

JOB DESCRIPTION

TELETHON KIDS INSTITUTE



Why is this Job Description being written?		<input type="checkbox"/> New Position <input type="checkbox"/> Replacement Position <input checked="" type="checkbox"/> Position re-designed <input type="checkbox"/> Position not previously described	
POSITION DETAILS:	Position Title:	RESEARCH ASSISTANT – THE ORIGINS PROJECT	
RFA:	Early Environment	Research Group:	The ORIGINS Project
Position reports to: (role)	Study Recruitment Coordinator		
Location: <i>include all possible locations</i>	Joondalup Health Campus		
POSITION PURPOSE: In one or two sentences briefly summarise the overall purpose of this role, i.e. broadly, what this role does and why			
<p>To recruit participants in the ORIGINS Project and sub-projects and to follow-up participants at a range of time points during the Project implementation. This position will contribute to the day to day running of The ORIGINS Project, along with a team of other Research Assistants. This position will support the collection of quality research data, including specimens, as part of the Project.</p>			
KEY RESPONSIBILITY AREAS <i>(Please list in order of importance)</i>			
Key Position Accountabilities What are the main areas for which the position is accountable	% of Total Role	Inputs: What are the key activities or tasks to be carried out?	Outputs: What are the expected end results? Measures: How it is measured

Participant engagement	40%	<ul style="list-style-type: none"> • Screening and recruitment of participants including, but not limited to, participant phone calls, letters and emails, including appointment bookings and regular contact or compliance post-randomisation calls • Developing and maintaining study-specific protocols, participant information sheets, participant consent forms and master files • Performing research clinic activities including: obtaining written consent, skin prick testing, venepuncture, and administering study specific questionnaires 	<ul style="list-style-type: none"> • Tasks to be completed to a high standard of quality • Completed research tasks according to project timelines and requirements 	<ul style="list-style-type: none"> • Positive feedback from team members • Research tasks completed with high quality in a timely manner
Sample collection	25%	<ul style="list-style-type: none"> • Prepare visit packs with sample collection tubes, datasheets, informed consent forms, and other relevant materials • Confirm patient suitability and eligibility regarding protocol inclusion and exclusion criteria • Ensure Informed Consent is obtained according to the Guidelines for Good Clinical Practice (GCP) • Collection of study related data and any specimens required for studies according to each study protocol with adherence to GCP guidelines • Communicate and liaise with JHC clinical staff and ORIGINS researchers regarding participant care 	<ul style="list-style-type: none"> • Data and sample collection from participants • Informed consent is documented in accordance with GCP requirements • Collection and processing of biological samples is performed correctly and in a timely manner • Optimal care, advocacy and support given to participants and their families 	<ul style="list-style-type: none"> • Correctly completed informed consent documents filed in appropriate participant records • Biological samples are collected at appropriate time points using the correct materials • Biological samples are delivered to laboratory staff for processing within the required time period for stability • Feedback from team members • Feedback from participants/families

Data collection	25%	<ul style="list-style-type: none"> • Collection of participant visit data in accordance with Good Clinical Practice and research standards • Entry of participant data into the research database • Follow up on outstanding clinical results and participant information as required. 	<ul style="list-style-type: none"> • Paper participant records are correctly completed and up to date • Participant data is entered into the database in a timely manner • Incomplete data is followed up in a timely manner 	<ul style="list-style-type: none"> • Review of participant paper records • Review of database for missing/incomplete/flagged data • Paper and electronic records are complete and up to date within required timeframe
Team Membership	10%	<ul style="list-style-type: none"> • Attend team meetings • Other administrative activities as required • Working cohesively and collaboratively with others – both internal and external 	<ul style="list-style-type: none"> • Tasks completed according to project timelines and deadlines • Effective, harmonious teamwork 	<ul style="list-style-type: none"> • Positive feedback from team members
ESSENTIAL SKILLS, KNOWLEDGE AND EXPERIENCE:				
Qualifications: what are the minimum educational, technical or professional qualifications required to competently perform role		<ul style="list-style-type: none"> • Relevant undergraduate science degree 		
Skills, Knowledge & Experience:		<ul style="list-style-type: none"> • Ability to work as a part of a team • Ability to obtain Working With Children Check • Right to live and work in Australia • Availability to work on scheduled clinic days • Experience in working with infants and children • High level of interpersonal, verbal and written communication skills • High personal motivation and ability to work independently • Possession of a current WA drivers licence and your own transportation • Strong computer skills using Microsoft Office, data management and analysis programs • Attention to detail • Demonstrate excellent team working skills as well as ability to work using own initiative • Time management skills/ability to prioritise workload 		

DESIRABLE SKILLS, KNOWLEDGE AND EXPERIENCE:

Qualifications: what are the minimum educational, technical or professional qualifications required to competently perform role

- Registered nurse with current AHPRA registration

Skills, Knowledge & Experience:

- Previous laboratory experience including blood processing and sterile technique
- Previous experience in research
- Experience in statistical and data analysis
- Paediatric experience

SCOPE:

Financial accountability: Does this role have accountability for a budget?

No

People responsibility: Does this role have any direct reports or indirect reports (through direct reports)?

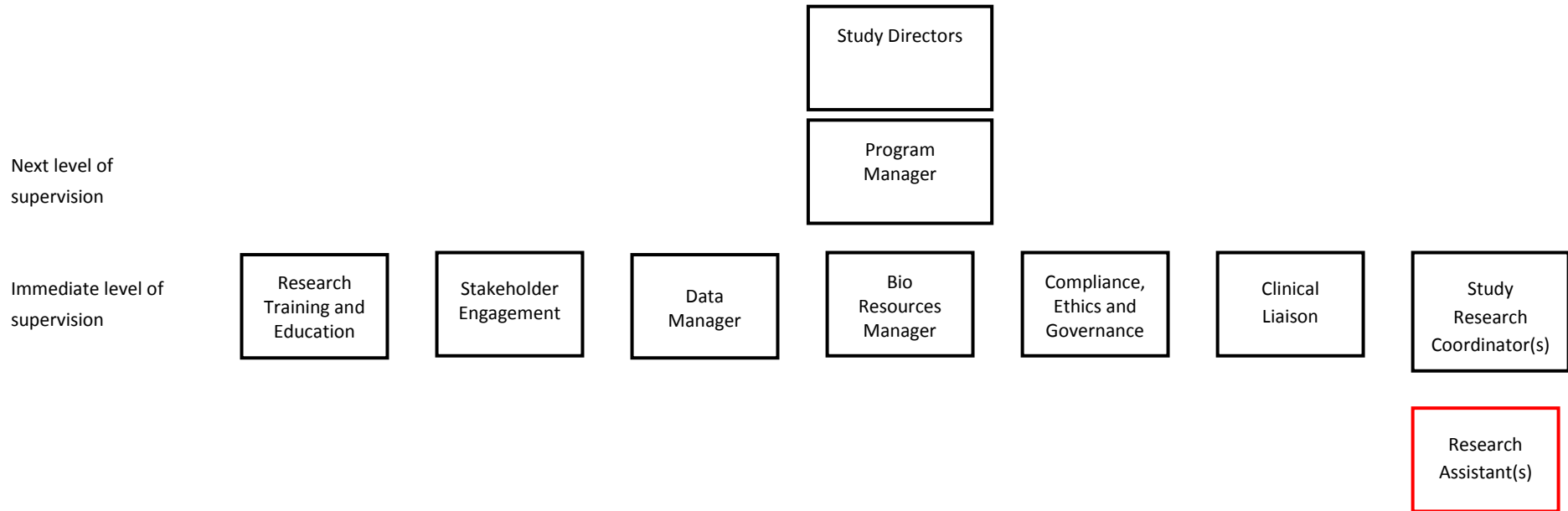
No. of direct reports

None

No. of indirect reports

None

ORGANISATIONAL CHART: (please complete using position titles or insert diagram below)



ADDITIONAL INFORMATION: is there any additional information that needs to be understood to explain this role?

This position is subject to a successful Criminal Record Screening Check and a Working with Children (WWC) Check. This is a compulsory check for people who are involved with child-related work in Western Australia.