

# **Regenerative Medicine and Magnetic Field in Neurodegenerative Pathologies from The Electromagnetic Point of View**

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## **ABSTRACT**

The use of regenerative medicine in combination with electromagnetic fields is opening new therapeutic possibilities for neurodegenerative diseases, such as Parkinson's and Alzheimer's. By energizing stem cells with electromagnetic fields and subsequently using them in damaged brain areas, it is possible to promote regenerative processes and improve neural repair capabilities. This article explores the electromagnetic principles underlying these interactions, focusing on diamagnetic therapy as a potential non-invasive therapeutic approach to support neurogenesis and synaptic plasticity.

**Keywords:** Stem cells; Diamagnetic; Brain; Parkinson; Alzheimer's

#### **INTRODUCTION**

Neurodegenerative diseases represent a significant medical challenge, given the lack of effective treatments capable of restoring damaged neuronal function [1]. In recent years, regenerative medicine has opened new avenues through the use of stem cells, whose ability to differentiate into various cell types offers tremendous therapeutic potential while [2], the use of electromagnetic fields like Diamagnetism, has shown significant promise in CNS diseases at the experimental level [3].

#### **BACKGROUND ON DIAMAGNETIC THERAPY**

Diamagnetic therapy is an approach that uses pulsed magnetic fields to induce biological responses in tissues without causing direct cellular depolarization. This technique leverages the sensitivity of biological tissues to magnetic fields to modulate cellular activity and support regeneration  $[4]$ . Several studies have demonstrated that pulsed electromagnetic fields, including the diamagnetic effect, can influence ion channel activity, cell membrane fluidity, and biochemical processes also correlated to the Intensity and the gradient of induced Magnetic Field all of which are potentially key factors in neurodegenerative diseases [5,6].



## **METHODOLOGY: DESIGNED METHOD S IN DIAMAGNETIM**

#### **Energizing and Administration of Stem Cells**

Preliminary unpublished data report of Stem cells that are exposed to low-frequency electromagnetic fields in a controlled environment to stimulate their proliferation and regenerative capacity. This energization induces metabolic changes, enhancing the production of neurotrophic growth factors that promote neuronal regeneration. Once energized, the cells are administered nasally, allowing them to cross the blood-brain barrier and migrate toward damaged brain areas (Figure 1).





#### **Magnetic Stimulation of the Brain**

After the stem cells reach the central nervous system, localized electromagnetic fields are applied in target brain areas. This stimulation increases cell penetration into damaged regions, promotes their differentiation into functional neurons, and stimulates the formation of new synaptic connections, which are crucial for functional recovery (Figure 2).



**Figure 2:** Localized electromagnetic fields are applied in target brain areas.



## **PRELIMINARY EXPECTED RESULTS**

The application of electromagnetic fields has shown positive effects on both stem cells and damaged brain tissue. Preliminary exposure of stem cells to electromagnetic fields increased their vitality and proliferative capacity, making them more effective in neuronal regeneration once they migrated into the brain. Furthermore, cerebral stimulation via magnetic fields facilitated cell integration into damaged areas, promoting a significant regenerative response.

#### **ELECTROMAGNETIC MECHANISMS OF INTERACTION AND MATHEMATICAL MODELS**

#### **Electromagnetic Induction and Induced Electric Field.**

One primary effect of electromagnetic fields on cells is the induction of electric fields through time-varying magnetic fields. Using Faraday's law, the induced electromotive force (E) in a loop is given by: ε=-(d∅)/dt

where  $\emptyset = B \cdot A$  is the magnetic flux, with B as the magnetic field and A as the area of the loop. For an oscillating magnetic field  $B(t) = B0 \sin(\omega t)$ , the time derivative of the flux results in an induced electric field:

$$
E_{ind}(\mathsf{t}) = \frac{dB}{dt} * r = B_0 \omega \cos(\omega t) * r
$$

where r is the distance from the loop's center. This induced field can alter the cell membrane potential, modulating ion channel activity and facilitating ion exchange across the membrane [7].

#### **Effect on Ion Channels and Membrane Potential.**

Ion channels are highly sensitive to changes in membrane potential induced by oscillating electric fields. The membrane potential Vm can be approximated by the Goldman-Hodgkin-Katz equation:

$$
V_m = \frac{RT}{F} \ln \frac{P_{Na}[Na^+]_{out} + P_K[K^+]_{out} + P_{Cl}[Cl^-]_{in}}{P_{Na}[Na^+]_{in} + P_K[K^+]_{in} + P_{Cl}[Cl^-]_{out}}
$$

where P represents the permeability of the respective ions  $(Na+, K+, Cl-)$ , and ion concentrations are indicated by location (intra- or extracellular). Interaction with electromagnetic fields can modulate these permeability values P, thereby altering membrane potential and facilitating neuronal signaling [8].

#### **Larmor Precession and Stability of Bound Ions.**

Another important effect is Larmor precession. Ions bound near the cell membrane undergo precession induced by the static magnetic field B0, described by the Larmor frequency omega L:

$$
\omega L = \frac{q B_0}{2m}
$$

where q is the ion's charge, and m is its mass. Precession at this frequency can influence the configuration of bound ions, affecting biochemical reactions at the membrane level and enhancing signal transmission.

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This phenomenon can be described by the motion equation for an ionic oscillator in the presence of a magnetic field and thermal noise:

$$
m\frac{d^2 r}{dt^2} + \beta \frac{dr}{dt} + kr = qv * B + n(t)
$$

where beta is the viscous damping coefficient, k is the elastic constant of the restoring force, and n(t) represents thermal noise. This model describes the balance of magnetic and thermal forces that stabilize or destabilize ion positions, potentially influencing neurotransmission [9, 10].

#### **Electrochemical Transfer Model for Stem Cells**

The electromagnetic interaction with stem cells can be represented by an electrochemical model based on charge transfer. The current Ib across the membrane is given by:

$$
I_b(S) = q_a \Delta \tau_a(S)
$$

where  $\Delta\Gamma$ a(s) is the change in surface concentration of the bound ion, and qa is a coefficient representing interfacial charge dependence. The impedance  $Z_A(s)$  of this electrochemical pathway can be written as:

$$
Z_A(s) = \frac{1}{q_a s(\alpha + v_a)}
$$

where alpha represents binding potential dependence, and va is the first-order binding rate constant. This model reflects how electromagnetic fields influence ion absorption and release, which are crucial for the differentiation and proliferation of stem cells in regenerative applications [11].

#### **Magnetic Field Distribution and Tissue Penetration Potential.**

The spatial distribution of the magnetic field in tissues is essential for ensuring adequate penetration into damaged brain regions. Using the Biot-Savart law, the magnetic field  $B(r)$  generated by a coil can be expressed as:

$$
(r)=\frac{\mu_0 I}{2\pi r}
$$

where I is the current in the coil, r is the distance from the coil's center, and  $\mu_0$  is the magnetic permeability of free space. This field can be modeled numerically to optimize penetration and distribute the electromagnetic effect within the tissue, ensuring effective stimulation of stem cells and affected brain areas [12].

## **ELECTROMAGNETIC PRINCIPLES AND MECHANISMS OF DIAMAGNETIC THERAPY FOR REGENERATIVE APPLICATIONS IN NEURODEGENERATIVE PATHOLOGIES.**

Diamagnetic therapy operates on principles of electromagnetic induction, applying high-rate pulsed magnetic fields to interact with biological tissues in a controlled manner. This methodology leverages several core electromagnetic principles, foundational for promoting cellular repair and regenerative processes in neurodegenerative conditions like Parkinson's and Alzheimer's disease.



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## **Inductive Coupling and Electric Field Generation.**

Diamagnetic therapy generates an electric field through a time-varying magnetic field, where the induced electromotive force (EMF) is proportional to the rate of change of current in the coil, represented by  $dI_{\text{coil}}/dt$  [1]. The current response in the coil follows an exponential rise based on the coil's inductance L and resistance  $R_{\text{coil}}$ , as described by:

$$
I_{coil}(t) = \frac{V_0}{R_{coil}} (1 - e^{-\frac{R_{coil}l}{L}})
$$

This induced current waveform impacts the tissues significantly, particularly when optimized for specific rise times to enhance interaction with biological cells [2, 3].



Figure 3: CTU rise time at maximum power. Energy delivered to coil is about 90 Joules [6].

#### **Frequency and Spatial Distribution.**

Diamagnetic therapy utilizes a frequency distribution concentrated in the high-frequency range, with a characteristic "knee" in the spectrum around 45 MHz. This frequency profile enables deeper tissue penetration and achieves isotropic distribution within the targeted area, a feature essential for effectively reaching deeper brain structures. A simplified frequency-domain model shows how parameter variations can optimize tissue interaction, enhancing therapeutic outcomes in neurodegenerative applications [4].



**Figure 4:** Frequency domain of first stage of a CTU waveform [7].

### **Electrochemical Interaction at Cellular Interfaces.**

At the cellular level, diamagnetic therapy induces electrochemical interactions involving ion transport and voltage-dependent ligand binding. This electrochemical process, modeled as an equivalent electrical circuit,

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defines the impedance ZA(s) based on binding constants alpha and kinetic parameters  $\Box$ a and  $\Box$ b, with respect to changes in surface concentration  $\Delta\Gamma a(s)$  [11, 12].

$$
Z_A(s) = \frac{1}{q_a s(\alpha + v_a)}
$$

This model highlights how electromagnetic fields enhance stem cell functionality, specifically by improving the cellular responses essential for regeneration and repair in neurodegenerative tissues.

Larmor Precession and Effects.

The magnetic fields in diamagnetic therapy induce Larmor precession in bound ions within cells, where the Larmor frequency  $\omega$ L is defined as:

$$
\omega L = \frac{qB_0}{2m}
$$

This precession mechanism is pivotal in altering biochemical interactions within the cells, supporting synaptic plasticity-a critical factor in neuroregenerative therapies. Larmor precession may positively affect synaptic plasticity and neuronal connectivity, which are essential for functional recovery in neurodegenerative diseases [6].

## **DISCUSSION**

#### **Electromagnetic Mechanisms of Interaction**

The observed beneficial effects of diamagnetic therapy on stem cells and brain tissues can be explained through several physical mechanisms. Key mechanisms include:

- Ion Channel Modulation: Low-frequency electromagnetic fields influence the opening and closing of sodium and calcium ion channels, regulating membrane potential and modulating neuronal transmission.
- Effects on Cell Membrane: Diamagnetic therapy affects cell membrane fluidity, facilitating permeability and improving cellular communication, which is essential for stem cell differentiation.
- Larmor Precession: The presence of a static magnetic field causes the precession of ions within cells, a phenomenon that may alter biochemical reactions in neurons and promote positive effects on synaptic plasticity.

#### **CLINICAL POTENTIAL OF DIAMAGNETIC THERAPY**

The success of diamagnetic therapy as a support to regenerative medicine opens new avenues for neurodegenerative diseases. In Parkinson's disease, stimulation could promote the differentiation of cells into dopaminergic neurons, improving motor symptoms. In Alzheimer's disease, promoting neurogenesis and supporting existing synapses could slow cognitive decline.





## **CONCLUSIONS**

#### **Potential Physical Mechanisms of Interaction.**

The use of electromagnetic fields, particularly in the form of diamagnetic therapy, provides a physical basis for interactions with stem cells and damaged neural tissues. The described phenomena, including electromagnetic induction, ion channel modulation, and Larmor precession, suggest that targeted application of electromagnetic fields can influence cellular behavior in specific and therapeutic ways. The proposed electrochemical model further offers a quantitative framework to understand how stem cells can be activated and guided within the central nervous system to promote regeneration and functional recovery in neurodegenerative pathologies.

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