

UQ Winter Research Scholarship Projects in Faculty of Medicine 2018

Read about the program on the <https://employability.uq.edu.au/node/215/0#0> page, and apply online from 5 March – 3 April 2018 via <https://employability.uq.edu.au/node/215/2#2>

Please take note of where each project is located. Projects are listed under the unit names on the application page (CareerHub).

Scholars can select from one of the following to see the associated projects:

- [Centre for Health Service Research](#)
- [Child Health Research Centre](#)
- [Princess Alexandra Hospital Southside Clinical Unit](#)
- [QIMR Berghofer Medical Research Institute](#)
- [School of Biomedical Sciences](#)
- [UQ Centre for Clinical Research](#)
- [UQ Diamantina Institute](#)

Important: These projects are located at multiple sites at St Lucia and Herston campuses and hospitals in Brisbane, Ipswich, and a number of rural and remote area facilities throughout the rest of the state.

Find out more about our [research sites](#) and research in our [clinical schools](#) and hospital sites.

[Centre for Health Service Research](#)

- Project 01** [RESCUE: Reduce sitting to improve cognitive function in elders](#)
Project 02 [Quality indicators for the aged](#)
Project 03 [Healthy ageing and well-being in older Indigenous adults](#)
Project 04 [Scoping review: Lifecourse factors associated with frailty](#)

[Child Health Research Centre](#)

- Project 05** [Bacterial interference: An alternative to antibiotics for the treatment of middle ear infections in Indigenous children?](#)

[Princess Alexandra Hospital Southside Clinical Unit](#)

- Project 06** [Mental health outcomes for people with Treatment Refractory Schizophrenia following clozapine withdrawal](#)

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- Project 15** [Pathogenesis and therapy for type 1 diabetes](#)

Project Details

Centre for Health Service Research

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| Project title: | RESCUE: Reduce sitting to improve cognitive function in elders |
| Project duration: | 30 hours per week for 6 weeks |
| Description: | This project will assist with a randomised controlled trial to examine whether a 12-week lifestyle program is beneficial for physical and brain health. The student will assist with assist with data management and processing. This will involve checking self-report questionnaires for completion and scanning and uploading data into a database, initialising monitors used to record sitting and movement and downloading data from those monitors, and preparing data for analyses. Other tasks may include searching the literature and formatting manuscripts for publication. |
| Location: | Building 33, Princess Alexandra Hospital, Woolloongabba |
| Expected outcomes and deliverables: | Participants will gain skills in the conduct of randomised controlled trials, including data collection, cleaning and analyses. They may have the opportunity to be involved in writing peer-reviewed publications. |
| Suitable for: | This project would be suitable for students in a health-related discipline. |
| Primary Supervisor: | Dr Paul Gardiner |
| Supervisor's contact details: | p.gardiner@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | Quality indicators for the aged |
| Project duration: | Up to 36 hours per week for 6 weeks |
| Description: | Quality of Care is an international priority in health service delivery. Our Centre provides a unique methodology for the development of quality indicators. We aimed to develop outcome, process and structure quality indicators in relation to common geriatric syndromes and function for the care of the frail aged hospitalised in acute general medical wards and the emergency department. A formal approach was taken for the development of quality indicators, including expert opinion, field study data and a formal voting process. We are at the concluding end of this project where involvement provides unique insight into the methodology for developing quality indicators and manipulating complex datasets. |
| Location: | Building 33, Level 2, Princess Alexandra Hospital, Woolloongabba |
| Expected outcomes and deliverables: | Small literature searches will be completed to update the evidence on geriatric syndromes relating to the quality indicators to assist in finalising the documentation for publication. Some data checking will be carried out. There will be an opportunity to manipulate the dataset using SPSS to provide some frequency data and prepare some tables. A sophisticated voting system has been used with the expert panels to finalise these QIs. A round of voting will be undertaken during this period. The scholar will have the opportunity to facilitate the voting which will be a unique learning experience. |
| Suitable for: | An individual with a keen eye for detail, and a willingness to learn new skills. All information will be explained on the job so no prior experience is required. |
| Primary Supervisor: | Dr Melinda Martin-Khan |
| Primary contact, if not supervisor: | Dianna Ang |

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| Supervisor's contact details: | m.martinkhan@uq.edu.au |
| Note before application: | The supervisor CAN be contacted by students prior to submission of an application |

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| Project title: | Healthy ageing and well-being in older Indigenous adults |
| Project duration: | 20 - 36 hours per week for 4 weeks |
| Description: | Projects are currently underway investigating the prevalence of dementia and healthy ageing in Indigenous communities in the Torres Strait and Far Northern Queensland. A winter scholar is required to assist with work associated with these projects. Tasks may include: assistance with the development and piloting of a healthy ageing assessment tool; working with data from the Australian Aboriginal and Torres Strait Islander Health Survey, or literature searching and the writing up of a scoping review. |
| Location: | Princess Alexandra Hospital, Woolloongabba |
| Expected outcomes and deliverables: | Literature search to identify assessment items for inclusion in the healthy ageing assessment tool. Extraction of data into spreadsheets. Mapping of data items to different lifestyle components: diet, physical activity, social activities, access to health care services etc. Evaluation around the suitability of data for inclusion in the tool. Review of data items in the Health Survey. Extraction of variables of interest. Basic analysis of health Survey data. Writing up of results. Systematic literature searches to identify literature for inclusion in the scoping review. Assisting with the writing up of the review. |
| Suitable for: | An individual with a keen eye for detail, and a willingness to learn new skills. Experience in quantitative data analysis and/or systematic literature searching would be highly regarded. |
| Primary Supervisor: | Dr Yvonne Hornby-Turner |
| Supervisor's contact details: | (07) 3176 6636 |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | Scoping review: Life course factors associated with frailty |
| Project duration: | 20-36 hours per week for 4 weeks |
| Description: | <p>Frailty is a common clinical syndrome in older adults that carries an increased risk for poor health outcomes including falls, incident disability, hospitalization, and mortality. Identifying factors across the life course that can delay the onset of frailty or improve the quality of life for those living with frailty is therefore a key public health priority.</p> <p>This project will complete a scoping review to identify life course associated with frailty in community dwelling older persons. A review of peer-reviewed published literature has already commenced. The work for this project will involve updating this literature search and the assistance with all other tasks associated with getting the paper ready for publication (see below). This will result in a co-author opportunity with one of the world's leading experts in frailty research.</p> |
| Location: | Princess Alexandra Hospital, Woolloongabba |
| Expected outcomes and deliverables: | Systematic literature searches will be performed to identify papers reporting on health assets that are associated with a lower frailty risk in community dwelling persons. Review of papers for relevant data and the extraction of this data into a spreadsheet. The mapping of evidence according to six key domains: Social, behavioural, economic, environmental, personal, and health and social services. Evaluation of data for quality and reliability of evidence. Assist in finalising the documentation for publication. Preparation of tables for inclusion in publication. |

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| Suitable for: | An individual with a keen eye for detail, and a willingness to learn new skills. All information will be explained on the job so no prior experience is required. |
| Primary Supervisor: | Associate Professor Ruth Hubbard |
| Supervisor's contact details: | y.hornbyturner@uq.edu.au |
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Child Health Research Centre

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| Project title: | Bacterial interference: An alternative to antibiotics for the treatment of middle ear infections in Indigenous children? |
| Project duration: | 20 hours per week for 4-6 weeks |
| Description: | <p>Australia's Indigenous children have some of the highest rates of middle ear infections (otitis media) in the world. This can result in hearing loss and have a devastating impact on language development, education and future employment. Despite antibiotic treatment and widespread vaccination programs, there has been little change in the prevalence of the disease.</p> <p>The aim of this project is to develop a probiotic specifically to treat otitis media in Australian Indigenous children using strains of 'friendly' bacteria from the upper airways of Indigenous children from remote communities. This winter project will contribute to this aim by assisting in the bacterial interference studies, where we determine which of these 'friendly' bacteria have the ability to kill the disease-causing bacteria in the lab.</p> |
| Location: | Centre for Children's Health Research, South Brisbane |
| Expected outcomes and deliverables: | The scholar will gain hand-on skills in the laboratory including culture-based microbiological work, writing laboratory reports, preparing a range of media and broths. The scholar will have the opportunity to present the results of their project via oral presentation within CHRC. |
| Suitable for: | This project is suitable for students with a microbiological background/ pre-medical students with an interest in the MD-HRD pathway. There is a possibility for expanding the project into a HRD for the appropriate candidate. |
| Primary Supervisor: | Dr Seweryn Bialasiewicz |
| Primary contact, if not supervisor: | Dr Andrea Coleman |
| Supervisor's contact details: | seweryn@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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Princess Alexandra Hospital Southside Clinical Unit

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| Project title: | Mental health outcomes for people with Treatment Refractory Schizophrenia following clozapine withdrawal |
| Project duration: | 20-36 hours per week for 4-6weeks |
| Description: | Clozapine is reserved for people with treatment refractory schizophrenia who have failed two trials of other antipsychotics. However, it has the most severe adverse events of all antipsychotics including agranulocytosis and myocarditis. This can lead to precipitous cessation of clozapine, and the risk of severe rebound psychosis. |

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| | This project will involve a Cochrane-like systematic review of the literature relating to the mental health outcomes of people who are ceased on clozapine. It will include data from RCTs, observational studies and case reports. This information will help inform both clinicians and mental health consumers as to the expected outcomes associated with clozapine cessation, and guide clinical management. |
| Location: | Princess Alexandra Hospital, Woolloongabba |
| Expected outcomes and deliverables: | The expectation is that the student will be a co-author on a manuscript for submission to a peer reviewed journal. |
| Suitable for: | Students with an interest in or experience in systematic reviews of medical literature |
| Primary Supervisor: | Associate Professor Dan Siskind |
| Supervisor's contact details: | d.siskind@uq.edu.au |
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QIMR Berghofer Medical Research Institute

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| Project title: | The D-Health trial |
| Project duration: | Flexible according to student's availability, however, expected workload is a minimum of 20 hours and maximum of 36 hours per week for a period between 4 and 6 weeks. |
| Description: | The D-Health Trial: The D-Health Trial is one of the largest trials of vitamin D ever conducted. Over 21,000 Australian adults aged over 65 years were enrolled between 2014 and 2015 and randomised to take either monthly vitamin D or placebo for five years. They are being followed to ascertain health events through self-report and linkage with Australian health databases. |
| Location: | QIMR Berghofer Medical Research Institute |
| Expected outcomes and deliverables: | Students will have the opportunity to learn how a trial as large as the D-Health trial operates. They will contribute to most aspects of the project, including telephoning participants to administer interviews, cleaning data, and dispensing tablets. Students interested in data analysis may have the opportunity to contribute to preliminary analyses. |
| Suitable for: | This project is suitable for students who have an interest in research into public health using epidemiological approaches. They must have an excellent telephone manner and extremely high attention to detail. |
| Primary Supervisor: | Associate Professor Rachel Neale |
| Supervisor's contact details: | Rachel.neale@qimrberghofer.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | Projects in melanoma research |
| Project duration: | Flexible according to student's availability, however, expected workload is a minimum of 20 hours and maximum of 36 hours per week for a period between 4 and 6 weeks. |
| Description: | Treatment for late-stage melanoma is currently challenging due to the frequent occurrence of drug resistance. To identify new treatment strategies, we have used unbiased genome-wide functional experiments (shRNA and CRISPR) to identify unique melanoma gene dependencies (genes that when knocked out will kill a majority of melanomas, or that will kill drug-resistant melanoma cells). In |

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| | <p>this project, we are exploring the role of these novel genes to determine their significance as potential drug targets. Students will be responsible for helping characterize one of our identified candidate genes. Potential techniques to be used in this project include (but likely not all) cell culture, proliferation and or other related functional assays, protein analysis via western blots, genetic manipulation via CRISPR gene knock-out, or pharmacological inhibition through small molecule drugs.</p> <p>Project 2: The difference between a benign (safe) mole and a malignant (cancerous) melanoma can be difficult to determine. In most instances, suspect lesions are usually surgically removed, a process which can be both invasive and costly. Furthermore, these lesions frequently end up being classified as benign, resulting in unnecessary procedures. To address this, we have been developing a new, non-invasive approach to determine if lesions are benign or malignant. Potential techniques that students will perform include basic molecular biology techniques, DNA and RNA extraction, transcriptomic and genomic analyses with RT-PCR and PCR, and active trouble shooting to determine the sensitivity and specificity of these approaches.</p> |
| Location: | QIMR Berghofer Medical Research Institute |
| Expected outcomes and deliverables: | Applicants will gain knowledge surrounding the biology and current therapeutic intervention strategies of melanoma; gain expertise in a variety of molecular biology techniques and; may have the opportunity to generate data that will contribute toward a publication. Applicants will be expected to maintain a lab book, follow directions as given by supervisor, analyse results, and generate appropriate figures or graphs relating to their findings. A final report or oral presentation summarising their results may be required at the conclusion of the project. |
| Suitable for: | This project is open to applications from students with a background in Biomedical Science, Biochemistry and Molecular Biology and have an interest in Cancer Biology |
| Primary Supervisor: | Professor Nicholas Hayward |
| Primary contact, if not supervisor: | Dr Ken Dutton-Regester |
| Supervisor's contact details: | ken.dutton-regester@qimrberghofer.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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School of Biomedical Sciences

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| Project title: | Prenatal alcohol programs offspring disease |
| Project duration: | 20 hours per week for 4 weeks |
| Description: | <p>Alcohol consumption amongst women of reproductive age is widespread in Australia (~80%). Given that ~50% of all pregnancies are unplanned, there is the potential to expose the early embryo to alcohol before the pregnancy has even been detected. Although the teratogenic effects of alcohol on the fetus are well-documented, particularly the impacts on the brain and subsequent behavioural deficits, the programming of other aspects of offspring health by more moderate exposures to alcohol are less well defined.</p> <p>The Moritz laboratory has developed several preclinical models of alcohol exposure that are relevant to common patterns of alcohol consumption in reproductive age women. This includes a chronic, low dose throughout pregnancy; an acute, binge exposure later in pregnancy; and a moderate exposure just around the periconceptual period (PCP) in which the dam is</p> |

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| | <p>exposed during ovulation, conception and pre-implantation embryo development. Two of these models have been well characterised in terms of impacts on the metabolic health of adult offspring (the PCP and chronic model) and indicate that prenatal alcohol can result in a tendency to develop a 'taste' for a high fat diet, an increased propensity for obesity, and results in glucose intolerance and insulin insensitivity. However, the binge model has been relatively understudied in these areas. This model exposes the fetus to a relatively high level of alcohol at embryonic day 13.5 and 14.5, a period well within the 1st trimester equivalent in human pregnancies (~5th-7th week of pregnancy), when women often don't know that they are pregnant and have not yet altered their drinking habits. This is a peak period of organogenesis and thus a critical window during development where alcohol exposure has the potential for profound effects on offspring health. Recent studies of Australian women of reproductive age indicate that this high level, 'special occasion', drinking is a prominent mode of alcohol consumption and thus can pose a potential risk to the health of the fetus in an undiagnosed pregnancy.</p> <p>Given the already high rates of obesity and metabolic disease in the Australian population, which at current projections are likely to rise, it is important to understand how this prevalent lifestyle factor can potentially contribute to this burden of disease.</p> |
| Location: | Sir William MacGregor Building (64), UQ St Lucia Campus |
| Expected outcomes and deliverables: | By participating in this project, students can expect to gain experience in using a preclinical (animal) model to understand how a maternal insult can program disease in offspring. They will gain hands-on experience in experiments looking at metabolism and food preference and will assist with data collection and analysis. This project is expected to contribute to a publication on the metabolic outcomes of offspring exposed to a 'binge' alcohol exposure. |
| Suitable for: | This project would be suitable for a student with a background in physiology or endocrinology, or a pre-medical provisional student. |
| Primary Supervisor: | Professor Karen Moritz |
| Primary contact, if not supervisor: | Dr Lisa Akison |
| Supervisor's contact details: | k.moritz@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | The effects of placental function on pregnancy complications |
| Project duration: | 24 hours per week for 6 weeks |
| Description: | The Placenta is the central regulator of pregnancy success. A range of pregnancy complications result from placental dysfunction leading to long term health complications for both the mother and child. This project will investigate placental stress signalling in a range of pregnancy disorders including gestational diabetes and pre eclampsia to understand pathways which can be targeted to treat these conditions. |
| Location: | Sir William MacGregor Building (64), UQ St Lucia Campus |
| Expected outcomes and deliverables: | This project will allow students to gain skills required for a successful career in research. Students will be offered the opportunity to learn multiple laboratory techniques from an experienced research team. Students will be asked to participate in laboratory meetings and present their research findings to the lab group in a friendly interactive format. |
| Suitable for: | Students interested in a career in research or with a hope to pursue further study in a field which would benefit from understanding complications of pregnancy. |
| Primary Supervisor: | Dr James Cuffe |

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| Supervisor's contact details: | j.cuffe1@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | Metabolic treatments for epilepsy |
| Project duration: | 30 hours per week for 6 weeks |
| Description: | Although many people with epilepsy can be successfully treated with drugs or surgery, about 45% still suffer from seizures. To this end the laboratory is seeking new metabolic treatments to improve energy metabolism in the epileptic brain and to prevent seizures. |
| Location: | Skerman Building (65), UQ St Lucia Campus |
| Expected outcomes and deliverables: | Increased understanding of energy deficiency in epilepsy models. Progress regarding development of new metabolic treatments. |
| Suitable for: | Any 2nd or 3rd yr student, BIOC2000 a plus |
| Primary Supervisor: | Associate Professor Karin Borges |
| Supervisor's contact details: | k.borges@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | Harnessing the immune system to battle ovarian cancer: A novel approach using non-coding RNAs |
| Project duration: | 36 hours per week for 5-6 weeks |
| Description: | <p>We are interested in developing novel nano-therapeutic methods to overcome immune suppression in ovarian cancer. Ovarian cancer is the most deadly type of gynaecologic disease with more than 1500 new cases being diagnosed each year in Australia. The high recurrence rate is a major challenge in the clinical management of high grade serous ovarian cancer. While stimulating our own immune system to recognize and attack tumour cells represents an attractive means to facilitate complete elimination of tumours, emerging data suggest that many of the immunotherapy tools, such as immune checkpoint inhibitors, are minimally active in ovarian cancer. We aim to develop effective strategies to enhance the infiltration and function of cytotoxic T lymphocytes in ovarian tumours and to develop clinically feasible means to monitor T-lymphocytes activity in tumours following therapy. Ultimately, strategies developed in this project could harness the power of the immune system to eliminate tumours and significantly increase the survival of patients with ovarian cancer.</p> <p>We are seeking a motivated undergraduate student who is interested in contributing to a large project involving nanotechnology and cancer biology, and who is eager to learn how to develop effective strategies to enhance anti-tumour immunity. The student will learn critical laboratory skills and knowledge needed to develop new strategies to enhance the infiltration and function of cytotoxic T lymphocytes in ovarian tumours. In addition, the student will gain experience in developing novel nanoparticle platforms for tumour-targeted delivery. He/She will gain experience in working in a multidisciplinary environment, obtain hands-on training from the lab head and a postdoctoral fellow, and contribute to an exciting project in the area of cancer nanomedicine and immunology. Students are strongly encouraged to contact the primary supervisor (Dr. Sherry Wu) prior to applying for this scholarship opportunity.</p> |
| Location: | Sir William MacGregor Building (64), UQ St Lucia Campus |
| Expected outcomes and deliverables: | The student will learn critical laboratory skills and knowledge needed to develop new strategies to enhance the infiltration and function of cytotoxic T lymphocytes in ovarian tumours. In addition, the student will gain experience in |

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| | developing novel nanoparticle platforms for tumour-targeted delivery. He/She will gain experience in working in a multidisciplinary environment, obtain hands-on training from the lab head and a postdoctoral fellow, and contribute to an exciting project in the area of cancer nanomedicine and immunology. The student will be required to contribute to other research activities in the laboratory, be a good lab citizen, and present his/her work in written or oral presentation format. |
| Suitable for: | This project is open to applications from students with a background in biomedical sciences, pharmacy, or biomedical engineering, who is interested in exploring research as a career path. |
| Primary Supervisor: | Dr. Sherry Wu |
| Supervisor's contact details: | sherry.wu@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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| Project title: | Sensory processing in zebrafish |
| Project duration: | 36 hours per week for 6 weeks |
| Description: | Our lab builds light-sheet microscopes, combines them with genetically-encoded indicators of neural activity, and uses them to image activity in all of the neurons in zebrafish brains while they respond to stimuli. We are interested in the pathways that information follows through the brain as the outside world is perceived and interpreted, and on the ways that these pathways change in disease states such as autism and schizophrenia. We have positions for biologists, physicists, mathematicians, and neuroscientists, especially those interested in spanning discipline boundaries. |
| Location: | School of Biomedical Sciences, UQ St Lucia Campus |
| Expected outcomes and deliverables: | Experience in microscopy, optical physics, neuroscience, and bioinformatics |
| Suitable for: | Particularly suitable for optical physicists and computational biologists who are interested in transitioning to neuroscience |
| Primary Supervisor: | Associate Professor Ethan Scott |
| Supervisor's contact details: | ethan.scott@uq.edu.au |
| Note before application: | The supervisor CAN be contacted by students prior to submission of an application |

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UQ Centre for Clinical Research

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| Project title: | Controlability of complex networks in the brain |
| Project duration: | 36 hours per week for 5 weeks |
| Description: | We are interested in the analysis of complex networks using computational and mathematical techniques. Such complex network can be inferred from the human brain. The goal of this project is to develop a theoretical-computational technique to assess the formation and evolution of brain network in the newborn infants. |
| Location: | UQ Centre for Clinical Research, UQ Herston Campus |
| Expected outcomes and deliverables: | The student will gain mathematical skills in brain data analysis, and have an opportunity to publish the outcomes of his/her project. |
| Suitable for: | Second/third year undergraduate students in school of mathematics and physics are highly encouraged to apply for this project. Third year undergraduate students in Electrical Engineering are also encouraged to apply |

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| Primary Supervisor: | Dr Steve Mehrkanoon |
| Supervisor's contact details: | s.mehrkanoon@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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UQ Diamantina Institute

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| Project title: | Pathogenesis and therapy for type 1 diabetes |
| Project duration: | 36 hours per week for 6 weeks |
| Description: | <p>i) Immunotherapy in type 1 diabetes Our laboratory focuses on investigating the immune defects leading the destruction the insulin producing cells of the pancreas and type 1 diabetes and studying immunotherapeutic strategies to correct these defects. We are investigating using a nanoparticle delivery system containing islet antigen and an immunomodulatory drug to reinstate T cell tolerance in a model of type 1 diabetes. The project will utilise animal models, in vitro assays of cellular function and flow cytometry to study immunological tolerance in this system.</p> <p>ii) Gut microbiota - host interactions in type 1 diabetes The gut microbiota has been shown to be disturbed in type 1 diabetes and this is linked to genetic risk and the environment. We are investigating the link between inflammation in the gut and the pancreas and the microbiota. We use human samples and animal models to characterise gut and pancreas derived proteins as well as sequencing the bacteria present in the gut. This project combines wet and dry lab techniques and may include proteomics, sequencing and bioinformatics.</p> |
| Location: | UQ Diamantina Institute, TRI Building, Woolloongabba |
| Expected outcomes and deliverables: | The student will be exposed to an active research environment and learn techniques that may include multi-colour flow cytometry, animal disease model of type 1 diabetes, proteomics, sequencing or histology. They will also be involved with experimental design, data analysis and interpretation. |
| Suitable for: | Students with a background in at least one of these subjects: immunology, microbiology or bioinformatics. |
| Primary Supervisor: | Dr Emma Hamilton-Williams |
| Supervisor's contact details: | e.hamiltonwilliams@uq.edu.au |
| Note before application: | The supervisor MUST be contacted by students prior to submission of an application |

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