## Transcriptional Analysis of ASCL1 and the Effects of Neurogenesis in GABAergic Neurons [Logbook]

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**Supervisor: Dr Frank Schubert** 

Date	Object	ive(s)	Next Steps
10/06/2020	•	Meeting with	NEXT STEP: Start research on
		supervisor Dr Schubert	the topic of neurogenesis
		to discuss ideas for the	NEXT STEP: Consider planning
		project and the next	to change the project structure
		steps	from a lab project to a
			computer-based project
25/07/2020	•	Decide on what The	<b>NEXT STEP</b> : Consider the
		approaches to use for	approaches to computer-based
		the project idea and	projects and look at past
		provide advantages	projects
		and disadvantages to	
		each approach	
	Litorot	uro roviou	
	Literati	ure review (Advantage) easy to	
		conduct	
		(Disadvantage) very	
		hard to get the	
		standard of a first-class	
		and requires critical	
		analysis and as require	
		a lot of journals and	
		papers to review	
	Biointo	ormatics	
	•	(Advantage) Easier to	
		get the first class as the supervisor is more	
		experienced	
		(Disadvantage)	
		Requires learning	
		bioinformatics	
		programs and requires	
		a lot more analysis	
	Data Submitted by		
	transcr	iption of RNA molecules	
	•	(Advantage) is in line	
		with my project based	
		on the comparative	
		analysis of the	
		expression of genes among different and	
		similar cell clusters.	
		The idea is dissociation	
		from the cells to	
		obtain a data sheet	
		which is	
		multidimensional	
		which requires	

07/10/2020	mathematical methods to reduce the complexity of the data. This will result in the 2D graphical representation between the gene cluster and the clustering of the gene themselves  Refine my introduction	<b>NEXT STEP</b> : Define my aims
	·	·
15/10/2020	<ul> <li>The first group meeting with Dr</li> <li>Schubert on how to organise your project</li> </ul>	NEXT STEP: Refine my introduction and how clear understanding of my aims
28/10/2020 - 21/11/20	<ul><li>Refine my introduction</li><li>Defined my aims</li></ul>	NEXT STEP: Begin to research the methodology that is going to be used in my project
22/11/2020	<ul> <li>Research on the methodology that is going to be used; RNA-seq analysis and choose my three primary datasets</li> </ul>	NEXT STEP: Begin to learn how to use Galaxy for my project and to further define my methodology and include the use of Gene ontology in the methods  NEXT STEP: Submit my project to Dr Schubert for feedback
30/11/2020	<ul> <li>Group meeting with Dr Schubert discussing how to deal with the technical aspects for gene lists and GO term analysis and STRING analysis</li> </ul>	NEXT STEP: Practice how to use GO term analysis and practice string analysis
18/12/2020	<ul> <li>Submit my project to Dr Schubert for feedback</li> </ul>	NEXT STEP: N/A
13/01/2021	<ul> <li>Created a defined title for the project. "The transcription analysis of ASCL1 and the effect of neurogenesis in GABAergic Neurons"</li> </ul>	NEXT STEP: Reorganise the project into clearer sections with the result section, discussion and the conclusion being added
10/02/2021	Refined my introduction	NEXT STEP: Plan to add diagrams and images into my project showing tangential migration, bHLH proteins and stem cell division
17/02/2021 - 18/02/2021	<ul> <li>Added diagrams of bHLH proteins and stem cell division</li> </ul>	NEXT STEP: Plan to add diagrams and images into my project showing tangential migration

18/02/2021	<ul> <li>Added diagrams showing tangential migration</li> </ul>	<b>NEXT STEP</b> : Sort out the methods section
19/02/201	The first training session for data techniques with Dr Schubert. Using GEO2R. Comparing two or more samples. usually derived from microarray or NGS data, and typically is used for differential gene expression analysis	NEXT STEP: Obtain two more datasets to supplement the existing data NEXT STEP: Process the previous microarray data acquired early
5/03/2021	<ul> <li>Training session         exploring with Dr         Schubert through GO         term analysis and         String first and         introducing Galaxy</li> </ul>	<b>NEXT STEP</b> : Perform a preliminary GO term analysis
19/02/2021 - 10/03/2021	<ul> <li>Obtain preliminary results from the Venn diagram and the STRING analysis</li> <li>Constructed the methodology pipeline and accompanying diagram</li> <li>Project meeting with Dr Schubert discussing the poster and how to prepare for it</li> </ul>	NEXT STEP: Sort out the methods section; replacing GeneVenn with InteriVenn NEXT STEP: Obtain two more datasets NEXT STEP: Sort out the methods section NEXT STEP: Finalising my poster NEXT STEP: Replace AmiGo2 with g:profiler
11/03/2021	<ul> <li>Preliminary Principal component analysis plot (PCA) and uniform manifold approximation and projection (UMAP) results</li> <li>Preliminary volcano plots</li> </ul>	NEXT STEP: Reanalyse the STRING data NEXT STEP: Make a table in the methodology section which details of the datasets NEXT STEP: Refine the preliminary Principal component analysis plot (PCA) and uniform manifold approximation and projection (UMAP) results and the preliminary volcano plots

12/02/2024 24/02/2024	Company and a collection	NEVT CTED. Ctant the
12/03/2021 - 24/03/2021	<ul> <li>Constructed a table in the methodology section which details of the datasets used in the present study and the biological function</li> <li>Construct a GO Term enrichment analysis visualised as a Manhattan-like-bubble graph.</li> <li>Replace AmiGo2 with g:profiler for the GO term analysis. Results of GO enrichment analysis for A, molecular function B, biological process and C, cellular component categories</li> </ul>	NEXT STEP: Start the discussion based on the experiment data NEXT STEP: Construct a diagram displaying adult neurogenesis
16/03/2021 - 31/03/2021	<ul> <li>Project meeting with         Dr Schubert discusses         possible points for             areas for the             discussion area     </li> <li>Expanded the             discussion section to             include "limitations of             experiment design"             and "future research"             section</li> </ul>	NEXT STEP: Being to research various discussion points. NEXT STEP: Reanalyse the STRING data
05/04/2021 - 28/04/2021	<ul> <li>Identified specific factors associated with ASCL1 expression:         Pixt2, FGFR1, GOT2,         NeuroD6 and NOTCH2         were found and play roles in the brain development</li> <li>Finalise the STRING analysis</li> </ul>	NEXT STEP: Decided what genes to focus on my discussion
19/04/2021 - 23/04/2021	<ul> <li>Project group meeting with Dr Schubert discussing and focusing on the preparation of the actual project write-up</li> </ul>	NEXT STEP: Reanalyse UMAP and volcano for GSE78949 NEXT STEP: To fine tune the methodology and results (and explain and summarise the meaning of the results)

02/05 2021	<ul> <li>Reanalyse UMAP and volcano for GSE78949</li> <li>Add details to Table 1</li> </ul>	NEXT STEP: Annotate my bibliography NEXT STEP: Rework my stem cell diagram and clarify the terminology NEXT STEP: Create a second table to explain the justification of the used database being selected
13/05/2021	<ul> <li>Project meeting with         Dr Schubert discussing         and finalising the         discussion and         amending the         diagrams and their         captions parts of the         results and general         queries</li> <li>Complete my         annotated         bibliography</li> </ul>	NEXT STEP: Finalise conclusion and abstract NEXT STEP: explain and summarise the meaning of the of the STRING analysis NEXT STEP: Perform a final count