# Position Details

## Research Scientist/Engineer- CSOF5

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| The following information is for applicants |
| Advertised Job Title | Research Scientist – GMP Upstream Lead |
| Job Reference | * 75901
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| Tenure | IndefiniteFull-time  |
| Salary Range | * AU$100,710 to AU$108,985 pa (pro-rata for part-time) + up to 15.4% superannuation
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| Location(s) | CSIRO Clayton, Victoria |
| Relocation Assistance | Will be provided to the successful candidate if required |
| Applications are open to | Australian/New Zealand Citizens and Australian Permanent Residents Only |
| Position reports to the | GMP Team Leader, CSIRO Biomedical Manufacturing |
| Client Focus – Internal | 10% |
| Client Focus – External | 90% |
| Number of Direct Reports | 0 |
| Enquire about this job | Contact Dr John Power via email at john.power@csiro.au or phone +61 3 9545 2472 |
| How to apply | Apply online at <https://jobs.csiro.au/> Internal applicants please apply via **Jobs Central**If you experience difficulties when applying, please email careers.online@csiro.au or call 1300 984 220. |

### Role Overview

The role of Research Scientist Staff in CSIRO is to conduct innovative research leading to scientific achievements that are aligned with CSIRO’s strategies. You may be engaged in scientific activity ranging from fundamental research to the investigation of specific industry or community problems. You will have the opportunity to build and maintain networks, play a lead role in securing project funds, provide scientific leadership and pursue new ideas and approaches that create new concepts.

The purpose of this position is to provide hands-on technical leadership in the field of Upstream (bioreactor) operations for manufacture of biotherapeutic proteins and vaccines within a GMP setting. The position reports to the GMP Team Leader in CSIRO Biomedical Manufacturing and is responsible for provision of advanced knowledge in the discipline of bioreactor operations. The role plays a key function in facilitating transfer of new bioreactor protocols into the GMP laboratory. The successful candidate will be a team player with highly developed written and verbal communication skills.

The position is based at Clayton, Melbourne but in the first year the appointee may be required to work at Parkville, or at non-CSIRO sites elsewhere in Melbourne to gain experience in additional aspects to the cGMP manufacture of biologics.

### Duties and Key Result Areas:

* Provide hands-on leadership of upstream cell culture activities in a GMP setting and serve as a point of accountability for transfer and implementation of upstream manufacturing methods.
* Lead by expertise and influence to ensure that new manufacturing protocols are established in accord with the cGMP quality framework and are completed within the agree timeframes and budget.
* Generate and analyse upstream bioprocess data to identify opportunities for continuous improvement in the operation of bioprocess systems.
* Assume ownership of upstream bioprocess equipment and maintain the same equipment in a GMP-ready state.
* Accurately write and review GMP batch manufacturing records. Communicate research results to clients and the scientific community through oral and written reports, which may include the preparation of documents for patent applications.
* Engage with clients as an authority on upstream cell culture methods. Act as a trusted advisor, utilising knowledge of client’s business and understanding of their underlying needs.
* Participate in project scoping, planning, and budgeting activities that shape the future direction of the upstream function. Anticipate industry needs and market direction through client liaison/networking and identify and adapt quickly to changes.
* Communicate openly, effectively and respectfully with all staff, clients and suppliers in the interests of good business practice, collaboration and enhancement of CSIRO’s reputation.
* Work collaboratively as part of a multi-disciplinary, regionally dispersed research team, and business unit to carry out tasks in support of CSIRO’s scientific objectives.
* Adhere to the spirit and practice of CSIRO’s Code of Conduct, Health, Safety and Environment procedures and policy, Diversity initiatives and Making Safety Personal goals.
* Other duties as directed

## **Required Competencies:**

* **Teamwork and Collaboration:** Cooperates with others to achieve organisational objectives and may share team resources in order to do this. Collaborates with other teams as well as industry colleagues.
* **Influence and Communication:** Uses knowledge of other party's priorities and adapts presentations or discussions to appeal to the interests and level of the audience. Anticipates and prepares for others reactions.
* **Resource Management/Leadership:** Allocates activities, directs tasks and manages resources to meet objectives. Provides coaching and on the job training, recognises and supports staff achievements and fosters open communication in the team.
* **Judgement and Problem Solving:** Investigates underlying issues of complex and ill-defined problems and develops appropriate response by adapting/creating and testing alternative solutions.
* **Independence:** Plans, sets and works to meet challenging standards and goals for self and/or others. Recognises where endeavours will make the most impact or difference, decides on desired outcome and sets realistic goals to reach this target.
* **Adaptability:**Copes with ambiguity or situations that lack clarity. Adapts readily to changing circumstances and new responsibilities (which may include activities outside own preferences) in the interests of achieving team objectives. Recognises the need for and undertakes personal development as a result of changes.

## **Selection Criteria**

#### Essential

*Under CSIRO policy only those who meet all essential criteria can be appointed.*

1. A degree in Biotechnology, Biochemical Engineering, or related field.
2. Industrial experience in the development and application of cell culture methods for manufacture of biotherapeutic proteins and vaccines for regulated markets.
3. Theoretical and practical knowledge of bioreactor unit operations including process development, optimisation, transfer, and troubleshooting.
4. Extensive experience working under the framework for Good Manufacturing Practices (GMP) gained through industrial experience.

## **Desirable:**

1. An advanced degree in Biotechnology, Chemical Engineering, or other relevant field.
2. Direct experience and working knowledge of the GMP frameworks for manufacture of human vaccines and biotherapeutics with global regulators such as the Therapeutic Goods Administration, US Food and Drug Administration, or European Medicines Authority.

Special Requirements

* The successful candidate will be asked to obtain and provide evidence of a National Police Check or equivalent. Please note that people with criminal records are not automatically deemed ineligible. Each application will be considered on its merits.
* The position is based at Clayton, Melbourne but in the first year the appointee may be required to work at Parkville, or at non-CSIRO sites elsewhere in Melbourne to gain experience in additional aspects to the cGMP manufacture of biologics.

## **About CSIRO:**

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Find out more about CSIRO [Manufacturing](https://www.csiro.au/en/Research/MF)