

## Opis delovnega mesta mladega raziskovalca/ke (*Description of the Young Researcher's position*)

1. Članica UL (*UL member*):

Univerza v Ljubljani, Medicinska fakulteta

2. Ime, priimek in elektronski naslov mentorja/ice (*Mentor's name, surname and email*):

akad.prof.dr. Robert Zorec; robert.zorec@mf.uni-lj.si

3. Raziskovalno področje (*Research field*):

Nevrobiologija, program »Celična fiziologija« - razvoj novih zdravil

4. Opis delovnega mesta mladega raziskovalca/ke (*Description of the Young Researcher's position*):

Vključuje morebitne dodatne pogoje, ki jih mora izpolnjevati kandidat/ka za mladega raziskovalca/ko, ki niso navedeni v razpisu za mlade raziskovalce.

*slo:*

Raziskovalno delo mlade(ga) raziskovalke(ca) bo potekalo v Laboratoriju za nevroendokrinologijo-molekulska celična fiziologija, na Inštitutu za patološko fiziologijo Medicinske fakultete v Ljubljani. Začetek raziskovalnega dela je predviden v jeseni 2022. Kandidate(ke), ki bodo do predvidenega roka septembra 2022 zaključili magistrski študij na 2. stopnji naravoslovnih smeri, kot so biologija, biokemija, medicina, biotehnologija, mikrobiologija, farmacija, laboratorijska biomedicina, kemija in so motivirani za raziskovanje vabimo, da se prijavijo prek spletnne aplikacije ter pošljejo življenjepis in motivacijsko pismo na naslov: [robert.zorec@mf.uni-lj.si](mailto:robert.zorec@mf.uni-lj.si). Prednost pri izbiri bodo imeli kandidati(ke) z visoko povprečno oceno študija in izkušnjami z delom v celični biologiji/fiziologiji in biokemiji/molekularni biologiji/vedenskih poskusih /razvoju zdravil.

### **Vsebina raziskovalnega dela: *Uravnavanje presnove astroglijie***

Astrociti so številčne in heterogene celice glije v možganih z mnogimi homeostatskimi funkcijami, med drugim uravnavajo presnovo. Privzemajo glukozo iz krvi in jo shranjujejo v obliki glikogena kot rezervno možgansko gorivo. Glikogen se ob povečani aktivnosti nevronov razgradi do glukoze, ki se v procesu aerobne glikolize presnovi v laktat. Laktat se sprosti iz astrocitov in kot emergent prenese v nevrone. Presnova v astrocitih je uravnana, prek z G-proteini sklopljenih receptorjev (GPCR) na površini astrocitov. Napake v presnovni povezanosti med astrociti in nevroni lahko privedejo do bolezni, kot je kognitivni upad zaradi nevrodegeneracije. Najpogostejsa oblika demenc je povezana z Alzheimerjevo boleznijo (AB), ki pa je povezana s hipometaboličnim stanjem, kar se redno izmeri klinično Eden od razlogov za hipometabolično stanje je degeneracija jedra *Locus coeruleus* (LC) v možganskem deblu, kjer je sicer zelo malo nevronov, ki prek difuznega pleteža aksonov inervirajo praktično vse dele možganov in hrbtnače in na teh mestih izločajo noradrenalin. Ker nevroni, ki izločajo noradrenalin propadejo, je količina noradrenalina zmanjšana. S tem pa tudi upade stimulacija astrocitne produkcije laktata v astrocitih. Ena od strategij je zmanjšane količine noradrenalina nadomestiti s transplantacijo nevronov, kar je zelo zahteven in kompleksen postopek. Alternativa je stimulirati aerobno glikolizo z ligandi, ki delujejo stimulativno na aerobno glikolizo, podobno kot noradrenalin. Ker smo pred kratkim odkrili molekule, ki aktivirajo ta proces in tudi odkrili nov receptor (gprB Dolanc in sod. Cells; 2022), ki se aktivira s temi molekulami, bo predmet raziskave določiti sekundarne prenašalce ( $\text{Ca}^{2+}$ , cAMP), ki se aktivirajo prek novega receptorja gprB in tudi prek katerih G-proteinov je učinek povečane produkcije laktata povezan (Gs, Gi Gq). Cilj projekta je ugotoviti ali je uravnavanje prek novega receptorja modulirano z zunajceličnim laktatom, ki naj bi na eni strani stimuliral proizvodnjo in izločanje laktata iz

astrocitov

Rezultati projekta bodo zagotovili nov vpogled v celično uravnavanje produkcije laktata v astrocitih, kar je pomembno za razvoj novih zdravil.

### Metode dela

Mladi(a) raziskovalec(ka) bo izvajal(a) meritve znotrajceličnih sekundarnih prenašalcev in metabolitov v astrocitih po aktivaciji receptorjev GPCR. Pri tem bo uporabljal(a) fluorescenčne označevalce in pa genetsko kodirajoče nanosenzorje in visokoločljivo fluorescenčno mikroskopijo v realnem času. Meritve bo izvajal v posameznih astrocitih v i) kulturi (*in vitro*) in tudi na izoliranih celicah iz transgenih živali in celičnih linij z izbitimi geni. Delo bo potekalo tudi ob sodelovanju s partnerji pri vedenjskih vzorcih transgenih živali in s specifičnimi lezijami v možganih.

eng:

The research work of the selected young researcher will be carried out at the Laboratory of Neuroendocrinology-Mol. Cell Physiology, Institute of Pathophysiology of the Faculty of Medicine in Ljubljana. The beginning of the research work is planned for autumn 2022. Interested candidates, who hold a Master degree in Biology, Microbiology, Pharmacy, Medicine, Chemistry, Physics are invited to apply via the web and send a motivation letter to: [robert.zorec@mf.uni-lj.si](mailto:robert.zorec@mf.uni-lj.si). Priority will be given to the candidates with a high average grade of study, with motivation and working experience in the field of cell biology, biochemistry/molecular biology/animal behaviour/drug development..

### Content of the research work: *Regulation of astroglial metabolism*

Astrocytes are an abundant and heterogeneous subtype of glial cells in the brain with many homeostatic functions, including the regulation of brain metabolism. They store blood-derived glucose in the form of glycogen as the brain fuel reserve, which is used during intense neuronal activity. It is degraded to glucose and metabolized in aerobic glycolysis to lactate. The latter is released from astrocytes and distributed as an energy fuel to neurons. Metabolism in astrocytes is regulated, via G-protein coupled receptors (GPCRs) on the surface of astrocytes. Defects in the metabolic connection between astrocytes and neurons can lead to diseases such as cognitive decline due to neurodegeneration. The most common form of dementia is associated with Alzheimer's disease (AD), which is associated with a hypometabolic condition that is regularly measured clinically. One reason for the cognitive decline is the demise of the brainstem *Locus coeruleus* neurons, which through diffuse axon plexus, innervate virtually all parts of the brain and secrete norepinephrine at these sites. As norepinephrine-secreting neurons are degraded in neurodegenerative diseases, the amount of norepinephrine is reduced. This also reduces the stimulation of astrocyte lactate production in astrocytes. One of the strategies is to replace the reduced amounts of norepinephrine with neuronal transplantation, which is a very demanding and complex procedure. An alternative is to stimulate aerobic glycolysis with ligands that have a stimulating effect on aerobic glycolysis, similar to norepinephrine. As we have recently discovered molecules that activate this process and also discovered a new receptor (gprB; Dolance et al., Cells, 2022), activated by these molecules, the subject of the study will be to identify secondary messengers ( $\text{Ca}^{2+}$ , cAMP) that are activated through the new gprB receptor and also through which type G-protein is the increased lactate production is mediated associated (Gs, Gi Gq). Moreover, the project will address the question whether extracellular lactate modulates the production of lactate as well as its release through channels.

The results of this project will provide a new insight into the cellular regulation of lactate production in astrocytes.

### Methods

The young researcher will perform measurements of intracellular second messengers and metabolites in astrocytes upon activation of GPCR receptors, using fluorescence markers, genetically encoded nanosensors and real-time high-resolution fluorescence microscopy. Measurements will be performed in individual rodent and human astrocytes in culture and from transgenic animals. Moreover, animal behavior with transgenic and lesion animals will be performed in cooperation with a partner lab.