The chapter represents a vision about how DNA might act as a topological

quantum computer). TQC means that the braidings of braid strands define TQC  $\,$ 

programs and M-matrix (generalization of S-matrix in zero energy ontology)

defining the entanglement between states assignable to the end points of

strands define the TQC usually coded as unitary time evolution for Schr\"odinger equation.

Before a representation of the model of TQC general vision about what

happens in quantum jump, which at least in formal sense can be regarded as

quantum computation (TQC), is represented. Included is also a section about

possible modification of thermodynamics required by the possibility of

negentropic entanglement. The modification corresponds simply to the replacement \$S\rightarrow S-N\$ for the entropy in standard thermodynamics.

The implications of this replacement are however highly non-trivial. The

\blockquote{pessimistic} generalization of the second law allows to
understand the

thermodynamical aspect of TQC. One can understand why living matter is

so effective entropy producer as compared to inanimate matter and also the

characteristic decomposition of living systems to highly negentropic and

entropic parts as a consequence of generalized second law. ADP-ATP process

of metabolism provides a concrete application for the generalized thermodynamics and allows to see this process as a transfer of negentropic

entanglement. Also DNA double strand for which sugar-phosphate backbone

consists of XMPs, X= A,T,C,G containing negentropy carrying phosphate bonds

can be seen as analogous to conscious brain with DNA strands representing

right and left hemispheres.

One can end up to the model of TQC in the following manner.

\begin{enumerate}

\item Darwinian selection for which the standard theory of self-organization provides a model, should apply also to TQC

programs. Tqc

programs should correspond to asymptotic self-organization patterns selected by dissipation in the presence of metabolic energy feed. The

spatial and temporal pattern of the metabolic energy feed characterizes

the TQC program - or equivalently - sub-program call.

\item Since braiding characterizes the TQC program, the self-organization

pattern should correspond to a hydrodynamical flow or a pattern of magnetic

field inducing the braiding. Braid strands must correspond to magnetic flux  $% \left( 1\right) =\left( 1\right) +\left( 1\right)$ 

tubes of the magnetic body of DNA. If each nucleotide is transversal magnetic dipole it gives rise to transversal flux tubes, which can also

connect to the genome of another cell. As a matter fact, the flux tubes

would correspond to what I call wormhole magnetic fields having pairs of

space-time sheets carrying opposite magnetic fluxes.

\item The output of TQC sub-program is probability distribution for the

outcomes of state function reduction so that the sub-program must be

repeated very many times. It is represented as four-dimensional patterns

for various rates (chemical rates, nerve pulse patterns, EEG power distributions,...) having also identification as temporal densities of zero

energy states in various scales. By the fractality of TGD Universe there is

a hierarchy of TQCs corresponding to p-adic and dark matter hierarchies.

Programs (space-time sheets defining coherence regions) call programs in

shorter scale. If the self-organizing system has a periodic behavior each

TQC module defines a large number of almost copies of itself asymptotically. Generalized EEG could naturally define this periodic pattern and each period of EEG would correspond to an initiation and halting of TQC. This brings in mind the periodically occurring solgel

phase transition inside cell near the cell membrane. There is also a connection with hologram idea: EEG rhythm corresponds to reference wave and

nerve pulse patters to the wave carrying the information and interfering

with the reference wave.

\item Fluid flow must induce the braiding which requires that the ends of

braid strands must be anchored to the fluid flow. Recalling that lipid

mono-layers of the cell membrane are liquid crystals and lipids of interior mono-layer have hydrophilic ends pointing towards cell interior,

it is easy to guess that DNA nucleotides are connected to lipids by magnetic flux tubes and hydrophilic lipid ends are stuck to the flow.

\item The topology of the braid traversing cell membrane cannot be affected

by the hydrodynamical flow. Hence braid strands must be split during TOC.

This also induces the desired magnetic isolation from the environment.

Halting of TQC  $\,$  reconnects them and make possible the communication of the

outcome of TQC.

\end{enumerate}

There are several problems related to the details of the realization.

\begin{enumerate}

\item How nucleotides A,T,C,G are coded to the strand color and what this color corresponds to physically? There are two options

which could be characterized as fermionic and bosonic.

\begin{enumerate}

\item Magnetic flux tubes having quark and anti-quark at their ends with

\$u\$,\$d\$ and \$u\_c\$, \$d\_c\$ coding for A,G and T,C. CP conjugation
would

correspond to conjugation for DNA nucleotides.

\item Wormhole magnetic flux tubes having wormhole contact and its CP

conjugate at its ends with wormhole contact carrying quark and antiquark

at its throats. The latter are predicted to appear in all length scales in

TGD Universe.

\end{enumerate}

\item How to split the braid strands in a controlled manner? High \$T\_c\$

super conductivity suggests a possible mechanism: braid strand can be

split only if the supra current flowing through it vanishes. A suitable

voltage pulse induces the supra-current and its negative cancels it. The

conformation of the lipid could control whether it it can follow the flow or

not. The absence of both genuine magnetic monopoles and boundaries

however demands that the monopole flux tubes must be closed. One manner to

achieve this is to assume that the magnetic flux returns back along second

space-time sheet.

A more realistic variant of this model is based on pairs of flux tubes going through

the membrane and carrying opposite currents and parallel (opposite) magnetic

fields. Reconnection for the members of the pair occurring the cell membrane effectively cuts both. This conforms with the identification

of Cooper pairs as S=0 or S=1 states of electrons at the two flux tubes.

The reconnection occurs naturally at the limit when the velocity of electrons

and thus current goes to zero.

\item How magnetic flux tubes can be cut without breaking the conservation

of the magnetic flux? The notion of wormhole magnetic field could save the

situation now: after the splitting the flux returns back along the second

space—time sheet of wormhole magnetic field. An alternative solution is

based on reconnection of flux tubes. Since only flux tubes of same color

can reconnect this process can induce transfer of color:
\blockguote{color

inheritance}: when applied at the level of amino-acids this leads
to a

successful model of protein folding. Reconnection makes possible breaking

of flux tube connection for both the ordinary magnetic flux tubes and

wormhole magnetic flux tubes.

\item How magnetic flux tubes are realized? The interpretation of flux

tubes as correlates of directed attention at molecular level leads to

concrete picture. Hydrogen bonds are by their asymmetry natural

correlates

for a directed attention at molecular level. Also flux tubes between

acceptors of hydrogen bonds must be allowed and acceptors can be seen as

the subjects of directed attention and donors as objects. Examples of

acceptors are aromatic rings of nucleotides, \$0=\$ atoms of phosphates,

etc.. A connection with metabolism is obtained if it is assumed that various phosphates \$XMP,XDP,XTP\$, \$X=A,T,G,C\$ act as fundamental acceptors

and plugs in the connection lines. The basic metabolic process \$ATP\rightarrow ADP+P\_i\$ allows an interpretation as a reconnection splitting flux tube connection, and the basic function of phosphorylating

enzymes would be to build flux tube connections as also of breathing and

photosynthesis. \end{enumerate}

The rest of the article represents a more concrete vision about how

DNA might act as a topological quantum computer (TQC). The topics discussed are following.

# \begin{enumerate}

\item How the basic gates are realized concretely? Gates can be identified

as basic braid operations so that the question reduces to how braidings of

magnetic flux tubes represent gates and what kind of particles represent

the quantum states. The identification of the particles is in terms of

quarks: TGD indeed predicts a hierarchy of scaled variants of hadron physics.

\item How the braiding is realized? What do braid strands identified as magnetic flux tubes look like? How the braiding operation is

induced? The tentative answer is that color magnetic flux tubes connecting

DNA nucleotides to the lipids of nuclear and cell membrane define braid

strands and that braiding operations are induced by hydrodynamic flow

around membrane generating 2-D flow of liquid crystal defined by the lipids. Also nerve pulse propagation can induced this kind of 2-D flow.

\item How magnetic flux tubes are realized? The interpretation of

flux

tubes as correlates of directed attention at molecular level leads to

concrete picture. Hydrogen bonds are by their asymmetry natural correlates

for a directed attention at molecular level. Also flux tubes between

acceptors of hydrogen bonds must be allowed and acceptors can be seen as

the subjects of directed attention and donors as objects. Examples of

acceptors are aromatic rings of nucleotides, \$0=\$ atoms of phosphates,

etc.. A connection with metabolism is obtained if it is assumed that various phosphates \$XMP,XDP,XTP\$, \$X=A,T,G,C\$ act as fundamental acceptors

and plugs in the connection lines. The basic metabolic process \$ATP\rightarrow ADP+P\_i\$ allows an interpretation as a reconnection splitting flux tube connection, and the basic function of phosphorylating

enzymes would be to build flux tube connections as also of breathing and photosynthesis.

### \end{enumerate}

The model is certainly very speculative and heavily relies on the new

physics predicted by TGD. One can also imagine alternative scenarios.

The model makes however strong predictions and is therefore testable.

#### \begin{enumerate}

\item The model makes several testable predictions about DNA itself. In particular, matter—antimatter asymmetry and slightly broken

isospin symmetry have counterparts at DNA level induced from the breaking

of these symmetries for quarks and antiquarks associated with the flux

tubes. DNA cell membrane system is not the only possible system that could

perform TQC like activities and store memories in braidings: flux tubes

could connect biomolecules and the braiding could provide an almost definition for what it is to be living. Even water memory might reduce to braidings.

\item The model leads also to an improved understanding of other roles of

the magnetic flux tubes containing dark matter. Phase transitions changing

the value of Planck constant for the magnetic flux tubes could be key

element of bio-catalysis and electromagnetic long distance communications

in living matter. For instance, one ends up to what might be called code

for protein folding and bio-catalysis. There is also a fascinating connection with Peter Gariaev's work suggesting that the phase transitions

changing Planck constant have been observed and wormhole magnetic flux

tubes containing dark matter have been photographed in his experiments.

\item In the proposed vision genes define the hardware and TQC programs

the software responsible for what becomes cultural evolution at the higher

levels of evolutionary hierarchy. This vision explains also the mystery of

introns. The quite recent findings challenging genetic determinism expressed using the term \blockquote{genetic dark matter} provide support for an

existence of new information carrying level at the level of genome identifiable in terms of TQC programs.

## \end{enumerate}

It must be emphasized that this model of DNA as TQC is only one option among many. There is large flexibility concerning the identification of

fermions involved. For instance A,T,C,G could be represented also in terms

of 4 states assignable to two spin half fermions at parallel flux tubes.

This would give rise to high  $T_c$  superconductor with both S=0 (S=1)

Cooper pairs assignet to flux tubes with opposite (parallel) magnetic

fields. The spin-spin interaction energy for the Cooper pair would be

negative and proportional to  $h_{eff}\$  and same for all fermion pairs if

\$h\_{eff}=h\_{gr}\$ hypothesis holds true at microscopic level.

#### %\end{abstract}