

Chapter 8



***Endosymbiotic Archaeal Generated RNA Viroids
Can Regulate Cell Function and Contribute to
Autoimmunity – Systemic Lupus Erythematosus,
Multiple Sclerosis and Rheumatoid Arthritis***

Introduction

Actinides like rutile producing intracellular magnesium deficiency due to rutile-magnesium exchange sites in the cell membrane has been implicated in the etiology of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis^{1,2}. Organisms like phytoplasmas and viroids have also been demonstrated to play a role in the etiology of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis^{3,4}. RNA viroids could contribute to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis². The possibility of generation of RNA viroids by actinide based primitive organism like archaea with a mevalonate pathway and cholesterol catabolism was considered⁵⁻⁸. An actinide dependent shadow biosphere of archaea and viroids in systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis is described⁶. The role of RNA viroids generated by actinidic archaea in regulation of body functions and the pathogenesis of human disease is discussed.

Materials and Methods

Informed consent of the subjects and the approval of the ethics committee were obtained for the study. The following groups were included in the study: – systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis. There were 10 patients in each group and each patient had an age and sex matched healthy control selected randomly from the general population. The blood samples were drawn in the fasting state before treatment was initiated. Plasma from fasting heparinised blood was used and the experimental protocol was as follows (I) Plasma+phosphate buffered saline, (II) same as I+cholesterol substrate, (III) same as II+rutile 0.1 mg/ml, (IV) same as II+ciprofloxacin and doxycycline each in a concentration of 1 mg/ml. Cholesterol substrate was prepared as

described by Richmond¹⁰. Aliquots were withdrawn at zero time immediately after mixing and after incubation at 37 °C for 1 hour. The following estimations were carried out: – Cytochrome F420, free RNA and free DNA¹¹⁻¹⁴. Cytochrome F420 was estimated fluorimetrically (excitation wavelength 420 nm and emission wavelength 520 nm). The statistical analysis was done by ANOVA.

Results

The parameters checked as indicated above were: – cytochrome F420, free RNA and free DNA. Plasma of control subjects showed increased levels of the above mentioned parameters with after incubation for 1 hour and addition of cholesterol substrate resulted in still further significant increase in these parameters. The plasma of patients showed similar results but the extent of increase was more. The addition of antibiotics to the control plasma caused a decrease in all the parameters while addition of rutile increased their levels. The addition of antibiotics to the patient's plasma caused a decrease in all the parameters while addition of rutile increased their levels but the extent of change was more in patient's sera as compared to controls. The results are expressed in tables 1-2 as percentage change in the parameters after 1 hour incubation as compared to the values at zero time.

Table 1 *Effect of rutile and antibiotics on cytochrome F420.*

Group	CYT F420 % (Increase with Rutile)		CYT F420 % (Decrease with Doxy+Cipro)	
	Mean	±SD	Mean	±SD
Normal	4.48	0.15	18.24	0.66
MS	22.12	1.81	61.33	9.82
SLE	22.29	1.66	59.02	7.50
RA	22.06	1.61	57.81	6.04
F value	306.749		130.054	
P value	< 0.001		< 0.001	

Table 2 *Effect of rutile and antibiotics on free RNA and DNA.*

Group	DNA % change (Increase with Rutile)		DNA % change (Decrease with Doxy+Cipro)		RNA % change (Increase with Rutile)		RNA % change (Decrease with Doxy+Cipro)	
	Mean	±SD	Mean	±SD	Mean	±SD	Mean	±SD
Normal	4.37	0.15	18.39	0.38	4.37	0.13	18.38	0.48
MS	22.62	1.38	63.82	5.53	23.29	1.98	67.46	3.96
SLE	22.56	2.46	62.70	4.53	23.32	1.74	65.67	4.16
RA	23.30	1.42	65.07	4.95	23.11	1.52	66.68	3.97
F value	337.577		356.621		427.828		654.453	
P value	< 0.001		< 0.001		< 0.001		< 0.001	

Discussion

There was increase in cytochrome F420 indicating archaeal growth. The archaea can synthesize and use cholesterol as a carbon and energy source^{15, 16}. The archaeal origin of the enzyme activities was indicated by antibiotic induced suppression. The study indicates the presence of actinide based archaea with an alternate actinide based enzymes or metalloenzymes in the system as indicated by rutile induced increase in enzyme activities^{17, 18}. The archaea can undergo magnetite and calcium carbonate mineralization and can exist as calcified nanoforms¹⁹. This can lead on to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis.

There was an increase in free RNA indicating self replicating RNA viroids and free DNA indicating generation of viroid complementary DNA strands by archaeal reverse transcriptase activity. The actinides modulate RNA folding and catalyse its ribozymal action. The viroids are evolutionarily escaped archaeal group I introns which have retrotransposition and self splicing qualities²⁰. Archaea induced immune activation and redox stress can produce histone deacetylase inhibition resulting in endogenous retroviral (HERV) reverse transcriptase and integrase expression. This can integrate the RNA viroidal

complementary DNA into the noncoding region of eukaryotic non coding DNA using HERV integrase as has been described for borna and ebola viruses²¹. The noncoding DNA is lengthened by integrating RNA viroidal complementary DNA with the integration going on as a continuing event. The archaea genome can also get integrated into human genome using integrase as has been described for trypanosomes²². The integrated viroids and archaea can undergo vertical transmission and can exist as genomic parasites^{21, 22}. This increases the length and alters the grammar of the noncoding region producing memes or memory of acquired characters²³. The viroidal complementary DNA can function as jumping genes producing a dynamic genome important in storage of synaptic information, HLA gene expression and developmental gene expression. The RNA viroids can regulate mRNA function by RNA interference²⁰. The phenomena of RNA interference can modulate T cell and B cell function, insulin signaling lipid metabolism, cell growth and differentiation, apoptosis, neuronal transmission and euchromatin/heterochromatin expression. This can lead on to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis.

The RNA viroids and its complementary DNA developed into cholesterol enveloped RNA and DNA viruses like herpes, retrovirus, influenza virus, borna virus, cytomegalo virus and Ebstein Barr virus by recombining with archaeal, eukaryotic and human genes resulting in viral speciation^{24, 25, 26}. The RNA viroids can also recombine with endogenous commensal RNA and DNA viruses producing speciation. Viral species are ill defined and fuzzy with all of them forming one common genetic pool with frequent horizontal gene transfer and recombination. Thus the multi and unicellular eukaryote with its genes serves the purpose of viral speciation. These viruses that are generated can lead on to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis.

The multicellular eukaryotes are like archaeal biofilms. The archaea with a mevalonate pathway uses the extracellular RNA viroids for quorum sensing and in the generation of symbiotic biofilm like structures which develop into multicellular eukaryotes^{27, 28}. The endosymbiotic archaea and bacteria with mevalonate pathway still uses the RNA viroids for the regulation of multicellular eukaryote. Pollution is induced by the primitive nanoarchaea and mevalonate pathway bacteria synthesized PAH and methane leading on to redox stress. Redox stress leads to sodium potassium ATPase inhibition, inward movement of plasma membrane cholesterol, defective SREBP sensing, increased cholesterol synthesis and nanoarchaeal/mevalonate pathway bacterial growth²⁹. Redox stress leads on to viroidal and archaeal multiplication. Redox stress can also lead to HERV reverse transcriptase and integrase expression. The noncoding DNA is formed of integrating RNA viroidal complementary DNA and archaea with the integration going on as a continuing event. The archaeal pox like dsDNA virus forms evolutionarily the nucleus. The integrated viroidal, archaeal and mevalonate pathway bacterial sequences can undergo vertical transmission and can exist as genomic parasites. The genomic integrated archaea, mevalonate pathway bacteria and viroids form a genomic reserve of bacteria and viruses which can recombine with human and eukaryotic genes producing bacterial and viral speciation. The change in the length and grammar of the noncoding region produces eukaryotic speciation and individuality³⁰. The integration of nanoarchaea, mevalonate pathway prokaryotes and viroids in to the eukaryotic and human genome produces a chimera which can multiply producing biofilm like multicellular structures having a mixed archaeal, viroidal, prokaryotic and eukaryotic characters which is a regression from the multicellular eukaryotic tissue. This results in a new neuronal, metabolic, immune and tissue phenotype leading to autoimmune disease.

The archaea and viroids can regulate the nervous system including the NMDA receptor³¹. NMDA receptors can be modulated by viroid induced RNA interference². The dipolar viroids combined with actinides in the setting of digoxin induced sodium potassium ATPase inhibition can produce a pumped phonon system mediated Frohlich model superconducting state inducing quantal perception with nanoarchaeal sensed gravity producing the orchestrated reduction of the quantal possibilities to the macroscopic world^{2, 31}. This leads to the perception of low level EMF and the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis. The viroids can regulate limbic lobe transmission by RNA viroid mediated RNA interference modulating norepinephrine, dopamine, serotonin and acetyl choline receptors¹⁸. The higher degree of integration of the archaea and viroids into the genome produces increased digoxin synthesis producing right hemispheric dominance and lesser degree producing left hemispheric dominance². Right hemispheric dominance is important in the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis. The viroid RNA interference mediated altered monoamine and NMDA transmission contributes to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis. Archaea and RNA viroid can bind the TLR receptor induce NFkB producing immune activation and cytokine TNF alpha secretion^{2, 32}. The archaea and viroid induced chronic immune activation and generation of superantigens can lead on to autoimmune disease. Archaea and viroids can induce the host AKT PI3K, AMPK, HIF alpha and NFkB producing the Warburg metabolic phenotype³³. The increased glycolytic hexokinase activity, decrease in blood ATP, leakage of cytochrome C, increase in serum pyruvate and decrease in acetyl CoA indicates the generation of the Warburg phenotype. There is induction of glycolysis, inhibition of PDH activity and mitochondrial dysfunction resulting in inefficient energetics and metabolic syndrome. The archaea and viroid generated cytokines can lead to TNF alpha induced insulin

resistance. Insulin resistance is important in the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis. The accumulated pyruvate enters the GABA shunt pathway and is converted to citrate which is acted upon by citrate lyase and converted to acetyl CoA, used for cholesterol synthesis³³. The pyruvate can be converted to glutamate and ammonia which is oxidised by archaea for energy needs. The increased cholesterol substrate leads to increased archaeal growth and digoxin synthesis leading to metabolic channeling to the mevalonate pathway³⁴. Viroid induced RNA interference can modulate the mRNAs concerned with insulin receptor function and lipid metabolism contributing to insulin resistance. Insulin resistance is important in autoimmunity. The RNA viroids can recombine with HERV sequences and get encapsulated in microvesicles contributing to the retroviral state. The prion protein conformation is modulated by RNA viroid binding producing prion disease. Prions and HERV sequences can contribute to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis.

Thus the actinidic archaea generated RNA viroids can regulate cell function and produce neuro-immuno-genetic-endocrine-metabolic integration. The RNA viroids and their complementary DNA can serve the purpose of viral speciation. The RNA viroids also contribute to the pathogenesis of systemic lupus erythematosus, multiple sclerosis and rheumatoid arthritis.

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