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How To Over Come Antibiotic Drug Resistance? Drug Resistance is Disorder of Second Order Of Universe Either By Drug Itself Or By Alteration In Implementation Due To Changed Behavior (Mutation) Of Target B.B.Bs. This Second Order of Universe Could be Reset by First Order Of Universe (AGE).

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Abstract: *Any disorder in implementation of second order of universe carried by drug by alteration in behavior of target B.B.Bs (mutations) , drug resistance develop . Drugs or antibiotics are set of information's (Code PCPs bactericidal / bacteriostatic activity) that trigger (second order carried by photons) bactericidal / bacteriostatic effects by giving specific order to bacterial cell . If this order (Code PcPs bactericidal / bacteriostatic activity) is failed to deliver (drug half life or dose related) to bacterial cell or failed to implement due to changed behavior by target B.B.Bs (due to mutation) , drug resistance would be there . This disorder could be reset by certain drugs when added with antibiotics . Resistance-modifying agents are capable of partly or completely suppressing bacterial resistance mechanisms like Beta lactase inhibitor - Cluvanic Acid or sulbactum . There are many thoughts that design the effects. These are fed thoughts and feeding was done in pre creation era by Highest center of universe . Similarly this disorder could be reset by First order (quantum entanglement) of the universe by using Atomic Genetic Engineering (AGE) technology as adjuvant therapy . The entire study comes under pharmacogenetics drug science [1]. Atomic Genetic Engineering (Quantum Entanglement - First order of Universe) could reset Divine Mechanics of drug and bacteria that cause drug resistance which has failed to follow second order of universe by virtue of mutation in bacterial cell . It is time mindness (TM) that triggers and regulate the entire phenomenon . It is prediction of theory .*

Keywords: *Basic Building Blocks , Mind and Mass , Atomic genetics , Atomic transcription and Translation , Tachyons*

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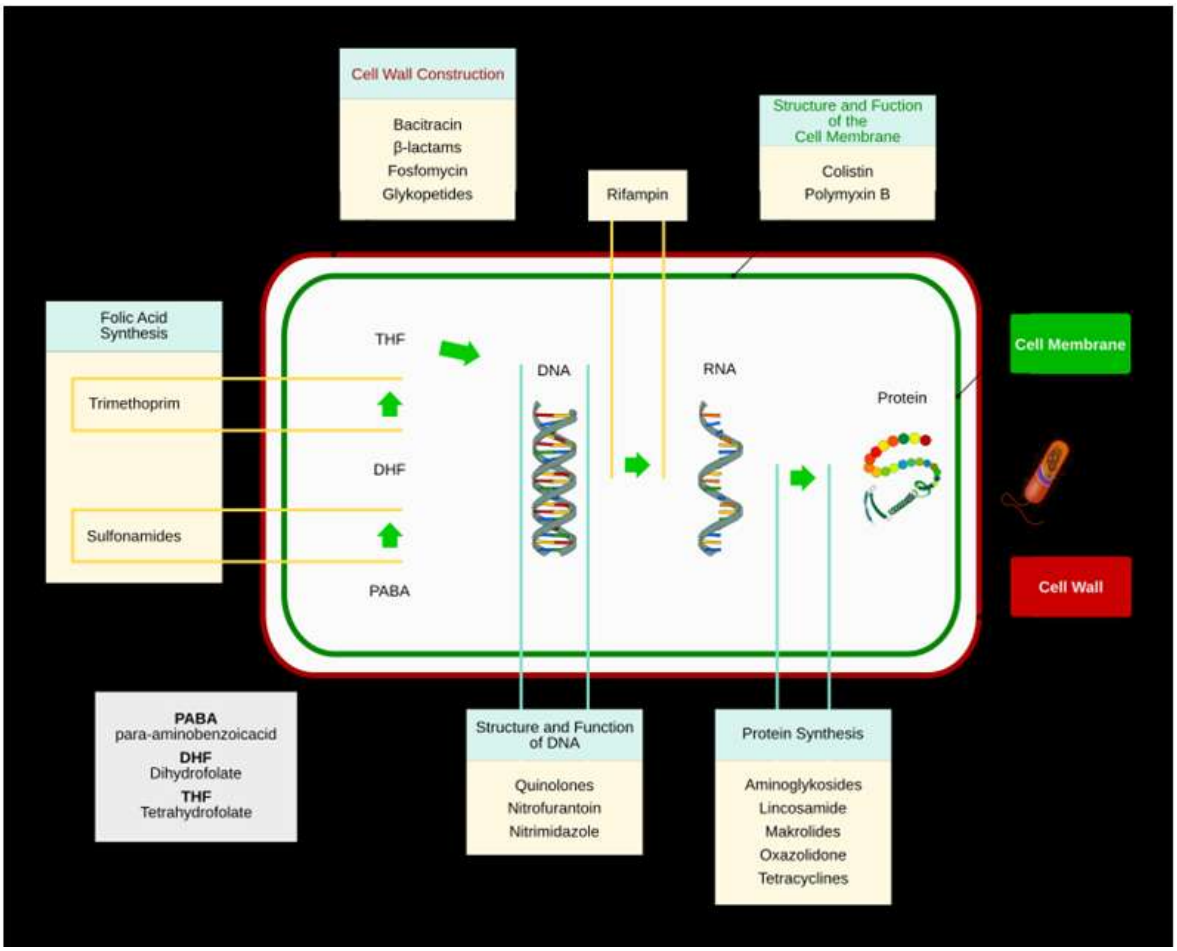
I. Introduction

1.1 Pharmacogenetics [1]

Pharmacogenetics [1] is new drug knowledge where we investigated how drug triggers divine mechanics of cell to control / uncontrol (side effects) its functions (metabolism - both normal and abnormal) and finally to set cell at changed function by second order of the universe. To understand pharmacogenetics of drug and to understand phenomenon of body structure and function (Both normal and abnormal and life and death phenomena) , we have to understand Basic Building Blocks (B.B.B – Mind – CCP , Code PcPs and CP and Mass – B.Bit -- Fig 1). Life effects are higher thought expressions (activated different Code PcPs – informations) of B.B.Bs. Formations of molecules , atoms and particles are due to lower thought expressions (Thought - CCP – Programming – Code PcPs –Interaction – Effects) . Drugs are divine units with fixed activated Code PcPs (structural Code PcPs , physical Code PcPs , chemical CodePcPs and spectral Code PcPs) . When drug enters in body it becomes life molecule and apart from physical nature it posses life consciousness that obey not only pharmacokinetics discipline rather it obeys pharmacodynamics discipline i.e drug knows where to go and where to fix itself to the cell (receptor) and how to trigger second orders of universe to modify cell functions (Pharmacogenetics). Pharmacogenetics starts after pharmodynamics (fixing with cell receptor) is over and before effects come. Pharmacogenetics is conditioned phenomenon (Dose dependent orders) or Unconditioned phenomenon (No effect seen) – i.e drug fails to order cell's Higher center . It happens only when death order – First order of universe is triggered and it orders higher center (say S,A, Node) to stop cardiac rhythmicity thoughts to trigger cardiac arrest or death . It is Time mindness (TM) that regulates and controls all orders of the universe . Without pharmacogenetics , drug science is incomplete ..

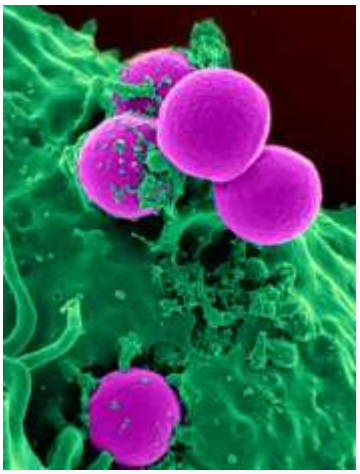
Antibiotics are commonly classified based on their mechanism of action, chemical structure, or spectrum of activity. Most target bacterial functions or growth processes.[56] Those that target the bacterial cell wall (penicillins and cephalosporins) or the cell membrane (polymyxins), or interfere with essential bacterial

enzymes (rifamycins, lipiarmycins, quinolones, and sulfonamides) have bactericidal activities. Protein synthesis inhibitors(macrolides, lincosamides and tetracyclines) are usually bacteriostatic (with the exception of bactericidal aminoglycosides).[57] Further categorization is based on their target specificity. "Narrow-spectrum" antibiotics target specific types of bacteria, such as gram-negative or gram-positive, where as broad-spectrum antibiotics affect a wide range of bacteria. Following a 40-year break in discovering new classes of antibacterial compounds, four new classes of antibiotics have been brought into clinical use in the late 2000s and early 2010s: cyclic lipopeptides (such as daptomycin), glycylcyclines(such as tigecycline), oxazolidinones (such as linezolid), and lipiarmycins (such as fidaxomicin).[58][59]



Molecular targets of antibiotics on the bacteria cell

Resistance



Scanning electron micrograph of a human neutrophil ingesting methicillin-resistant *Staphylococcus aureus*(MRSA)

The emergence of resistance of bacteria to antibiotics is a common phenomenon. Emergence of resistance often reflects evolutionary processes that take place during antibiotic therapy. The antibiotic treatment may select for bacterial strains with physiologically or genetically enhanced capacity to survive high doses of antibiotics. Under certain conditions, it may result in preferential growth of resistant bacteria, while growth of susceptible bacteria is inhibited by the drug.[62] For example, antibacterial selection for strains having previously acquired antibacterial-resistance genes was demonstrated in 1943 by the Luria–Delbrück experiment.[63] Antibiotics such as penicillin and erythromycin, which used to have a high efficacy against many bacterial species and strains, have become less effective, due to the increased resistance of many bacterial strains.[64]

Resistance may take the form of biodegredation of pharmaceuticals, such as sulfamethazine-degrading soil bacteria introduced to sulfamethazine through medicated pig feces.[65] The survival of bacteria often results from an inheritable resistance,[66] but the growth of resistance to antibacterials also occurs through horizontal gene transfer. Horizontal transfer is more likely to happen in locations of frequent antibiotic use.[67]

Antibacterial resistance may impose a biological cost, thereby reducing fitness of resistant strains, which can limit the spread of antibacterial-resistant bacteria, for example, in the absence of antibacterial compounds. Additional mutations, however, may compensate for this fitness cost and can aid the survival of these bacteria.[68]

Paleontological data show that both antibiotics and antibiotic resistance are ancient compounds and mechanisms.[69] Useful antibiotic targets are those for which mutations negatively impact bacterial reproduction or viability.[70]

Several molecular mechanisms of antibacterial resistance exist. Intrinsic antibacterial resistance may be part of the genetic makeup of bacterial strains.[71][72] For example, an antibiotic target may be absent from the bacterial genome. Acquired resistance results from a mutation in the bacterial chromosome or the acquisition of extra-chromosomal DNA.[71] Antibacterial-producing bacteria have evolved resistance mechanisms that have been shown to be similar to, and may have been transferred to, antibacterial-resistant strains.[73][74] The spread of antibacterial resistance often occurs through vertical transmission of mutations during growth and by genetic recombination of DNA by horizontal genetic exchange.[66] For instance, antibacterial resistance genes can be exchanged between different bacterial strains or species via plasmids that carry these resistance genes.[66][75] Plasmids that carry several different resistance genes can confer resistance to multiple antibacterials.[75] Cross-resistance to several antibacterials may also occur when a resistance mechanism encoded by a single gene conveys resistance to more than one antibacterial compound.[75]

Antibacterial-resistant strains and species, sometimes referred to as "superbugs", now contribute to the emergence of diseases that were for a while well controlled. For example, emergent bacterial strains causing tuberculosis that are resistant to previously effective antibacterial treatments pose many therapeutic challenges. Every year, nearly half a million new cases of multidrug-resistant tuberculosis (MDR-TB) are estimated to occur worldwide.[76] For example, NDM-1 is a newly identified enzyme conveying bacterial resistance to a broad range of beta-lactam antibacterials.[77] The United Kingdom's Health Protection Agency has stated that "most isolates with NDM-1 enzyme are resistant to all standard intravenous antibiotics for treatment of severe infections."[78] On 26 May 2016 an E coli bacteria "superbug" was identified in the United States resistant to colistin, "the last line of defence" antibiotic.[79][80]

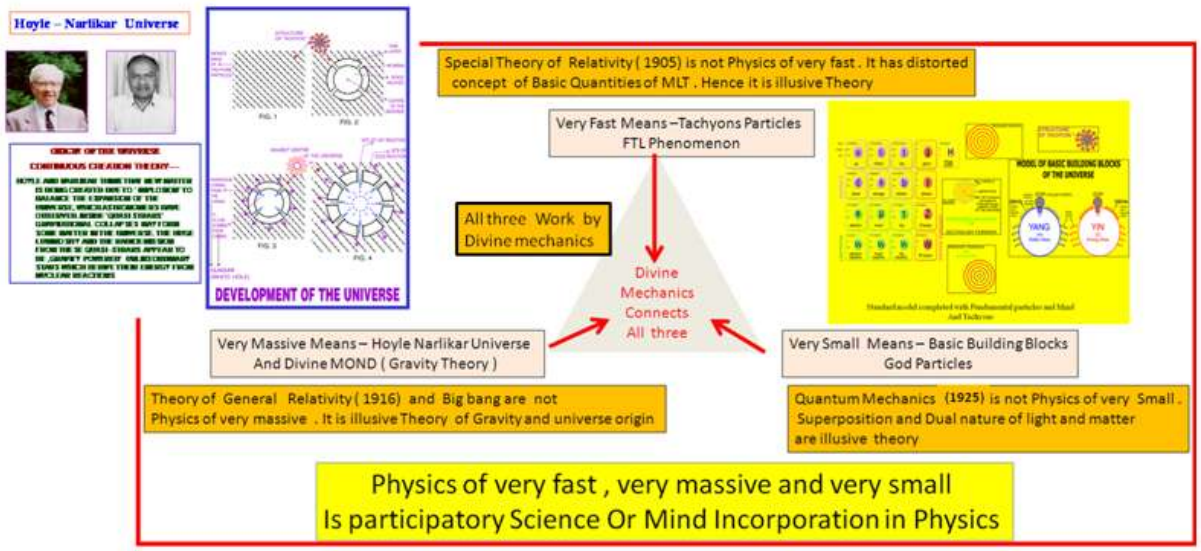
Resistance-modifying agents[edit]

One strategy to address bacterial drug resistance is the discovery and application of compounds that modify resistance to common antibacterials. Resistance modifying agents are capable of partly or completely suppressing bacterial resistance mechanisms.[139] For example, some resistance-modifying agents may inhibit multidrug resistance mechanisms, such as drug efflux from the cell, thus increasing the susceptibility of bacteria to an antibacterial.[139][140] Targets include:

- The efflux inhibitor Phe-Arg-β-naphthylamide.[140]
- Beta-lactamase inhibitors, such as clavulanic acid and sulbactam[141]

Metabolic stimuli such as sugar can help eradicate a certain type of antibiotic-tolerant bacteria by keeping their metabolism active.[142]

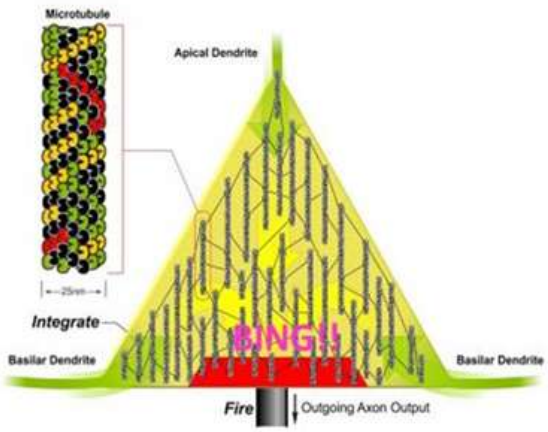
What is New Physics or Physics of Mind ?

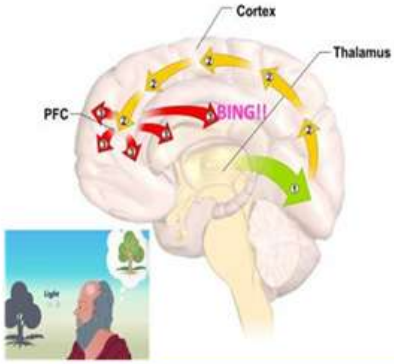


Physics of Mind

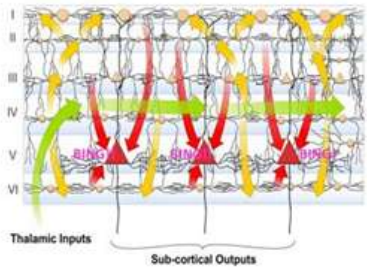
Quantum Consciousness – According to Physicists and Biologist [1]

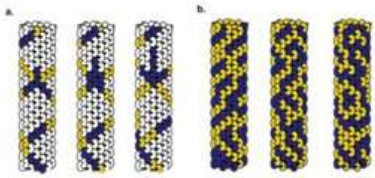
The nature of consciousness remains deeply mysterious and profoundly important, with existential, medical and spiritual implication. We know what it is like to *be* conscious – to have awareness, a conscious 'mind', but who, or what, are 'we' who know such things? How is the subjective nature of phenomenal experience – our 'inner life' - to be explained in scientific terms? What consciousness actually *is*, and how it comes about remain unknown. The general assumption in modern science and philosophy - the 'standard model' - is that consciousness emerges from complex computation among brain neurons, computation whose currency is seen as neuronal firings ('spikes') and synaptic transmissions, equated with binary 'bits' in digital computing. Consciousness is presumed to 'emerge' from complex neuronal computation, and to have arisen during biological evolution as an adaptation of living systems, extrinsic to the makeup of the universe. On the other hand, spiritual and contemplative traditions, and some scientists and philosophers consider consciousness to be intrinsic, 'woven into the fabric of the universe'. In these views, conscious precursors and Platonic forms preceded biology, existing all along in the fine scale structure of reality.





Wrong depiction of visual realization





My research involves a theory of consciousness which can bridge these two approaches, a theory developed over the past 20 years with eminent British physicist Sir Roger Penrose. Called 'orchestrated objective reduction' ('Orch OR'), it suggests consciousness arises from quantum vibrations in protein polymers called microtubules inside the brain's neurons, vibrations which interfere, 'collapse' and resonate across scale, control neuronal firings, generate consciousness, and connect ultimately to 'deeper order' ripples in spacetime geometry. Consciousness is more like music than computation.

Colleagues Travis Craddock and Jack Tuszynski and I also study how anesthetics act in microtubules to erase consciousness, and with Jay Sanguinetti, John JB Allen and Sterling Cooley, we are studying how transcranial ultrasound (TUS) can be used noninvasively to resonate brain microtubules and treat mental, cognitive and neurological disorders.

Many thanks to my assistant Abi Behar-Montefiore and Ed Xia for maintaining this website.

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1.2 Mind and Mass Realities [2]

Fig. 1 Divine Mechanics Unit – CCP, CP and information s – Code PcPs with B-Bit – Mass

Atomic genetics is the branch of science where we investigate about fundamental interactions of the universe i.e. atomic transcription and translations . New words have been coined to understand hidden science of mind part of reality. Mind reality has been recognized as different faces by "I" about 5000 years back to Arjuna in Mahabharata.as shown in Fig 1. It is just like to understand any language through Alphabets. These are (different faces) Alphabets of mind reality. One Mind reality has one face identity and the second mind reality has second face identity and so on. The facial expression represents phenomenon of intelligence and different faces represent different types of properties carrying property. The open eyes means property is activated while close eye means property is inactivated. In spite of carrying properties conscious ness they also know how to conduct not only origin of universe but also how to create two different universe i.e. next creation could be different from this creation . In all, it is automatic system of the universe . The mind realities which are of good properties have devtas face identity (first five faces on both side) and those mind realities which are of bad properties have demons face identity (last four faces on both side) . These are named as code PCPs or messenger atomic genes . The central face is CCP or Thought script where all thoughts of the universe are banked . It is bank of data of all information s of the universe It is face identity of Anti mind particles as data of all information's of the universe are stored as anti mind particles . It is the Time mind ness (biological clock) that keeps on expressing different thoughts from this thought script (CCP) . There are four more faces (black bodies) shown on extreme left and right floating in fire are CPs (translating Atomic genes) . That translates the messages and realizes it and reacts accordingly.[2]

1.3. Messages From Biological world to understand B.B.B world as shown in Fig 2 [2]

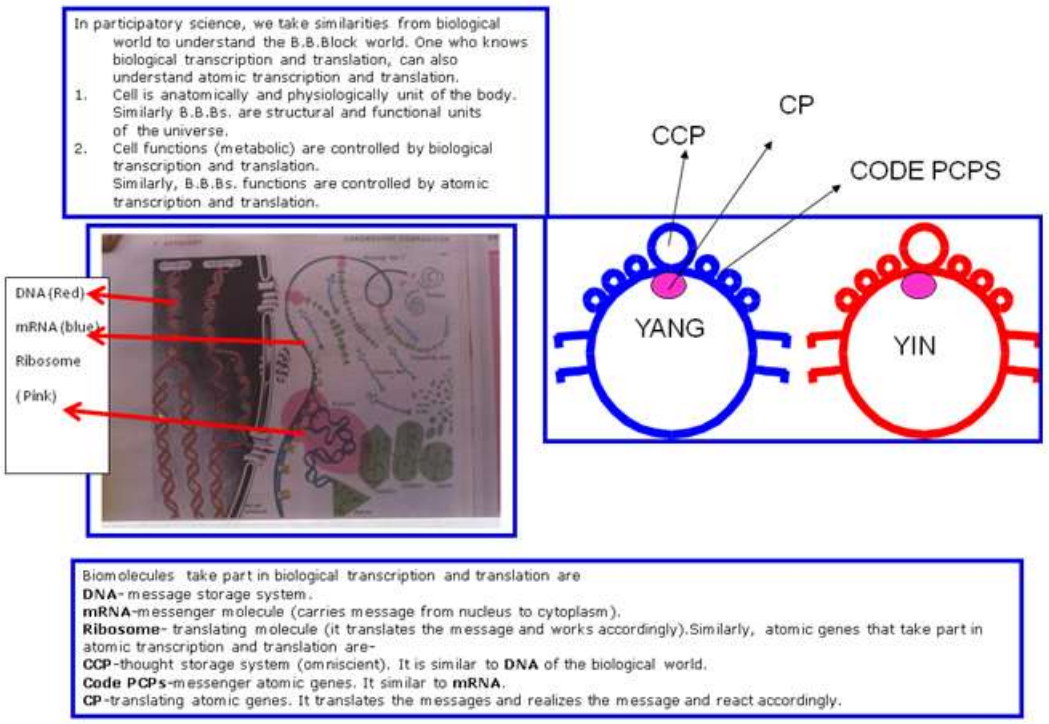
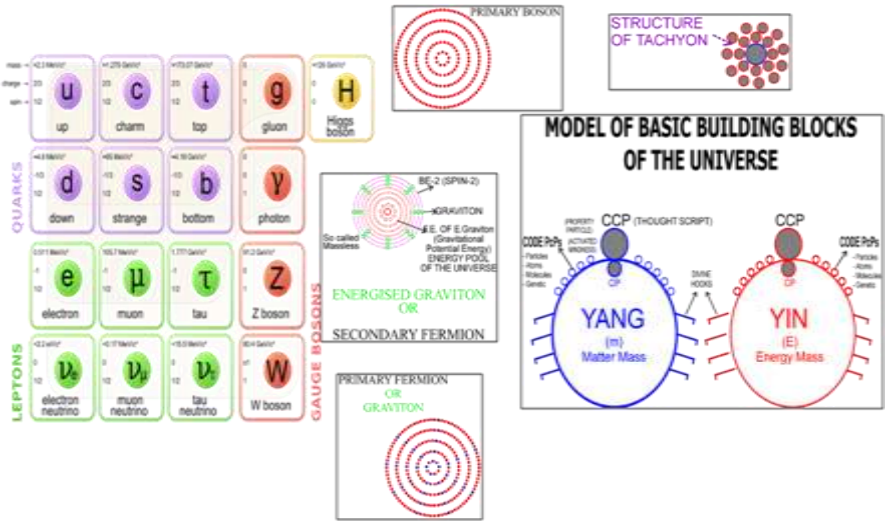


Fig. 2 Parallel teaching by participatory science

The standard model not only modified rather it has been completed [3] with introduction of energized gravitons, primary fermions, primary bosons, Basic Building Blocks, Mind and Tachyons as shown in Fig 3



Standard model completed with Fundamental particles and Mind And Tachyons

Fig. 3 standard Model chart [3]

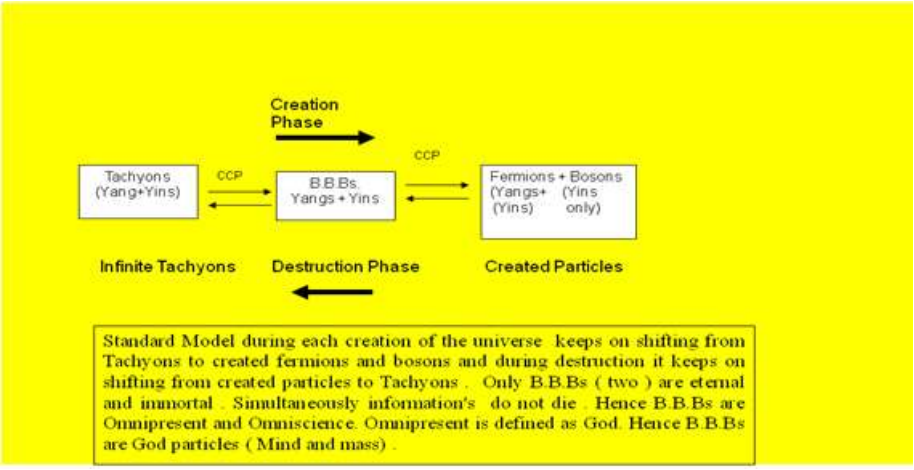


Fig .3.1 one creation and destruction cycle [4]

Participatory science has coined [5] some new words and there are some words which already exist in physics. Both these words are defined here and the definitions are according to participatory science. It has no relation with the definitions given in modern physics.

1.Mass - The part which gives shape to the smallest mass unit i.e. basics building blocks (B. B. Bs.) is called mass.

2.Matter :-Mixture of fermions and bosons or only bosons which are made up of mass (B.B.Bs) and that is why we realize their shape is called matter.

3.Inertial mass :-Mass (smallest mass units i.e. B. B. Bs) having inertial properties (classical inertia) either absolute rest or uniform motion in straight line is called inertial mass. Or Number of B. B. Bs. Per unit space present in bigger units is called inertial mass. Or Total matter contained by the bigger units or total number of fermions and bosons contained by the bigger units is called inertial mass. Therefore it never changes from place to place.

4. Gravitational mass :- The mass (interacting surfaces) which takes part in gravity (divine energized gravitons theory) interaction (which is due to mind) is called gravitational mass. When gravity interaction increases (number of divine energized gravitons increases) the interacting mass (interacting surface) also increases or when interacting mass (interacting surfaces) increases, gravity interaction (number of divine energized gravitons) also increases. Therefore gravitational mass (interacting mass or surfaces) changes from place to place. It is the fed mind that decides gravity interaction. We shall discuss it in gravity chapter and in atomic genetics.

5. Pure m (matter) mass :-matter mass (smallest mass unit or B.B.B) which have got inertial property of absolute rest.

6. Energy mass:-Energy mass (smallest energy mass unit B.B.B) which have got inertial property of uniform motion in straight line and which also gives shape to the bigger energy mass (Higgs Bosons etc) units.

7.m (impure matter) of E=mC^2 :-Matter particles (fermions) which have got spin property are called impure matter particles.

1.4 Prayer message formation in brain.

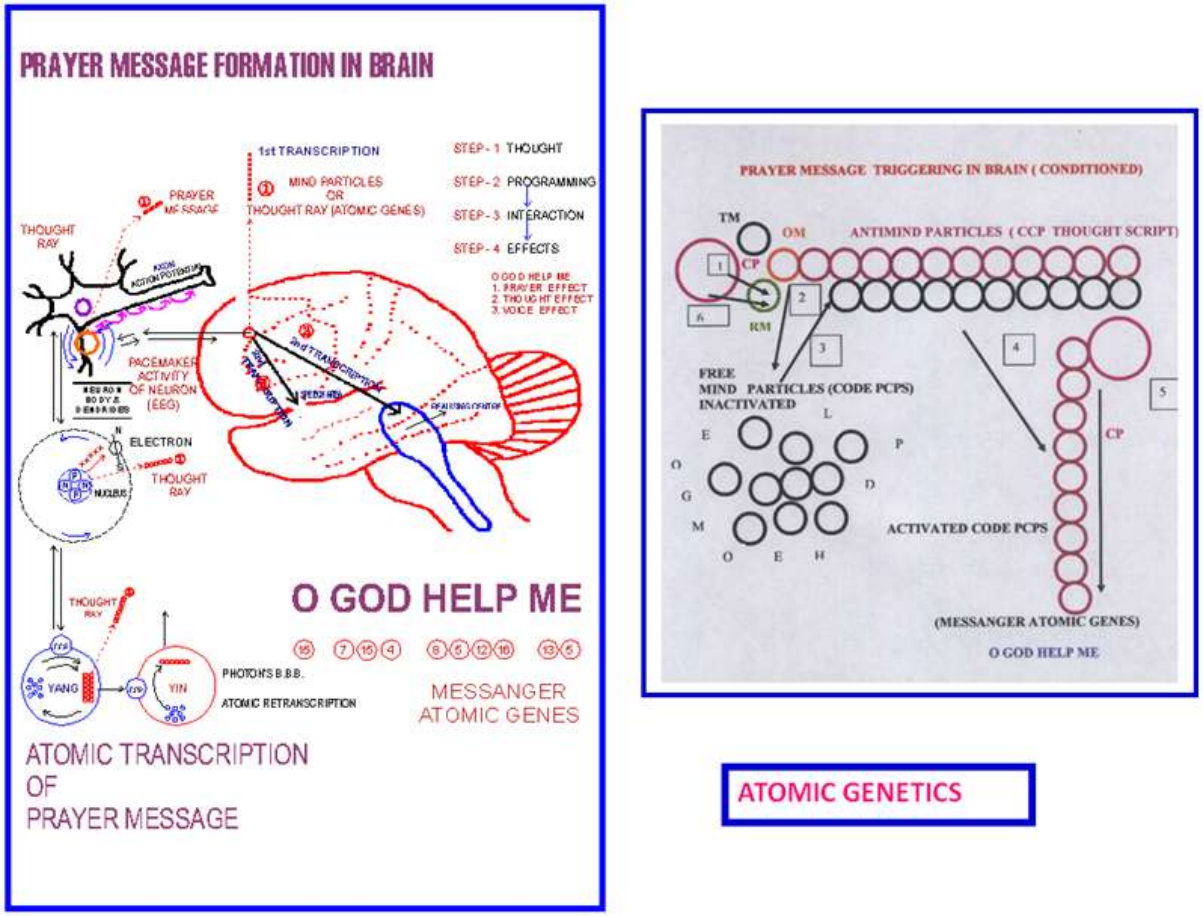


Fig 4 Divine Mechanics - Prayer message formation in brain

In atomic transcription and translation of prayer [5], following steps take place on Yang B.B.B – B-Bit as shown in Figure 4.

1.CP removes RM (repressor mindness-green) from OM (operating mindness -orange) thus induction of atomic transcription triggers.

2.OM triggers activation of free mind particles (black -inactivated code PCPs) of that thought script (magenta) of "o god help me ".

3.Free mind particles (black -inactivated code PCPs) get attached to anti mind particles script (magenta one) to form messenger thought script of "o god help me ".

4.Messenger atomic genes (black) get activated by anti mind particles thought script and further they get detached from anti mind particles thought script to form activated messenger atomic genes (activated code PCPs) (magenta) of " o god help me "

5.CP carries phenomenon of splicing by translating the messenger activated atomic genes (activated code PcPs) and finally there is activated message of " o god help me" is formed .

6.CP represses atomic transcription by adding RM (green) to OM (orange) . Thus atomic transcription gets halt.

Having formed the message it comes out in three forms .

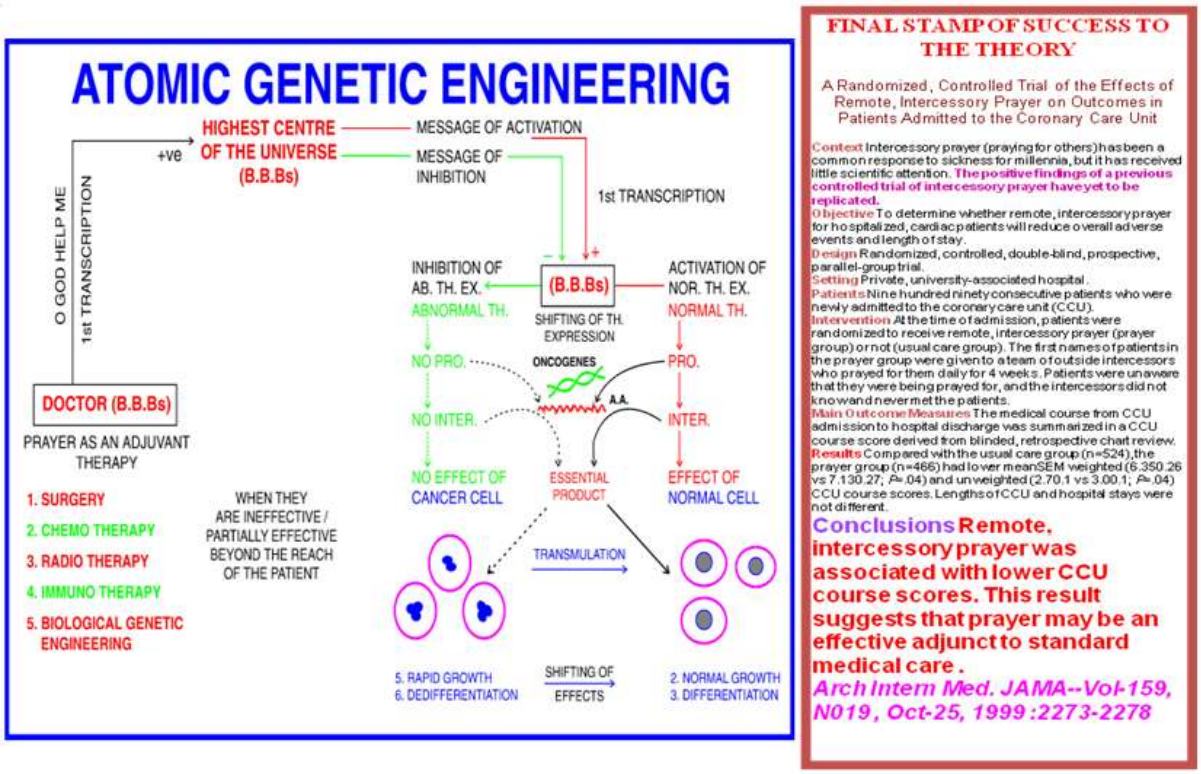
In atomic genetic engineering (prayer) we use our basic power i.e. power of B.B.Bs. Our B.B.B. (higher center) talks with highest center of the universe by sending the message by first transcription. Till today nobody knows how does the brain generate thoughts. I am going to tell you that mystery too. In the frontal lobe the neurons are responsible for thought generation. In the neuron there is electrical activity called pacemaker activity which is occurring between dendrites and the body of the neuron. The membrane of the cell is made up of atoms and atom is made up of B.B.Bs. At the level of B.B.B. say thought of **'O GOD HELP ME'** is expressed. As a result programmed messages of O GOD HELP ME (code PCPs) are formed. Out of three programmed messages, one is carried by atomic genes to highest center of the universe. It is called **THOUGHT RAY (Quantum entanglement)** which is made up of pure atomic genes and then the message goes through

phenomenon called first transcription. They come out from brain directly. The other two messages are carried by photons from nucleus of atom to electrons. Here they are modulated on electrical activity of the cell called pacemaker activity. Further they are modulated on actions potentials going towards **REALIZING CENTER** situated in brain stem (RAS) and from RAS to speech area situated in the frontal area. Target B.B.Bs. of the realizing center finally realizes thought effect of O GOD HELP ME. While from speech area message goes to motor cortex again via RAS and from there to vocal cords and finally it comes out as a speech effect of O GOD HELP ME. In layman's terminology formation of the thought ray means PRAYER as shown in Fig 4

Where Does Prayer Message go ?

Prayer message goes as shown in Figure 5 to highest center of the universe via first transcription where it is realized and it is accepted, the highest center sends two messages to B.B.Bs working as higher center in cancer cell. These messages are message of inhibition of abnormal thought expression and message of activation of normal thought expression. Having received the messages, higher center stops expressing the abnormal thoughts and it starts expressing the normal thoughts. As a result, there are no more abnormal programmed messages and in place of that normal programmed messages are there. Now the messages have shifted from abnormal (5and 6) to normal (2 and 3). This shifting of thought expression is called **ATOMIC GENETIC ENGINEERING** as shown in **Figure 4.1**

The changed messages reach to target B.B.Bs. through same route. Having received the changed messages, target B.B.Bs. stop expressing the previous programmng and they start expressing the normal programming. As a result the cancer cells transmutate into normal cells. **Or diseased cell gets cured [6]** as shown in **Fig 4.1**



FINAL STAMP OF SUCCESS TO THE THEORY

A Randomized, Controlled Trial of the Effects of Remote, Intercessory Prayer on Outcomes in Patients Admitted to the Coronary Care Unit

Context Intercessory prayer (praying for others) has been a common response to sickness for millennia, but it has received little scientific attention. **The positive findings of a previous controlled trial of intercessory prayer have yet to be replicated.**

Objective To determine whether remote, intercessory prayer for hospitalized, cardiac patients will reduce overall adverse events and length of stay.

Design Randomized, controlled, double-blind, prospective, parallel-group trial.

Setting Private, university-associated hospital.

Patients Nine hundred ninety consecutive patients who were newly admitted to the coronary care unit (CCU).

Intervention At the time of admission, patients were randomized to receive remote, intercessory prayer (prayer group) or not (usual care group). The first names of patients in the prayer group were given to a team of outside intercessors who prayed for them daily for 4 weeks. Patients were unaware that they were being prayed for, and the intercessors did not know and never met the patients.

Main Outcome Measures The medical course from CCU admission to hospital discharge was summarized in a CCU course score derived from blinded, retrospective chart review.

Results Compared with the usual care group (n=524), the prayer group (n=466) had lower mean±SEM weighted (6.350.26 vs 7.130.27; P=.04) and unweighted (2.70.1 vs 3.00.1; P=.04) CCU course scores. Lengths of CCU and hospital stays were not different.

Conclusions Remote, intercessory prayer was associated with lower CCU course scores. This result suggests that prayer may be an effective adjunct to standard medical care.

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Fig . 4.1 A.G.E and Final stamp of success to New Theory[6]

1.5 Message system of the Universe

Before the origin of the universe [**2**] nature had only one type of message systems which is called FIRST TRANSCRIPTION. Messages (Code PcPs) used to go from one B.B.B. to another B.B.B. by atomic transcription. Messages were carried by atomic genes (Code PcPs) with very very high velocity. It is the fundamental message system and it is called **Quantum Entanglement .**

After the origin of the universe, nature created atoms. It also created one more message system called SECOND TRANSCRIPTION. Here the message (code PcPs) are carried by photons from one atom to another atom with velocity of light. Thus atoms ,molecules, cells, and even individuals talk with one another

After the formation of the cell , nature created one more system called THIRD TRANSCRIPTION .Here there is a message storage system formed by DNA. There are messenger molecules called mRNA that carry message from DNA script to cytoplasm where the message (code PCPs) is read or translated by ribosome and they work accordingly. Thus the messages reach to enzymes and hormones and finally messages reach to

target units. Having received the messages, target units work accordingly. Finally life effects (metabolic) are observed.

These three types of message systems are working in the nature. These message system are being used by the nature according to nature's need. as shown in Fig 5

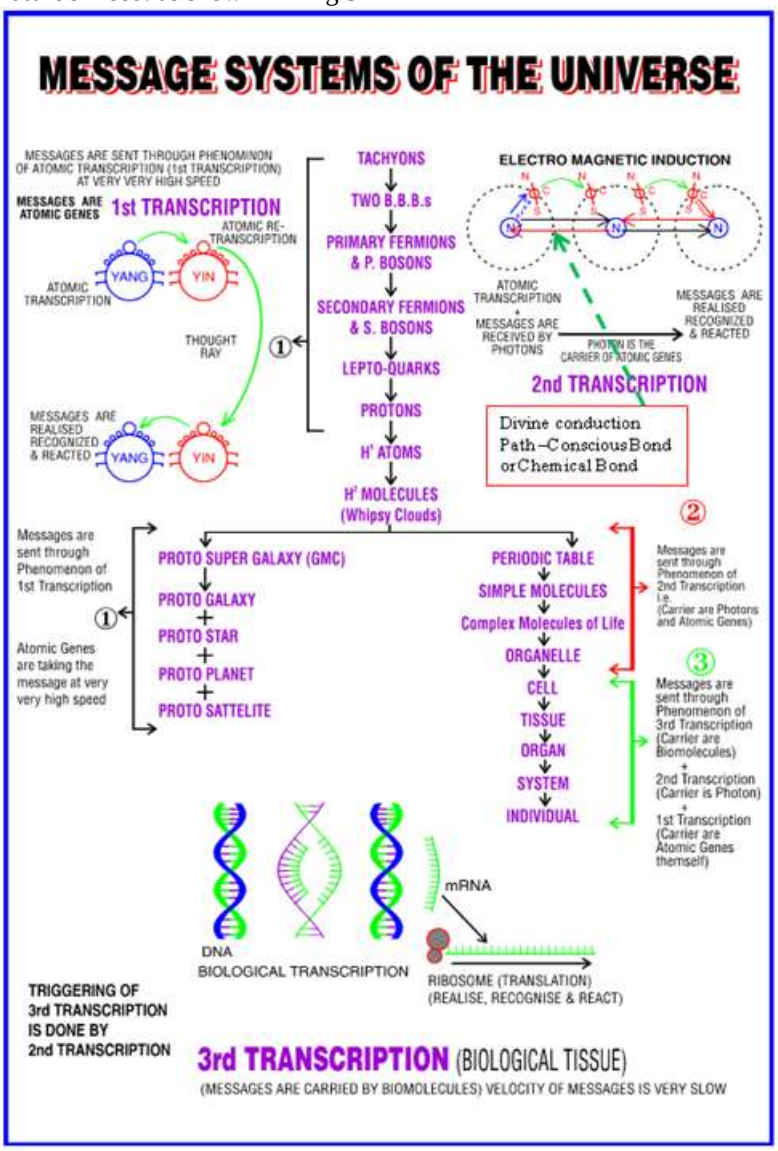

Fig. 5 Messages system of the universe

1.6 How does nature work & triggering of normal & abnormal effects

To understand creation physics as shown in Fig 6 and Fig 7. . There are two types of thought stimulation [2] . One is CONDITIONED THOUGHT STIMULATION and other one is UNCONDITIONED THOUGHT STIMULATION.

STIMULATION OF THOUGHT EXPRESSION --- There are two types of thought expressions one is CONDITIONED STIMULATION of thought expression, and other one is self stimulation of thoughts i.e. UNCONDITIONED STIMULATION of thought expression .

At the time of the origin of the universe, all effects got created. The cause of all effects of the universe is THOUGHT expression. These thought expressions were triggered by UNCONDITIONED OR SELF STIMULATED WAY . It is the first step and it is followed by PROGRAMMING or formation of programmed messages by code PCPs. This programmed message moves from higher centers to target B.B.Bs. it is called INTERACTION. Having received the messages, the mind and mass of the target B.B.Bs. work in a synchronized way so as to produced the effects as thought by a the higher center. If the thought expression by higher center is normal, the shapes, properties and laws produced by target B.B.Bs. would be normal and if the

thought expressions are abnormal, the shapes, properties and laws would be abnormal. This is the basic concept of transmutation phenomenon. Finally what we observe is called EFFECT.

Appearance of new shapes. properties and laws is called TRANSMUTATION. The first three steps are collectively called CCP. During transmutation process if CCP is written, it does mean that unless the thought, programming and interaction take place, nature cannot transmutate. Transmutation phenomenon is seen in particles, atoms, molecules and even in cells. The basic steps of any transmutation remain the same except that the thought expressions differ.

The subatomic particle are made up of more fundamental particles called Basic Building Blocks (B.B.Bs) which are made up of mind and mass. These B.B.Bs are divine in nature with the result they talk with each other by phenomenon called atomic transcription and translation (thought expressions). The triggering of broken symmetry is caused by atomic transcriptions. Unless the atomic transcriptions occur, subatomic particles could never exhibit phenomenon of broken symmetry . So the broken symmetry is never spontaneous. It is being mis understood that sub atomic particles do have spontaneous activities as far as broken symmetry is concerned. Hence the Nobel prize physics 2008 awarded to this work is too early to give prize.

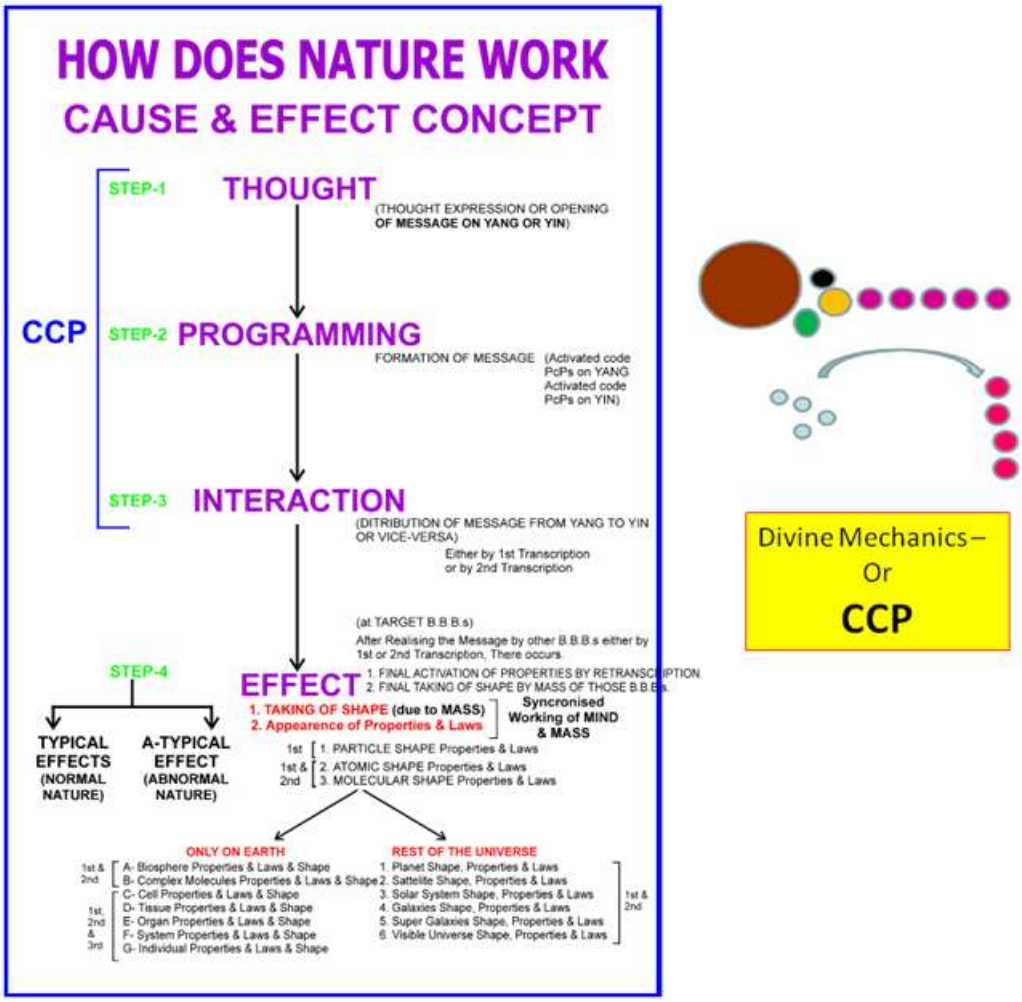

Fig. 6 Divine Mechanics – How Does Nature work ?

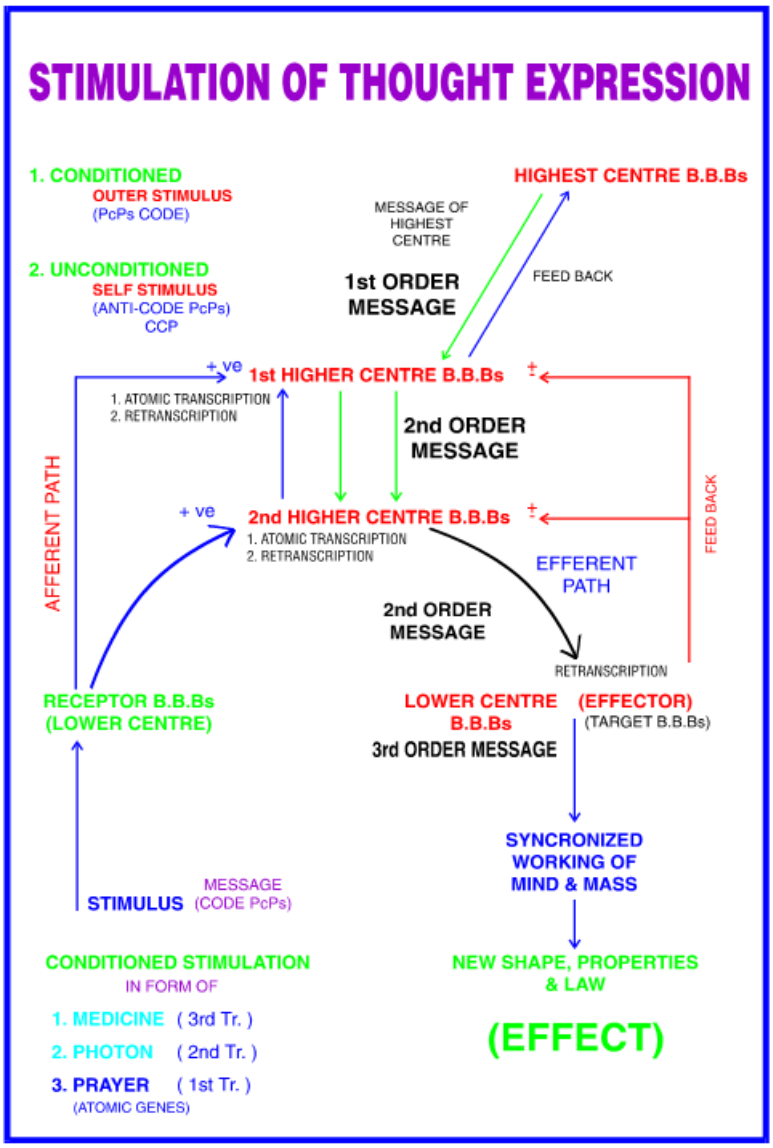

Fig. 7 Conditioned and Unconditioned thought expressions

1.7 Message network of the Universe (Feed Back Mechanism and different centers of the Universe)

With the origin of universe, nature first created primary units i.e. primary fermions (gravitation) and primary boson, these primary units are equipped with one higher center (one B.B.B.) and rest of the B.B.Bs. are working as lower centers or target B.B.Bs.[2] After primary units ,nature created secondary units i.e. secondary fermions and secondary bosons. similarly nature created tertiary units (lepto-quarks) and then quaternary units (protons& neutrons).

Each unit is equipped with higher centers, lower centers and target B.B.Bs. After quaternary units nature created atomic units, molecular units, complex molecules of life units, organelle units, cell units, tissue units, organ units, system units and individual units. Each unit is equipped wit**h** higher centers, lower centers, and target B.B.Bs. Similarly nature created satellite units, planet units, solar system units, galaxy units, super galaxy units, dark matter layer unit. These units are also equipped with higher centers, lower centers and target B.B.Bs. Thus our universe is divided into different units and each unit is equipped with higher and lower centers.

All higher centers are under control of highest center of the universe by efferent paths. This efferent path is made up of first transcription. Higher centers can send messages to highest center of the universe by afferent path or feed back path. Thus highest center of the universe is well informed about all effects of the universe. Messages can come from lower centers to higher centers and from higher centers to highest center of the universe via afferent path. The highest center of the universe can send messages to higher centers and from higher center to lower centers. There is an inter unit message network also which is made up of first, second and third transcription depending upon the nature's need. Thus the entire universe is under control of highest center

of the universe. Highest center can change any programming programmed by it during pre creation era as shown in Fig 8

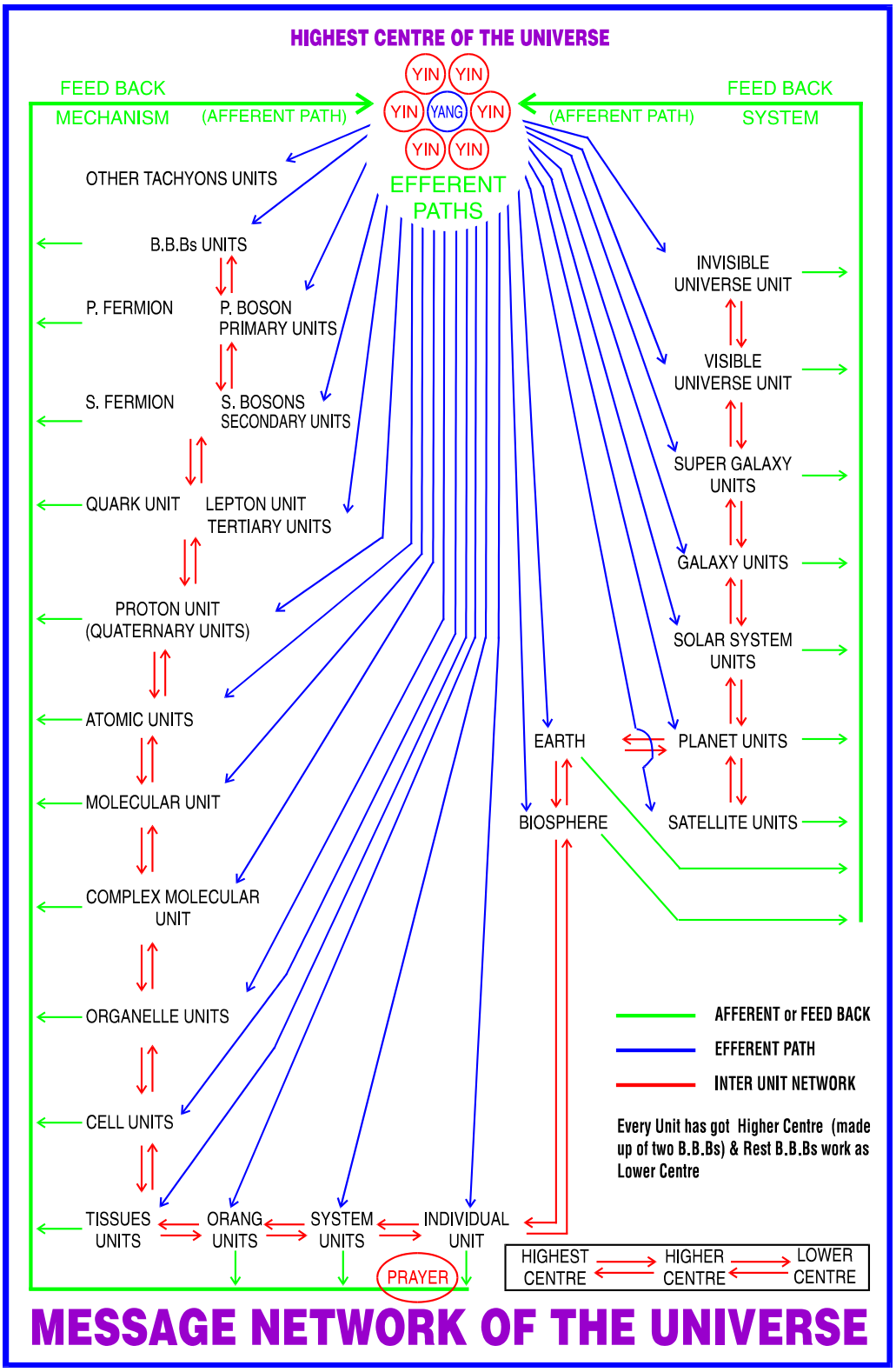

Fig. 8 Messages Net work of the Universe

1.8 Origin of the universe [7]

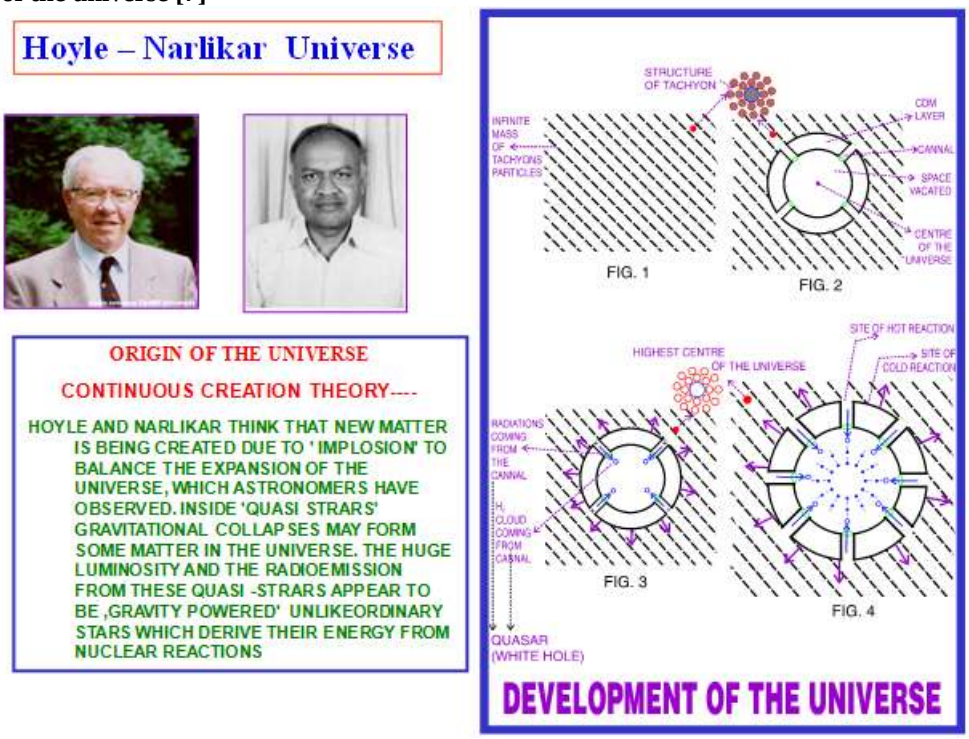

Fig. 9 Development of the Universe

Before the origin of the universe, these Basic Building Blocks (B.B.Bs) as shown in **Fig-3** were in the form of tachyons as shown in **Fig- 9 [7].** It means that at that time the tachyons were everywhere in the universe. Let us look at the structure of tachyons; it is made up of one matter B.B.B. (YANG) and many energy (**YINs**) B.B.Bs. Initially out of the infinite tachyons, one became the **highest center** of the universe. Messages used to go from highest center to rest of the universe and messages could come from rest of the universe to highest center of the universe by atomic transcription. Thus highest center had fed its thought to rest of the B.B.Bs. that would take part in creation - that they would express only those thoughts to give desired effect as wished by the highest center of the universe. So all B.B.Bs were informed about their role before creation of the universe. In pre-creation era programming of the future universe was done by highest center of the universe. Our universe is oscillating and it is a divine universe. It means that it has a creation phase and a destruction phase. During creation phase tachyons break into their B.B.Bs. and from these B.B.Bs, formation of fermions and bosons take place as shown in **Fig 3.1**. After the creation phase, destruction would start and in this phase all created particles would again break into their B.B.Bs and finally tachyons would form.

At the time of origin of the universe, all the effects got created. These effects are taking of different shapes and appearance of properties and laws. All these effects are studied in various branches of science.

With the origin of the universe, nature first created a sphere of **COLD DARK MATTER (C.D.M)** and canals in it. With the result space got created. At the other end of the canals, hot reaction started **(the relics are back ground radiations 2.7 degree K of our hydrogen clouds)**. As a result hydrogen clouds and lot of radiations were created. The empty canals were filled by these hydrogen clouds and radiations and thus **QUASARS** appeared in the universe. Simultaneously C.D.M. layer started expanding and clouds and radiations kept on coming in this closed universe as shown in **Fig 9 .** With the passage of time more and more C.D.M. layer formed, more and more quasars formed. The hydrogen cloud came in this closed universe. They started running towards C.D.M. layer as they were attracted by the gravity of C.D.M. layer. Those clouds, which were nearer, moved faster than those, which were away from CDM Layer. The **HUBBLE LAW**, can thus be explained. With some more passage of time, clouds were joined to form **GMC** (giant molecular clouds). Later by self-gravitation different proto stars, proto planets, proto satellites were formed. Finally stars became bright and thus bright galaxies appeared in this universe. Our universe is still in expansion phase and creation is still going inside quasars. It is to be remembered that highest center of the universe does not come in the visible universe. It keeps on receiving the messages by atomic transcription and it has power to change any programming programmed by it during pre-creation era.

It is the atomic genes which constitutes mind part of reality. Mind incorporation in physics is awaited as theory of every thing is not yet investigated . I have investigated theory of every thing (ToE) and I found that while studying unified theory at the time of origin of the universe by Hoyle Narlikar universe , it was mind

reality that triggered symmetry breaking phase . It is mind reality that triggers oscillation phenomenon of the universe. It is the mind reality which is behind perfect cosmological principle. There is nothing like artificial intelligence in the universe. These mind particles constitute intelligence of the universe that controls the deterministic order of the universe. It is the mind reality behind all effects of the universe. But this is not our mind rather it is mind of Almighty B.B.B (Yang B.B.B or matter B.B.B. or Male B.B.B working as Highest center of the universe) . His mind could be manipulated by prayer only. as shown in figure 9.1and 9.2

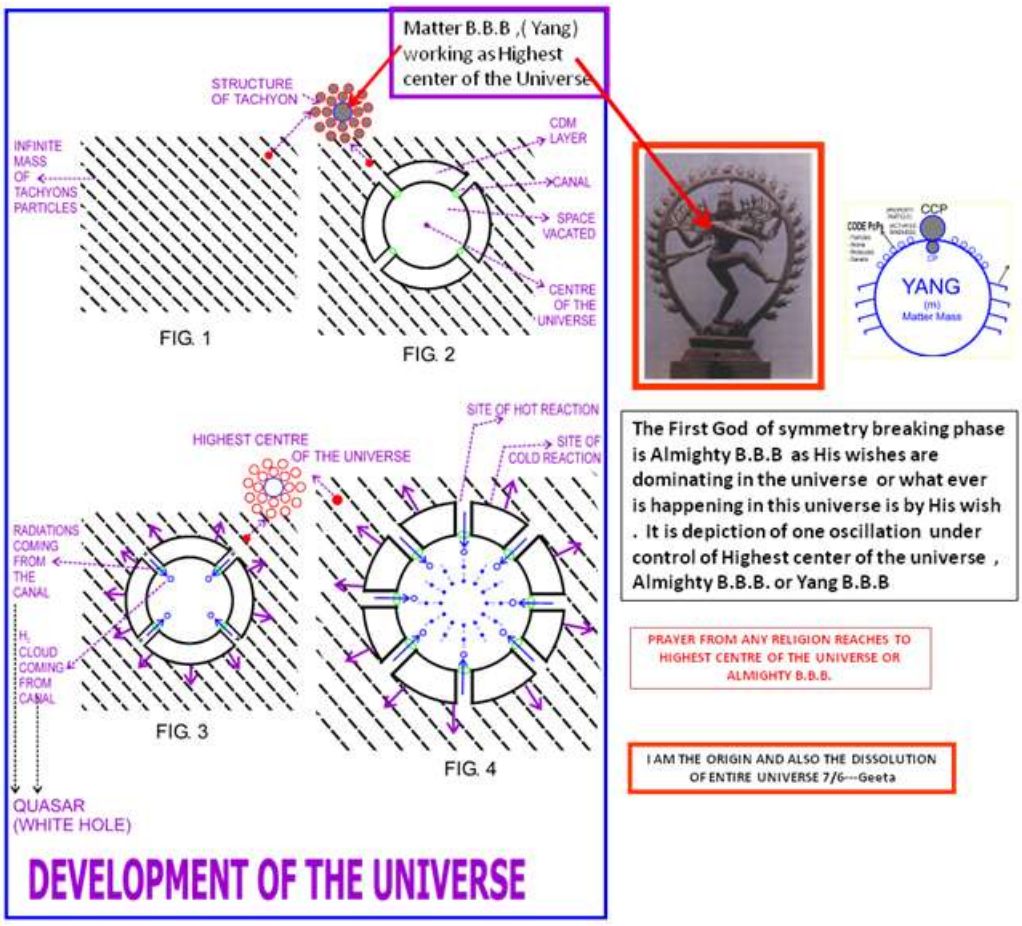

Figure 9.1 Hoyle Narlikar Universe

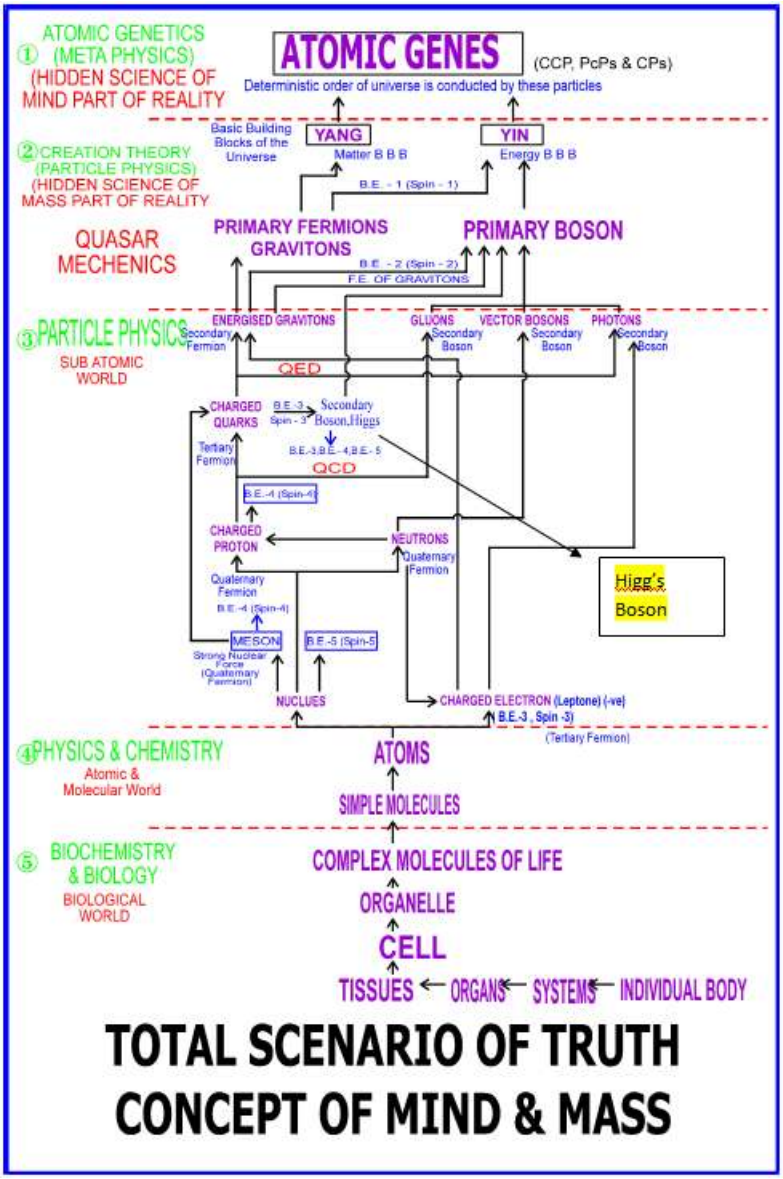

Fig 9.2 Total scenario of Truth Mind and Mass

1.9 DM and DE i.e. energized gravitons and energy pool of the universe as shown in Fig 10 and Fig 11 [8]

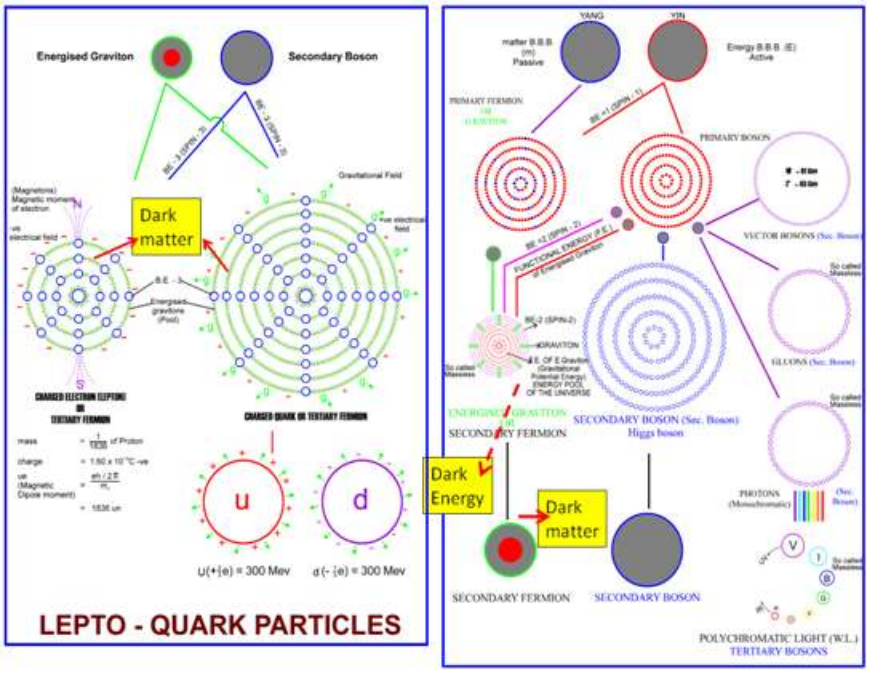

Fig. 10 Structural configuration of Leptons and Quarks as regard DM and structural configuration of DM and DE at micro level

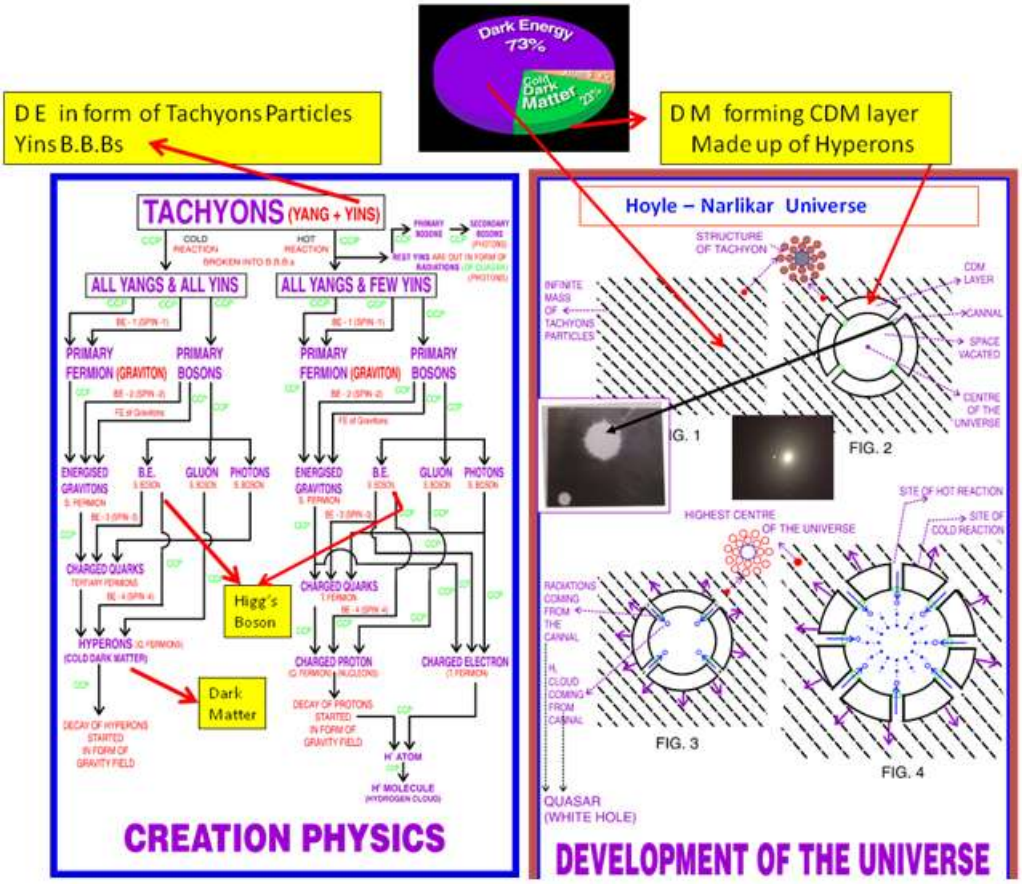

Fig 11 structural configuration of DM and DE at macro level [9]

1.10 Divine Mechanics and Brain Computation

Learning and Memory and information s (Code PcPs) Net work in Brain [10]

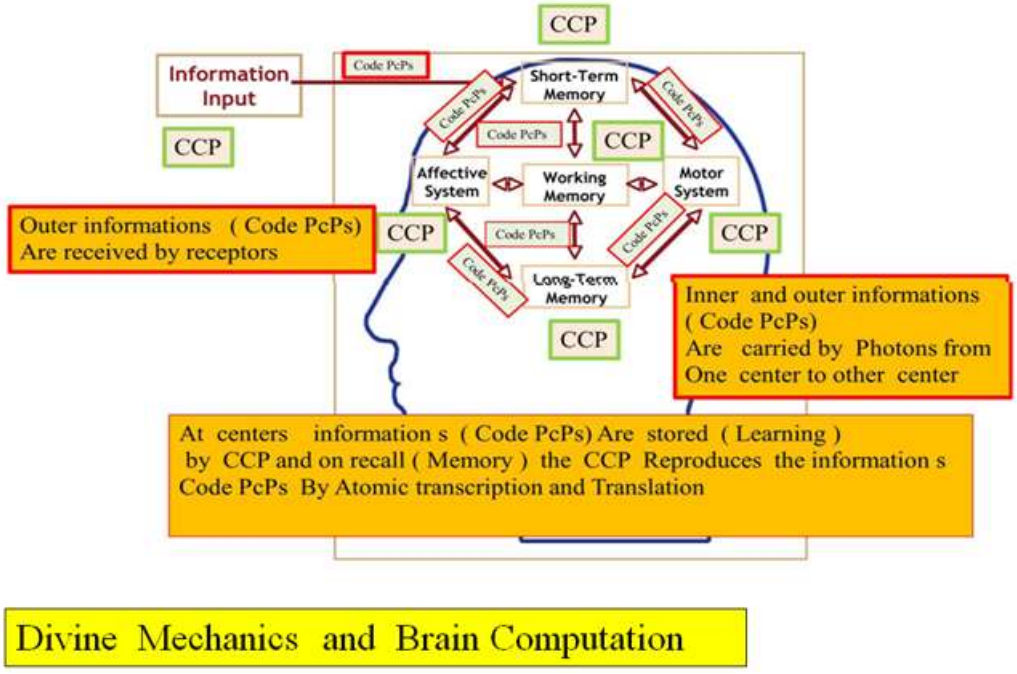

Fig 12 Divine Mechanics of Brain Computing System – Learning and memory

At all steps, these are B.B.Bs (B-Bit) that are responsible for information's control and triggering . If CCP is written it means unless atomic transcription and translation occur, brain cannot have learing and memory triggering and control and it is called Divine mechanics of Biology and it is life. These are higher thought expressions of B.B.Bs. In Bits of classical computers and qubits of QC, the thought expressions are lower and different but the mechanics (DIVINE) remains the same. as shown in Fig 12

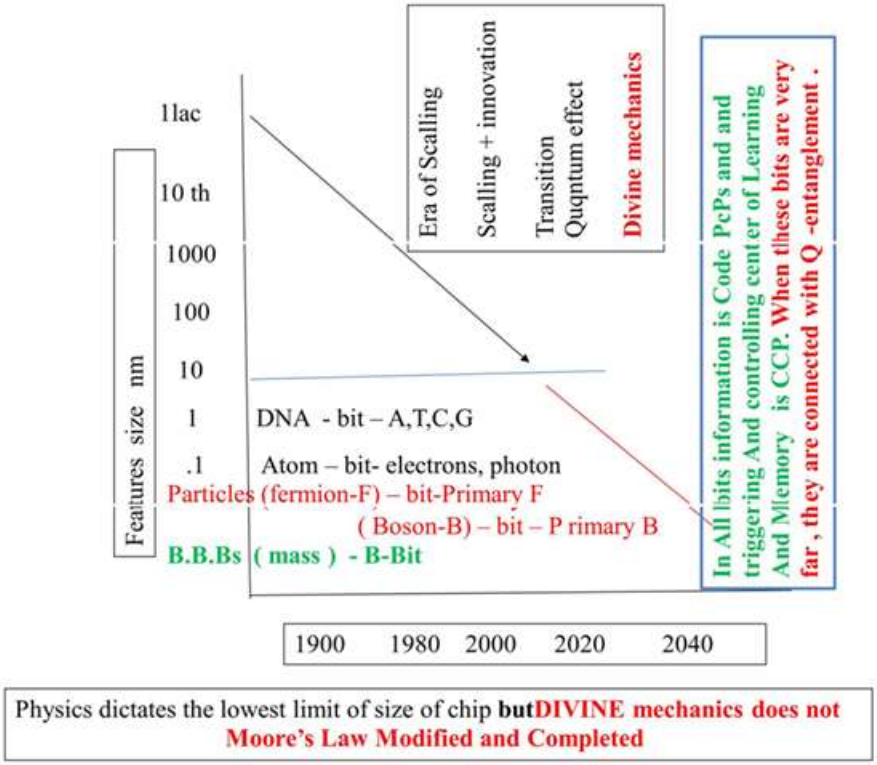

Fig 13- The Theory of Computers and Brain are same – DIVINE MECHANICS

(Structural and Functional smallest Bit (mass) is B-Bit of the universe and the Divine Mechanical Unit is CCP , Code PcPs and CP.)

DIVINE MECHANICS The Theory of Computers and Brain is SAME as shown in Fig 13

Physics dictates the lowest limits of the size of chips [**11**]. But divine mechanics does not . Hence the bit is a typical unit of information. ((Classical and Quantum Bit) is wrong and to describe informations by adding qubit is also illusion as in complicate system ,it is the different thoughts expressions that describe informations . Informations are separate (Code PcPs) and bits (Electron of transistor) qubits (electron of diamond crystal) are separate. At DNA level the bit is nucleotide (A,T,C,G) and at Atomic level the bit is photon and electron , at particle level (secondary fermions and secondary bosons) the bit is primary fermions and primary bosons and informations are still code PcPs and at B.B.B level the bit is B.B.B (Yang mass and Yin mass - B- bit) it self and the information is carried by Code PcPs but at all level of all bit units the information storage system is CCP (Physiological arrow of Time) it never dies even after destruction of the universe. Hence information s are eternal . In computers, Bit is a structural configuration that describes information s of the system and information s are in form of Code PcPs. The working of the bit is triggered and controlled by thoughts of higher centers that form that bit. If the bit is classical (transistors of chips) the effect is different . But if the bit is qubit with entanglement the working is triggered and controlled by thoughts of higher center that form that crystal . In both the effect of describing informations would be different.

One is classical (thought expressions are fixed and limited)and other one is quantum (thought expressions are changing and more wide to describe the more wide complicated system to have fast results .) . In life sciences in Brain the bit is again Photon and the information s are Code PcPs and they are triggered and controlled by CCP of higher centers that is triggered by DIVINE MECHANICS .

In life sciences the CCP (cryptography) of B-bit expresses informations (Code PcPs) which were not fed like speaking Lie and unlike Bit of classical computer and qubit of QC it shows phenomenon of forgetfulness and IQ phenomenon (mental age /real age) . It is Higher functions of B.B.B or B Bit and it is life . Hence there is difference in some aspect of DIVINE MECHANICS \of Computers computation and Brain computation

II. Structure

2.1 Modified theory of atomic structure as Niels Bohr's theory has limitations in explaining different series of hydrogen spectra.

New assumptions of structure of hydrogen atom. – Divine Modified Niels Bohr 's Theory (**Divine MONB**) as shown in Fig 15

1. Hydrogen atom is like a miniature solar system with the negatively charged electron moving in circular orbit around positively charged nucleus having positively charged particle proton. In this circular orbit or stationary orbits , electron is being held by single string of energized gravitons . This energy level is called ground state.
2. When Hydrogen atom is excited, the electron moves from ground state to excited state energy levels making a bigger circular orbit and having reached in this new energy levels , it also emits radiations . Hence absorption and emission of energy quanta occur simultaneously. Shifting is understood in inter orbital shift phenomenon.
3. When excitation is over, the electron once again come back to ground state energy level . All above mentioned events are triggered by different thoughts or atomic transcriptions from the nucleus or proton in case of hydrogen atom.
4. These atomic transcriptions are being triggered by conditioned stimulation of CCP by the message system made up of photons (second transcription or –ve charged photons or afferent path) coming from electron to proton and further they are transmitted by photons (second transcription made up of +ve charged photons, or efferent path) to electron. Having received the messages , electron works accordingly. Unless the thought is expressed, nature cannot work . as shown in Fig 15
5. When electron is in ground state, its interaction with energized graviton and electromagnetic photon is more hence IP -13.59ev (n=1) is required to remove electron from this energy level. During excited stage the interaction is different hence I.P. to remove electron from this state is -3.4ev (n=2), -1.511ev (n=3),-0.850ev (n=4) , -0.544ev (n=5) respectively. It the changing thought of interaction with energized graviton and electromagnetic photons that gives changing effect .
6. Electronic configuration of higher atoms could be known by studying I.P. curve . Working of hydrogen atom as shown in Fig 14 .

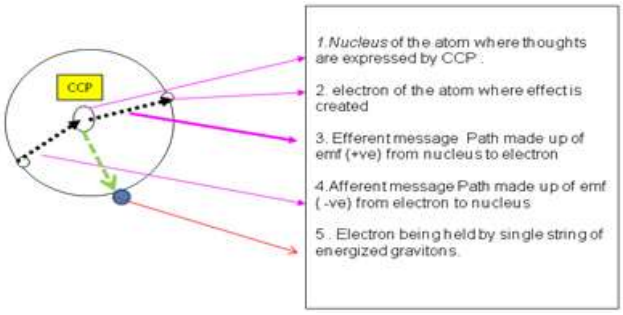

Fig 14 working of hydrogen atom – Divine MONB Theory

2.2 Conditioned stimulation of CCP and phenomenon of life.

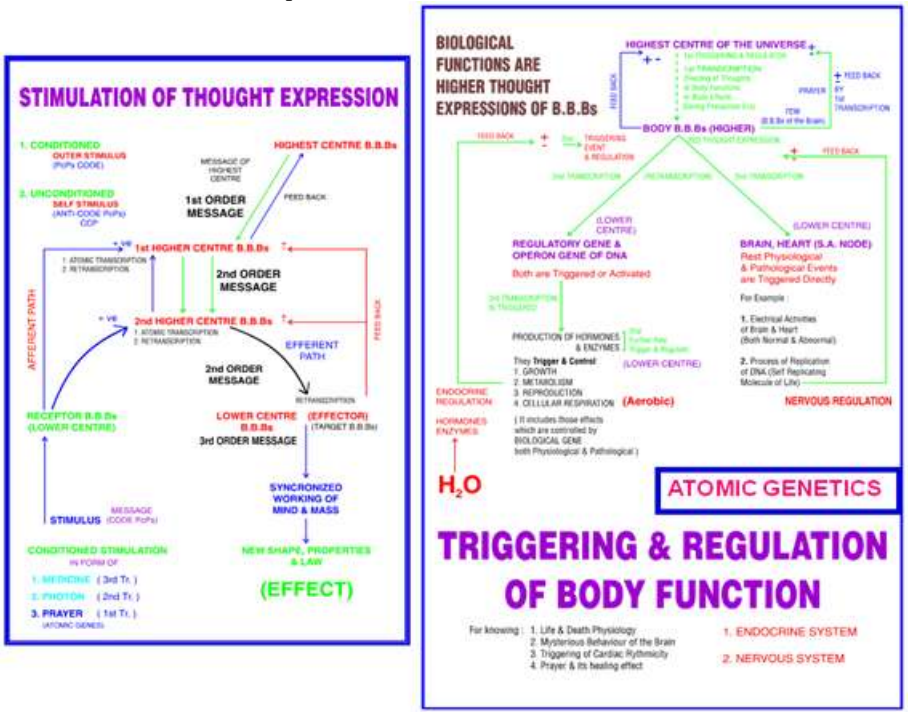

Fig 15 Stimulation of thought expression by conditioned way

Few body functions are triggered by unconditioned thought expressions like mysterious behavior of brain i.e. creative thoughts, few dreams and triggering of cardiac cycle i.e first systole in utro. Few other functions are triggered by conditioned stimulation of thoughts like life and death physiology i.e. if you add water, you would see life effect and if you remove water, you would see death effect and prayer and its healing effect.

2.3 unconditioned and conditioned stimulation of CCP in life phenomenon

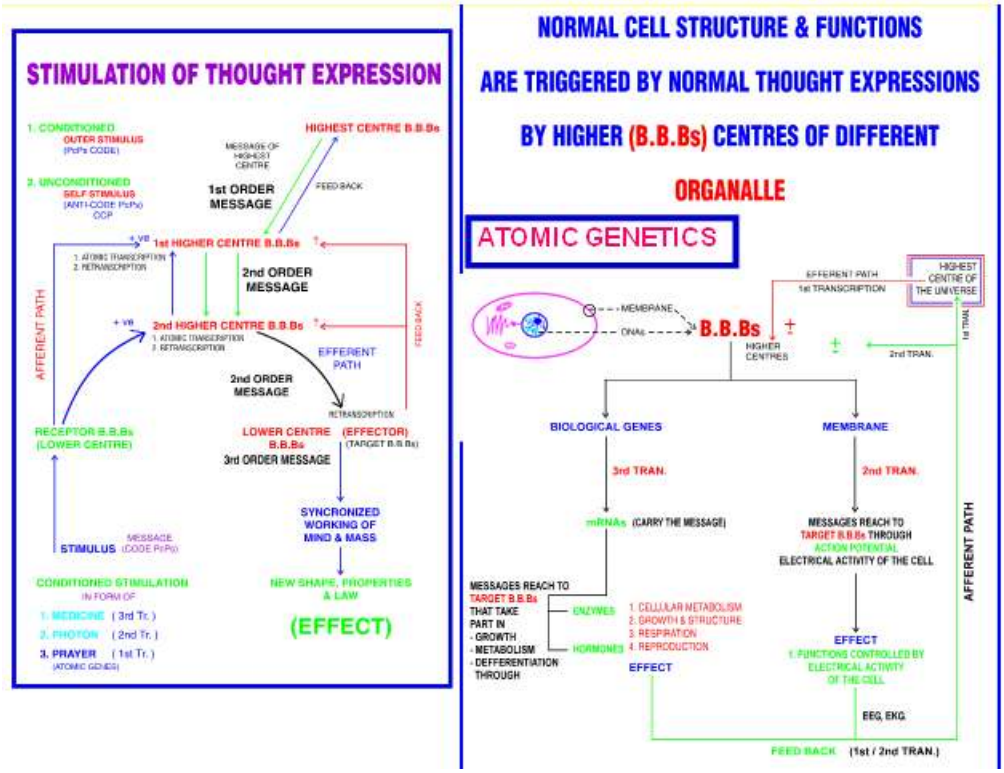

Fig 16 Stimulation of thought expression by unconditioned or conditioned way in our body

Triggering and regulation of cell functions or cell physiology as shown in Fig 16

Normal cell structure and function are triggered by atomic transcriptions or thought expressions by higher centers (B.B.Bs.) present inside the cell. These higher centers are found in **DNAs** and in the membrane. Higher centers in **DNA** express thoughts. The programmed messages (code PCP) are carried by photons from nucleus of the atom to electrons. Thus messages come on the surface of **DNA** molecule. These messages shift to mRNA during third transcription. **mRANs** carry the massages (code PCPs) from nucleus to cytoplasm where messages are translated by ribosome. Simultaneously, messages shift to peptide chains (enzymes &hormones) These enzymes and hormones carry the messages to target units (B.B.Bs.). Having received the messages, target units work accordingly and life effects like metabolism, cellular respiration, growth and structure formation, and other metabolic effects are observed. There is feedback of the effects to the higher centers. Thus triggering and regulation of metabolic functions are carried out.

Similarly, the higher centers of membranes express thoughts. These thoughts are carried by photons form nucleus to the electron of the atom. Thus messages (code PCPs) come on the surface of the membrane. Here they are modulated on pace maker electrical activity and further on action potential and thus action potential (second transcription) carries the messages from one part of the cell to the other part of the cell. This is called electrical effects (EEG and ECG) of the cell. There is feed back of these effects to higher center and thus these effects are triggered and regulated.

Both the effects (metabolic and electrical) have there feed back not only to higher centers but also to highest centers via first transcription. So highest centers is well informed about all effects of the cell. Higher centers are not autonomous centers rather they are also under control of highest centers of the universe.

2.4 Body functions and their regulation by different centers

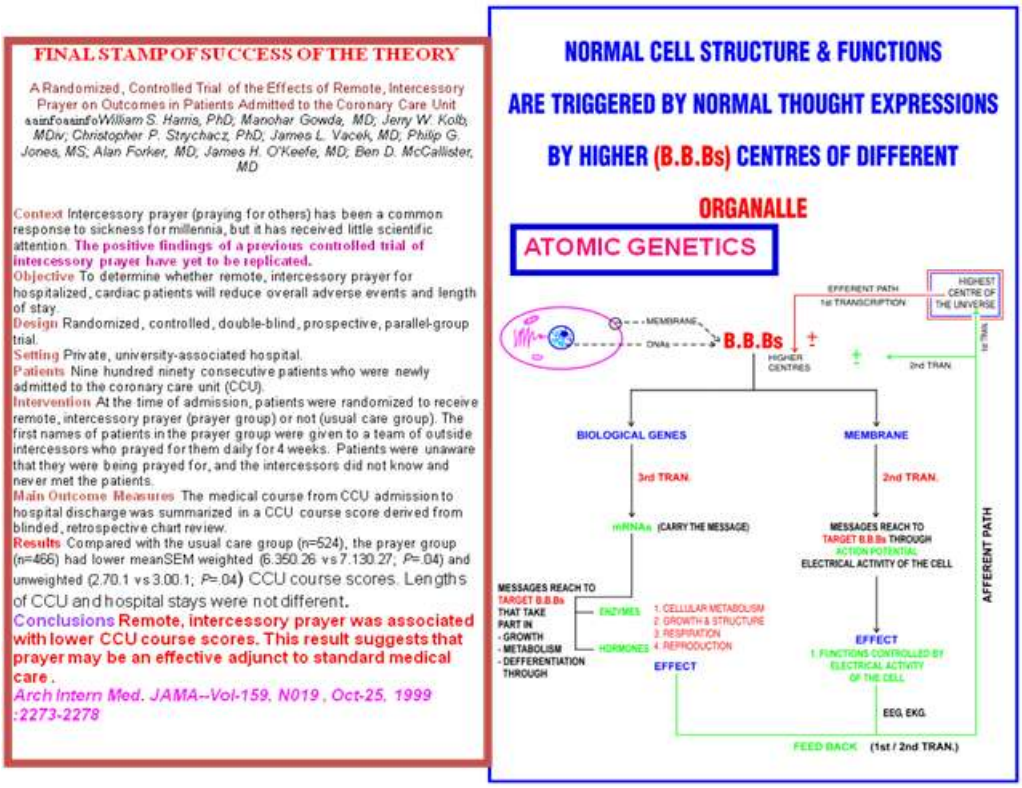

Fig 17 Unconditioned and conditioned stimulation of thought expression

Our all body functions are triggered by different thoughts expressions by different higher centers present in biological genes or DNA or in membrane of the cell. Thus different life effects are seen. There is feed back to Highest center of the universe. That controls higher centers of our body. Prayer triggers this highest center to have its healing effect. Highest center is situated in invisible universe beyond CDM layer. Concept of SOUL is myth.

2.5 Conditioned stimulation of CCP phenomenon of Germination

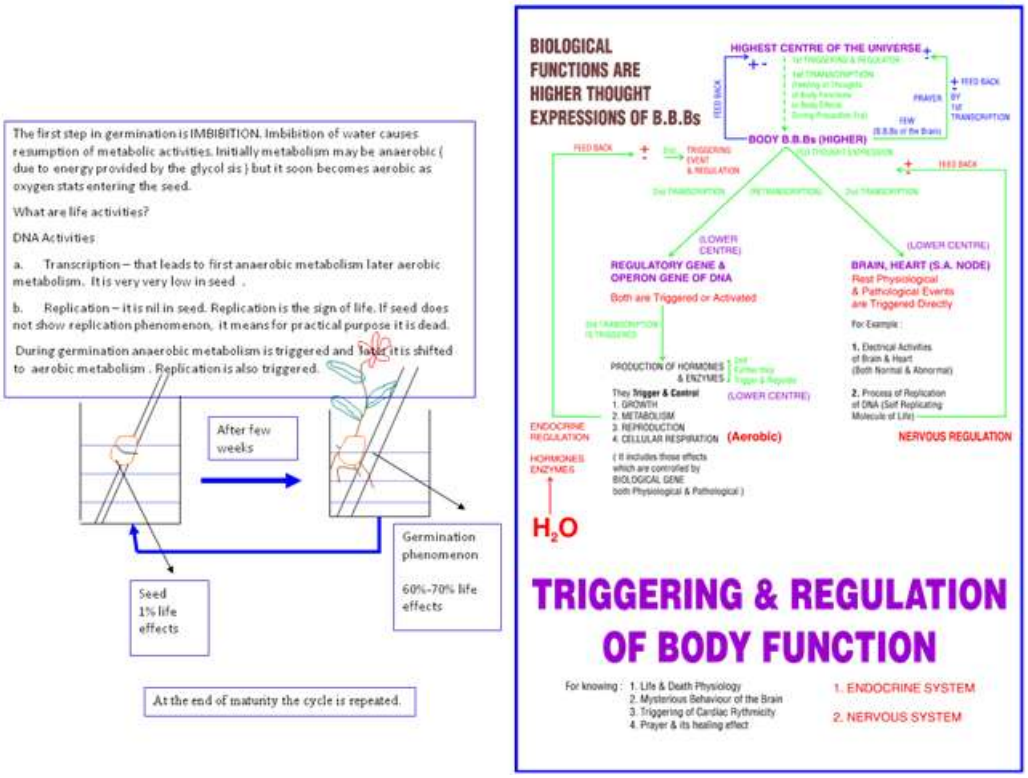

Fig18 Conditioned stimulation of thought expression

Few functions are triggered by conditioned stimulation of thoughts like life and death physiology i.e. if you add water, you would see life effect and if you remove water, you would see death effect.

Seed is alive (say having 1% of life activity) but hardly show any sign of life. It has low water content and exhibits virtually no metabolic activity. Such quiescent seeds can live for many years but germinate when soaked in water under suitable temperature and in presence of oxygen. Metabolic activity (anaerobic) are very low in seeds. Metabolic activities come to visually standstill as the seed coat becomes increasable impermeable to oxygen and moisture.

The first step in germination is IMBIBITION. Imbibition of water causes resumption of metabolic activities. Initially metabolism may be anaerobic (due to energy provided by the glycol sis) but it soon becomes aerobic as oxygen stats entering the seed.

What are life activities?

DNA Activities

a. Transcription – that leads to first anaerobic metabolism later aerobic metabolism. It is very very low in seed .
b. Replication – it is nil in seed. Replication is the sign of life. If seed does not show replication phenomenon, it means for practical purpose it is dead.

During germination anaerobic metabolism is triggered and later it is shifted to aerobic metabolism . Replication is also triggered. The triggering of both the activities is onset of atomic transcriptions of replication as well as onset of transcriptions of aerobic metabolism. With the result messages come on the surfaces of DNAs and during 3rd transcriptions they are shifted to different mRNAs and finally they reach to different enzymes and hormones. These enzymes and hormones carry messages to target units . With the result we observe phenomenon of germination. The entire working has been depicted by line diagram **(Fig –19)**. These atomic transcriptions are stimulated by water that is why it is **CONDITIONED STIMULATION of CCP.** The percentage of thought expressions increase with the time and we observe increase in the number of effects. At present we can say that plant is showing from 1 % to 20% or 40% of its effect till it reaches its maturity . At maturity the plant exhibits all effects and at that time we can say it is expressing 100% life atomic transcriptions. With the formation of new seed life atomic transcriptions once again reduced to 1% only. This cycle i.e. going from 1% life effects or atomic transcription to 100% life effects and coming back to 1% again is being visible

to us at present. When water is withdrawn, it leads to suppression of life thought expressions and death thought expressions are triggered with the result we observe death effects of plant.

CONCLUSION OF THE EXPERIMENT- The phenomenon of life effect is triggered by atomic transcription of life. Unless life atomic transcriptions are triggered, life effects are not visible. So life effects are nothing but higher thought expressions of basic building blocks. Phenomenon of death is triggered by death atomic transcription. At the time of death life thought expressions are inhibited and death thought expressions are triggered with the result we observe death effects. Water only stimulates life thought expressions that leads to triggering of different life activities (metabolic , replication and other electrical activities) in side the cell.

Being a scientist, one must know how do life effects come about. Life effects are higher thought expressions of B.B.Bs. Formation of particles, atoms and molecules are due to lower thought expressions of B.B.Bs. But their higher thought expression lead to appearance of all life effects. One who knows properties of B.B.Bs. and atomic genetics, can understand how life effects are triggered. There is nothing like **SOUL**. **It is a myth** that when soul goes inside we get life and when it moves out we are dead. When thought expressions of life are suppressed and thought expressions, of death are triggered, we observe death effects. So life effects are basically triggered by atomic transcription occurring on B.B.Bs.

2.6 Laws on which atomic genes work.

Atomic Genetics and genetic damage as shown in figure 19

Having read **Atomic genetics and Basic etiology of the cancer,** now we discuss how atomic genetics trigger molecular effects like genetic damage etc during carcinogenesis. **ATOMIC GENETICS** is a new branch of science in which we study about **Laws**, **PROPERTIES** and **FUNCTIONS** of atomic genes. Now I shall highlight the laws on which atomic genes work. As we have seen that Gregor Mendel had made three laws of inheritance known as ---

1.The principle of Dominance---- Mendel therefore concluded that what were transmitted from parent to offspring were discrete factors. Each factor contained information about the form of the trait. The factor associated with the form which was expressed in the hybrid offspring (F1) was **DOMINANT**. For example, the factor for yellow seed color was a dominant factor. The factor associated with the form which remained hidden in the **F1** but reappeared in the **F2** was **RECESSIVE**, Thus the factor for the green seed color was recessive. Mendel's factor is now recognized as the gene.

In atomic genetics this law is interpreted like this ---- During atomic transcription or thought expressions few thoughts are expressed and these are called dominating thoughts and rest thoughts which are not expressed are called recessive thoughts. Thus particles, atoms and molecules show only those properties which are triggered by dominating thought expressions. **During transmutation, dominating thoughts get recessive while recessive thoughts get dominant.** Single effect or property is triggered by single dominating thought expression or atomic transcription.

For example-- Normal genetic arrangements are triggered by normal arrangement genetic dominating thought expressions while genetic damage is triggered by abnormal arrangement genetic dominating thought expressions. Abnormal arrangement genetic thought expression is triggered by carcinogens only after normal arrangement genetic thought expressions get recessive. Or we can say normal genetic arrangement are triggered by normal arrangement genetic thought expressions and when carcinogens come in contact with the cell, they shift the thought expression from normal to abnormal genetic arrangement. With the result we observe genetic damage. Thus carcinogens suppress dominating normal arrangement genetic thoughts and they trigger abnormal arrangement recessive genetic thoughts. With the result genetic damage is seen.

2.The Principle of Segregation---The principle of segregation states that allele pairs separate or segregate during gamete formation, and the paired condition is restored by random fusion of gametes during fertilization. In atomic genetics it will be interpreted like this --- There is separate atomic transcription for separate effects. So if there are hundred effects, they all are triggered by hundred separate thought expressions or atomic transcriptions.

For example --- Genetic damage is triggered by separate thought expression and rapid growth is triggered by separate thought expression and dedifferentiation is triggered by separate thought expression. And all these thought expressions are triggered by carcinogens. **But the timing of thought expressions is different.** Genetic damage thought expression is first to trigger and later rapid growth and dedifferentiation thought expressions.

3. The Principle of independent assortment---- The principle of independent assortment states that if we consider the inheritance of two or more genes at a time, there distribution in the gametes and in the progeny of subsequent generations is independent of each other.

In atomic genetics it will be interpreted like this--- In one phenomenon, if two or more than two transcriptions or thought expressions are expressed that will give rise to two or more than two effects, **it dose not mean that there expression is DEPENDENT ON EACH OTHER.**

For example --- In phenomenon of carcinogen sis, there is effect of genetic damage, there is effect of rapid growth and there is effect of dedifferentiation. All these effects are triggered by separate thought expressions. The simultaneous expression of these atomic transcriptions is independent of each other. Or these thoughts expressions are the part of the carciogenesis phenomenon but their expressions are independent of one another. **It does mean that for rapid growth thought expression, genetic damage thought expression is NOT essential.**

Having informed about the laws and working of atomic genes, Now we discuss phenomenon of carcinogen sis, which is triggered by, outer stimuli like physical, chemical carcinogens or virus or they are self stimulated i.e. hereditary factors.

When Genetic damage is not essential for cancer transmutation then what is the role of Genetic damage? . **Why do carcinogens trigger genetic damage first? How do carcinogens trigger carcinogenesis?**

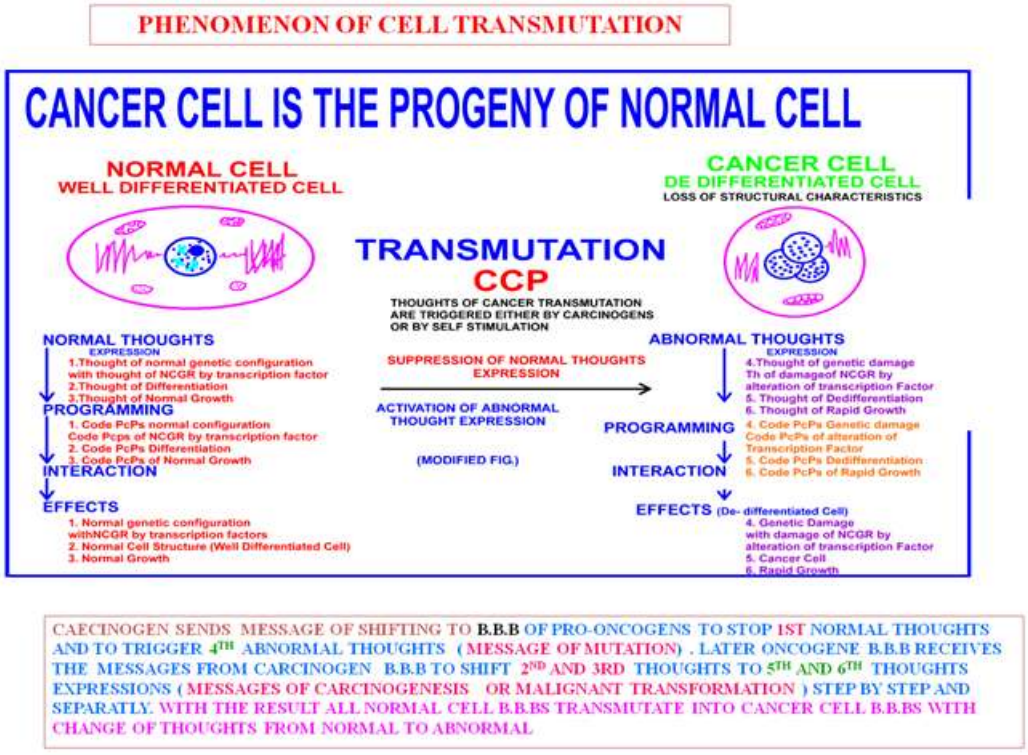

Fig 19 phenomenon of cell transmutation into cancer

STIMULATION OF THOUGHT EXPRESSION --- Shifting of thought expressions from normal to abnormal could be triggered by either from outer stimuli i.e. **CONDITIONED STIMULATION** of thought expression, which is caused by physical and chemical carcinogens or viruses etc or it is self stimulate i.e. **UNCONDITIONED STIMULATION** of thought expression , which is caused by hereditary factors.

2.7 Unconditioned and conditioned stimulation of CCP in life phenomenon

Why do patients die during treatment ? We doctors do not know physiology of Life and Death till date. We are made up of B.B.Bs (Basic Building Blocks). S.A node That controls heart rhythms is under control of Almighty B.B.B (Highest center of the universe). During death, First order of universe comes from Highest center to Higher center of S A Node to stop thought expression of cardiac rhythms (Thought of SDD, Depolarization, Repolarization) with the result heart stops and patients die .

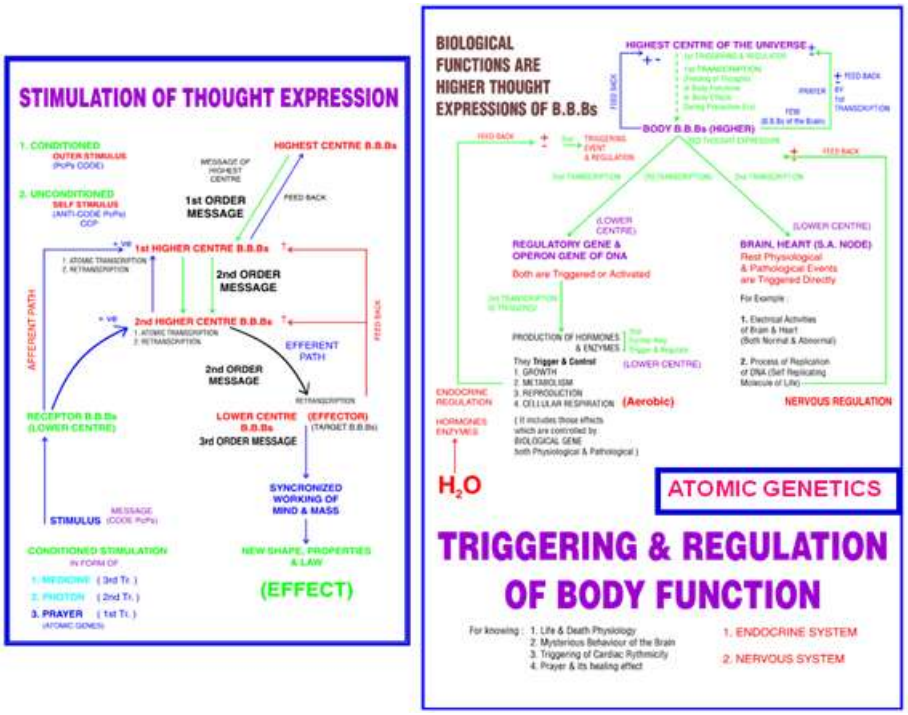

Fig 20 Stimulation of thought expression by conditioned way

2.8 Body functions and their regulation by different centers

If we know science of prayer, we could stop this message of death and we could prevent death during treatment.

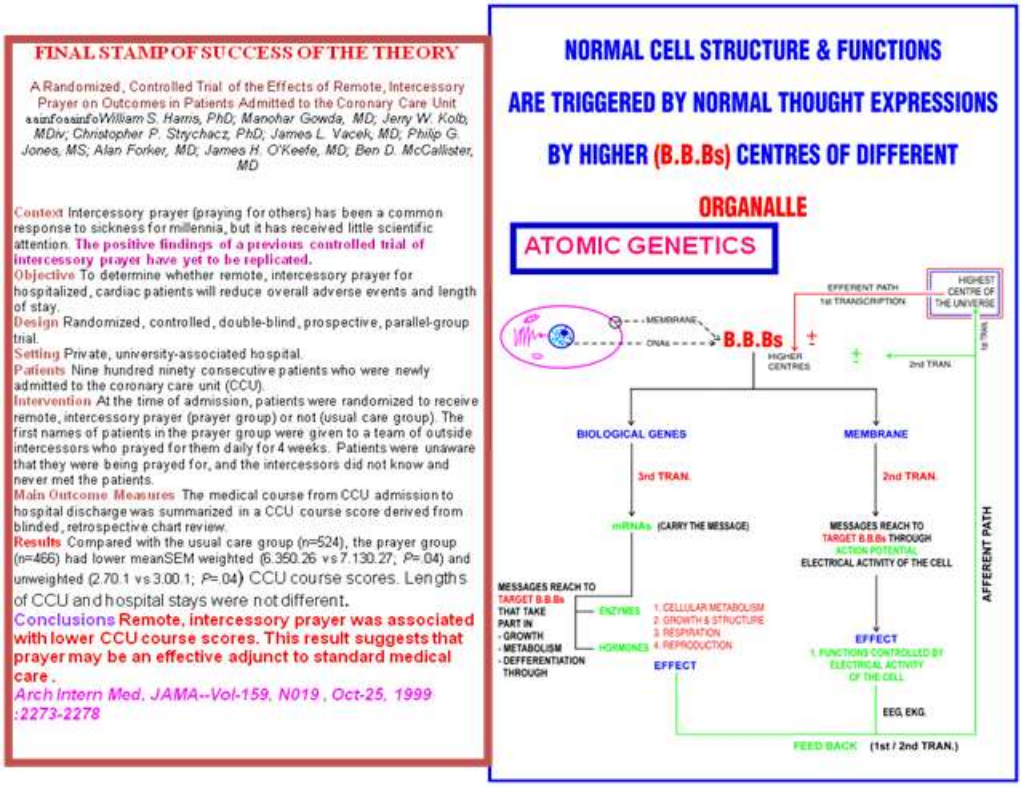

Fig 21 Stimulation of thought expression by unconditioned and conditioned ways

Our all body functions are triggered by different thoughts expressions by different higher centers present in biological genes or DNA or in membrane of the cell. Thus different life effects are seen. There is feed back to Highest center of the universe. That controls higher centers of our body. Prayer triggers this highest

center to have its healing effect. Highest center is situated in invisible universe beyond CDM layer. Concept of SOUL is myth.

It looks that there is lack of facilities (say oxygen) given by doctor to the patients but it is all illusions . We live in a tank of Oxygen (atmosphere) and if it does not short then what ever is happening to body (fall in Po2 %) is due to First order of universe that causes death of Patients (multi factorial causes by order of Highest center of the universe). To prevent triggering of death by First order of the universe, Almighty B.B.B has made prayer (Quantum entanglement - talking of two B.B.Bs) to talk to Him . If prayer is accepted , death could be prevented during treatment despite of lack of facilities .

2.9 In atomic transcription and translation of bactericidal / bacteriostatic Activity

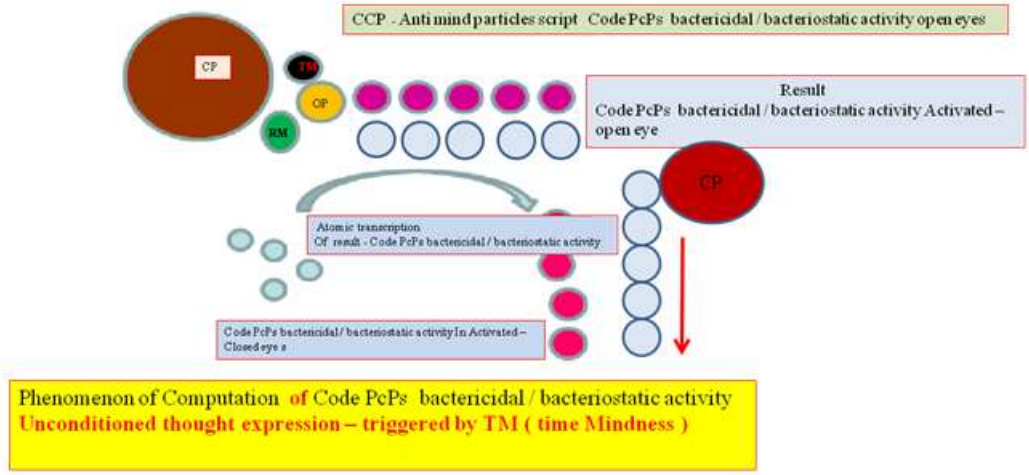

Fig 22 In atomic transcription and translation of bactericidal / bacteriostatic activity (micro level)

In atomic transcription and translation of bactericidal / bacteriostatic activity , following steps take place on Yang B.B.B – B-Bit (Figure 22, Fig 23) of Antibiotic drug .

1.CP removes RM (repressor mindness-green) from OM (operating mindness -orange) thus induction of atomic transcription triggers.

2.OM triggers activation of free mind particles (black -**inactivated code PCPs**) of that thought script (magenta) of **"**bactericidal / bacteriostatic activity ".

3.Free mind particles (black -**inactivated code PCPs**) get attached to anti mind particles script (magenta one) to form messenger thought script of **"** bactericidal / bacteriostatic activity ".

4.Messenger atomic genes (black) get activated by anti mind particles thought script and further they get detached from anti mind particles thought script to form activated messenger atomic genes (activated code PCPs) (magenta) of "bactericidal / bacteriostatic activity "

5.CP carries phenomenon of splicing by translating the messenger activated atomic genes (activated code PcPs) and finally there is activated message of " bactericidal / bacteriostatic activity " is formed .

6.CP represses atomic transcription by adding RM (green) to OM (orange). Thus atomic transcription gets **halt**.

7. Activated Code PcPs of bactericidal / bacteriostatic activity move from yang B-Bit to Yin B-Bit .

8. Yin B-Bit re translate the message of Code PcPs bactericidal / bacteriostatic activity and retranscripts the message of Code PcPs bactericidal / bacteriostatic activity .

9. this yin B-Bit carries in information (code PcPs - bactericidal / bacteriostatic activity) to target B.B.B to trigger effect of bactericidal / bacteriostatic activity .

Pharmaco Genetics - Divine Mechanics and drug action on bacterial cell by second order of Universe .

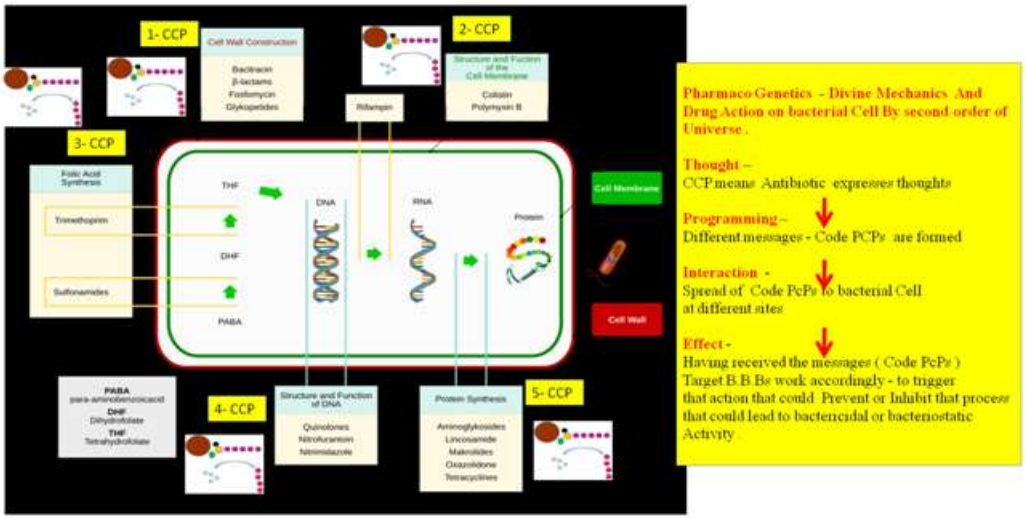

Fig 23 Pharmacogenetics - Divine Mechanics And Drug Action on bacterial Cell by second order of Universe .(Macro level)

Higher center(divine mechanics) in drug expresses thought that trigger the process that cause bactericidal and bacteriostatic activity of the bacteria . Having expressed the thoughts , programmed messages (Code PcPs) are formed . These messages spread (Code PcPs bactericidal or bacteriostatic activity) via second order (Photons) in the entire bacterial unit. The target B.B.Bs of bacteria having received the messages (Code PcPs bactericidal or bacteriostatic activity), they work accordingly and we observe the effect either bactericidal or bacteriostatic activity in vivo and vitro both .

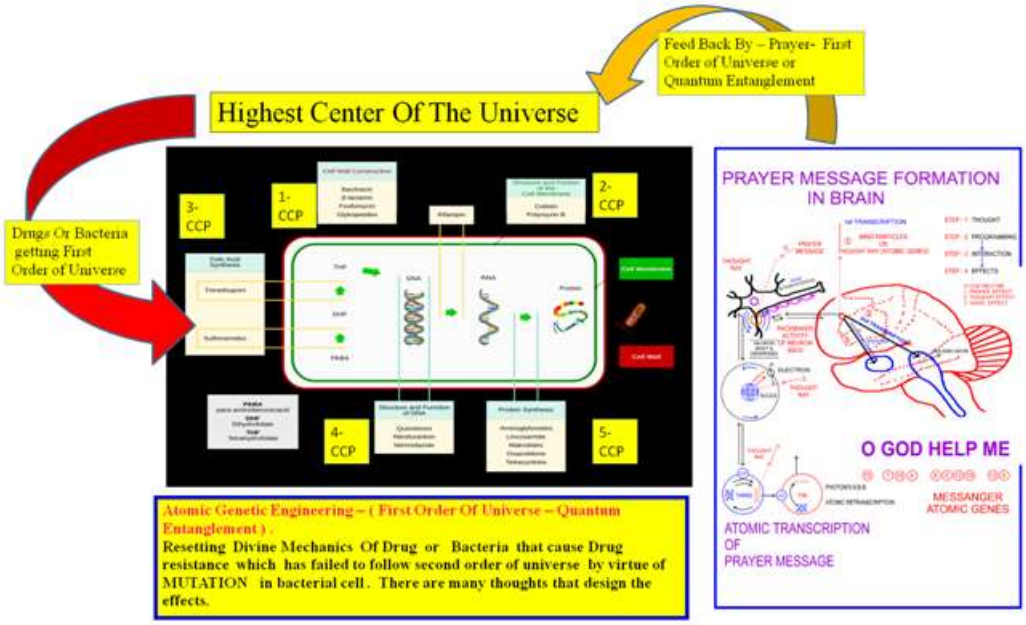

Fig 24 Atomic Genetic Engineering (First order of Universe) as adjuvant therapy could reset Divine Mechanics (CCP) of drug and Bacteria .

Atomic Genetic Engineering (First order of Universe) could reset Divine Mechanics of drug and bacteria that cause drug resistance which has failed to follow second order of universe by virtue of mutation in bacterial cell. There are many thoughts that design the effects. It is prediction of theory.

III. Conclusion

Any disorder in implementation of second order of universe carried by drug by alteration in behavior of target B.B.Bs (mutations), drug resistance develop. Drugs or antibiotics are set of information's (Code PCPs bactericidal / bacteriostatic activity) that trigger (second order carried by photons) bactericidal / bacteriostatic

effects by giving specific order to bacterial cell . If this order (Code PcPs bactericidal / bacteriostatic activity) is failed to deliver (drug half life or dose related) to bacterial cell or failed to implement due to changed behavior by target B.B.Bs (due to mutation) , drug resistance would be there . This disorder could be reset by certain drugs when added with antibiotics. Resistance-modifying agents are capable of partly or completely suppressing bacterial resistance mechanisms like Beta lactase inhibitor - Cluvanic Acid or sulbactum . There are many thoughts that design the effects. These are fed thoughts and feeding was done in pre creation era by Highest center of universe . Similarly this disorder could be reset by First order (quantum entanglement) of the universe by using Atomic Genetic Engineering (AGE) technology as adjuvant therapy . The entire study comes under pharmacogenetics drug science . [1]. Atomic Genetic Engineering (Quantum Entanglement - First order of Universe) could reset Divine Mechanics of drug and bacteria that cause drug resistance which has failed to follow second order of universe by virtue of mutation in bacterial cell . It is time mindness (TM) that triggers and regulate the entire phenomenon . It is prediction of theory .

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