

combination with resonant acoustic energy. Using the resonant acousto-EM and acoustic energy together, allows for the use of lower power levels of both types of energy, lessening the potential adverse affects of electromagnetic energy and/or acoustic energy on nearby or adjacent nontargeted structures.

5 Electromagnetic energy may interact with and complement an acoustic energy wave in a system in at least four ways: via the piezoelectric effect, intrinsic dissipation of electromagnetic energy and via the acoustoelectric or magnetoacoustic effect.

10 In the piezoelectric effect, acoustic vibratory energy is converted interchangeably with EM energy by a transducer. Biologic piezoelectric structures can modulate the same conversion of energy, thereby acting as living transducers. When an EM field is applied to a biologic piezoelectric structure, an acoustic wave is produced. Likewise, when an acoustic wave is applied to a biologic piezoelectric structure, EM energy is produced. The piezoelectric effect in biologic structures has many useful applications (see below.) This effect becomes even more useful when principles of acoustic resonance are applied. In the  
15 present invention specific biologic structures can be targeted with an acoustic wave or EM energy at power levels that dramatically affect the target structure, but have virtually no effect on adjacent, nonresonant structures. Although not previously postulated by others, biologic structures functioning as living, resonant piezoelectric transducers which modulate the conversion of mechanical and EM energy is undoubtedly one of the major underlying  
20 mechanisms responsible for the interaction of EM fields with biologic structures.

In the acoustoelectric effect, the passage of an acoustic wave through a semiconductor induces an electric current. The passage of an acoustic wave through the material is postulated to cause a periodic spatial variation of the potential energy of the charge carriers. This results in an electric field across the ends of the semiconductor as long  
25 as the acoustic wave is traversing the semiconductor. Free electrons carriers are bunched in the potential-energy troughs, and as the acoustic wave having a specific frequency propagates, it drags the bunches along with it, resulting in an electric field such as a DC field pulsing at the specific acoustic frequency or an AC field having a frequency equal to the specific acoustic frequency. The effect is enhanced where there are both positively and  
30 negatively charged carriers, and where there are many different groups of carriers - conditions which are frequently found in biologic systems. The attributes of the current

produced depend on the unique molecular and atomic components of the structure in question. This aspect alone provides a means to perform acoustoelectric spectroscopy on biologics many of which are semiconductors, and depending on the selected frequency, the acoustoelectric effect in biological structures has many potentially useful applications. Thus  
5 understood, a targeted structure can be irradiated or exposed to acoustic energy having non-resonant frequency and an electromagnetic energy pattern of the acoustoelectric effect in the structure can be detected. This detected electromagnetic energy pattern can be used as a signature to detect and identify the targeted structure.

However, the acoustoelectric effect becomes even more useful when principles of  
10 acoustic resonance are applied. Augmentation, detection, and/or disruption of biologics can be targeted to specific structures at power levels that dramatically affect the target structure, but have virtually no effect on nearby, nonresonant structures. The current produced by the acoustoelectric effect in a resonant structure will be much stronger than any current produced by neighboring non-resonant structures, and may be of an alternating nature. The large signal  
15 to noise ratio obtained from a resonant structure improves accuracy of acoustic and EM pattern identification and detection. Similar to reversal of the piezoelectric effect and acoustic resonance intrinsic energy dissipation pathway (see above), application of the resonant acoustoelectric EM pattern to a targeted structure will amplify the acoustic wave (acoustoelectric gain which peaks at the frequency for which the acoustic wavelength is the  
20 Debye length, where bunching is optimum). Thus, combined use of the resonant acoustic and acoustoelectric EM fields can allow for greater tissue penetration of high frequency acoustic energy that would otherwise be highly attenuated and have poor tissue penetration. Using the resonant acoustic frequency and acoustoelectric EM fields together also allows for the use of lower power levels of both types of energy, lessening the potential effects on other  
25 nontargeted and nonresonant structures.

The magnetoacoustic effect is the magnetic-field-dependent attenuation of an acoustic field in a monotonic, oscillatory, or resonant manner, depending on the electronic properties of the substance in question. This variability in result, depending on structural composition, provides a further enhancement of resonant acousto-EM spectroscopy in relation to biologics  
30 and other structures, via addition of a magnetic field. Also, the addition of a magnetic field provides the means to amplify or attenuate an acoustic field, thus improving or modulating

the penetration of the acoustic field in biologic tissues.

Similarly, resonant acoustics combined with acoustic cyclotron resonance (ie. resonant acoustic cyclotron resonance) and Doppler-shifted resonant acoustic cyclotron resonance presents a powerful, and precise means of selectively causing augmentation,  
5 detection and/or disruption of structures.

The present invention provides a method that applies the principles of acoustic resonance to biologic structures for the purpose of disruption and/or augmentation of functions of the specifically targeted biologic structure. The resonant acoustic frequency of a biologic structure may be determined by performing resonant acoustic spectroscopy using  
10 methods and systems well know in the art. Particularly, a resonant acoustic frequency of a biologic structure may be determined by the steps of:

- a) applying acoustic energy to the biologic structure and scanning through a range of acoustic energy frequencies; and
- b) detecting at least one specific frequency which causes a maximum signal output  
15 from the biologic structure indicating the biologic structure being induced into acoustic resonance by the at least specific frequency.

The specific frequencies causing the maximum signals are the resonant acoustic frequencies of the biologic structure which are defined and used herein as the acoustic signature of the biologic structure. Once determined, at least one resonant acoustic  
20 frequency may be applied to the biologic structure to affect functioning therein and/or to determine its acousto-EM signature.

The acoustic energy including the resonant acoustic frequencies is applied at a power level sufficient to affect functioning of the biologic structure. Depending on the power intensity of the acoustic energy and the type of targeted structure that is induced into acoustic  
25 resonance, the structure may have its functions affected, such as disruption and/or augmentation.

At lower power levels functions of the biologic structure can be augmented while at higher power levels disruption of the structure may occur. Augmentation as used herein encompasses beneficial effects on the biologic structure. Such augmenting of functions or  
30 enhancing effects include but are not limited to enhancement of growth, reproduction, regeneration, embryogenesis, metabolism, fermentation, and the like. The results of such

enhancement include but are not limited to increase in bone mass or density, increase in number and maturation of eggs, increase in number and/or function of leukocytes, increase in fermentation products in beer, wine and cheese manufacturing, increase in plant germination and growth and the like.

5           There are some situations where the ability to selectively disrupt a structure with resonant acoustic energy is very useful as disclosed in the present invention. As stated above, disruption as used herein refers to deleterious effects on the biologic structure. Such deleterious effects include but are not limited to structural failure of the biologic structure resulting in lysis, shattering, rupture or inactivation of the biologic or of one or more  
10 components of the biologic structure. Disruption as used herein also includes within its ambit inhibition of vital processes required for growth, reproduction, metabolism, infectivity and the like. Components which may be targeted for disruption include, but are not limited to DNA, RNA, proteins, carbohydrates, lipids, lipopolysaccharides, glycolipids, glycoproteins, proteoglycans, chloroplasts, mitochondria, endoplasmic reticulum, cells, organs and the like.  
15 In the case of virulent organisms, the virulence factors may be specifically targeted for disruption to prevent or inhibit the growth, infectivity or virulence of the organism. Such virulence factors include but are not limited to endotoxins, exotoxins, pili, flagella, proteases, ligands for host cell receptors, capsules, cell walls, spores, chitin, and the like.

          Organics, biologics or one or more targeted portions thereof which are amenable to  
20 disruption using the methods of the present invention include but are not limited to viruses, bacteria, protozoans, parasites, fungi, worms, mollusks, arthropods, tissue masses, and the like. The organics or biologics to be disrupted may be isolated, present in a multicellular organism or portion thereof, or other complex environment.

          It is postulated that disruption of the targeted biologic structure without affecting  
25 nearby tissue or structures occurs due to acoustic resonance being induced only in the targeted structure which until now has not been considered a mechanism to affect a biologic structure. This is very different from that disclosed in the prior art which contemplates only three mechanisms for affecting a biologic structure which include cavitation, thermal and mechanical.

30           At specific power levels, such as in lower levels, that do not cause the actual disruption of a structure, resonant acoustic energy can intrinsically dissipate within the

structure and to the adjacent medium. This intrinsically dissipated energy can be converted by the structure into an electromagnetic energy having specific properties and/or fields that may be manifested as direct and alternating currents, electric and magnetic fields, electromagnetic radiation and the like. The pattern of the electromagnetic energy represents  
5 an acousto-EM signature of the structure.

The present invention provides a method to determine an acousto-EM signature of a structure which comprises irradiating the structure with acoustic energy having a frequency at or near a previously determined resonant acoustic frequency of the structure to induce resonance therein and detecting the electromagnetic energy pattern caused by the intrinsic  
10 dissipation of energy.

Once an acousto-EM signature is determined for a specific structure, this structure can be induced into acoustic resonance by applying EM energy equivalent to the acousto-EM signature of the structure, such as equivalent to the acousto-EM signature of the structure. direct and alternating current, electric and magnetic fields, and electromagnetic radiation and  
15 the like.

As such, the present invention applies the principles of acoustic resonance by applying resonant acoustic frequencies and electromagnetic energy equivalent to the predetermined acousto-EM signature of a targeted structure individually or in combination to affect the targeted structure, the method comprising the steps of:

20 a) applying at least one resonant acoustic frequency of the targeted structure and/or introducing electromagnetic energy equivalent to part or all of the acousto-EM signature of the targeted structure; and

c) applying (a) and/or (b) each at a power intensity level to induce acoustic resonance within the targeted structure and affect functioning of the structure.

25 Both the resonant acoustic frequency of the targeted structure and the acousto-EM signature must be predetermined, as discussed above, to provide the applicable energy for inducing acoustic resonance in the structure. The electromagnetic energy can be introduced into the targeted structure in the form of a direct or alternating current have a specific frequency that is equivalent to the electromagnetic energy pattern detected when the  
30 structure is induced into acoustic resonance. Furthermore each type of energy can be applied at a power level less than used individually and this allows for inducing acoustic resonance