MANUFACTURING TECHNICIAN

Company Description

VIVEbiotech is a GMP Contract Development and Manufacturing Organisation (CDMO) specialized in lentiviral vectors. As specialists in the development and manufacture of lentiviral vectors, we aim to contribute to the development of gene therapy treatments which provide an improvement in/cure of diseases in the field.

Information

- Deadline: 2020-06-30
- Category: Business
- Province: Gipuzkoa
- Country: Basque Country
- City: Donostia - San Sebastian

Company

VIVEBIOTECH SL

Main functions, requisites & benefits

Main functions

Execute Lentiviral vectors manufacturing processes according to GMP manufacturing guides. Drafting of manufacturing protocols, standardized operating procedures (SOPs) and other relevant manufacturing guides. Comply with and ensure compliance with internal manufacturing standards. Be part of a multidisciplinary team to carry out process optimizations and researches. Carry out maintenance of manufacturing areas to guarantee their status and compliance with GMP regulations.

Requisites

Background: Degree related to research in biomedicine (Science, Biology, Biochemistry, Chemistry, Pharmacy ...) or related engineering. Higher degree related. Ability to propose potential process improvements and essays. Results analyses. Previous experience in GMP manufacturing and/or bioprocesses. Spoken and written communication skills (both in Spanish and English). Team working skills and ability to work under deadlines.

Benefits

You will join a young company, in constant growth and with a highly qualified team. VIVEbiotech develops its activities in an atmosphere of honesty and ethics, promoting rigor, effort, perseverance, participation and teamwork, while remaining committed to the development of its personnel.