**Regd. Office:** CM-05, SINE office, 3rd Floor, CSRE building, IIT Bombay, Powai, Mumbai - 400076

**GMP Facility:** PLOT NO. R-977, TTC Industrial Area, MIDC Rabale, Navi Mumbai, India - 400 701



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| **Job Description** |
| **Job Title:** | QC Analyst |
| **Function:** | Quality Control | **Report to:** | QC Head  |
| **Location:** | Rabale, Navi Mumbai | **Position Type:** | Full Time |
| **About Immunoadoptive Cell Therapy Private Limited.** |
| ImmunoACT is a pioneering cell & gene therapy company, currently in Phase II trials for India’s first indigenously developed CAR-T therapy for the treatment of refractory/relapsing B-cell malignancies (HCAR19), with an aggressive pipeline to treat liquid and solid tumors, With strong, strategic Research (IIT-Bombay) and Clinical (Tata Memorial Hospital) collaborations, ImmunoACT is paving the way towards affordable and accessible gene-modified cell therapies for resource-constrained economies. Our first state-of-the-art GMP facility has been operational since mid-2022, as we scale-up and scale-out our production capabilities to serve the clinically unmet needs of patients across India.  **Our Vision:** To be a Leader in Cell & Gene Therapy for the Long-Term Cure of Patients, through Translational Research in India & beyond **Our Values:*** Accessible and affordable to all
* Cutting edge research
* Transforming ideas to reality

*Become a part of this revolution in the healthcare industry - grow your career with us.***Website** – <https://www.immunoact.com/> - Visit our website, understand about us.**LinkedIn Profile** - <https://www.linkedin.com/company/immunoact/>  |
| **Roles and Responsibilities** |
| * Plan and execute activities of Environment Monitoring and Personnel Monitoring activities and its associated activities.
* Conduct routine and non-routine analyses of in-process materials, raw materials, environmental samples, finished goods, or stability samples.
* Interpret test results, compare them to established specifications and control limits, and make recommendations on appropriateness of data for release.
* Compile laboratory test data and perform appropriate analyses.
* Complete documentation needed to support testing procedures.
* Calibrate, validate, or maintain laboratory equipment.
* Supply quality control data necessary for regulatory submissions.
* Receive and inspect raw materials.
* Ensure that lab cleanliness and safety standards are maintained.
* Support cGMP stability/IPQC testing activities for CGT products
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| **Qualifications and Educational Requirement** |
| * Master’s degree in microbiology, or relevant field.
* 1 to 2 years of experience in Environmental Monitoring in a cGMP area.
* Shall have experience in other microbiological related assays related to Sterility testing of Drug product, Water testing etc.
* Should be familiar with cGMP practices.
* Able to write technical reports/protocols/SOP with regulatory understanding
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| **Desired Qualities** |
| * Organizing skills
* Problem solving approach
* Multitasking
* Reporting skills
* Excellent Coordinating skills
* Flexibility
* A Team player
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| **Compensation** | Compensation would not be a constraint for the suitable candidate |
| **Mail CVs to Email Id** | *jobs@immunoact.com* |

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