



CURRICULUM VITAE

Date personale

Nume, prenume: Csépp Katalin
Email instituțional: katalin.csep@umfst.ro
Titlu academic: Conferențiar universitar
Disciplină, departament: Genetică, ME1

Domeniul profesional de activitate

Specialitate/supraspecializări: Medicină internă, Genetică medicală

Activitate de cercetare

Teme și direcții de cercetare: boli genetice rare, sindromul metabolic și tulburări comportamentale, tulburări cognitive

Număr ORCID: 0000-0003-4802-096X

Lucrări publicate în extenso (5 lucrări reprezentative):

1. **Csep K**, Sebesi Sz, Gabos Grecu I, Torok A, Baki G, Sardi K. Influence of BDNF-Val66Met and PPARGC1A-Gly482Ser Polymorphisms on Sedentary Lifestyle and Stress Associated Risk of Metabolic Syndrome and Depression. Proceedings of 5th Medical Genetics Congress with International Participation, Filodiritto Editore-Proceedings. 2019. 2:115-120
2. Gábos G, Moldovan D, Dobru D, Mihály E, Bara N, Nădășan V, Hutanu A, **Csép K**. Mutational spectrum and genotype-phenotype relationships in a cohort of Romanian hereditary angioedema patients caused by C1 inhibitor deficiency. Revista Romana de Medicina de Laborator-Romanian Journal of Laboratory Medicine. 2019. 27:255-269
3. Szabo M, Mate B, **Csep K**, Benedek T. Epigenetic Modifications Linked to T2D, the Heritability Gap, and Potential Therapeutic Targets. Biochem Genet. 2018. 56(6):553–574
4. **Csep K**, Szigeti E, Szalman K. MTHFR-Ala222Val effects on metabolic syndrome progression Acta Medica Marisiensis. 2018. 64(2):62-69
5. **Csép K**, Szigeti E, Vitai M, Korányi L. The PPARGC1A - Gly482Ser Polymorphism (rs8192678) and the metabolic syndrome in a central Romanian population. Acta Endo (Buc) 2017. 13(2): 161-167

