Spanish SME pharma GMP-certified lab manufacturer (CMO) offers production capacity, facilities & expertise (as outsourcing) on sterile biological medicines for human use in Advanced Therapies & Blood Products (PRP). Seeking Int´/EU manufacturing agreements

Summary

Profile type
Business Offer

Company's country
Spain

POD reference
BOES20230613024

Profile status
PUBLISHED

Type of partnership
Commercial agreement
Outsourcing agreement
Investment agreement

Targeted countries
• World

Contact Person
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Term of validity
13 Jun 2023
12 Jun 2024

Last update
21 Jun 2023

General Information

Short summary

Spanish SME pharma GMP-certified lab manufacturer and developer (CDMO) located in Madrid offers production capacity, facilities, and expertise as outsourced service.
Proven expertise on sterile biological medicines for human use in the field of advanced therapies & Blood Products (e.g.: Platelet-Rich Plasma). Authorized and certified by the Spanish Drug Agency.
Likewise, established experience on trading of own branded blood derivative for regenerative medicine. Open to International/EU investo

Full description

Spanish SME lab founded in 2018 and located in Madrid dedicated to the manufacture and development (CDMO) of sterile biological medicines for human use on Cell Therapies, as for instance, blood derivatives such as Plasma-Rich Platelet (PRP) with Good Manufacturing Practices (GMP) certification and authorization issued by Spanish Drug Agency (AEMPS). We offer production capacity and facilities under sterile environment according to GMP and leading experience in Advanced Therapies.
Seeking, preferably EU/International customers and investors interested in Cell Therapy manufacture to outsource

their production capacity by hiring our services as CDMO. Similarly, open to EU/International R&D collaboration on hemoderivative projects.

Advantages and innovations

Advantages from our certified and safe manufacturing and developing service we offer:
- GMP certification granted by Spanish Drug Agency (AEMPS) for Cell Therapies.
- Issuance of certificate of analysis (COA).
- Drug safety by certificates of sterility and endotoxins.
- Control and monitoring of the whole process, including cold chain by assuring quality, efficacy, and safety of the treatment.
- Guaranteed ready-to-use packaging by direct application for treatment.
- Additionally, due to Advanced Therapies expertise under GMP, comprehensive consulting and training.

Technical specification or expertise sought

Stage of development

Already on the market

Sustainable Development goals

- **Goal 9: Industry, Innovation and Infrastructure**
- **Goal 3: Good Health and Well-being**

IPR Status

Secret know-how

Partner Sought

Expected role of the partner

Partners interested in manufacturing and/or developing their Cell Therapies or investing in our GMP facilities, as the SME offers their production capacity as a CDMO. Open as well to partners interested in R&D collaborative projects/consortia.

Type of partnership

Type and size of the partner

Commercial agreement

Outsourcing agreement

Investment agreement

- **Big company**
- **R&D Institution**
- **SME 50 - 249**
- **SME <=10**
- **Other**
- **University**
- **SME 11-49**

Dissemination

Technology keywords

- **06002002 - Cellular and Molecular Biology**

Market keywords

- **05007007 - Other medical/health related (not elsewhere classified)**
- **05003001 - Therapeutic services**
- **05004003 - Laboratory equipment**
- **04006 - Cellular and Molecular Biology**
- **09004001 - Business products and supplies**

Targeted countries

- **World**

Sector groups involved

- **Health**