard deviation
Antibiotics susceptibility profile for selected Gram-positive bacteria as read in mean diameter of zones of inhibition (mm) after exposure to acoustic treatment of 5000 Hz/90 dB compared with control

| Bacteria            | PEF | CN  | APX | Z    | AM  | R   | CPX | S   | SXT | E   |
|---------------------|-----|-----|-----|------|-----|-----|-----|-----|-----|-----|
| B. subtilis ATCC6633|     |     |     |      |     |     |     |     |     |     |
| a) 14 ± 1.4         | a) 16 ± 1.4 | a) 12 ± 2.8 | a) 12 ± 1.4 | a) 12 ± 1.4 | a) 16 ± 2.1 | a) 18 ± 1.4 | a) 20 ± 2.8 | a) 12 ± 1.4 | a) 14 ± 2.8 |
| b) 26 ± 1.4         | b) 20 ± 2.8 | b) 27 ± 3.5 | b) 21 ± 0.7 | b) 22 ± 3.5 | b) 15 ± 2.8 | b) 28 ± 2.1 | b) 30 ± 0.7 | b) 21 ± 2.1 | b) 30 ± 3.5 |
| c) 16 ± 1.4         | c) 16 ± 2.1 | c) R  | c) R  | c) 16 ± 2.1 | c) 14 ± 1.4 | c) 19 ± 1.4 | c) 14 ± 5.7 | c) 16 ± 0  | c) 17 ± 0.7 |
| d) 16 ± 0.7         | d) 12 ± 2.1 | d) R  | d) R  | d) 14 ± 0  | d) 19 ± 0  | d) 16 ± 2.1 | d) R  | d) R  | d) R  |
| S. pyogenes ATCC19613|     |     |     |      |     |     |     |     |     |     |
| a) 14 ± 2.8         | a) 16 ± 2.1 | a) 12 ± 2.1 | a) 14 ± 1.4 | a) 16 ± 1.4 | a) 18 ± 4.2 | a) 18 ± 1.4 | a) 18 ± 1.4 | a) 18 ± 1.4 | a) 12 ± 2.8 |
| b) 20 ± 1.4         | b) 17 ± 28 | b) 18 ± 21 | b) 17 ± 14 | b) 17 ± 21 | b) 15 ± 1.4 | b) 19 ± 1.4 | b) 19 ± 1.4 | b) 19 ± 21  | b) 21 ± 0.7 |
| c) 18 ± 1.4         | c) 15 ± 21 | c) R  | c) 12 ± 0.7 | c) 16 ± 0  | c) 18 ± 21 | c) 18 ± 21 | c) 15 ± 0  | c) R  | c) 14 ± 1.4 |
| d) 16 ± 0.7         | d) 19 ± 28 | d) 17 ± 28 | d) 19 ± 0.7 | d) 17 ± 14 | d) 16 ± 28 | d) 19 ± 21 | d) 16 ± 28 | d) R  | d) R  | d) R  | d) R  |
| S. aureus ATCC25923 |     |     |     |      |     |     |     |     |     |     |
| a) 12 ± 1.4         | a) 14 ± 14 | a) 12 ± 2.8 | a) R  | a) 16 ± 2.1 | a) 16 ± 1.4 | a) 16 ± 1.4 | a) 18 ± 1.4 | a) 18 ± 2.8 | a) 14 ± 1.4 |
| b) 12 ± 5.7         | b) 14 ± 49 | b) R  | b) R  | b) 15 ± 1.4 | b) 19 ± 0.7 | b) R  | b) R  | b) R  | b) R  |
| c) 12 ± 0.7         | c) 12 ± 2.1 | c) R  | c) R  | c) R  | c) 17 ± 2.1 | c) 19 ± 14 | c) R  | c) R  | c) R  | c) R  |
| d) R                | d) 15 ± 14 | d) R  | d) R  | d) R  | d) 15 ± 0  | d) 18 ± 35 | d) R  | d) R  | d) R  | d) R  |

a) Control b) 24 h c) 48 h d) 72 h R—Resistance PEF: Penicillin CN: Gentamicin APX: Ampicillin Z: Cefuroxime AM: Amoxacillin R: Ceftriaxone CPX: Ciprofloxacin S: Streptomycin SXT: Cotrimoxazole E: Erythromycin
changing its conformity between an open and closed state and this response is dependent on the frequency of sound, type of microorganism and exposure duration (Sarvaiya and Kothari 2017).

Non-ionising, non-thermal, extremely low frequency and electromagnetic fields (ELF-EMF) have been reported to produce a number of biological effects during their interaction with biological systems (Belyaev et al. 2011; Nguyen et al. 2015; Taheri et al. 2017a; Movahedi et al. 2019). In this study, there exists a decrease in growth rate after exposure to electromagnetic field of 5mT. This observation is in accordance with Inhan-Garip et al. (2011) who carried an experiment on three Gram-negative and three Gram-positive bacteria and observed a decrease in growth of all test cultures after exposure to electromagnetic flux. This reduced cell growth observed could be a result of increased level of reactive oxygen species prompted by exposure to electromagnetic field which may have affected macromolecular synthesis and caused protein injury, ultimately leading to a decrease in bacterial growth rate as suggested by Souvik et al. (2018).

After exposure to electromagnetic field, some of the bacteria were found to developed resistance to antibiotics they were previously susceptible to while some became more susceptible. Chiefly among the several factors affecting antimicrobial susceptibility are the physical properties of the magnetic field, such as frequency, duration of radiation, the structure of the bacterial cell wall and the nature of the peptidoglycan properties in the cell wall of the Gram-positive and Gram-negative bacteria (specifically with changes in the bacterial cell wall protein as suggested by Taheri et al. 2017a) and Said-Salman et al. 2019). Mechanisms of acquired susceptibility or acquired resistance

Table 3 Antibiotics susceptibility profile for selected Gram-negative bacteria as read in mean diameter of zones of inhibition (mm) after exposure to acoustic treatment at 1125 Hz/80 dB compared with control

| Bacteria            | Exposure | CAZ | CIP | TET | IMP | AMP |
|---------------------|----------|-----|-----|-----|-----|-----|
| S. marcescens ATCC14756 | a) Control  | 26±1.4 | 28±2.8 | R | 26±1.4 | R |
| b) 24 h | 23±1.4 | 21±1.4 | R | 29±1.4 | R |
| c) 48 h | 23±0  | 29±2.8 | R | 32±1.4 | R |
| d) 72 h | 20±2.9 | 25±0.7 | 12±2.1 | 26±5.7 | 19±4.2 |
| P. aeruginosa ATCC9027 | a) Control  | 26±1.4 | 34±3.5 | a) R | 26±2.8 | a) R |
| b) 24 h | 20±9.2 | 23±3.5 | b) R | 17±4.2 | b) R |
| c) 48 h | 26±2.1 | 22±0  | c) R | 27±2.8 | b) R |
| d) 72 h | 30±0.7 | 27±1.4 | d) R | 31±4.9 | d) R |
| G. oxydans ATCC19357 | a) Control  | 27±2.8 | 26±0.7 | a) R | 26±1.4 | a) 23±0.7 |
| b) 24 h | 20±0  | 17±2.1 | b) R | 24±3.5 | b) R |
| c) 48 h | 14±2.1 | 19±1.4 | c) R | 26±2.8 | b) R |
| d) 72 h | R      | 22±0  | d) R | 18±1.3 | d) R |
| R. sphaeroides ATCC17023 | a) Control  | 24±0.7 | 24±0.7 | a) R | 24±0.7 | a) R |
| b) 24 h | 19±0.7 | 23±0  | b) R | 23±3.5 | b) R |
| c) 48 h | 24±1.4 | 24±3.5 | c) R | 28±0.7 | c) R |
| d) 72 h | 28±0.7 | 23±3.5 | d) R | 34±0.7 | d) R |
| C. freundii ATCC33128 | a) Control  | 14±0.7 | 26±0.7 | a) R | 29±4.2 | a) 30±2.1 |
| b) 24 h | 14±0  | 20±0  | b) R | 25±0.7 | b) R |
| c) 48 h | 20±7.1 | 31±2.1 | c) R | 12±0.7 | c) R |
| d) 72 h | R      | 30±0  | d) R | 11±1.4 | d) R |
| Y. pestis ATCC11953 | a) Control  | 13±1.4 | 23±0  | a) R | 23±1.4 | a) 15±0 |
| b) 24 h | 23±0.7 | 22±2.1 | b) R | 20±4.9 | b) 14±2.1 |
| c) 48 h | 18±0  | 20±0.7 | c) R | 26±0.7 | c) 15±0.7 |
| d) 72 h | 16±2.8 | 18±2.8 | d) R | 23±0.7 | d) 17±1.4 |
| K. pneumoniae ATCC25955 | a) Control  | 26±1.4 | 31±2.1 | a) R | 22±2.8 | a) 24±0.7 |
| b) 24 h | 27±1.4 | 21±1.4 | b) R | 30±0.7 | b) R |
| c) 48 h | 26±0.7 | 23±0.7 | c) R | 25±5.7 | c) R |
| d) 72 h | 26±2.8 | 25±3.5 | d) R | 31±0.7 | d) R |
| A. aceti ATCC15973 | a) Control  | 19±4.2 | 21±9.9 | a) R | 17±1.4 | a) 16±4.2 |
| b) 24 h | 26±0  | 19±2.1 | b) R | 34±7.1 | b) 13±1.4 |
| c) 48 h | 14±4.2 | 28±4.9 | c) R | 21±14.1| c) 22±2.8 | c) R |
| d) 72 h | 19±0.7 | 20±0  | d) R | 10±0  | d) 27±1.4 | d) R |
| E. coli ATCC25922 | a) Control  | 29±1.4 | 22±1.4 | a) R | 33±4.2 | a) 26±1.4 |
| b) 24 h | 14±1.4 | 20±0.7 | b) R | 20±0  | b) R |
| c) 48 h | 17±0.7 | 21±1.4 | c) R | 20±4.2 | c) R |
| d) 72 h | 18±1.4 | 21±0.7 | d) R | 27±1.4 | d) R |

a) —Control b) —24 h c) —48 h d) —72 h R —Resistance CAZ—Ceftazidime CIP—Ciprofloxacin TET—Tetracycline IMP—Impinem AMP—Ampicillin *Data are mean of replicate ± S.D
Table 4  Antibiotics susceptibility profile for selected Gram-positive bacteria read in mean diameter of zones of inhibition (mm) at 1125 Hz/80 dB compared with control

| Bacteria          | PEF     | CN     | APX     | Z       | AM     | R       | CPX     | S       | SXT     | E       |
|-------------------|---------|--------|---------|---------|--------|---------|---------|---------|---------|---------|
| B. subtilis ATCC6633 |         |        |         |         |        |         |         |         |         |         |
| a) 14 ± 1.4       | a) 16 ± 1.4 | a) 12 ± 1.4 | a) 12 ± 1.4 | a) 16 ± 2.1 | a) 18 ± 1.4 | a) 20 ± 2.8 | a) 12 ± 1.4 | a) 14 ± 2.8 |
| b) 12 ± 1.4       | b) R    | b) R   | b) R    | b) R    | b) R   | b) 16 ± 4.2 | b) 21 ± 2.1 | b) R    | b) R    |
| c) 18 ± 4.2       | c) 18 ± 2.8 | c) R   | c) R    | c) R    | c) 14 ± 0.7 | c) 17 ± 0 | c) 13 ± 1.4 | c) R    | c) 15 ± 4.9 |
| d) 16 ± 5.7       | d) R    | d) R   | d) R    | d) R    | d) R   | d) 14 ± 0.7 | d) 13 ± 0 | d) R    | d) 15 ± 3.5 |
| S. pyogenes ATCC19613 |        |        |         |         |        |         |         |         |         |         |
| a) 14 ± 2.8       | a) 16 ± 2.1 | a) 12 ± 2.1 | a) 14 ± 1.4 | a) 16 ± 1.4 | a) 18 ± 4.2 | a) 18 ± 1.4 | a) 18 ± 1.4 | a) 12 ± 2.1 |
| b) 14 ± 4.1       | b) 12 ± 4.9 | b) 12 ± 1.4 | b) 13 ± 1.4 | b) 14 ± 2.8 | b) R   | b) 17 ± 1.4 | b) 12 ± 1.4 | b) R    | b) R    |
| c) 13 ± 0.7       | c) R    | c) 12 ± 1.4 | c) 15 ± 1.4 | c) 15 ± 2.8 | c) R   | c) 16 ± 3.5 | c) 15 ± 4.9 | c) R    | c) 12 ± 0.7 |
| d) 17 ± 0.7       | d) R    | d) R   | d) 12 ± 2.1 | d) R    | d) 12 ± 3.5 | d) 12 ± 1.4 | d) 12 ± 0 | d) R    | d) 16 ± 2.1 |
| S. aureus ATCC25923 |        |        |         |         |        |         |         |         |         |         |
| a) 12 ± 1.4       | a) 14 ± 1.4 | a) 14 ± 2.8 | a) R    | a) 16 ± 2.1 | a) 16 ± 1.4 | a) 20 ± 1.4 | a) 18 ± 1.4 | a) 14 ± 2.8 | a) 14 ± 1.4 |
| b) R              | b) R    | b) R   | b) R    | b) R    | b) 12 ± 0 | b) 17 ± 1.4 | b) R    | b) R    | b) 15 ± 4.9 |
| c) R              | c) R    | c) R   | c) R    | c) R    | c) 16 ± 3.5 | c) 16 ± 2.8 | c) R    | c) R    | c) R    |
| d) R              | d) R    | d) R   | d) R    | d) R    | d) 13 ± 2.1 | d) 14 ± 2.8 | d) R    | d) R    | d) R    |

a) —Control  b) —24 h  c) —48 h  d) —72 h  R—Resistance  PEF—Pefloxacin  CN—Gentamycin  APX—Ampicillin  + Cloxacillin
Z—Cefuroxime  AM—Amoxicillin  R—Ceftriaxone  CPX—Ciprofloxacin  S—Streptomycin  SXT—Cotrimoxazole  E—Erythromycin
Table 5  Antibiotics susceptibility profile of selected Gram-negative bacteria as read in mean diameter of zones of inhibition (mm) after treatment to electromagnetic flux at 5mT intensity compared with control

| Bacteria culture | CAZ | CIP | TET | IMP | AMP |
|------------------|-----|-----|-----|-----|-----|
| *S. marcescens* ATCC14756 | a) 26 ± 1.4 b) 15 ± 0.1 | a) 28 ± 2.8 b) 29 ± 0.1 | a) R b) R | a) 26 ± 1.4 b) 21 ± 2.8 | a) R |
| *P. aeruginosa* ATCC9027 | a) 26 ± 1.4 b) 16 ± 0.1 | a) 34 ± 3.5 b) 26 ± 8.5 | a) R b) R | a) 26 ± 1.4 b) 26 ± 0 | a) R |
| *G. oxydans* ATCC19357 | a) 27 ± 2.8 b) 18 ± 1.4 | a) 26 ± 0.7 b) 23 ± 3.5 | a) R b) 18 ± 1.4 | a) 26 ± 1.4 b) 28 ± 3.5 | b) R |
| *R. sphaeroides* ATCC17023 | a) 24 ± 0.7 b) R | a) 24 ± 0.7 b) 24 ± 0 | a) R b) R | a) R b) 30 ± 0.7 | b) R |
| *C. freundi* ATCC33128 | a) 14 ± 0.7 b) R | a) 26 ± 0.7 b) 26 ± 0 | a) 25 ± 0 b) 24 ± 0 | a) 29 ± 4.2 b) 30 ± 2.1 | b) R |
| *Y. pestis* ATCC11953 | a) R b) 19 ± 1.4 | a) 25 ± 0 b) 19 ± 1.4 | a) R b) 21 ± 1.4 | a) 23 ± 1.4 b) R | a) 15 ± 0 |
| *K. pneumoniae* ATCC25955 | a) 26 ± 1.4 b) 20 ± 0 | a) 31 ± 2.1 b) 19 ± 1.4 | a) R b) 19 ± 1.4 | a) 22 ± 2.8 b) 23 ± 1.4 | b) R |
| *A. aceti* ATCC15973 | a) 9 ± 4.2 b) R | a) 21 ± 9.9 b) 16 ± 0.7 | a) R b) 13 ± 4.2 b) 17 ± 1.4 | a) 16 ± 4.2 b) 23 ± 2.8 | b) 18 ± 0.7 |
| *E. coli* ATCC25922 | a) 29 ± 1.4 b) R | a) 22 ± 1.4 b) 18 ± 0.7 | a) 19 ± 0 b) 14 ± 2.1 | a) 33 ± 4.2 b) R | b) R |

Table 6  Antibiotics susceptibility profile for selected Gram-positive bacteria as read in mean diameter of zones of inhibition (mm) after treatment to electromagnetic flux at 15 mT intensity compared with control

| Bacteria | PEF | CN | APX | Z | AM | R | CPX | S | SXT | E |
|----------|-----|----|-----|--|----|--|-----|--|-----|---|
| *B. subtilis* ATCC6633 | a) 14 ± 1.4 b) 17 ± 3.5 | a) 16 ± 1.4 b) 12 ± 2.1 | a) 12 ± 1.4 b) 12 ± 1.4 | a) 12 ± 1.4 b) 12 ± 1.4 | a) 16 ± 2.1 b) 13 ± 0.7 | a) 12 ± 1.4 b) 12 ± 1.4 | a) 16 ± 1.4 b) 19 ± 0.7 | a) 18 ± 1.4 b) 21 ± 1.4 | a) 20 ± 2.8 b) 22 ± 2.8 | a) 12 ± 1.4 b) 14 ± 2.8 |
| *S. pyogenes* ATCC19613 | a) 14 ± 2.8 b) 20 ± 2.8 | a) 16 ± 1.4 b) 17 ± 2.1 | a) 12 ± 2.1 b) 13 ± 1.4 | a) 14 ± 1.4 b) 14 ± 1.4 | a) 16 ± 1.4 b) 16 ± 1.4 | a) 18 ± 1.4 b) 18 ± 1.4 | a) 18 ± 1.4 b) 18 ± 1.4 | a) 18 ± 1.4 b) 18 ± 1.4 | a) 18 ± 1.4 b) 18 ± 1.4 | a) 14 ± 2.8 |
| *S. aureus* ATCC25923 | a) 12 ± 1.4 b) 16 ± 6.7 | a) 14 ± 1.4 b) 12 ± 2.8 | a) R b) R | a) R b) R | a) 16 ± 2.1 b) 16 ± 2.1 | a) 16 ± 1.4 b) 18 ± 1.4 | a) 20 ± 1.4 b) 20 ± 0 | a) 18 ± 1.4 b) 18 ± 1.4 | a) 14 ± 2.8 b) 14 ± 1.4 | a) 14 ± 2.8 |

a) —Control b) —24 h c) —48 h d) —72 h R—Resistance PEF—Pefloxacin, CN—Gentamycin, APX—Ampicillin + Cloxacillin, Z—Cefuroxime, AM—Amoxicillin, R—Ceftriaxone, CPX—Ciprofloxacin, S—Streptomycin, SXT—Cotrimoxazole, E—Erythromycin

(after exposure to EMF) are reported to be largely due to the importance of the membrane potential of the bacterial cell wall, these waves can have an effect on the antibiotic susceptibility (Torgomyan et al. 2011). Furthermore, increased antibiotic susceptibility may be due to the interaction of electromagnetic fields on the water molecules in the cell. An increase in the permeability of the bacteria which might make the ion channels to be kept open in the bacterial wall, efflux pumps and ion channels in the cell wall, could also be playing an important role in the absorption of antibiotics. Interaction between the electromagnetic waves and the molecules of the antibiotics (such interaction affecting the Hydrophilicity or hydrophobicity of the antibiotic, antibiotic activity and antibiotic molecule size) thereby altering the charge distribution on antibiotics molecule has also been reported to affect the transportation of antibiotics across the cell membrane (Taheri et al. 2017). So far, the process of antibiotic resistance (with accelerated increase through various anthropogenic activities) seems to proceed unabated. Results of this study can further help in understanding the various mechanisms of antibiotic resistance thereby adding valuable angles in the development of effective antibiotic therapy for treatment of infectious diseases. Based on findings in this work, the roles of mechanosensitive channels present on bacterial membranes with respect to permissively or selectively admitting antibiotic molecules crossing the membrane barriers can be further studied. The effects of electromagnetic flux and sound waves demonstrated in this study could be incorporated in the evaluation of mechanosensitive channels as pharmacological targets for novel antibiotics design. It has recently been identified that the antibiotic streptomycin can act as an unlocking agent of
large mechanosensitive channels in bacteria, as the drug uses the channels as a route of entry into the cytoplasm (Blount and Iscla 2020). Further research has also identified the potential promiscuity of mechanosensitive channels in some species in allowing access to multiple antibiotics into the cytoplasm depending on the size and vibrational effect (Wray et al. 2019).

Earlier reports have identified bacteria to possess potentials to respond specifically to vibrational patterns that induce ionic flow and electrical charges on the membrane which most likely are unique to each species (Norris and Hyland 1997). It was shown that sound-induced extracellular vibrations could be detected as unique acoustic signals that influence intracellular vibrations linked to the movements of intracellular organelles like molecular motors, cytoskeletons, and chromosomal package in a specific pattern (Reguera 2011). Electromagnetic flux has also been identified as particularly important in modulating microbial metabolism in a method that can be detrimental or beneficial to cellular proliferation depending on the specifics of the flux frequency and intensity (Beretta et al. 2019). Specific examples with respect to Pseudomonas and Enterobacter species showed that zero magnetic fields induced antibiotic resistance in these bacteria in a magnetosensitive pattern (Creanga et al. 2004). This clearly means that understanding of the exact electromagnetic and sound frequencies that control membrane mechanosensation and intracellular vibrations can be utilised in species-specific microbial responses to sound and electromagnetic flux. To consolidate on the effects of sound, there have been practical observations of acoustic frequencies emitted by a B. subtilis strain detected at a range of 8–43 kHz (Matsumashi et al. 1998), while a lower range of 0.9–1.6 kHz was observed for Saccharomyces cerevisiae cells (Pelling et al. 2004). Apart from cellular vibrational patterns, it is understood that sound waves are propagated by mere vibration of particles within a medium (Reguera 2011), this therefore means that there can also be distinct sound patterns as observed within certain microbial ecological environments. These sound patterns could also be further studied as signals and frequencies and built upon to determine the specific responses of such bacteria to modulated sounds; a process that can be harnessed in designing therapeutics targeting sound-sensitivity in bacteria.
With the exposition of this knowledge, it could then be easier to design adjuvants or small sized frequency specific agonists that can be used to complement the existing antibacterial drugs in a synergistic fashion against target bacteria, especially drug resistant phenotypes. Data generated in this work can also potentially enrich aspects of sonobiology and sonogenetics for further studies of microbial membrane and intracellular behaviour. Studies involving recently applied biomolecular ultrasound techniques have elucidated species-specific acoustic modulation of bacterial membranes and intramolecular materials for higher resolution imaging and characterisation of intracellular activities (Maresca et al. 2018). Results from this work could be further expounded upon to provide a basis for deeper experimentation on variable species behaviour for improvement into direct sonar scanning and imaging of bacteria in the future.

Industrially, the information from this study can be useful for strain improvement of microorganisms for enhanced production of desired microbial metabolites where applicable. This current study can also be applied in advancing the techniques of sonoporation in bacteria, as a combined application of electromagnetic pulses and sound can be specifically made to target viable cells. Pores formed on the cells can be sites of DNA entry in transformation experiments, as well as points for seepage of viable intracellular metabolites of molecules (Song et al. 2007). These data could also be a basis for design of more effective sonobioreactors (Chisti 2003) and electrobioreactors (Mohtasham et al. 2016) for bioprocess optimisation. Ultimately this new research area should open a new frontier for multidisciplinary work at the interfaces of molecular biology, biochemistry and biophysics especially bioacoustics and sonobiology.

Despite the proper evaluations of potential application of the data in this study, it is important to note that, specific experiments linking measured cell membrane mechanosensation in line with the permissiveness to antibiotics as well as growth patterns were not fully elucidated here. We however, have extrapolated potential reasons explaining the behaviour of the bacterial cells under the influence of sound vibrations and electromagnetic flux. In line with this it is therefore necessary to further investigate mechanosensory biology, along with transcriptome and/or metabolomic profiling of sound and low electromagnetic stimulated cultures using different microorganisms in other to provide meaningful understanding in the area of cell-sound and cell-EMF interactions.

Conclusions
In this study, it has been shown that physical factors such as sound and electromagnetic flux (EMF) that abound in the environment interfere with the physiology of bacteria either by enhancing or inhibiting its growth as well as its response to antibiotics. This further point to the fact these factors can affect biomass accumulation and synthesis of intracellular molecules via a range of pathways, and that certain frequencies and amplitudes may favour some microbial species over others. This work also identifies the need for deeper investigations into the biomechanistic approaches microbial cells utilise in balancing membrane mechanosensitivity under the influence of sound and electromagnetic frequencies. The desired frequencies that could be applied in modulating cellular activities positively or negatively for applications in more efficient antimicrobial therapies and bioproduction processes are a major point of reference going forward.

Abbreviations
ATCC: American type culture collection; EMF: Electromagnetic field.

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Authors’ contributions
EOG and OMA designed the work, ANI-K collected the data and also prepared the manuscript together with EOG and OMA. EOG analysed some of the data. All authors read and approved the final manuscript.

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The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

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Competing interests
The authors declare that they have no competing interests.

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