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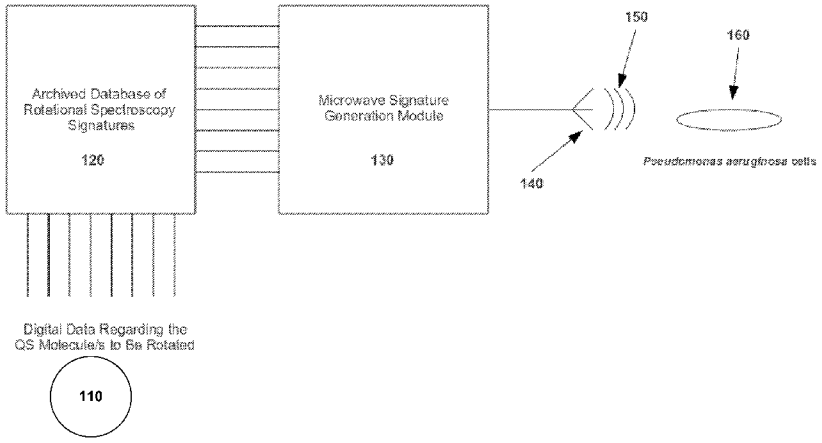
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(54) **Title**: SYSTEMS AND METHODS FOR MICROWAVE JAMMING OF MOLECULAR RECOGNITION



(57) **Abstract**: Systems and methods include directing limited frequencies of microwave signals toward target molecules, driving a motion of the target molecules to impact molecular recognition. In one implementation, a microwave spectra associated with the rotational modes of a target molecule is obtained. From peaks in the spectra, a mode of molecular movement is identified and a microwave signal profile is generated for driving a motion of binding portions of the molecule associated with the identified mode. Microwave signals are generated based on the signal profile for output by antennas. For example, the microwave signals can be can be used to impede the binding of quorum sensing molecules by receptors of P. aeruginosa. In one implementation, the antennas can be placed in a catheter for placement in a patient. In another implementation, the antennas can be placed in a sleeve or other device for use adjacent to the skin of a patient.

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SYSTEMS AND METHODS FOR MICROWAVE JAMMING OF MOLECULAR RECOGNITION

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority and the benefit under 35 U.S.C. § 119(e) of U.S. Provisional Patent Application No. 61/598,675, filed 14 February 2012, of which the entire contents and substance are hereby incorporated by reference as if fully set forth below.

BACKGROUND

[0002] The mechanism behind microwave spectroscopy is rooted in quantum mechanics, but can be understood from a classical perspective. For a molecule to exhibit a microwave spectrum it must have one or more dipole moments. In the presence of an electromagnetic wave the dipoles will be driven, resulting in the rotation of the molecule. The ubiquitous microwave oven is a kitchen appliance based on this principle. The electromagnetic field generated by the magnetron in the microwave oven induces the rotation of the H2O molecules in the food and this rotation results in a transfer of heat to the region surrounding each H2O molecule.

[0003] Microwave spectroscopy has been used as a means of identifying molecular content of far away planets. For example, the microwave spectra emitted by the molecules in the gaseous clouds of Jupiter are measured via radio telescopes on earth. The collected spectra are then analyzed and compared with the spectra of molecules measured in laboratories. Recently, there has been considerable work in identifying the microwave spectra of biologically relevant molecules such as DNA.

[0004] Another developing technological field is quorum sensing. As an evolutionary predecessor to the endocrine system, microbes communicate with one another by sending out small molecules. When these molecules are received by the other microbes in the community they trigger gene expression. For example, for a colony of *Pseudomonas aeruginosa* bacteria, the 3-oxo-C12 AHL ("C12") molecule signals that a bio-film should be constructed, in essence, a protective dome over the colony that can block chemical treatment.

[0005] So-called quorum-quenching mechanisms are currently being explored so as to minimize bio-film formation by microbes such as *P. aeruginosa*. Conventional approaches have been limited to antibodies which bind up the C12 molecules or drugs which are intended to do the

same. However, these conventional treatments are unable to pass through an established bio-film. Moreover, drug treatments can have deleterious side effects and undesirable systemic impact.

SUMMARY

[0006] Some or all of the above deficiencies may be addressed by certain implementations of the disclosed technology. Certain implementations include directing limited frequencies of microwave signals toward a target molecule, driving a motion of the target molecule to impact molecular recognition. For example, in some implementations, microwave signals can be used to impede the binding of the quorum sensing C12 molecule by receptors of *P. aeruginosa*. The relatively harmless energy levels for the prescribed microwave energy and the specificity of the microwave frequencies involved localize impact and reduce the risk of side effects.

[0007] According to an example implementation, a method for generating a microwave biological control signal is provided. The method can include obtaining a microwave spectrum associated with the rotational modes of a molecule. From one or more peaks in the spectrum, a mode of molecular movement can be identified. Then a microwave signal profile can be generated for driving a motion of one or more binding portions of the molecule associated with the identified mode. In a further implementation, microwave signals can be generated based on the signal profile. In another further implementation, the microwave signals can be output by one or more antennas. The method can also include identifying a molecule associated with a biological event.

[0008] According to an example implementation, system for generating a microwave control signal is provided. The system can include a processor and a memory operatively coupled to the processor. The memory can be configured for storing instructions that, when executed by the processor, cause the system to receive a microwave spectrum associated with rotational modes of a molecule, identify a mode of molecular movement associated with one or more peaks in the spectrum, and generate, by the processor, a microwave signal profile based on the identified mode for driving a motion of the molecule.

[0009] According to an example implementation, a computer-readable medium containing instructions that, when executed by at least one processor in a system, cause the system to perform a method including, receiving a microwave spectrum associated with the rotational

modes of a molecule. From one or more peaks in the spectrum, a mode of molecular movement can be identified. Then a microwave signal profile can be generated for driving a motion of one or more binding portions of the molecule associated with the identified mode. In a further implementation, microwave signals can be generated based on the signal profile. In another further implementation, the microwave signals can be output by one or more antennas.

[0010] Other implementations, features, and aspects of the disclosed technology are described in detail herein and are considered a part of the claimed disclosed technology. Other implementations, features, and aspects may be understood with reference to the following detailed description, accompanying drawings, and claims.

BRIEF DESCRIPTION OF THE FIGURES

[0011] Reference will now be made to the accompanying figures and flow diagrams, which are not necessarily drawn to scale, and wherein:

[0012] Fig. 1 depicts an illustration of a block diagram of the system, according to an example implementation.

[0013] Fig. 2A-B depict illustrations of the quorum sensing molecule, according to an example implementation.

[0014] Fig. 3 depicts an illustration of the rotational spectrum, according to an example implementation.

[0015] Fig. 4 depicts an illustration of a flow diagram of the method, according to an example implementation.

[0016] Fig. 5 depicts an illustration of a flow diagram of the method, according to an example implementation.

[0017] Fig. 6 depicts an illustration of a block diagram of the computing device system architecture, according to an example implementation.

[0018] Fig. L1 depicts an illustration of front, back, left, and right views of the linearly polarized patch antenna, according to an example implementation.

[0019] Fig. L2 depicts an illustration of top and bottom views of the linearly polarized patch antenna, according to an example implementation.

[0020] Fig. L3 depicts an illustration of a plot of S11 parameters for the linearly polarized patch antenna, according to an example implementation.

[0021] Fig. L4 depicts an illustration of a plot of realized gain for the linearly polarized antenna, according to an example implementation.

[0022] Fig. C1 depicts an illustration of front, back, left, and right views of the circularly polarized patch antenna, according to an example implementation.

[0023] Fig. C2 depicts an illustration of top and bottom views of the circularly polarized patch antenna, according to an example implementation.

[0024] Fig. C3 depicts an illustration of a plot of S11 parameters for the circularly polarized patch antenna, according to an example implementation.

[0025] Fig. C4 depicts an illustration of a plot of realized gain for the circularly polarized antenna, according to an example implementation.

DETAILED DESCRIPTION

[0026] The disclosed technology is based upon a radical concept — that a microwave signal of the proper frequency and character (e.g., polarization, temporal characteristics) can be used to alter the nature of biomolecular interactions.

[0027] In certain implementations, these microwave signals can cause target molecules to rotate billions of times per second and thus impede their binding. Such impedance will herein be referred to as microwave jamming, as in essence, certain implementations can provide a microwave signal which will jam cell-to-cell communication in an analogous fashion to which radar signals are jammed in electronic warfare. By jamming the microbe-to-microbe communication, certain implementations of the disclosed technology can impede the formation of bio-film by *P. aeruginosa*.

[0028] In describing example implementations, additional terminology will be resorted to for the sake of clarity. It is not intended that the disclosed technology be limited in scope to the details of construction and arrangement of components set forth in the following description or illustrated in the drawings. Rather, the disclosed technology is capable of other implementations and of being practiced or carried out in various ways.

[0029] Throughout the specification and the claims, the following terms take at least the meanings explicitly associated herein, unless the context clearly dictates otherwise. The term "connected" means that one function, feature, structure, or characteristic is directly joined to or in communication with another function, feature, structure, or characteristic. The term "coupled" means that one function, feature, structure, or characteristic is directly or indirectly joined to or in communication with another function, feature, structure, or characteristic. Relational terms such as "first" and "second," and the like may be used solely to distinguish one entity or action from another entity or action without necessarily requiring or implying any actual such relationship or order between such entities or actions. The term "or" is intended to mean an inclusive "or." Further, the terms "a," "an," and "the" are intended to mean one or more unless specified otherwise or clear from the context to be directed to a singular form. The term "include" and its various forms are intended to mean including but not limited to.

[0030] *Pseudomonas aeruginosa* is a life-shortening infection in the lungs of patients with Cystic Fibrosis (CF). *P. aeruginosa* sets up residence in healthy lungs as well as CF lungs. In CF patients, however, a genetic defect inhibits the body's ability to fight off the infection. Quorum sensing molecules transmitted between members of the microbial community are an indicator of what the microbes are up to. From a biological point of view, the quorum sensing molecules control gene regulation and inform the colony when to, among other things, replicate, sporulate, and when to construct a protective bio-film. Once the protective bio-film has been constructed, chemical treatment is unlikely to be effective in managing *P. aeruginosa* colonies.

[0031] At present there is considerable interest in the field of controlling bacterial infections by quenching their quorum sensing systems. Problematic in these approaches is the fact that conventional attempts to quench the quorum sensing systems involve chemical means. These techniques are futile once a biofilm has been constructed.

[0032] Certain implementations of the disclosed molecular jamming techniques can be effective against even microbial colonies with established bio-films. Although chemicals may not penetrate a bio-film, microwave energy can. For example, a typical bio-film thickness can be approximately 100μm. Assuming the conductivity of the biofilm to be roughly equal to that of seawater, at 10 GHz, an example electromagnetic signal may be reduced by only 0.073dB in propagating through the bio-film. Moreover, as certain implementations include placing a

microwave antenna in close proximity to the offending microbial colony, there can be very little loss of electromagnetic energy during propagation through the bio-film.

[0033] Virtually every biomolecular interaction involves molecular recognition as a critical component — disrupt molecular recognition and a cell process has been altered or stopped. Although this description details an exemplary application of microwave jamming for inhibiting bio-film formation among *P. aeruginosa* colonies, the techniques disclosed herein have a wide variety of applications.

[0034] For example, and not limitation, certain implementations of the disclosed technology can be applied to mold treatment. Currently, the accepted method for eradicating mold from a home is to use chemicals to kill the microbial communities associated with the mold. Using the disclosed technology, microwave energy can be transmitted at the proper frequency to jam the communication within the mold microbial community and ultimately resulting in cell death.

[0035] Another potential application is the supplement or replacement of herbicides and insecticides with microwave jamming. In an example implementation, rather than genetically engineer plants, such as soy beans, to be unaffected by highly toxic herbicides, non-genetically engineered soy beans could be planted and microwave transmitters placed in the crop field with the soy beans. In another example implementation, an unmanned air vehicle (UAV) equipped with a microwave transmitter over the field. In this example implementation and others, alternate means of radiation delivery will be apparent to one of skill in the art.

[0036] Yet another potential application of this technology is for use in cancer treatment. In an example implementation, a microwave signal or sequence of signals can be targeted at cancer cell DNA. Sustained rotational stressing on the DNA from the microwave energy can cause the cancer cell's DNA to fracture, similar to the mechanism behind some chemotherapy drugs. For localized treatment, an antenna on a catheter can be fed to a location near the tumor and the microwave signal transmitted for sufficiently long to weaken or kill the cancer cells. Because microwave energy at certain frequencies (e.g. 10 GHz) will not propagate very far in tissue, the microwave jamming signal is likely to destroy the tumor with reduced or nominal collateral damage to healthy surrounding tissue. In another example implementation, targeted microwave radiation can be used to block cell-to-cell communication for angiogenesis.

[0037] Other and additional applications of the disclosed technology are contemplated and within the scope of this disclosure.

[0038] Referring now to the figures, in which like reference numerals represent like parts throughout the views, various implementations of the disclosed technology will be described in detail.

[0039] In an example implementation, the disclosed technology can be used to treat infections of *P. aeruginosa*. **Fig. 1** depicts an illustration of a block diagram of the system, according to an example implementation. As shown in **Fig. 1**, a digital encoding **110** of the quorum sensing molecule C12 can be input to a database of rotational spectroscopy signatures **120** to obtain the appropriate microwave spectra. The signature retrieved from the database can be sent to the Microwave Signature Generation Module (MSGM) **130**. The output of the MSGM can drive an antenna **140** for outputting microwave signals.

[0040] Electromagnetic waves **150** output by the antenna **140** can propagate towards the colony of *P. aeruginosa* cells **160**, causing the C12 molecules within the colony to rotate at the frequencies of the microwave signal. Accordingly, a microwave signal in the GHz range can cause a molecule to rotate billions of times per second, thus impeding molecular recognition and blocking the message to form a bio-film. In a further implementation, a CF patient could be treated using a catheter containing the antenna. The catheter could be threaded into the lungs and positioned proximal to the location of the infection.

[0041] Microwave jamming of the quorum sensing communication system can be used in conjunction with conventional treatments unable to penetrate the bio-film such as antibiotics. Moreover, in some implementations, microwave jamming of cell-to-cell communication can lead directly to cell death. Thus, certain implementations of the disclosed technology can provide a treatment methodology which requires no drugs — only microwave energy of the correct frequencies — ushering a new class of targeted microwave pharmaceuticals.

[0042] Certain implementations of the disclosed technology are generally applicable to moderating molecular recognition in any biological interaction. **Fig. 5** depicts an illustration of a flow diagram of the method, according to an example implementation. The method **500** starts in block **502**, and according to an example implementation, includes identifying a molecule associated with a biological event. The biological event can be any event in which molecular

recognition, i.e., binding of the molecule, is implicated. For example, the biological event could include conversion of a substrate or the reuptake of a neurotransmitter. Moreover, the biological event can be a complex or multi-stage process involving the binding of several molecules, as in angiogenesis or apoptosis.

[0043] In certain implementations, the target molecule can be a quorum sensing molecule, such as C12 for instructing *P. aeruginosa* communities to form a bio-film, or any of various other molecules biologically interactive molecules, including, without limitation, ligands, substrates, hormones, neurotransmitters, cytokines, growth factors, toxins, sugars, etc. Moreover, in some implementations, the target molecule may moderate molecular recognition of a second molecule, for example, as with an enzyme inhibitor or activator.

[0044] Target molecules exhibiting one or more dipole moments can effectively respond to the microwave signals. For example, polar molecules have a permanent dipole moment, and thus can be suitable targets for implementations of the disclosed technology. However, in certain implementations, a dipole moment can be induced in a molecule that would otherwise be unresponsive to targeted microwave signals, for example by application of an electric field.

[0045] **Figs. 2A-B** depict illustrations of the quorum sensing molecule, C12 **200**, according to an example implementation. As shown in **Fig. 2A**, the elements hydrogen (H) **220,** oxygen (O) **230**, nitrogen (N) **240**, and carbon (C) **250** are represented by spheres of various sizes. In this molecule, the presence of the uneven charge distribution of the electron clouds leads to a net dipole moment **260**. There are other weaker dipole moments in the molecule related to the hydrogen bonding between the H atoms and the N and O atoms. The complexity of these dipoles and attendant quadrupole, as well as the molecule's moment of inertia tensor gives rise to the rich microwave spectra which can be specific to a particular conformer of a molecule.

[0046] In block **504**, the method **500** includes obtaining a microwave spectra associated with the rotational modes of the molecule. These spectral frequencies can be very narrow in bandwidth. Thus, the identification of the rotational resonances can help ensure that the microwave energy to be radiated into a targeted collection of cells will substantially affect only the targeted molecule of interest.

[0047] In certain implementations, the microwave spectra can be measured, using, for example, microwave spectroscopy. For astronomical spectroscopic studies, the thermal motion of the

molecules in a gas leads to rotation of these molecules which in turn generates the microwave spectra observed. In microwave spectroscopic techniques such as laser ablation-molecular beam Fourier Transform microwave (LA-MB-FTMW) spectroscopy, attenuation of specific frequencies is looked for through a carefully constructed gas of the target molecule surrounded by a select number of H2O molecules.

[0048] In another implementation, spectra can be obtained by modeling the molecule and analytically deriving the spectra. These calculations can consider the molecule of interest, e.g. C12, with the proper number of H2O molecules which will surround it in its biological environment. The modeled spectra can closely approximate experimentally measured spectra in some cases. For example, **Fig. 4** depicts an illustration of an experimentally measured spectrum **462** and theoretically derived spectrum **463** for a an example serine conformer **461**. There are a host of preexisting programs that can be used to predict the rotational resonances of a molecule, for example, the open source program, GAMESS, and the commercially available program, Gaussian09.

[0049] In certain implementations, an indication or signature of the spectra can be stored in or retrieved from a database. Accordingly, a previously measured or derived spectra for the target molecule can be obtained from the database, obviating the need to re-analyze the rotational spectra of the target molecule for each treatment session. For example, as shown in **Fig. 1**, a digital encoding **110** of the target molecule can be generated and input to an archived database of rotational spectroscopy signatures **120**. The database can provide the rotational spectra for the target molecule, in accordance with block **504**.

[0050] **Fig. 3** depicts an illustration of the rotational spectra **300** for C12, according to an example implementation. As shown in **Fig. 3**, the rotational frequencies of these example spectra are illustrated by the peaks **320**, **330** and **340**. These frequency ranges corresponding to these peaks can be substantially unique to the target molecule — so much so that different conformers of the same molecule can have distinct spectra.

[0051] In this example, the rotational spectra appear sufficiently unique and narrow band that they can be used to discriminate between conformers of the same molecule. Accordingly, it is unlikely that this signal, which drives the rotational modes of the molecule C12, will cause other molecules to rotate. (However, a cautious approach could also consider the spectra of other

molecules.) Moreover, microwave energy is strongly attenuated in organic tissue as, from an electromagnetic standpoint, tissue is very similar to sea water. Thus a catheter with an antenna for outputting targeted microwave signals that is introduced into the lungs of a CF patient may only affect the cells nearby. Hence, systemic effects can be reduced or minimized, in contrast to treatment with chemicals.

[0052] In block **506**, the method **500** includes identifying a mode of molecular movement associated with one or more peaks in the spectra. In certain implementations, this can include relating one or more peaks in the spectra to a mode of vibration of one or more moieties, or functional groups, of the molecule. By driving the motion of moieties that include binding portions, the rate of binding of the target molecule can be affected. For example, certain implementations can drive the rotation of the molecule C12 by irradiating the sample with microwave energy at the appropriate frequencies gleaned from molecular modeling and an experimental technique, such as LA-MB-FTMW.

[0053] In certain implementations, the induced motion can impede the binding of the target molecule. However, targeted microwave signals can be designed to induce molecular motion to achieve other effects. In various other implementations, the induced motion can promote binding of the target molecule, promote unbinding of the molecule, or trigger release of the molecule, for example, from a receptor. In yet another implementation, the induced motion can cause the target molecule, itself, to fracture from physical stress.

[0054] In block **508**, the method **500** includes identifying one or more binding portions associated with identified mode. In certain implementations, a binding portion can be all or a portion of an active site.

[0055] In block **510**, the method **500** includes generating, using a processor, a microwave signal profile for driving a motion of the one or more binding portions. In certain implementations, one or more microwave signals can be generated based on the signal profile. In another implementation, the generated signal profile can be stored, for example, on a database for later retrieval and use.

[0056] In certain implementations, blocks **506** to **510** can be performed by the MSGM **130**. In an example implementation, the MSGM can select a subset of the rotational spectroscopy signal which is most likely to be unique to the target molecule so as not to induce incidental rotation of

other molecules. Moreover, the MSGM can prescribe other microwave signal characteristics to achieve the desired motion and corresponding effect. One of skill in the art will be able to select microwave signal characteristics for achieving the desired effect.

[0057] For example, a basic microwave signal profile may prescribe outputting a fixed frequency microwave signal within a range corresponding to the one or more peaks in the rotational spectra. However, in some implementations, a more complicated signal profile can be used to exert greater control over the motion of a target molecule. In certain implementations, a microwave signal can be output that sweeps or chirps between frequencies within a frequency range corresponding to the one or more peaks of the spectra. In another example implementation, the frequency of the first microwave signal can "hop" between frequency ranges corresponding to the first peak and the second peak.

[0058] According to another example implementation, the microwave signal can be a substantially continuous signal. In another implementation, the microwave signal can be a discrete or pulsed signal.

[0059] According to another example implementation, the microwave signal can have a substantially narrow bandwidth, for example less than about 100 MHz. Narrowing the bandwidth can reduce side effects by reducing the possibility of exciting molecules other than the target molecule. In another implementation, the bandwidth of the microwave signal can be constrained so as not overlap frequency ranges corresponding to more than one peak of the associated spectra.

[0060] According to another example implementation, the amount of power of the microwave signal can be less than a microwatt. However, especially if significant attenuation is expected, the power may be increased such that the power received at the target molecule meets a certain threshold. In one implementation, the total amount of power of all microwave signals output can be less than a microwatt.

[0061] According to another example implementation, a microwave signal profile can prescribe outputting multiple microwave signals of the various qualities described above. In one implementation, the microwave signals can be a sequence of microwave signals. In another implementation, the microwave signals can be output concurrently.

[0062] In certain implementations, the generated microwave signals **150** can be output from one or more antennas **140**. The antennas can be designed and uniquely constructed for the frequency range of the desired rotational spectra and for efficiency in near-field generation. Moreover, the antennas can be constructed to emanate radiation with one or more of elliptical, linear, or circular polarizations. One of skill in the art will be able to select antenna properties to achieve the desired effect.

[0063] According to an example implementation, the antenna can be placed inside a catheter or similar device. The catheter can be placed inside a patient substantially proximal to a microbial infection or potential infection site. For example, a catheter could be threaded into the lungs of a CF patient. In another example implementation, the antenna can be placed in a sleeve, patch, or other suitable construct for placement or attachment substantially adjacent to the skin of a patient. For example, the antenna could be placed next to a burn site on a patient to manage infection. In another example implementation, a device containing the antenna could be surgically implanted, and coupled to an internal or external signal generation means. In yet another example implementation, a device containing the antenna could be ingested. Moreover, in another example implementation, the antenna could be incorporated into a device for cleansing. For example, targeted microwaves could be used to manage bio-film formation on contact lenses or other medical devices.

[0064] Two example antennas were designed and constructed to output radiation at 10.2 GHz, one of the important frequencies in the rotational spectra of C12, as shown in **Fig. 3**. The antennas were of two types, linearly polarized and circularly polarized antenna. The two example antennas are now described in detail.

[0065] A single-element linearly polarized 10.2 GHz antenna **400** was designed to create a template antenna that could be phased to produce a high-gain antenna. **Figs. L1-L2** depict illustrations of front **401** back **402** left **403** right **404** bottom and top views of the linearly polarized antenna **400**, according to an example implementation. The example antenna as shown in **Figs. L1-L2** can be used as a building-block to create 2-element, 8-element, and 16-element antenna arrays. The antenna arrays can be phased a half-wavelength apart to provide for constructive interference. This constructive interference can provide radiation with greater

intensity. If the radiation intensity is too low with the 1-element array, higher-element arrays can be used to provide greater exposure to 10.2 GHz radiation.

[0066] The example linearly polarized antenna **400** is a 10.2 GHz microstrip patch antenna. The patch **410** is 5.10 mm in length and 9.03 mm in width. The substrate **430** height is 1.524 mm and is made of FR4 substrate which has a relative permittivity of 4.3. The impedance of the antenna **400** is approximately 238.5 ohms. In order to match the antenna **470 400** to a 50 ohm transmission line **490**, a quarter wave transformer **470** was used. The quarter wave transformer **470** has a length of 7.35 mm and a width of .535 mm. The .535 mm width gives the quarter wave transformer an impedance of ~ 109 ohms. A 50 ohm microstrip transmission line **490** is connected to the 109 ohm quarter wave transformer **470**. The 50 ohm microstrip **490** line has a width of 2.97 mm and a length of 4.0 mm. Table 1 lists additional dimensions and specifications of the example linearly polarized antenna **400**.

[0067] The connection of the patch **410**, 109 ohm transmission line, and 50 ohm transmission line **490** is relatively reflection-less at 10.2 GHz, as illustrated in the S11 graphed in **Fig. L3**. As shown in **Fig. L3**, the antenna **400** is resonant at 10.229 GHz **310** and has a usable frequency range of approximately 9.7 GHz to 10.7 GHz.

[0068] **Fig. L4** depicts an illustration of a plot of realized gain for the linearly polarized antenna **400**, according to an example implementation. **Fig. L4** shows the azimuthal farfield realized gain pattern for the antenna. The antenna **400** has a realized gain of 5.1 dBi at 10.2 GHz.

Table 1. Dimensions of the 10.2 GHz 1-element Patch Antenna **400**

Reference	Parameter	Value
410	**Patch**	
	Patch Length	5.10 mm
	Patch Width	9.03 mm
	Patch Thickness	.038 mm
	Patch Material	Copper
430	**Substrate**	
	Substrate Thickness	1.524 mm
	Substrate Width	13.55 mm
	Substrate Length	13.55 mm
	Substrate Relative Permittivity	4.3
450	**Ground Plane**	
	Ground Plane Length	13.55 mm

	Ground Plane Width	13.55 mm
	Ground Plane Thickness	.038 mm
	Ground Plane Material	Copper
470	**Quarter Wavelength Transformer**	
	Quarter Wavelength Transformer Length	7.35 mm
	Quarter Wavelength Transformer Width	.535 mm
	Quarter Wavelength Transformer Thickness	.038 mm
	Quarter Wavelength Transformer Material	Copper
490	**50 Ohm Transmission Line**	
	50 Ohm Transmission Line Length	4.0 mm
	50 Ohm Transmission Line Width	2.97 mm
	50 Ohm Transmission Line Thickness	.038 mm
	50 Ohm Transmission Line Material	Copper

[0069] A circularly polarized patch antenna **800** was also designed and produced. **Figs. C1-C2** depict illustrations of views of the circularly polarized antenna **800**, according to an example implementation. One drawback of a linearly polarized antenna is in the linearity of is electric field. An electric field that is perpendicular to a molecule (or dipole moment) will not produce any induced EMF. In order to increase the induced EMF of the target molecule, a circularly polarized antenna **800** was designed. A circularly polarized antenna can create an electromagnetic field that has components in both the X direction and the Y direction, assuming the wave is travelling in the Z direction. Since the circularly polarized antenna contains equal E-field components in both the X and the Y direction, molecules that are possibly tangential to the X portion of the E-field will still be excited by Y-portion of the E-field. This notion can be of great importance when trying to excite molecules that are floating in an aqueous solution. The orientation of the molecules in the aqueous solution can be random. As such, using a circularly polarized electromagnetic field can raise the probability of exciting a molecule in the solution. The specifications for the 10.2 GHz circularly polarized antenna **800** are as follows:

[0070] The antenna **800** is a 10.2 GHz circularly polarized microstrip patch antenna. The patch **810** is 5.10 mm in length and 9.03 mm in width. The substrate **830** height is 1.524 mm and is made of FR4 substrate which has a relative permittivity of 4.3. There are cut-outs **820** in the upper left and bottom right corners. The cut-outs **820** are at approximately 45 degree angles to

the corner and form a 45, 45, 90 degree triangle with side lengths of 1.12 mm. The cut-outs can create the circular polarizing experienced in the farfield radiation.

[0071] The impedance of the antenna **800** is approximately 238.5 ohms. In order to match the antenna to a 50 ohm transmission line **890**, a quarter wave transformer **870** was used. The quarter wave transformer **870** has a length of 7.35 mm and a width of .535 mm. The .535 mm width gives the quarter wave transformer **870** an impedance of ~ 109 ohms. A 50 ohm microstrip transmission line **890** is connected to the 109 ohm quarter wave transformer. The 50 ohm microstrip line **89-** has a width of 2.97 mm and a length of 4.0 mm. Table 2 lists additional dimensions and specifications of the example circularly polarized antenna.

[0072] The connection of the patch **810**, 109 ohm transmission line, and 50 ohm transmission line **890** is relatively reflection-less at 10.2 GHz. This notion is illustrated in the S11 graphed in **Fig. C3**. **Fig. C3** depicts an illustration of a plot of S11 parameters for the circularly polarized patch antenna, according to an example implementation. As shown in **Fig. C3**, the example antenna is resonant at 10.342 GHz **710** and has a usable frequency range of approximately 10 GHz to 10.7 GHz

[0073] **Fig. C4** depicts an illustration of a plot of realized gain for the circularly polarized antenna **800**, according to an example implementation. **Fig. C4** shows the azimuthal farfield realized gain pattern for the antenna. The antenna **800** has a realized gain of 4.1 dBi at 10.2 GHz.

Table 2. Dimensions of the 10.2 GHz Circularly Polarized Antenna **800**

Reference	Parameter	Value
810	**Patch**	
	Patch Length	5.10 mm
	Patch Width	9.03 mm
	Patch Thickness	.038 mm
	Patch Material	45 degrees
820	**Cut Out**	
	Cut Out Width	1.12 mm
	Cut Out Length	1.12 mm
	Cut Out Angles	Copper
830	**Substrate**	
	Substrate Thickness	1.524 mm

	Substrate Width	13.55 mm
	Substrate Length	13.55 mm
	Substrate Relative Permittivity	4.3
850	**Ground Plane**	
	Ground Plane Length	13.55 mm
	Ground Plane Width	13.55 mm
	Ground Plane Thickness	.038 mm
	Ground Plane Material	Copper
870	**Quarter Wavelength Transformer**	
	Quarter Wavelength Transformer Length	7.35 mm
	Quarter Wavelength Transformer Width	.535 mm
	Quarter Wavelength Transformer Thickness	.038 mm
	Quarter Wavelength Transformer Material	Copper
890	**50 Ohm Transmission Line**	
	50 Ohm Transmission Line Length	4.0 mm
	50 Ohm Transmission Line Width	2.97 mm
	50 Ohm Transmission Line Thickness	.038 mm
	50 Ohm Transmission Line Material	Copper

[0074] Various implementations of the disclosed technology may be embodied all or in portion in non-transitory computer readable media for execution by a processor. An example implementation may be used in an application of a medical device, but other computing devices.

[0075] **Fig. 6** depicts a block diagram of an illustrative computer system architecture **600** according to an example implementation. Certain aspects of **Fig. 6** may be embodied in a computing device. It will be understood that the architecture **600** is provided for example purposes only and does not limit the scope of the various implementations of the communication systems and methods.

[0076] The architecture **600** of **Fig. 6** includes a central processing unit (CPU) **602**, where computer instructions are processed; a display interface **604** that acts as a communication interface and provides functions for rendering video, graphics, images, and texts on the display; a keyboard interface **606** that provides a communication interface to a keyboard; and a pointing device interface **608** that provides a communication interface to a pointing device, e.g., a touchscreen or presence-sensitive screen. Example implementations of the architecture **600** may include an antenna interface **610** that provides a communication interface to an antenna.

Example implementations may include a connection interface **612**. The connection interface may include one or more of a peripheral connection interface and network communication interface, providing a communication interface to an external device or network. In certain implementations, a camera interface **614** may be provided that acts as a communication interface and provides functions for capturing digital images from a camera. In certain implementations, a sound interface **616** may be provided as a communication interface for converting sound into electrical signals using a microphone and for converting electrical signals into sound using a speaker. According to example implementations, a random access memory (RAM) **618** may be provided, where computer instructions and data may be stored in a volatile memory device for processing by the CPU **602**.

[0077] According to an example implementation, the architecture **600** may include a read-only memory (ROM) **620** where invariant low-level system code or data for basic system functions such as basic input and output (I/O), startup, or reception of keystrokes from a keyboard are stored in a non-volatile memory device. According to an example implementation, the architecture **600** may include a storage medium **622** or other suitable type of memory (e.g. such as RAM, ROM, programmable read-only memory (PROM), erasable programmable read-only memory (EPROM), electrically erasable programmable read-only memory (EEPROM), magnetic disks, optical disks, floppy disks, hard disks, removable cartridges, flash drives), where the files include an operating system **624**, application programs **626** (including, for example, a web browser application, a widget or gadget engine, and or other applications, as necessary) and data files **628** are stored. According to an example implementation, the architecture **600** may include a power source **630** that provides an appropriate alternating current (AC) or direct current (DC) to power components. According to an example implementation, the architecture **600** may include a telephony subsystem **632** that allows the device **600** to transmit and receive sound over a telephone network. The constituent devices and the CPU **602** may communicate with each other over a bus **634**.

[0078] In accordance with an example implementation, the CPU **602** may have appropriate structure to be a computer processor. In one arrangement, the computer CPU **602** may include more than one processing unit. The RAM **618** may interface with the computer bus **634** to provide quick RAM storage to the CPU **602** during the execution of computing programs such as the operating system application programs, and device drivers. More specifically, the CPU **602**

may load computer-executable process steps from the storage medium **622** or other media into a field of the RAM **618** in order to execute computing programs. Data may be stored in the RAM **618**, where the data may be accessed by the computer CPU **602** during execution. In one example configuration, the device **600** may include at least 128 MB of RAM, and 256 MB of flash memory.

[0079] The storage medium **622** itself may include a number of physical drive units, such as a redundant array of independent disks (RAID), a floppy disk drive, a flash memory, a USB flash drive, an external hard disk drive, thumb drive, pen drive, key drive, a High-Density Digital Versatile Disc (HD-DVD) optical disc drive, an internal hard disk drive, a Blu-Ray optical disc drive, or a Holographic Digital Data Storage (HDDS) optical disc drive, an external mini-dual in-line memory module (DIMM) synchronous dynamic random access memory (SDRAM), or an external micro-DIMM SDRAM. Such computer readable storage media may allow the device **600** to access computer-executable process steps, application programs and the like, stored on removable and non-removable memory media, to off-load data from the device **600** or to upload data onto the device **600**. A computer program product, such as one utilizing a communication system may be tangibly embodied in storage medium **622**, which may comprise a machine-readable storage medium.

[0080] In an example implementation of the disclosed technology, the computing system architecture **600** may include any number of hardware and/or software applications **221**, **222** that are executed to facilitate any of the operations. In an example implementation, one or more I/O interfaces may facilitate communication between the computing system architecture **600** and one or more input/output devices. For example, a universal serial bus port, a serial port, a disk drive, a CD-ROM drive, and/or one or more user interface devices, such as a display, keyboard, keypad, mouse, control panel, touchscreen display, microphone, etc., may facilitate user interaction with the computing system architecture **600**. The one or more I/O interfaces may be utilized to receive or collect data and/or user instructions from a wide variety of input devices. Received data may be processed by one or more computer processors as desired in various implementations of the disclosed technology and/or stored in one or more memory devices.

[0081] One or more network interfaces may facilitate connection of the computing system architecture **600** inputs and outputs to one or more suitable networks and/or connections; for

example, the connections that facilitate communication with any number of sensors associated with the system. The one or more network interfaces may further facilitate connection to one or more suitable networks; for example, a local area network, a wide area network, the Internet, a cellular network, a radio frequency network, a Bluetooth enabled network, a Wi-Fi enabled network, a satellite-based network any wired network, any wireless network, a proximity network, etc., for communication with external devices and/or systems. As desired, implementations of the disclosed technology may include the computing system architecture **600** with more or less of the components illustrated in **Fig. 6**.

[0082] Certain implementations of the disclosed technology are described above with reference to block and flow diagrams of systems and methods and/or computer program products according to example implementations of the disclosed technology. It will be understood that one or more blocks of the block diagrams and flow diagrams, and combinations of blocks in the block diagrams and flow diagrams, respectively, may be implemented by computer-executable program instructions. Likewise, some blocks of the block diagrams and flow diagrams may not necessarily need to be performed in the order presented, or may not necessarily need to be performed at all, according to some implementations of the disclosed technology.

[0083] These computer-executable program instructions may be loaded onto a general-purpose computer, a special-purpose computer, a processor, or other programmable data processing apparatus to produce a particular machine, such that the instructions that execute on the computer, processor, or other programmable data processing apparatus create means for implementing one or more functions specified in the flow diagram block or blocks. These computer program instructions may also be stored in a computer-readable memory that may direct a computer or other programmable data processing apparatus to function in a particular manner, such that the instructions stored in the computer-readable memory produce an article of manufacture including instruction means that implement one or more functions specified in the flow diagram block or blocks. As an example, implementations of the disclosed technology may provide for a computer program product, comprising a computer-usable medium having a computer-readable program code or program instructions embodied therein, said computer-readable program code adapted to be executed to implement one or more functions specified in the flow diagram block or blocks. The computer program instructions may also be loaded onto a computer or other programmable data processing apparatus to cause a series of operational

elements or steps to be performed on the computer or other programmable apparatus to produce a computer-implemented process such that the instructions that execute on the computer or other programmable apparatus provide elements or steps for implementing the functions specified in the flow diagram block or blocks.

[0084] Accordingly, blocks of the block diagrams and flow diagrams support combinations of means for performing the specified functions, combinations of elements or steps for performing the specified functions and program instruction means for performing the specified functions. It will also be understood that each block of the block diagrams and flow diagrams, and combinations of blocks in the block diagrams and flow diagrams, may be implemented by special-purpose, hardware-based computer systems that perform the specified functions, elements or steps, or combinations of special-purpose hardware and computer instructions.

[0085] While certain implementations of the disclosed technology have been described in connection with what is presently considered to be the most practical and various implementations, it is to be understood that the disclosed technology is not to be limited to the disclosed implementations, but on the contrary, is intended to cover various modifications and equivalent arrangements included within the scope of the appended claims. Although specific terms are employed herein, they are used in a generic and descriptive sense only and not for purposes of limitation.

[0086] This written description uses examples to disclose certain implementations of the disclosed technology, including the best mode, and also to enable any person skilled in the art to practice certain implementations of the disclosed technology, including making and using any devices or systems and performing any incorporated methods. The patentable scope of certain implementations of the disclosed technology is defined in the claims, and may include other examples that occur to those skilled in the art. Such other examples are intended to be within the scope of the claims if they have structural elements that do not differ from the literal language of the claims, or if they include equivalent structural elements with insubstantial differences from the literal language of the claims.

CLAIMS

What is claimed is:

1. A method for generating a microwave biological control signal comprising:

identifying a molecule associated with a biological event;

obtaining a microwave spectrum associated with rotational modes of the molecule;

identifying a mode of molecular movement associated with one or more peaks in the spectrum;

identifying one or more binding molecule portions associated with the identified mode; and

generating, by a processor, a microwave signal profile for driving a motion of the one or more binding molecule portions.

2. The method of claim 1, wherein the molecule is polar.

3. The method of claim 1 further comprising inducing a dipole moment in the molecule.

4. The method of claim 1, wherein the molecule has an induced dipole moment.

5. The method of claim 1, wherein the molecule is a quorum sensing molecule.

6. The method of claim 1, wherein the molecule is a ligand.

7. The method of claim 1, wherein the molecule is an inducer.

8. The method of claim 1, wherein the molecule is an activator.

9. The method of claim 1, wherein the molecule is an enzyme substrate.

10. The method of claim 1, wherein the molecule is an enzyme activator.

11. The method of claim 1, wherein the molecule is an enzyme inhibitor.

12. The method of claim 1, wherein the molecule is a hormone.

13. The method of claim 1, wherein the molecule is a neurotransmitter.

14. The method of claim 1, wherein the molecule is a cytokine.

15. The method of claim 1, wherein the molecule is a growth factor.

16. The method of claim 1, wherein the molecule is a toxin.

17. The method of claim 1, wherein the molecule is a sugar.

18. The method of claim 1, wherein the biological event comprises binding of the molecule.

19. The method of claim 1, wherein the biological event comprises conversion of a substrate.

20. The method of claim 1, wherein the biological event comprises neurotransmitter re-uptake.

21. The method of claim 1, wherein the biological event comprises apoptosis.

22. The method of claim 1, wherein the biological event comprises angiogenesis.

23. The method of claim 1, wherein obtaining the microwave spectrum comprises measuring the spectrum using spectroscopy.

24. The method of claim 1, wherein obtaining the microwave spectrum comprises modeling the molecule and analytically deriving the spectrum.

25. The method of claim 1, wherein obtaining the microwave spectrum comprises receiving the spectrum from a database.

26. The method of claim 1, wherein identifying the mode of molecular movement comprises relating the one or more peaks to a mode of vibration of one or more moieties of the molecule and wherein the one or more binding portions of the molecule comprises at least part of the one or more moieties.

27. The method of claim 1, wherein identifying the mode of molecular movement comprises relating the one or more peaks to a mode of vibration of one or more functional groups of the molecule and wherein the one or more binding portions of the molecule comprises at least part of the one or more functional groups.

28. The method of claim 1, wherein the motion impedes binding of the molecule.

29. The method of claim 1, wherein the motion promotes binding of the molecule.

30. The method of claim 1, wherein the motion promotes the unbinding of the molecule.

31. The method of claim 1, wherein the motion triggers release of the molecule.

32. The method of claim 1, wherein the motion comprises a rotation of one or more portions of the molecule.

33. The method of claim 1, wherein the motion comprises a vibration of one or more portions of the molecule.

34. The method of claim 1 further comprising storing the signal profile in a database.

35. The method of claim 1 further comprising generating one or more microwave signals based on the signal profile.

36. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a fixed frequency signal within a frequency range corresponding to the one or more peaks.

37. The method of claim 35, wherein the one or more peaks comprises a first peak and a second peak and the one or more microwave signals includes a first microwave signal, the frequency of the first microwave signal hopping between frequency ranges corresponding to the first peak and the second peak.

38. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal sweeping between frequencies within a frequency range corresponding to the one or more peaks.

39. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a substantially continuous signal.

40. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a pulsed signal.

41. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a frequency bandwidth coinciding with only one peak of the spectrum.

42. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a frequency bandwidth of less than 101 MHz.

43. The method of claim 35, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a narrow frequency bandwidth.

44. The method of claim 35, wherein the one or more microwave signals is a sequence of microwave signals.

45. The method of claim 35, wherein the one or more microwave signals are concurrent microwave signals.

46. The method of claim 35, wherein the power of each of the generated signals is less than one microwatt.

47. The method of claim 35, wherein the total power of the generated signals is less than one microwatt.

48. The method of claim 35, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having an elliptical polarization.

49. The method of claim 35, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having a circular polarization.

50. The method of claim 35, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having a linear polarization.

51. The method of claim 35 further comprising outputting the one or more generated signals from one or more antennas.

52. The method of claim 51, further comprising placing a catheter including at least a portion of the one or more antennas into a patient.

53. The method of claim 51, further comprising positioning at least a portion of the one or more antennas substantially adjacent to the skin of a patient.

54. The method of claim 51, further comprising placing a sleeve including at least a portion of the one or more antennas substantially adjacent to the skin of a patient.

55. The method of claim 51, further comprising submerging at least a portion of the one or more antennas.

56. A system for generating a microwave control signal comprising:

at least one processor;

at least one memory operatively coupled to the at least one processor and configured for storing data and instructions that, when executed by the at least one processor, cause the system to:

receive a microwave spectrum associated with rotational modes of a molecule;

identify a mode of molecular movement associated with one or more peaks in the spectrum;

generate, by the processor, a microwave signal profile based on the identified mode for driving a motion of the molecule.

57. The system of claim 56, wherein the molecule is polar.

58. The system of claim 56 further comprising inducing a dipole moment in the molecule.

59. The system of claim 56, wherein the molecule has an induced dipole moment.

60. The system of claim 56, wherein the molecule is a quorum sensing molecule.

61. The system of claim 56, wherein the molecule is a ligand.

62. The system of claim 56, wherein the molecule is an inducer.

63. The system of claim 56, wherein the molecule is an activator.

64. The system of claim 56, wherein the molecule is an enzyme substrate.

65. The system of claim 56, wherein the molecule is an enzyme activator.

66. The system of claim 56, wherein the molecule is an enzyme inhibitor.

67. The system of claim 56, wherein the molecule is a hormone.

68. The system of claim 56, wherein the molecule is a neurotransmitter.

69. The system of claim 56, wherein the molecule is a cytokine.

70. The system of claim 56, wherein the molecule is a growth factor.

71. The system of claim 56, wherein the molecule is a toxin.

72. The system of claim 56, wherein the molecule is a sugar.

73. The system of claim 56, the instructions further comprising selecting a molecule associated with a biological event.

74. The system of claim 73, wherein the biological event comprises binding of the molecule.

75. The system of claim 73, wherein the biological event comprises conversion of a substrate.

76. The system of claim 73, wherein the biological event comprises neurotransmitter re-uptake.

77. The system of claim 73, wherein the biological event comprises apoptosis.

78. The system of claim 73, wherein the biological event comprises angiogenesis.

79. The system of claim 56, wherein receiving the microwave spectrum comprises measuring the spectrum.

80. The system of claim 56, wherein receiving the microwave spectrum comprises modeling the molecule and analytically deriving the spectrum.

81. The system of claim 56, wherein receiving the microwave spectrum comprises receiving the spectrum from a database.

82. The system of claim 56, wherein identifying the mode of molecular movement comprises relating the one or more peaks to a mode of vibration of one or more moieties of the molecule.

83. The system of claim 56, wherein identifying the mode of molecular movement comprises relating the one or more peaks to a mode of vibration of one or more functional groups of the molecule.

84. The system of claim 56, wherein the motion impedes binding of the molecule.

85. The system of claim 56, wherein the motion promotes binding of the molecule.

86. The system of claim 56, wherein the motion promotes the unbinding of the molecule.

87. The system of claim 56, wherein the motion triggers release of the molecule.

88. The system of claim 56, wherein the motion comprises a rotation of one or more portions of the molecule.

89. The system of claim 56, wherein the motion comprises a vibration of one or more portions of the molecule.

90. The system of claim 56, the instructions further comprising storing the signal profile in a database.

91. The system of claim 56, the instructions further comprising generating one or more microwave signals based on the signal profile.

92. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a fixed frequency signal within a frequency range corresponding to the one or more peaks.

93. The system of claim 91, wherein the one or more peaks comprises a first peak and a second peak and the one or more microwave signals includes a first microwave signal, the frequency of the first microwave signal hopping between frequency ranges corresponding to the first peak and the second peak.

94. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal sweeping between frequencies within a frequency range corresponding to the one or more peaks.

95. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a substantially continuous signal.

96. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a pulsed signal.

97. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a frequency bandwidth coinciding with only one peak of the spectrum.

98. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a frequency bandwidth of less than 101 MHz.

99. The system of claim 91, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a narrow frequency bandwidth.

100. The system of claim 91, wherein the one or more microwave signals is a sequence of microwave signals.

101. The system of claim 91, wherein the one or more microwave signals are concurrent microwave signals.

102. The system of claim 91, wherein the power of each of the generated signals is less than one microwatt.

103. The system of claim 91, wherein the total power of the generated signals is less than one microwatt.

104. The system of claim 91, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having an elliptical polarization.

105. The system of claim 91, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having a circular polarization.

106. The system of claim 91, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having a linear polarization.

107. The system of claim 91 further comprising one or more antennas and the instructions further comprising outputting the one or more generated signals from the one or more antennas.

108. A computer-readable medium containing instructions that, when executed by at least one processor in a system, cause the system to perform a method comprising:

receiving a microwave spectrum associated with rotational modes of a molecule;

identifying a mode of molecular movement associated with one or more peaks in the spectrum;

identifying one or more binding portions of the molecule associated with the identified mode; and

generating, by a processor, a microwave signal profile for driving a motion of the one or more binding portions.

109. The method of claim 108, wherein the molecule is polar.

110. The method of claim 108 further comprising inducing a dipole moment in the molecule.

111. The method of claim 108, wherein the molecule has an induced dipole moment.

112. The method of claim 108, wherein the molecule is a quorum sensing molecule.

113. The method of claim 108, wherein the molecule is a ligand.

114. The method of claim 108, wherein the molecule is an inducer.

115. The method of claim 108, wherein the molecule is an activator.

116. The method of claim 108, wherein the molecule is an enzyme substrate.

117. The method of claim 108, wherein the molecule is an enzyme activator.

118. The method of claim 108, wherein the molecule is an enzyme inhibitor.

119. The method of claim 108, wherein the molecule is a hormone.

120. The method of claim 108, wherein the molecule is a neurotransmitter.

121. The method of claim 108, wherein the molecule is a cytokine.

122. The method of claim 108, wherein the molecule is a growth factor.

123. The method of claim 108, wherein the molecule is a toxin.

124. The method of claim 108, wherein the molecule is a sugar.

125. The method of claim 108 further comprising identifying a molecule associated with a biological event.

126. The method of claim 125, wherein the biological event comprises binding of the molecule.

127. The method of claim 125, wherein the biological event comprises conversion of a substrate.

128. The method of claim 125, wherein the biological event comprises neurotransmitter re-uptake.

129. The method of claim 125, wherein the biological event comprises apoptosis.

130. The method of claim 125, wherein the biological event comprises angiogenesis.

131. The method of claim 108, wherein receiving the microwave spectrum comprises measuring the spectrum using spectroscopy.

132. The method of claim 108, wherein receiving the microwave spectrum comprises modeling the molecule and analytically deriving the spectrum.

133. The method of claim 108, wherein receiving the microwave spectrum comprises receiving the spectrum from a database.

134. The method of claim 108, wherein identifying the mode of molecular movement comprises relating the one or more peaks to a mode of vibration of one or more moieties of the molecule and wherein the one or more binding portions of the molecule comprises at least part of the one or more moieties.

135. The method of claim 108, wherein identifying the mode of molecular movement comprises relating the one or more peaks to a mode of vibration of one or more functional groups of the molecule and wherein the one or more binding portions of the molecule comprises at least part of the one or more functional groups.

136. The method of claim 108, wherein the motion impedes binding of the molecule.

137. The method of claim 108, wherein the motion promotes binding of the molecule.

138. The method of claim 108, wherein the motion promotes the unbinding of the molecule.

139. The method of claim 108, wherein the motion triggers release of the molecule.

140. The method of claim 108, wherein the motion comprises a rotation of one or more portions of the molecule.

141. The method of claim 108, wherein the motion comprises a vibration of one or more portions of the molecule.

142. The method of claim 108 further comprising storing the signal profile in a database.

143. The method of claim 108 further comprising generating one or more microwave signals based on the signal profile.

144. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a fixed frequency signal within a frequency range corresponding to the one or more peaks.

145. The method of claim 143, wherein the one or more peaks comprises a first peak and a second peak and the one or more microwave signals includes a first microwave signal, the frequency of the first microwave signal hopping between frequency ranges corresponding to the first peak and the second peak.

146. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal sweeping between frequencies within a frequency range corresponding to the one or more peaks.

147. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a substantially continuous signal.

148. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal being a pulsed signal.

149. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a frequency bandwidth coinciding with only one peak of the spectrum.

150. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a frequency bandwidth of less than 101 MHz.

151. The method of claim 143, wherein the one or more microwave signals includes a first microwave signal, the first microwave signal having a narrow frequency bandwidth.

152. The method of claim 143, wherein the one or more microwave signals is a sequence of microwave signals.

153. The method of claim 143, wherein the one or more microwave signals are concurrent microwave signals.

154. The method of claim 143, wherein the power of each of the generated signals is less than one microwatt.

155. The method of claim 143, wherein the total power of the generated signals is less than one microwatt.

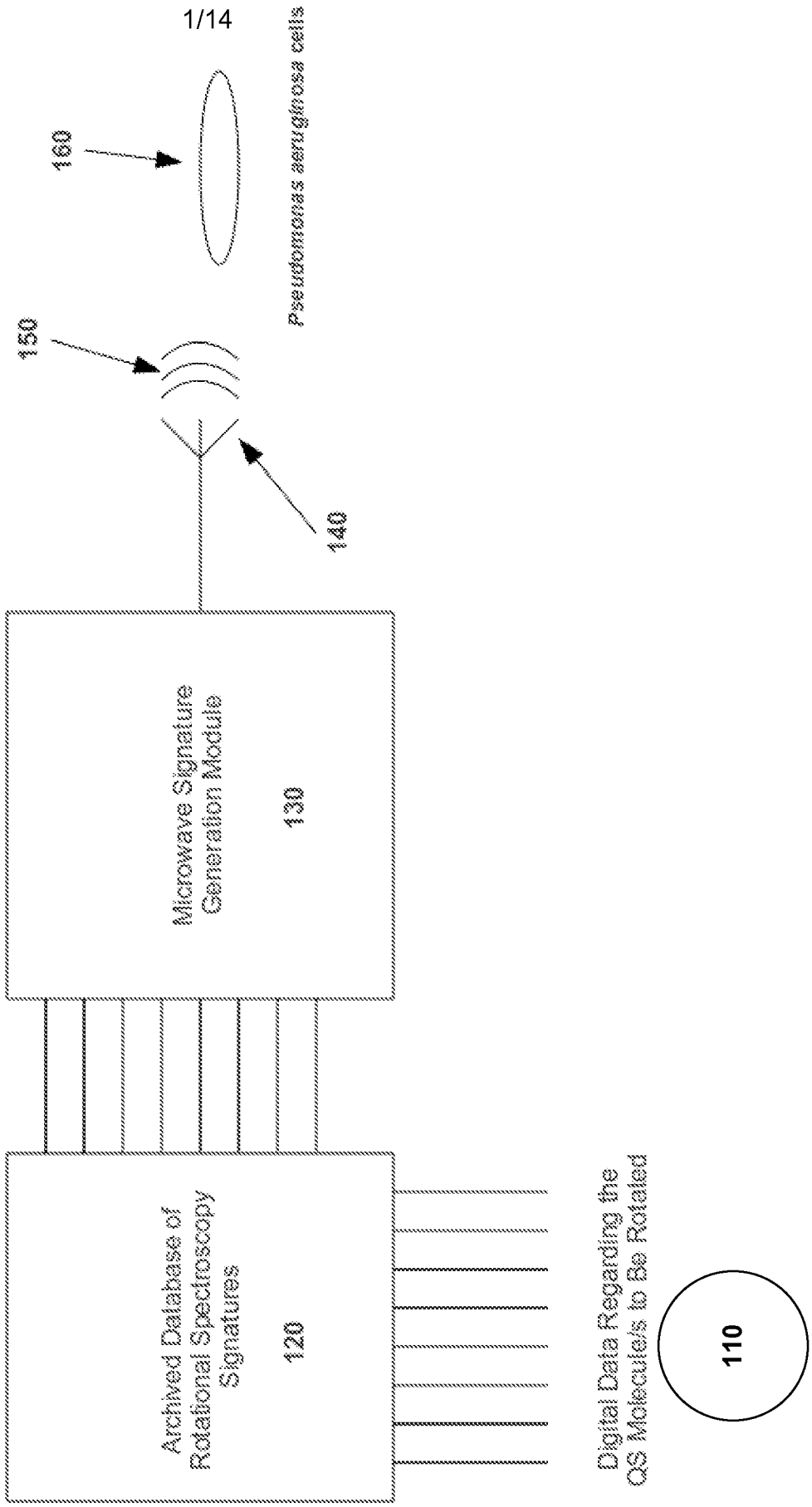
156. The method of claim 143, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having an elliptical polarization.

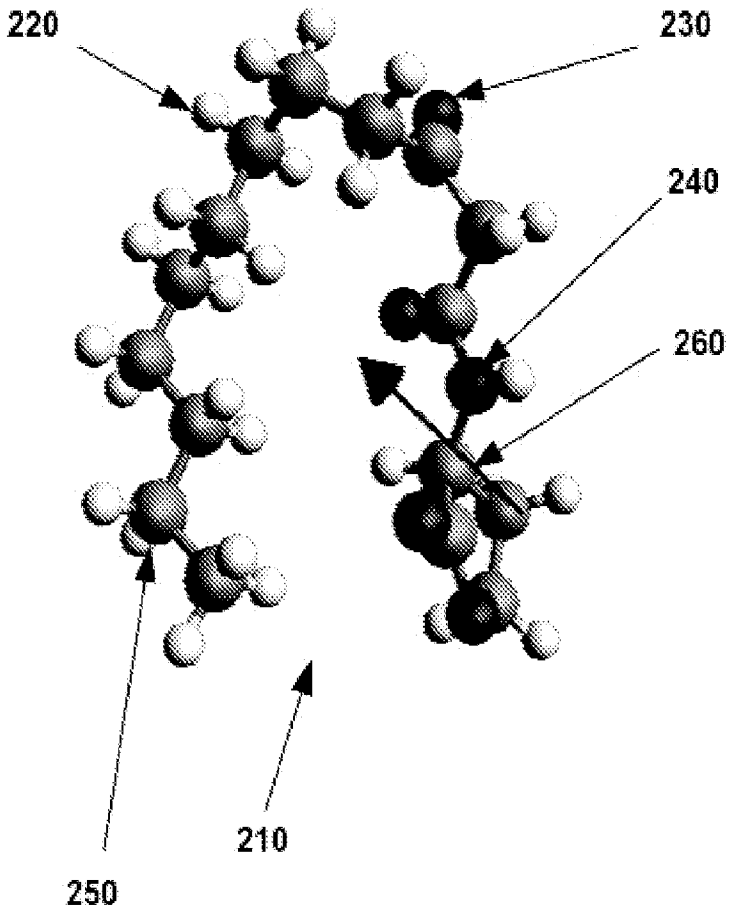
157. The method of claim 143, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having a circular polarization.

158. The method of claim 143, wherein the one or more generated signals includes a first microwave signal, the first microwave signal having a linear polarization.

159. The method of claim 143 further comprising outputting the one or more generated signals from one or more antennas.

Fig. 1

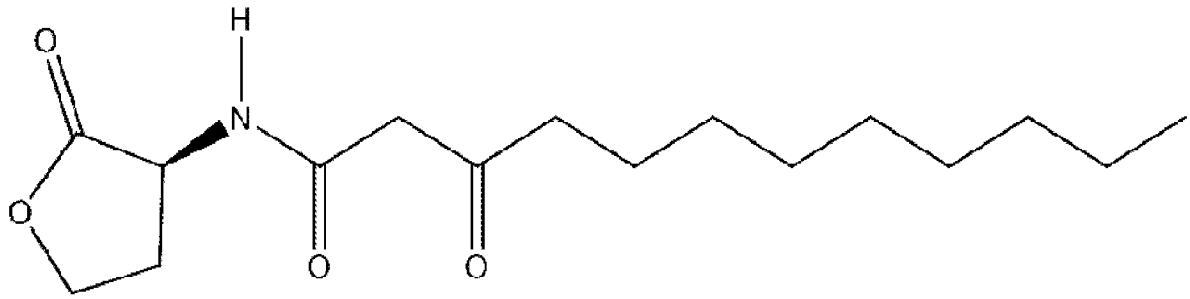

Pseudomonas aeruginosa cells
160
150
140
Microwave Signature Generation Module
130
Archived Database of Rotational Spectroscopy Signatures
120
Digital Data Regarding the QS Molecule/s to Be Rotated
110



N-dodecanoyl-L-Homoserine lactone ("C12")
Quorum Sensing Molecule

Fig. 2A

Fig. 2B



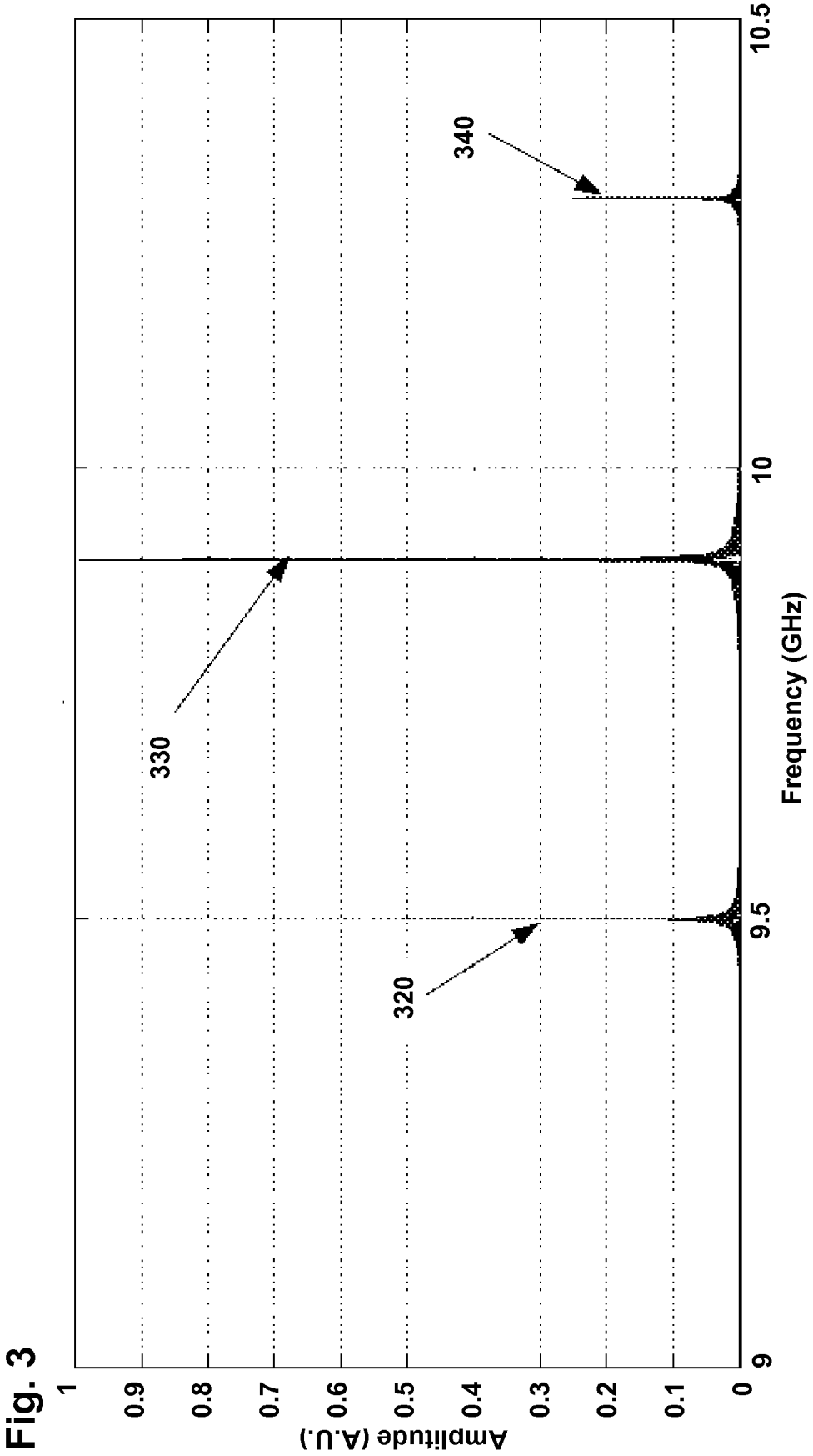

Amplitude (A.U.)
1
0.9
0.8
0.7
0.6
0.5
0.4
0.3
0.2
0.1
0
9
9.5
10
10.5
Frequency (GHz)
330
320
340

Fig. 3

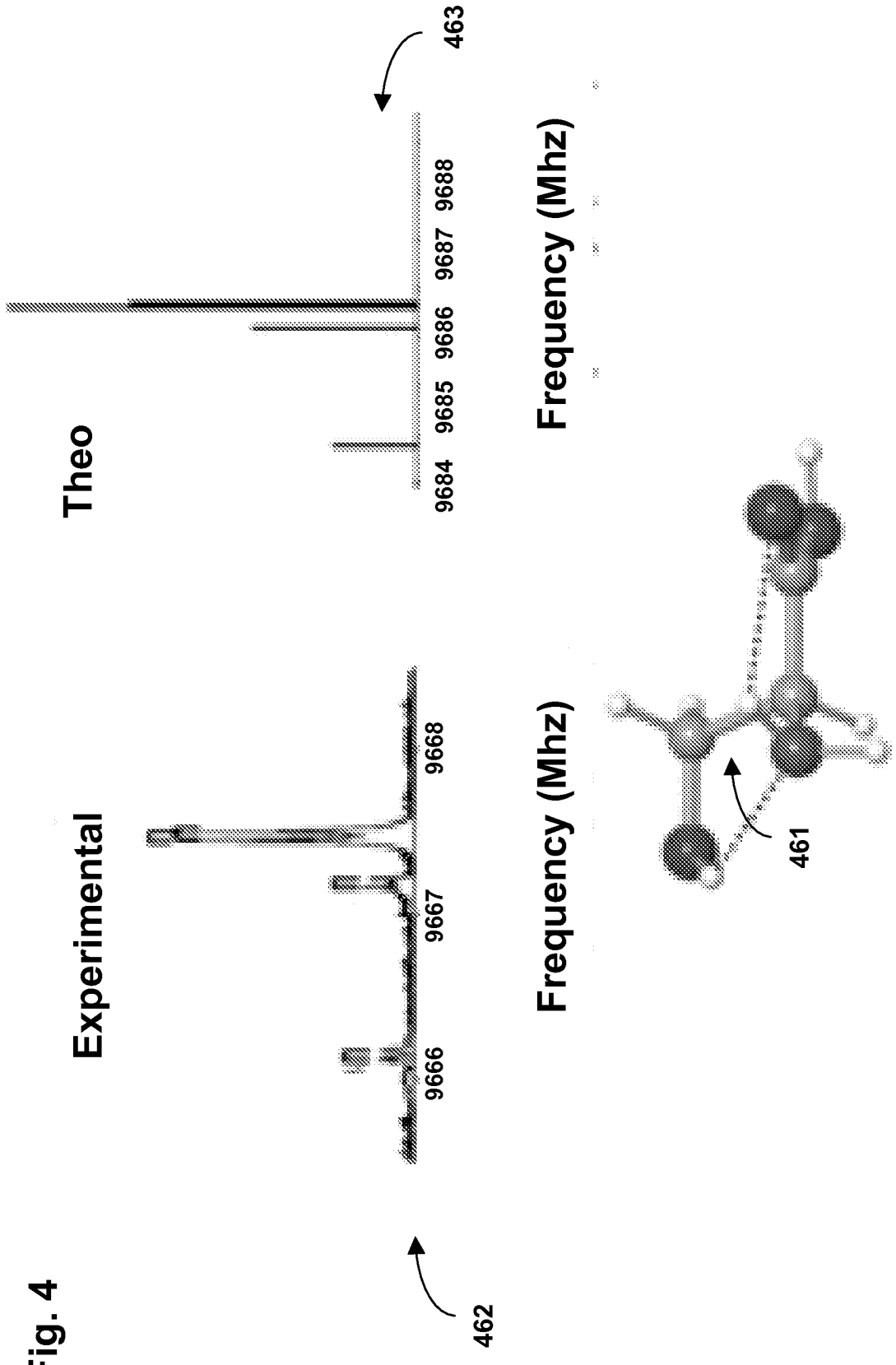

Theo
Experimental
Frequency (Mhz)
Frequency (Mhz)
9684
9685
9686
9687
9688
9666
9667
9668
461
462
463

Fig. 4

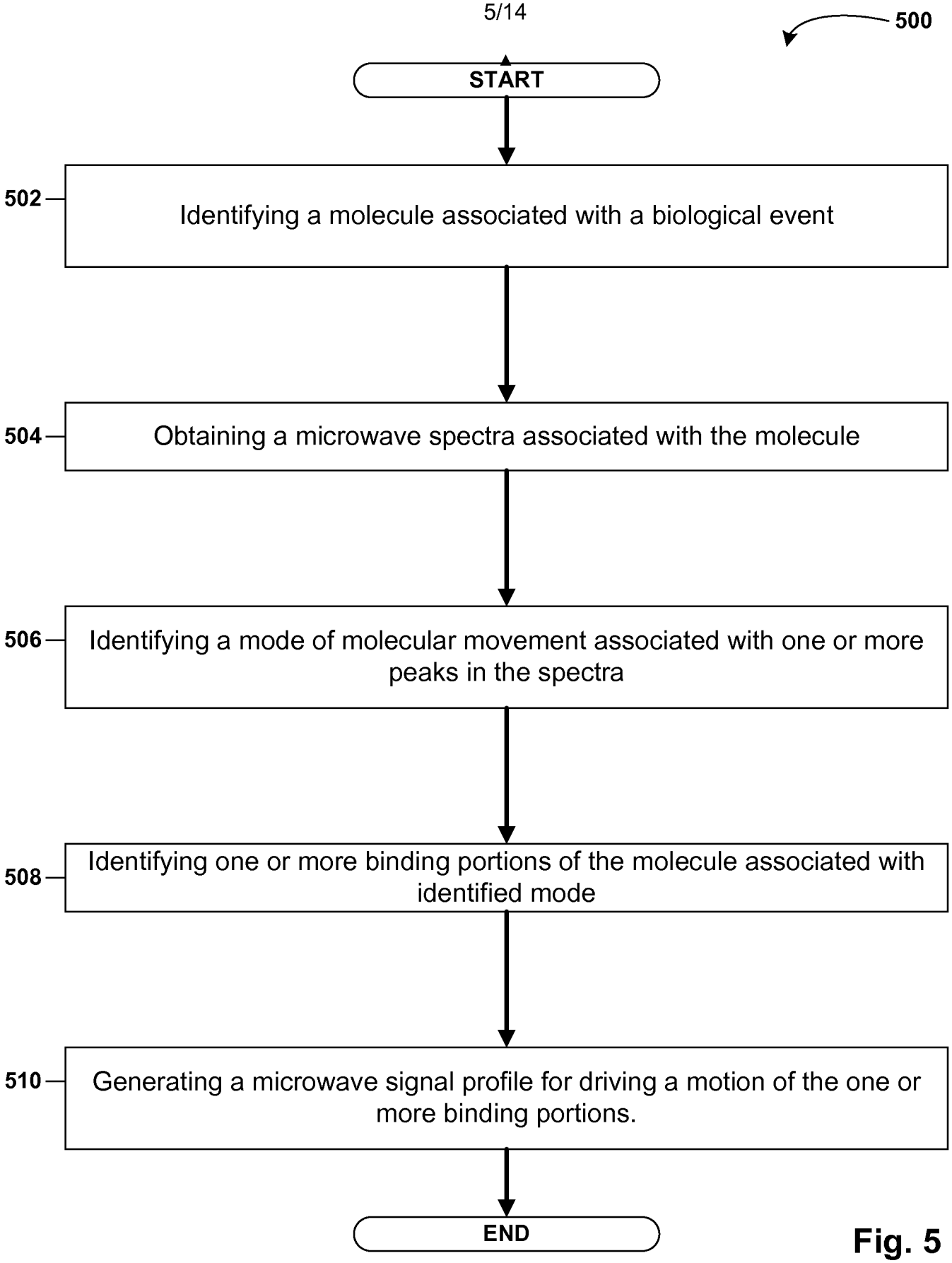

500
START
502
Identifying a molecule associated with a biological event
504
Obtaining a microwave spectra associated with the molecule
506
Identifying a mode of molecular movement associated with one or more peaks in the spectra
508
Identifying one or more binding portions of the molecule associated with identified mode
510
Generating a microwave signal profile for driving a motion of the one or more binding portions.
END

Fig. 5

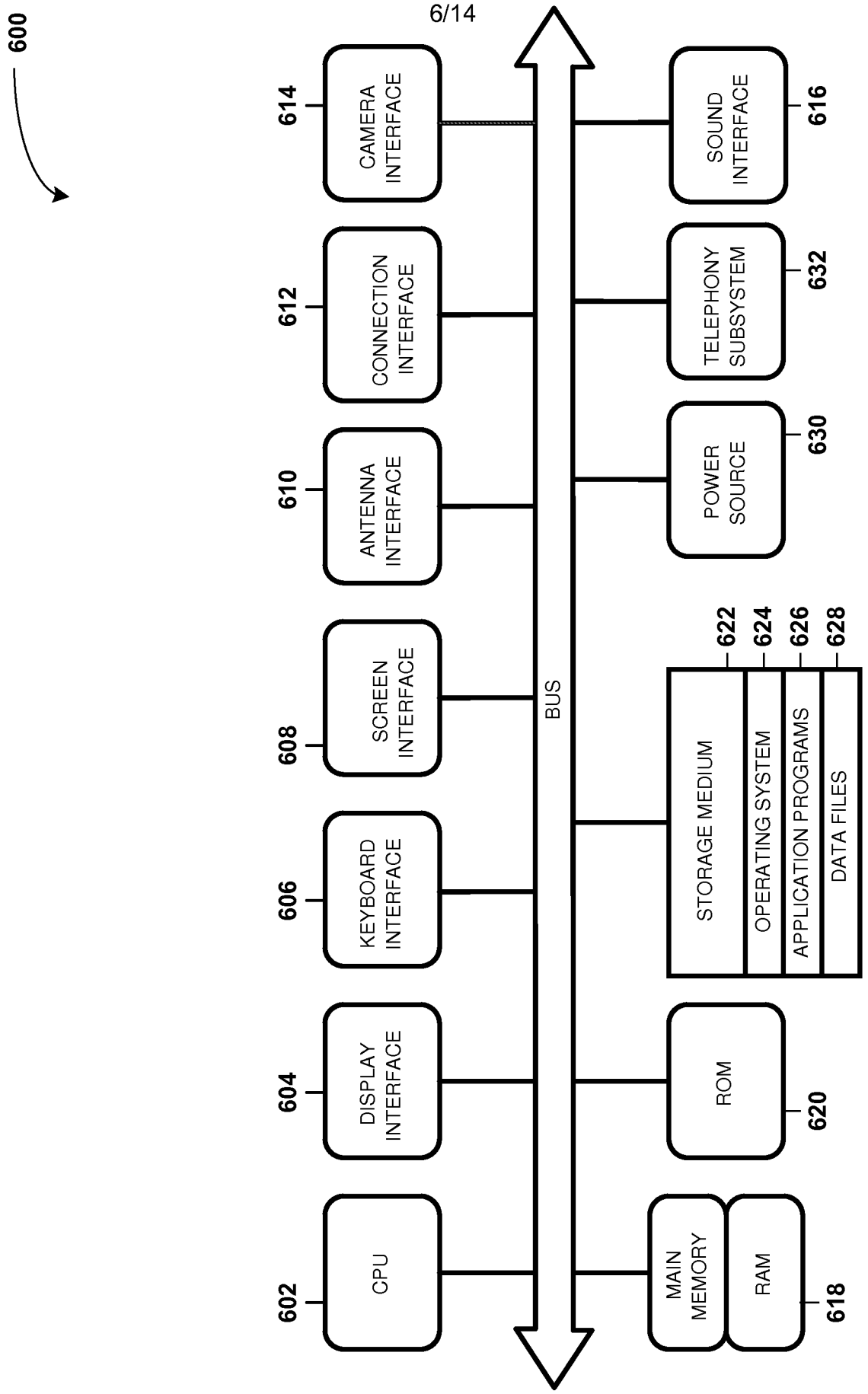
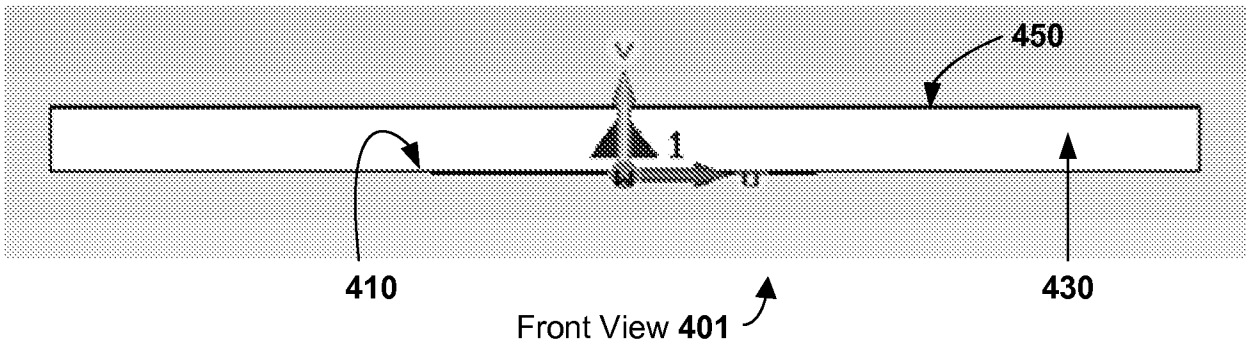
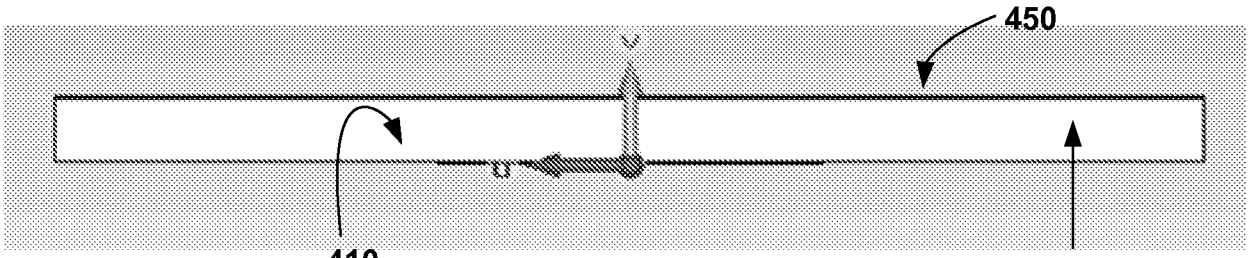

600
602
CPU
604
DISPLAY INTERFACE
606
KEYBOARD INTERFACE
608
SCREEN INTERFACE
610
ANTENNA INTERFACE
612
CONNECTION INTERFACE
614
CAMERA INTERFACE
BUS
618
MAIN MEMORY
RAM
620
ROM
622
STORAGE MEDIUM
624
OPERATING SYSTEM
626
APPLICATION PROGRAMS
628
DATA FILES
630
POWER SOURCE
632
TELEPHONY SUBSYSTEM
616
SOUND INTERFACE

Fig. 6



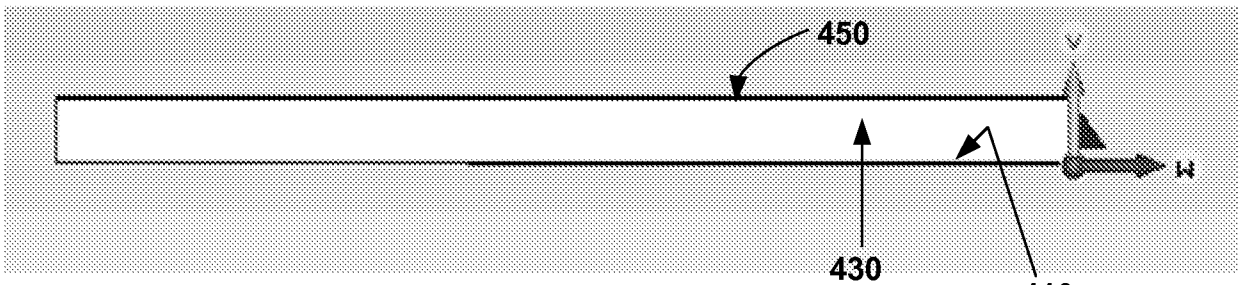
Front View **401**

Fig. L1

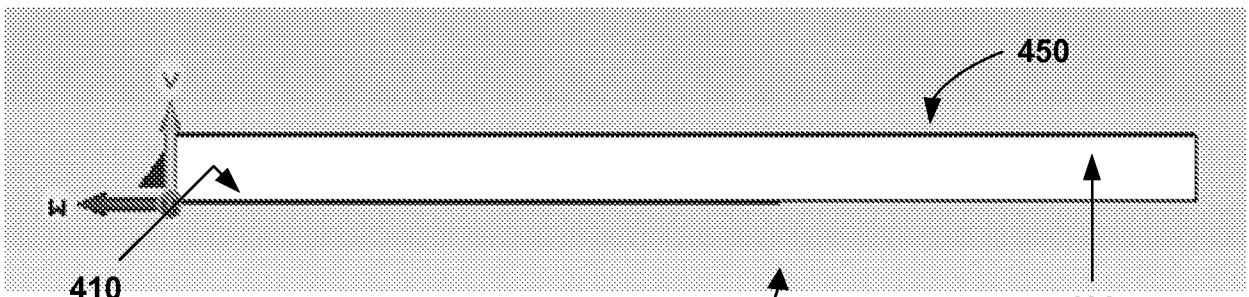


Back View **402**

400



Left View **403**



Right View **404**

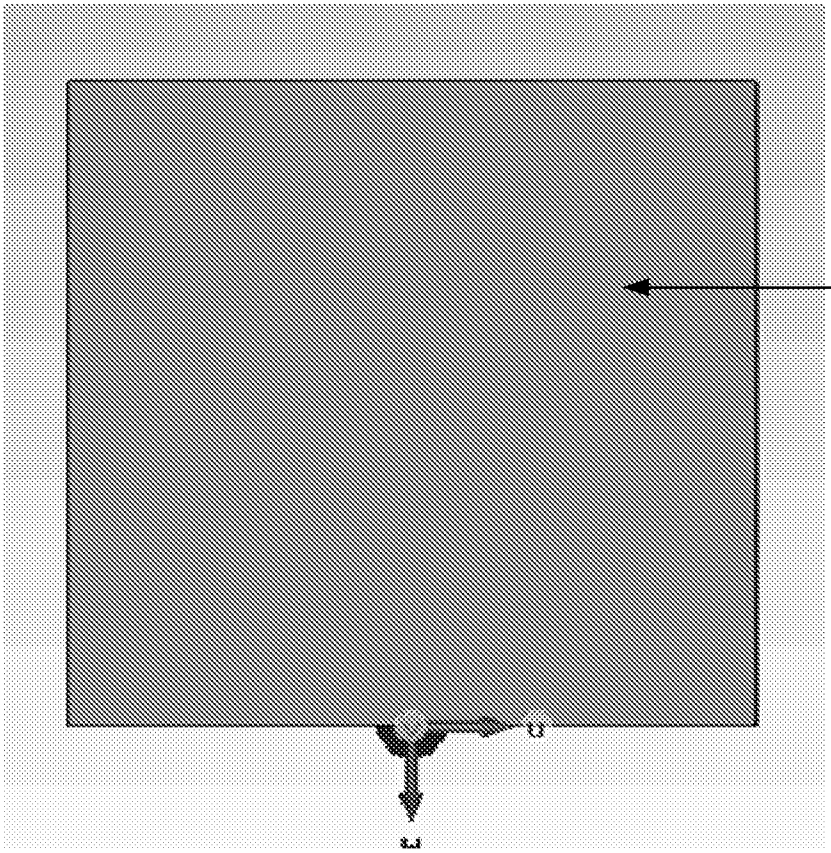
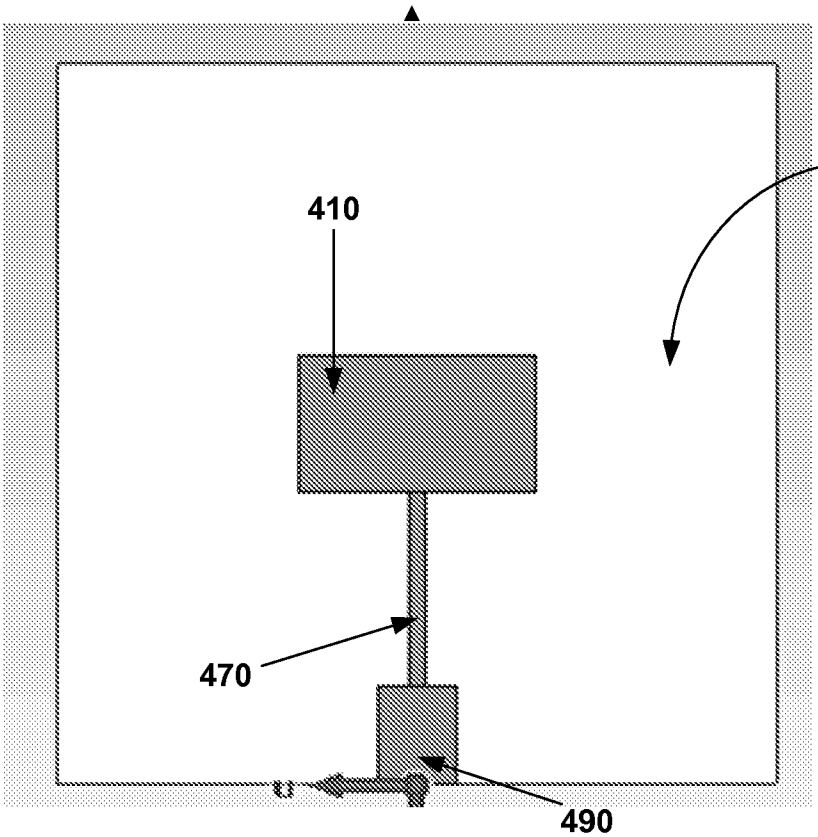


Fig. L2



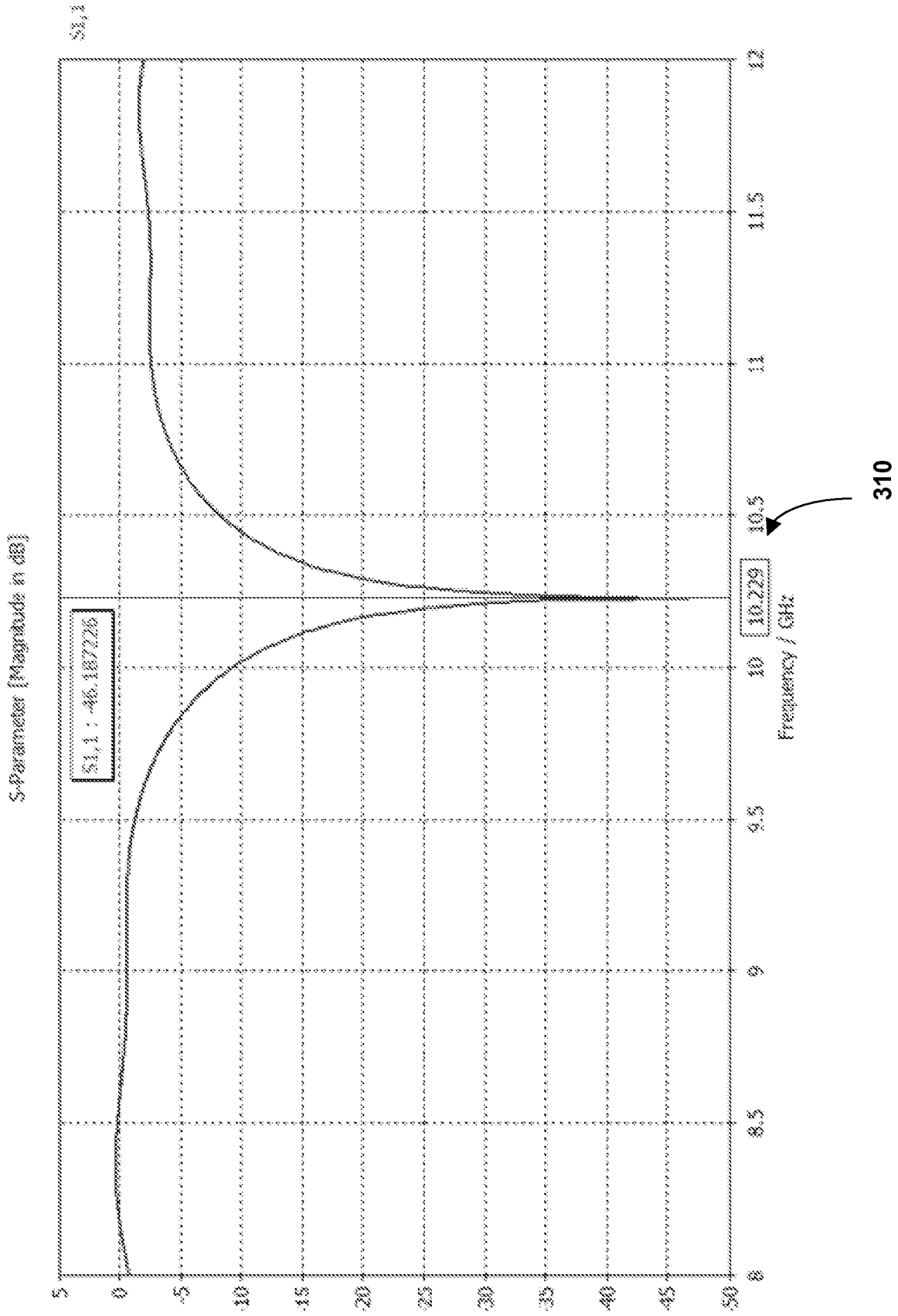

S1,1
S-Parameter [Magnitude in dB]
S1,1 : -46.187226
10.239
Frequency / GHz
310

Fig. L3

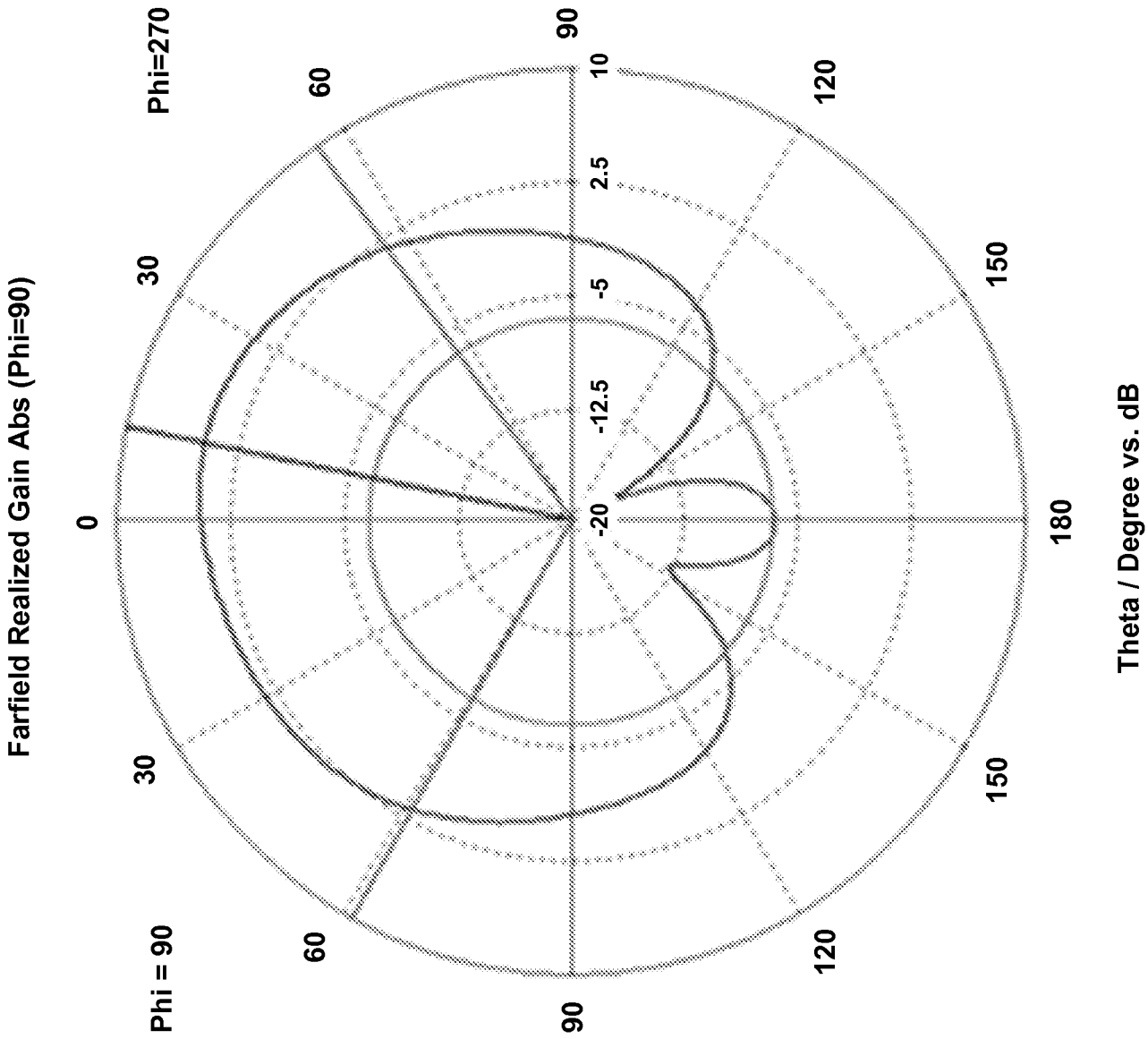
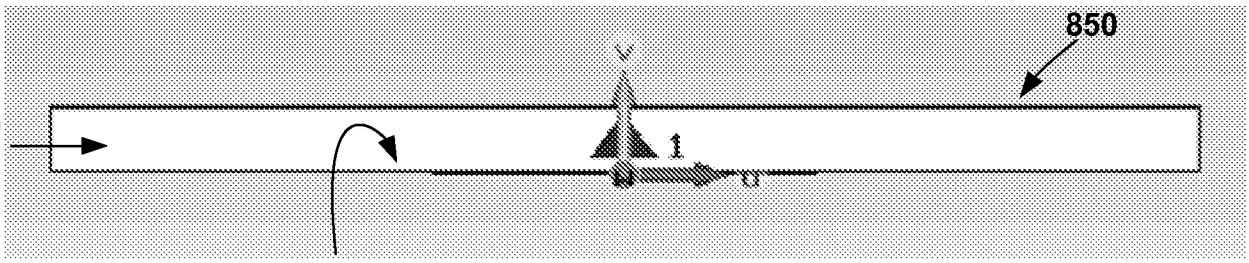
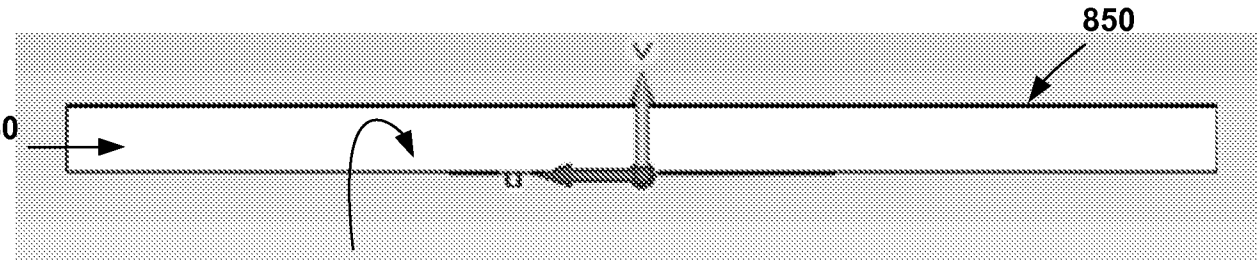

Farfield Realized Gain Abs (Phi=90)
Phi=270
Phi = 90
0
30
60
90
120
150
180
10
2.5
-5
-12.5
-20
Theta / Degree vs. dB

Fig. L4

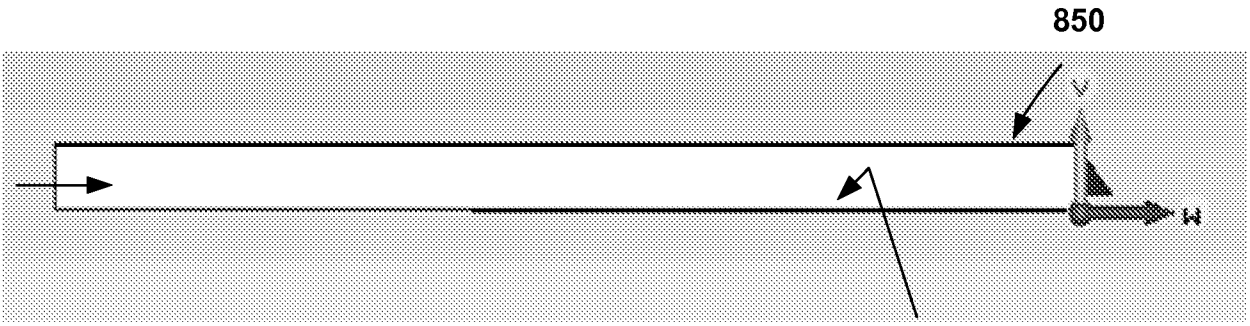


Front View **801**

Fig. C1

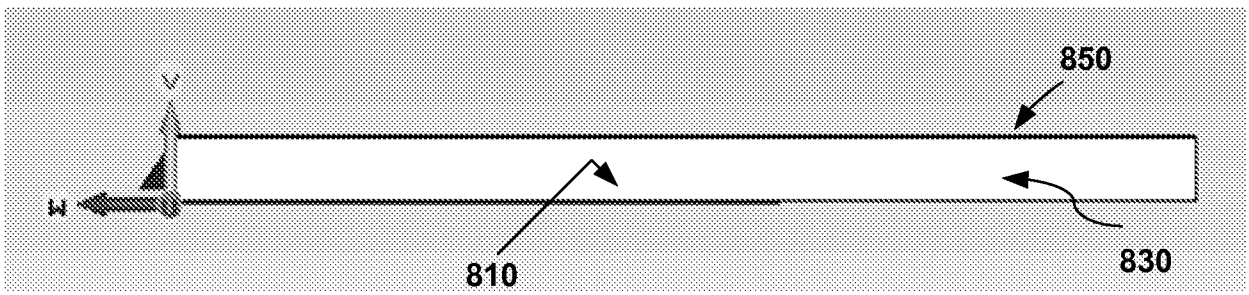


Back View **802**



Left View **803**

800



Right **804**

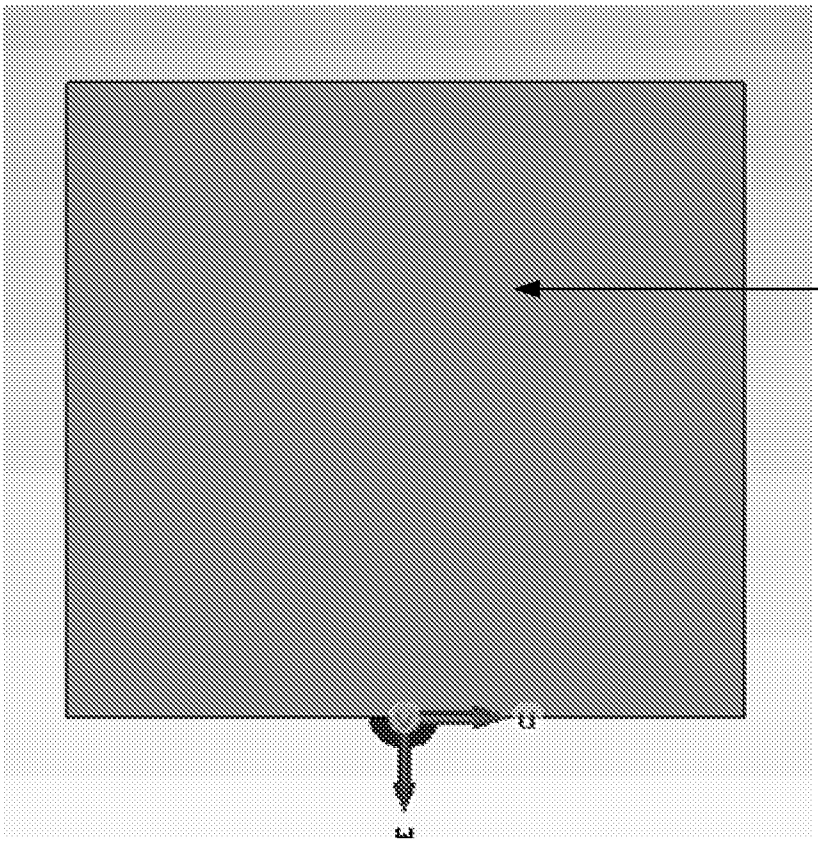
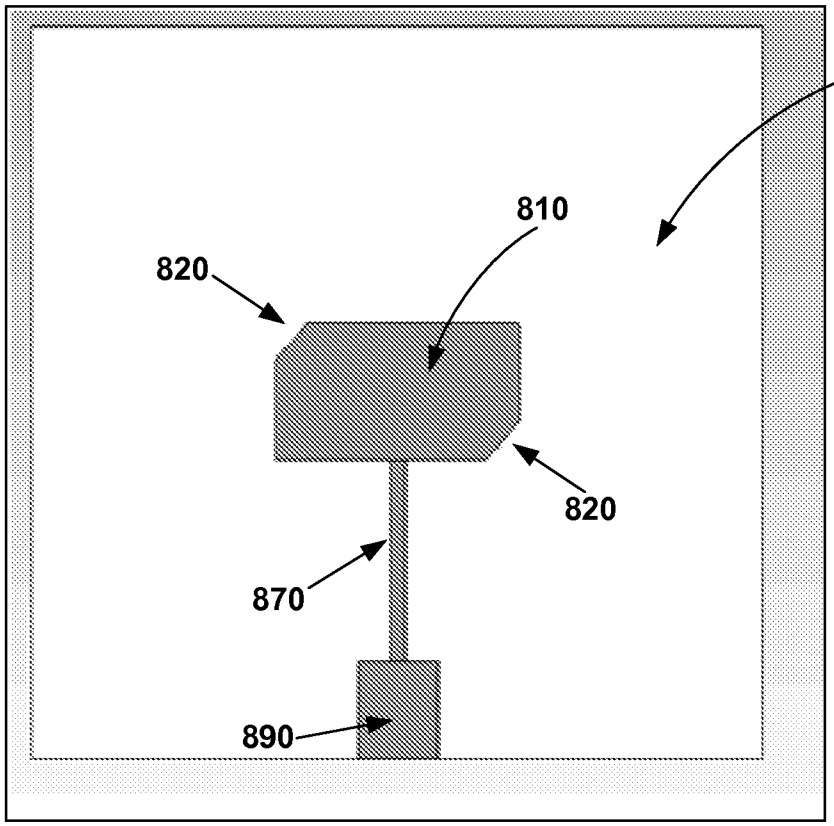


Fig. C2



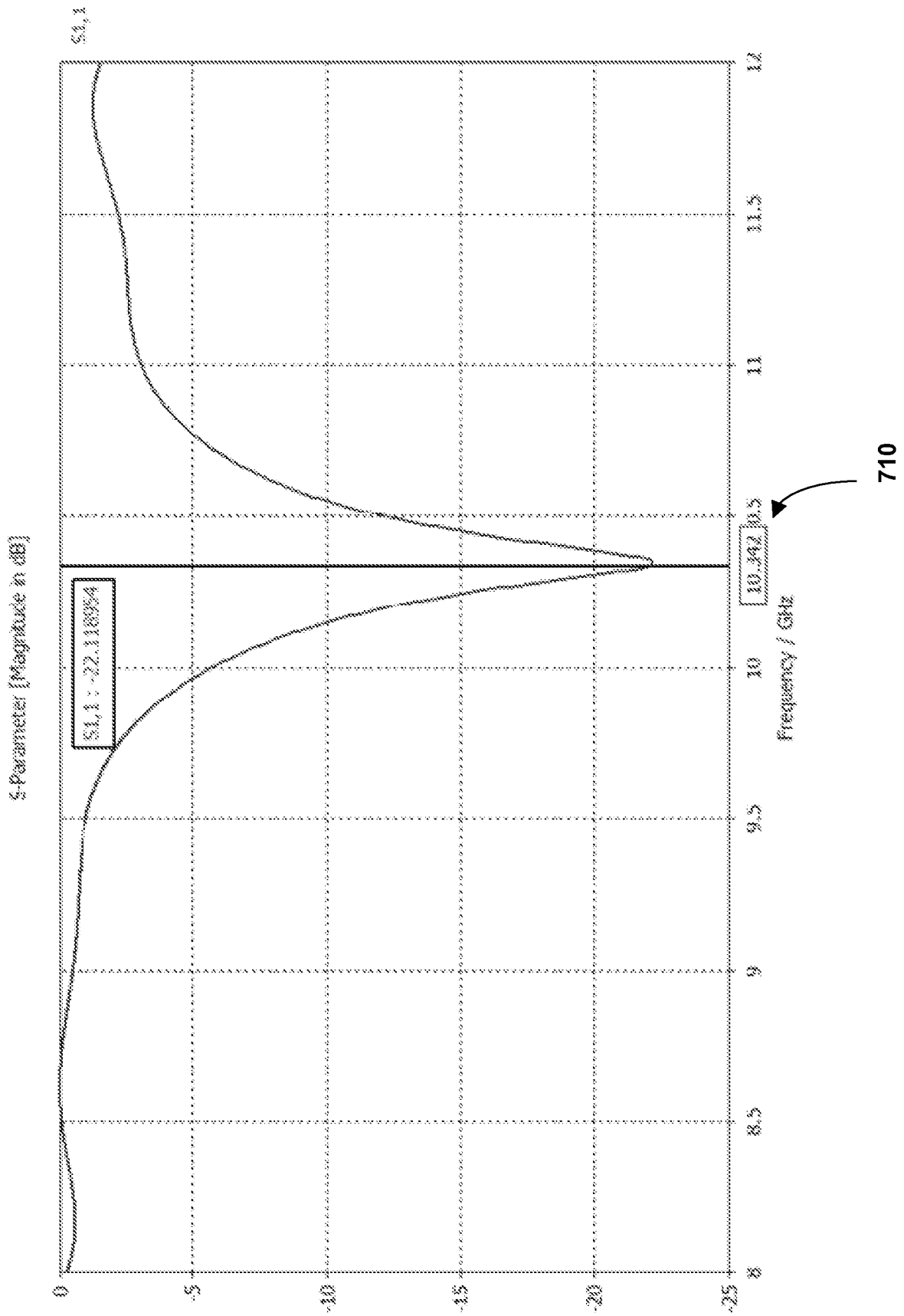

S1,1
S1,1 : -22.118094
10.342
710
S-Parameter [Magnitude in dB]
Frequency / GHz
0
-5
-10
-15
-20
-25
8
8.5
9
9.5
10
10.5
11
11.5
12

Fig. C3

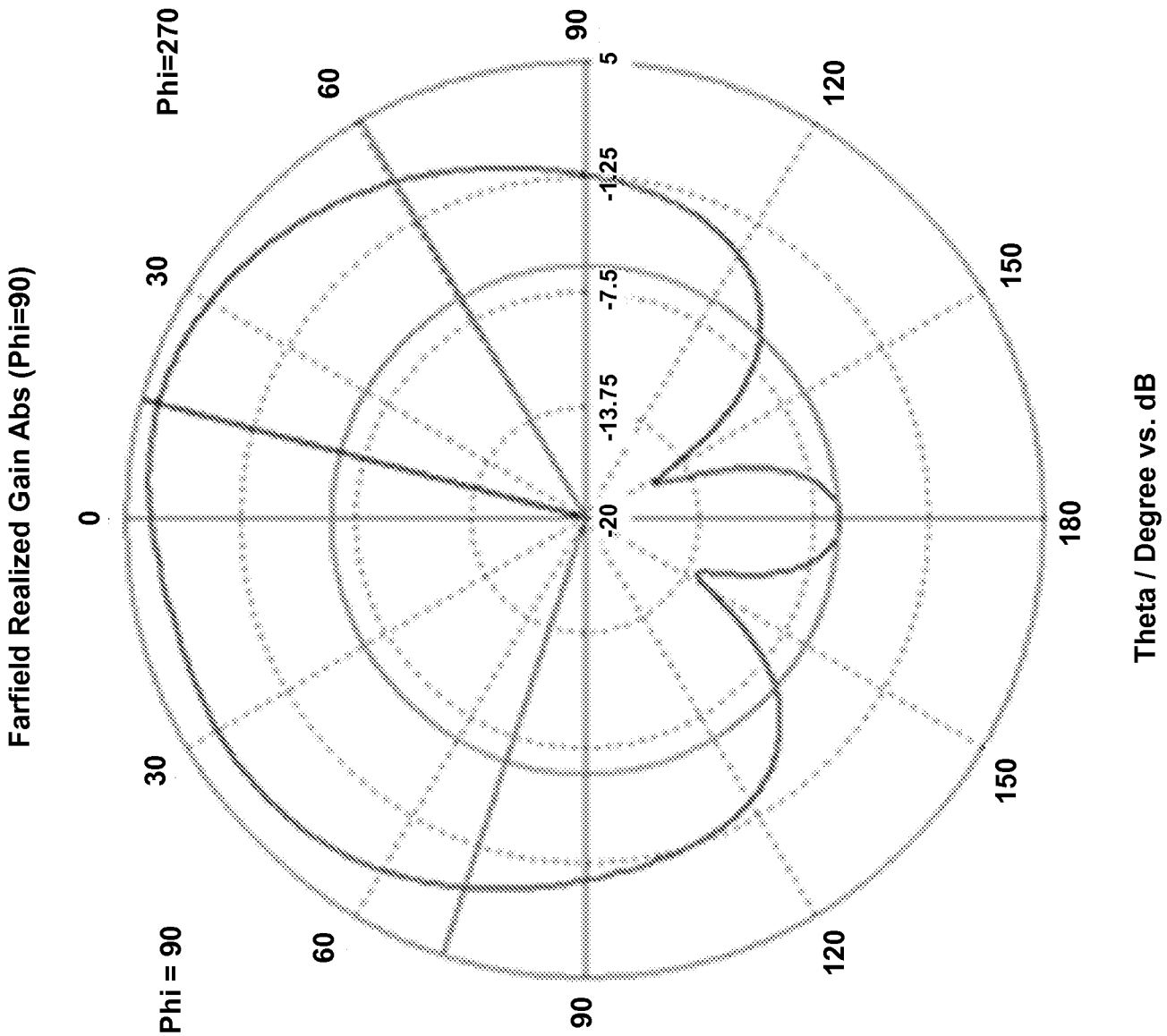

Farfield Realized Gain Abs (Phi=90)
Phi=270
Phi = 90
0
30
60
90
120
150
180
5
-1.25
-7.5
-13.75
-20
Theta / Degree vs. dB

Fig. C4

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US13/26193

A. CLASSIFICATION OF SUBJECT MATTER
IPC(8) - G01N 33/53, 22/00 (2013.01)
USPC - 204/157.15, 157.43; 702/38
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC(8): C07C 1/00, 2/00, 4/00, 5/00, 6/00; G01N 33/53, 22/00 (2013.01)
USPC: 204/157.15, 157.43; 702/19, 21, 1, 38; 435/4

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
MicroPatent (US Granted, US Applications, EP-A, EP-B, WO, JP, DE-G, DE-A, DE-T, DE-U, GB-A, FR-A); ScienceDirect; IEEE Xplore; Google/Google Scholar; 'microwave spectrum,' 'microwave spectroscopy,' fourier, 'binding molecules,' 'dipole moment,' 'quorum sensing,' 'microwave rotation'

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 2005/0139485 A1 (BROOKS, J et al.) June 30, 2005; paragraphs [0007], [0034], [0103], [0171], [0352]-[0358]	1-36, 38-92, 94-144, 146-159
Y	BIRGE, RR et al. Role of Calcium In The Proton Pump of Bacteriorhodopsin: Microwave Evidence For A Cation-Gated Mechanism. The Journal of Physical Chemistry. 1996. Vol.100, No. 23, pp 9990-10004. DOI: 10.1021/jp953669e.	1-36, 38-92, 94-144, 146-159
Y	OLIVER, CM et al. Microwave Synthesis And Evaluation of Phenacylhomoserine Lactones As Anticancer Compounds That Minimally Activate Quorum Sensing Patwhays In Pseudomonas aeruginosa. J. Med. Chem. 2009. Vol. 52, pp 1569-1575. DOI: 10.1021/jm8015377.	5, 60, 112
Y	WEINTRAUB, ST et al. Evaluation Of The Necessity For Rapid Inactivation Of Brain Enzymes Prior To Analysis of Norepinephrine, Dopamine And Serotonin In The Mouse. Life Sciences. 02 October 1975, Vol. 17, pp 1423-1428.	20, 76, 128
Y	SONG, X et al. Microwave Induces Apoptosis In A549 Human Lung Carcinoma Cell Line. Chinese Medical Journal. 2011. Vol. 124, No. 8, pp 1193-1198. DOI: 10.3760/cma.j.issn.0366-6999.2011.08.013.	21, 77, 129
Y	US 6421550 B1 (BRIDGES, JE et al.) July 16, 2002; column 6, lines 14-22, column 7, lines 3-5	22, 78, 130
Y	WO 2002/077607 A2 (MROSS, MR et al.) October 3, 2002; paragraph [42]; Claim 37	28, 41, 46, 47, 84, 97, 102, 103, 136, 149,154, 155
Y	US 5463226 A (MATSUZAKI, K et al.) October 31, 1995; column 7, lines 49-60	38, 94, 146

☒ Further documents are listed in the continuation of Box C. ☐

* Special categories of cited documents:
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"E" earlier application or patent but published on or after the international filing date
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"O" document referring to an oral disclosure, use, exhibition or other means
"P" document published prior to the international filing date but later than the priority date claimed
"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
30 April 2013 (30.04.2013)	13 MAY 2013
Name and mailing address of the ISA/US Mail Stop PCT, Attn: ISA/US, Commissioner for Patents P.O. Box 1450, Alexandria, Virginia 22313-1450 Facsimile No. 571-273-3201	Authorized officer: Shane Thomas PCT Helpdesk: 571-272-4300 PCT OSP: 571-272-7774

Form PCT/ISA/210 (second sheet) (July 2009)

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US13/26193

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 2007/0284241 A1 (KIBAR, O) December 13, 2007; paragraphs [0100], [0116]	48, 50, 104, 106, 156, 158
Y	US 2009/0131926 A1 (RUSIN, CT et al.) May 21, 2009; figures 1-5; paragraph [0021]; Claim 1	52, 55
Y	US 2010/0114086 A1 (DEEM, ME et al.) May 6, 2010; abstract; figures 5-11; paragraphs [0055], [0146]	53, 54
Y	US 2009/0001262 A1 (VISSER, E et al.) January 1, 2009; figures 1, 2; paragraphs [0034], [0035]	24, 25, 34, 56-92, 94-144, 146-159