



BioSynaptica S.A.

### Verticals

Biotechnology  
Human Health

### Contact

Matías Depetris  
mdepetris@biosynaptica.com  
+54 93424625309  
www.biosynaptica.com

### Team

**Milagros Bürgi**  
Co-founder- Researcher  
**Matías Depetris**  
Co-founder - Director  
**Marcos Oggero Eberhardt**  
Co-founder - Researcher  
**Ricardo Kratje**  
Co-founder - CTO

### Previous funding

**USD 124,000**  
Founders  
**USD 100,000**  
Aceleradora Litoral  
**USD 200,000**  
FONDCE - BICE

### Required investment

**USD 2M**  
Operative actions (production process development, scale-up and analytical development).  
Analysis and complementary tests (proof of concept in animal models).  
Technical consultancies.

### Institutions linked to IP

UNL  
CONICET  
UNSAM

### Strategic alliances

Aceleradora Litoral  
Biotechnological Center of Litoral (FBCB-UNL)  
Max-Planck-Institut für Multidisziplinäre Naturwissenschaften (Germany)  
ICIVET (UNL)  
CAB-Startups (Argentine Chamber of Biotechnology)

## An innovative medicine to treat neurodegenerative diseases

### Summary

We are a Technology-Based Company incubated at the School of Biochemistry and Biological Sciences of the Universidad Nacional del Litoral (UNL) in the city of Santa Fe, Argentina. Our development focuses on the generation of an innovative neuropharmaceutical from human erythropoietin (hEPO) derivatives designed to meet the needs of millions of people suffering from central nervous system conditions, such as Parkinson's and Alzheimer's diseases, amyotrophic lateral sclerosis, multiple sclerosis, neurotraumas, among others.

### Detected problem

Neurodegenerative diseases are disorders characterized by a progressive deterioration in motor function and/or cognition caused by a selective loss of neurons from the central nervous system. The best-known diseases such as Alzheimer's disease, Parkinson's disease, amyotrophic lateral sclerosis, among many others, are responsible of serious cognitive, physical, and psychosocial consequences and represent one of the main causes of morbidity in western countries. Besides, despite their significant worldwide increment, there are no current effective treatments to prevent or cure them.

### Technological solution

hEPO is a widely-used biotherapeutic product for the treatment of anemia, since it is responsible for the production and maintenance of red blood cells in circulation (erythropoiesis). This protein also protects neurons from the damage caused by neurotoxic agents (neuroprotection), reduces neuroinflammation and restores neuronal connections (neuroplasticity). However, when it is designed as a neuropharmaceutic for patients who do not suffer from anemia, the molecule displays erythropoiesis-associated side effects. At BioSynaptica, we modified the hEPO molecule by blocking the undesirable erythropoietic activity but preserving its neuroprotective and neuroplastic capacity and improving the properties and, as a result, we obtained a therapeutic candidate for the treatment of chronic diseases.

### Market

The global central nervous system treatment market size was USD 116 billion in 2021 and is expected to expand at a CAGR of 9.4% from 2021 to 2028. The neurodegenerative disease drug market size reached USD 40 billion in 2021 and is expected to register a CAGR of 3.1% over the same period.

The target market segment is biotechnology and/or pharmaceutical companies that may incorporate an innovative drug into their product portfolio. These companies will be able to commercialize the drug in a monopoly for 12 to 15 years in several countries from Asia, Europe, Oceania and America.

### Monetization model

#### **B2B: Licensing and royalties.**

Initial development results were protected by patent requests in Argentina and via PCT with WIPO. Based on the positive evaluation of this latest application, a patent nationalization phase was initiated in more than 20 countries in Asia, Europe, Oceania and America.

BioSynaptica will carry out successive rounds of investment to perform proof of concept tests in animal models of specific diseases, pre-clinical studies, and clinical trials phases I—II in the context of the medicinal species registration in Argentina and in other countries. Once these objectives are achieved, the technology will be sub-licensed to one or more pharmaceutical companies to perform clinical phase III and the drug introduction to the market. Initial payments, payments for fulfilling milestones, and royalties on sales will be collected.