

Search for methylation sites in the genome of *Baeotendipes noctivagus* chironomids based on nanopore sequencing data

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DNA methylation is one of the most significant epigenetic mechanisms for the regulation of gene expression in eukaryotes, involved in many biological processes. This process transfers a methyl group to a cytosine, mostly in positions 5' of a guanine (CpG sites).

While the most popular methods for detection of modified cytosines still are based on bisulfite sequencing data [1], a recent alternative is nanopore sequencing [2,3]. Here, nanopore sequencing data is used to investigate the existence of DNA methylation in the genome of chironomid *Baeotendipes noctivagus*. It is a non-biting midge from the Diptera order whose larva can survive in hypersaline lakes with the salinity rate of 80-240 g/l. It is the most halotolerant chironomid known [4]. It adapts to extreme conditions (hypoxia and osmotic stress) on the genomic level by extensive duplication of hemoglobin genes and responds to high salinity by high expression of the *Egfp1* gene encoding. That makes this midge an interesting model to study regulation gene expression.

The obtained results show the lack of methylation in *Baeotendipes noctivagus* genome. It can be explained by the absence of DNA methyltransferases 3 and 1 in Diptera insects, which are responsible for creating *de novo* methylation and maintaining existing.

It confirms that methylation is not a leading epigenetic mechanism in dipterans, and they may use other ways to regulate genes expression.

This study is a part of a large project, the goal of which is to understand how *Baeotendipes noctivagus* has adapted to hypersaline lakes.

References

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