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Article information

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Abstract

PSEUDO MORPHOLOGY AND COVID-19 CURE.o

COVID-19 is concluded as devoid of independent morphogenetic structure to develop itself but it inherits from host the distorted DNA signals to manifest its existence in basic cellular form with RNA superscripted during evaluation process. If the lapse or

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misinformation of hereditary features from genes lead to probagation of tumours instead of specialised cells whereas the suppression of such information in RNA signal transmission by COVID-19 induction will mutate virus

multiplication possibly. If so,the treatment should begin from

genetical correction of host rather than counteraction of virus initially.As future couldnot be predicted by anyway it is indeed not a neglecting factori if the evil virus leave damages due to its epigenetic action transmitted to subsequent generations!

https://en.wikiversity.org/wiki/WikiJournal_Preprints/COVID-19_ELIMINATION_AND_CELL_DIFFERENTIATION.

Introduction

An attempt to conceptualise the covid-19 development in contradiction with any other morphogenetic cell by virtue of its reactions and in the vague of its complications in arresting let us try using the method of eliminating its raising up from host without liquidising it directly but making itself to liquidate self is narrated.Early presumptions regarding the COVID-19 infection strategy went to attribute its silence prevailing over initial period to copyover of genetic transcriptions of hostcell through its RNA interaction with host cell DNA messages intercepted .This enables COVID=19 to maintain similarity in its RNA configuration found in host cell as well to protect itself against its identification distinguished from host cell immunity.Once it assumes integrity with hostcell, it just waits until the differentiation process starts up in hostcell as usual or by way of increasing immunity.When the differentiation starts up ,the nucleus of COVID-19 due to its fluid bubble nature spread as nano droplets keeping signal communication but seperate only physically through nano distance and by this action it gets access to more no of HOSTCELLS.The individual portions after entering individual hostcells raiseup to live as individual COVID-19 nucleus with disintegration of hostcell DNA into RNA of COVID-19 instead of transformation of same into mRNA of normal cell in regular differntiation supposed.It is deformed differentiation in support of COVID-19 rather than normal differentiation to form new fully developed hostcells. So, semi developed cell resume its contact with its associate of differntiated new cell to disintegrate its protein process by stopping some growthfactors leading to appearance of FAS-LIGANDS in cell clusters.Presence of FAS-LIGANDS initites APOTOSIS because the surrounding COVID-19 semi grown cell structure with decomposed proteins provide such gaps for well grown cells to shrink and rest themselves .So, white blood corpuscles and simultaneously the source of FAS-LIGAND generation i.e the new differentiated hostcell along with the COVID-19 encroached one gets demolished by attacking white cells sent by

Immune system. Eventual result is the both dying of white cells and new cells with the advantages reaped for proliferation of COVID-19. There is no surprise in COVID-19 choosing the period of cell differentiation and sharply launching its proliferation considering the Pseudomorphological Development associated with it.

PSEUDOMORPHOLOGY OF COVID-19

The basic concept of this argument has its root in the process of cell split which is the basic and essential phenomenon in the growth of tissue structure of living organisms. I am consentient to cell multiplication by splitting as far as fully developed cell structure is concerned but am reluctant to the multiplication of partially developed primordial cells like COVID 19 which lacks DNA configuration. This speculation leads to attribute a different method for the proliferation of COVID 19 which is to be confirmed experimentally by observation and verification through international testing labs.

The cell splitting process leads to the formation of new cells (with both DNA and RNA), eventually it is possible only in that kind of fully developed cells only and not in partially developed cells having RNA alone because splitting is the unique action that can be initiated if only DNA also is present in the cell according to the ancient past bio metamorphic activities information stored in DNA. For viruses like COVID-19 which is only semi grown organism without DNA no such initiation signal for cell split process hardly present as there is no storage spot possible without DNA. Its cell growth is only aggregation of already grown cells in the human or fully developed cells of living bodies cropped as smuggled and exists with continued supply of its past association with original cells.

It could be spelt out if the following development of COVID-19 affected cells exist virtually but not identified physically to an extent as to show critical symptoms. That is the distinct no of RNA cells of host body which do not merge with its associate DNA combination of which forms new developed host cells whereas the hanging RNA forms the COVID-19's semi developed cells eventually. It is evident that such cells do not act for DNA SIGNAL (Due to epigenicity and being target of COVID-19 particulate combination) and they look isolated once they arise initially in the healthy cells itself and does not do productive work as like original RNA and hence can be named pseudo RNA. The name is so given because in fact these nonintegrated RNA constituents may be later deployed as COVID-19 transformed portions during its sudden proliferation. If the presence of Pseudo RNA is identified and ascertained the infection of COVID-19 could well be recognised in advance before the spreading goes to dangerous extent which is explained below.

This Pseudo RNA form part of some newly split good cells and act in opposition to the signal of body immune system (due to epigenetic effect of COVID-19) sent from DNA of white corpuscles because it gets grouped in to RNA's of newly formed COVID-19 primordial cells which attacks host cells. How come the DNA information of new Cells having Pseudo RNA fail integrity with

body? Is it a ridicule? No. During splitting all the signals are halted within the cell and fission signal only active along the previously

assumed boundary to realise the formation of two new cells. This is the situation when the silent COVID-19 could also try to multiply its cells as it cannot initiate its own cell division due to its Pseudomorphological nature. COVID-19 interference in division prompts the Pseudo RNA to bring ahead splitting signals further even after termination of division. So some of the white corpuscles with Pseudo RNA and new host cells (Pseudo RNA linked) do not unilaterally act with good cells in protein synthesis and tend to form new boundary (like cancer cells) incorporating COVID-19 RNA to serve COVID-19 as a shield and supply platform from which it can proliferate until it multiplies to a large number of individual COVID-19 cells demolishing life organs. This trial of boundary forming is evident from experimental findings of a variety of configurations of COVID-19 in infected persons. Though it cannot form new boundary the time it associates in the formation process helps COVID-19 to grow its species. If the immune system goes on attacking infected cells without screening the negative trend by Pseudo RNA, the defence & counter attack ultimately turns into self-destruction. Please note that it is not necessary for apoptosis to complete but initialising it in an inappropriate time is as much as dangerous as it creates instability in cell development.

From the above correlations of proliferation of COVID-19 integrated with cell differentiation it is obvious its genetic structure is not stable one and is manifested in so many configurations corresponding to variations created during the process of cell differentiation which is critical for making morphological changes in cells. ([see illustration](#))

The virus transforms itself in different gene configurations in order to survive in the human body. The news reported mentioned that among two types of corona virus used for experiments one type transformed itself in 788 configurations and the other in 32. Definitely as per biological foundations an organism possessing definite morphological base will not, in short period migrate to different manifestations. This is the first coincidence I observe with the Pseudo morphology of COVID-19 I emphasized. Secondly, such transformation came as a result of alteration in SPIKE protein. This supports the supertranscription of encryptions in RNA disrupting NF- κ B protein formation leading to irregular differentiation of cells which I specified as cause of losing immunity and apoptosis.

WHETHER THIS GENETIC REFORMATION ACQUIRED BY COVID-19 COULD LIMIT ITS TERRITORY TO ITSELF ONLY WITHOUT HAVING ITS ROOT IN INFECTED CELL FROM WHERE ITS PROTEIN SYNTHESIS STEPS FORWARD?

I suspect strongly that perhaps once there was no specific modification supposed to have been preceded in the genomes of infected (host) cells there can possibly be no such strange insimilarities in configurations of virus morphology observed is possible.

This suspicion is not meant to puzzle any biological phenomenon but to make it transparent that like without inheriting mother cell the baby cell will not exhibit an attribute ,we should logically understand that unless any genetic phenomenon went to overwrite itself in host cell ,the RNA of the infecting COVID-19 would have been supposed to exhibit variety of manifestations it has been reported to come in over 5349 samples tested as on the date the above news was reported by researchers and confirmed by LSHTM,LONDON.

So,what this finding if its echo heard in days to come ,will alert the treatment of COVID-19 Infection?

DIFFERENTIATION THERAPY AS AN INEVITABLE TOOL

SCOPE OF ANTIBODIES

Normally vaccines and antiviral medicines act in support of fully differentiated cells resulting after successful differentiation and fighting germs. But they seldom act while differentiation going on and also there is meagre possibility their counteracting effect go against virus having imprinted same genetic configuration similar to host cells. Body immune system will initiate the action of drugs if only it finds contradiction of genetic structure associated with developing cells and as for COVID-19 development we already discussed how it has integrated itself with proteinsynthesis of host cells and so it will not expose any incidence for being attacked. So as for virus having definite morphogenetic configuration the drugs a, vaccines and antiviral medicines prove success as they are readily distinguished by immune system to initiate counteraction against them. But as for COVID-19 the immune system will never initiate any offence in association with drugs as the virus manifests as product of host cell differentiation with evidence of being participated in differentiation due to its proliferation closely united in differentiation. Under this circumstances there is no chance of coming out from covid-19 without removing the traces of differentiation in it . However without attacking virus the protective protein layer could be used to prevent virus from entering the healthy cells remaining away from its approach still. That is by way of supportive treatments came help in occasion.

The supportive treatments being offered worldwide as of today forms shielding protein layer around inside the life organs throat, lungs and breathing passages .So, though the COVID-19 cells are detached from their to contact with life cells and they (covid-19) disappear from their presence over human cells of life sensitive organs lying over chest, but due to the genetic changes the COVID-19 inherits ,is it can be ruled out from being rest itself with its new configuration as an associate cell member behind the observations of indigeneous equipments as the new configurations it acquire are not readily recognisable and only in timespan it may come to light just as like 788 now found? Because the presence of lifesaving protein shield just darkens the distinguishable micro constituents there exists every nano escape for such strange shapes come into picture suddenly.

I am aware that the researches went on and realised hitherto and will be would powerful anti-virus medicines which may get a breakthrough into COVID-19 gene with the fact that they too have to be brought into the action by immunesystem whose detection and response signals if already suppressed or influenced due to proliferation of COVID-19 or if the virus hide its identity through its evading RNA configuration simulated from host as already discussed, then definitely thete prompted a genetic rectification initialy before attempting such vaccines.

For safeside,let us not rule out the possibility projected in some reserches THAT THE IMMUNISATION RESTING IN CORONA healed patients will last only 1-2 years, and post treatment care shall periodically be maintained to protect.

On the grounds of assumption that supportive treatments just protect unaffected bodycells by only keeping outside the proliferated COVID-19 cells in live without removing them from stage but behind the closedscreen,This sidelined COVID-19 will not proliferate further or escape into active life regions due to protein wall but slowly induce acruing problems in further development of protected healthy cells by robbing the ingredients passing on the way through it to growing cells inside the besieged protein fort.

I am writing as a common man awayfrom active sects,day to day medical experiences and practices because no team from such engagements could come to this corner due to the fact that they have been tied with thousands of ropes which pull along millions of points diverting the route in this confiscation.It is evident from the scene that no differentiation therapy to treat this complication has been deployed as an experimental practice in order to study atleast the genetic data of the ailment reveals out.But time may come in future to force mission to utilise the differentiation tecniques in the event that no vaccine could ultimately be launched due to obstacles come across.It is no strange in the event of its interaction with genetical structure to consider the treatment in the grounds of retrofitication of cell bases rather than removing traces of covid-19 just recognised and identified in critical zones.Because considering the disastrous nature of this deep rooting virus,in order to protect order in humancell systems for ever such root decision,I feel is ambitious and shold be opted after analysing the affected for long period.

Poor differentiation is an important hallmark of cancer cells, and differentiation therapy holds great promise for cancer treatment. The restoration of I κ B kinase α (IKK α) leads to the differentiation of nasopharyngeal carcinoma cells with reduced tumorigenicity. The findings by Yan et al. validate the polycomb protein enhancer of zeste homologue 2 (EZH2) as a target for intervention

SCOPE OF DIFFERENTIATION THERAPHY

On the basis of above concluded facts it is obvious the DIFFERENTIATION THERAPIES practiced in the treatment of cancer in respiratory system spreading over throat, bronchus and lungs. In that treatments shall be deployed for COVID-19 eradication. It is used for removal of tumours, tumour to form matured fully grown cell out of poorly differentiated one but in current treatment COVID-19 is concerned the objective is not removal of tumour, but removal of contagious proteins created along with genetic infringement associated with virus. Afterwards when cells restored to original condition, the proliferated duplicated germ cells disappear completely and their further interaction genetic signals completely erased out. While writing this contradictory comparison, I sincerely admit that being non-medical, non-bio or physiological scientist I have not been associated anytime at diagnosis and treatment of such things but the discussions I have grasped in this line eventually made me come to this point due to following facts. 1. It is well known that medicines used in Differentiation therapy bring the poor differentiated tumour cell into an early stage from which the defective path started to rush it to severe irregular toxic development. What is to be understood is that in this method of curing when the tumour cell was brought to early stage, the growth processing route is diverted in correct path with the help of activators which prevent the silencing of IKK α (p45) kinase critical in protein production in DNA/NF- κ B signalling.

2. Though tumour development as a cause of poor differentiation is a practically distinguished fact, the realising factor to compare COVID-19 infection outcome as also same of distorted differentiation may well be comprehended by virtue of protein ubiquitination which almost halts respiratory tissues in contrast with any other severe virus infections which rather halt cell process functioning only but not interfering differentiation process. Secondly, the proliferated COVID-19 cluster appear suddenly as so many cases witnessed instant collapse in physical conditions during treatments eventually leading to deaths. Though the other factors like diabetes, kidney and immune weakness count it also implies that mutations of COVID-19 infections prove fatal due to incapability of life organs to withstand vibration resonated with operation revealing some underdevelopment in tissues rest or tissues became separately weakened. Finally the extreme victims witness badly damaged life organs as a result of extra proliferation of COVID-19 and such damages would not occur unless differentiation is bad. So, it is one which is closely associated with COVID-19 infection is differentiation and focus is to be diverted first on it. The same therapy when would be applied to current situation of COVID-19 proliferated cells only host cells which accompanied these virus cells during differentiation or the cell from which differentiation proceeded to yield new complete cell and partial COVID-19 inhabited cell will be brought to early stage. Here if once again the DNA/NF- κ B signalling pathway is made to go in normal way by providing the necessary activators prevent the silencing of IKK α kinase critical in protein production. Then naturally the COVID-19 inhibited cell gets devoid of such error factors and will no more support proteins necessary for COVID-19 survival. Because as the virus is out of DNA it cannot acquire growth signals from the RNA while the differentiation proceeds in correct manner. Only if differentiation gets halted due to breaking away of COVID-19 nucleus as

described earlier the virus has chance to overrule RNA. Before the complete destruction of cells due to APOTOSIS, if differentiation therapy administered in right time it will yield encouraging results. (SEE LINE SKETCH IN)

The above speculation is postulated by considering important morphological features of fully grown developed cell structure and semi grown undeveloped Pseudomorphological virus structure. Let us analyse that. The fully developed cell structure of human cells possess DNA and contains programs to be executed to complete protein synthesis and cell development. So, from the realised treatments of tumour observed it is found able to execute cell growth in correct formulae when the tumourised poor differentiated cell is brought to early stage during DIFFERENTIATION THERAPY. As such it can be reckoned well if the differentiation is applied to cell surrounded by contagious disintegrated DNA bound partially grown, halted proteination cell the same reversal process will not fail to take place. Because the differentiation process is not applied to completely DNA broken, transformed COVID 19 cell which is not possible as the Pseudomorphology does not have room for such kind of cellular activity. The feature of covid-19 cell is that once it started to grow with its encrypt residual in RNA that is capable to proliferate only once during its attempt to multiply. It cannot initiate second attempt of multiplication as like fully developed cell during therapy due to the fact that it lack DNA.

So, I have just combined the advantage of DNA in developed cell and disadvantage of null DNA in semi developed virus cell structure in order to proceed with differentiation therapy to destroy virus multiplied and regain host growth. Time will answer to this venturous prediction.

Additional information

- **^ Differentiation therapy** is an approach to treating advanced cancers in which malignant cells are encouraged to differentiate into more mature forms using pharmacological agents. The basis of the therapy stems from the tendency of malignant tumor cells to assume a less specialized, stem cell-like dedifferentiated state.
- **^ Phosphorylated IKK α (p45)** is a nuclear active form of the IKK α kinase that is induced by the MAP kinases BRAF and TAK1 and promotes tumor growth independent of canonical NF- κ B signaling. Insights into the sources of IKK α (p45) activation and its downstream substrates in the nucleus remain to be defined. Here, we discover that IKK α (p45) is rapidly activated by DNA damage independent of ATM-ATR, but dependent on BRAF-TAK1-p38-MAPK, and is required for robust ATM activation and efficient DNA repair. Abolishing BRAF or IKK α activity attenuates ATM, Chk1, MDC1, Kap1, and 53BP1 phosphorylation, compromises 53BP1 and RIF1 co-recruitment to sites of DNA lesions, and inhibits 53BP1-dependent fusion of dysfunctional telomeres. Furthermore, IKK α or BRAF inhibition synergistically enhances the therapeutic potential of 5-FU and irinotecan to eradicate chemotherapy-resistant metastatic human

tumors *in vivo*. Our results implicate BRAF and IKK α kinases in the DDR and reveal a combination strategy for cancer treatment.

- \wedge NF- κ B (nuclear factor kappa-light-chain-enhancer of activated B cells) is a protein complex that controls transcription of DNA, cytokine production and cell survival. NF- κ B is found in almost all animal cell types and is involved in cellular responses to stimuli such as stress, cytokines, free radicals, heavy metals, ultraviolet irradiation, oxidized LDL, and bacterial or viral antigens.[1][2][3][5][6] NF- κ B plays a key role in regulating the immune response to infection. Incorrect regulation of NF- κ B has been linked to cancer, inflammatory and autoimmune diseases, septic shock, viral infection, and improper immune development. NF- κ B has also been implicated in processes of synaptic plasticity and memory
- I consider comparison of TUMOUR GENERATION and COVID-19 complication because both are not possible unless there is lapse occurs in protein synthesis. The tissue damage and immunity loss could be attributed towards it. Whereas other infections not affecting tissue culture and immunity more while they may offer strong offensive to immunity. Because of damages in respiratory organ tissues sudden breathing syndromes witness.

Acknowledgements

I would like to thank wikipedia.org in this regard which contains relevant updated info available readily

Competing interests

Mysterious nature of COVID-19

Ethics statement

I WOULD LIKE TO DEDICATE MY CONSCIENTIOUS THINKING, WORK AND TIME TOWARDS COMPLETELY WIPING OUT COVID-19 EXISTENCE IN GLOBE.

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