

Glutathione deficiency and alterations in the sulfur amino acid homeostasis during early postnatal development as potential triggering factors for schizophrenia-like behavior in adult rats

Magdalena Górny¹, Agnieszka Wnuk², Adrianna Kamińska³, Kinga Kamińska², Grażyna Chwatko³, Anna Bilska-Wilkosz¹, Małgorzata Iciek¹, Małgorzata Kajta², Zofia Rogóż², Elżbieta Lorenc-Koci²

¹ The Chair of Medical Biochemistry, Jagiellonian University Medical College, 7 Kopernika Street, 31-034 Kraków, Poland

² Maj Institute of Pharmacology, Polish Academy of Sciences, 12 Smętna Street, 31-343 Kraków, Poland;

³ Department of Environmental Chemistry, University of Łódź, 163 Pomorska Street, 90-236 Łódź, Poland;

* Correspondence: lorenc@if-pan.krakow.pl

Supporting information

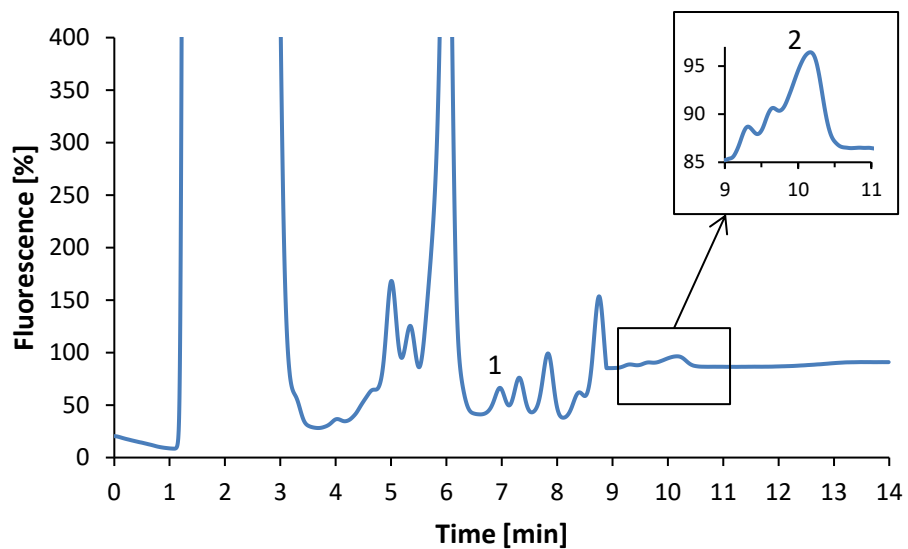


Figure S1. Representative chromatogram for methionine and homocysteine assay in the liver homogenate. Peaks: 1 - methionine, 2 - homocysteine.

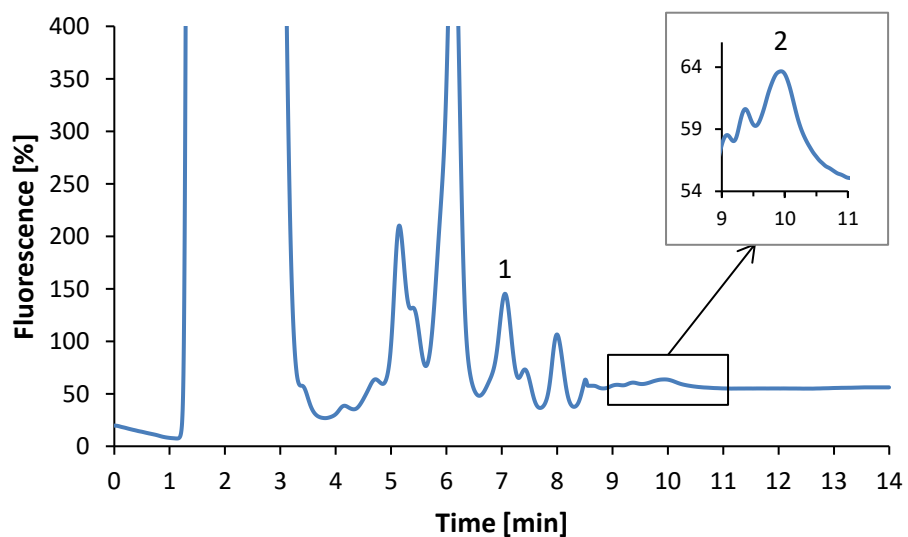


Figure S2. Representative chromatogram for methionine and homocysteine assay in the kidney homogenate. Peaks: 1 - methionine, 2 - homocysteine.

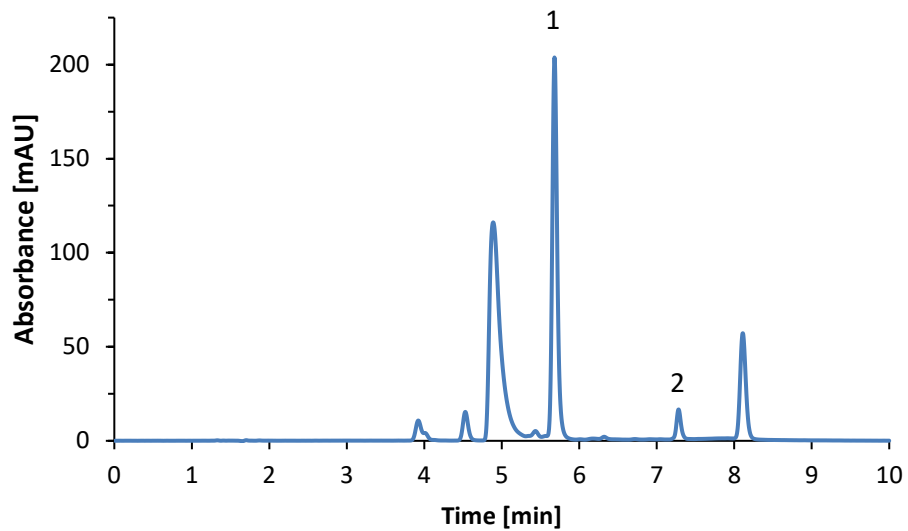


Figure S3. Representative chromatogram for glutathione and cysteine assay in the prefrontal cortex (PFC) homogenate. Peaks: 1 - glutathione, 2 - cysteine.

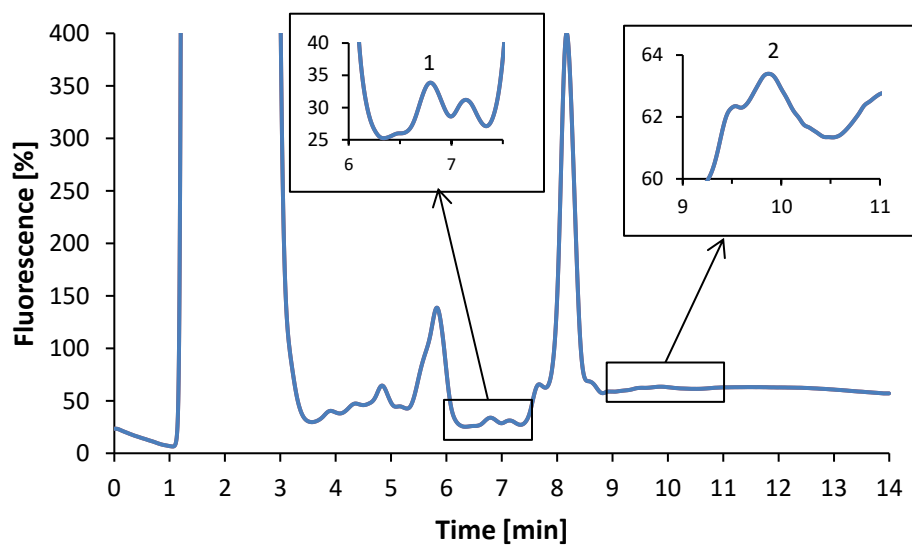


Figure S4. Representative chromatogram for methionine and homocysteine assay in the PFC homogenate. Peaks: 1 - methionine, 2 - homocysteine.

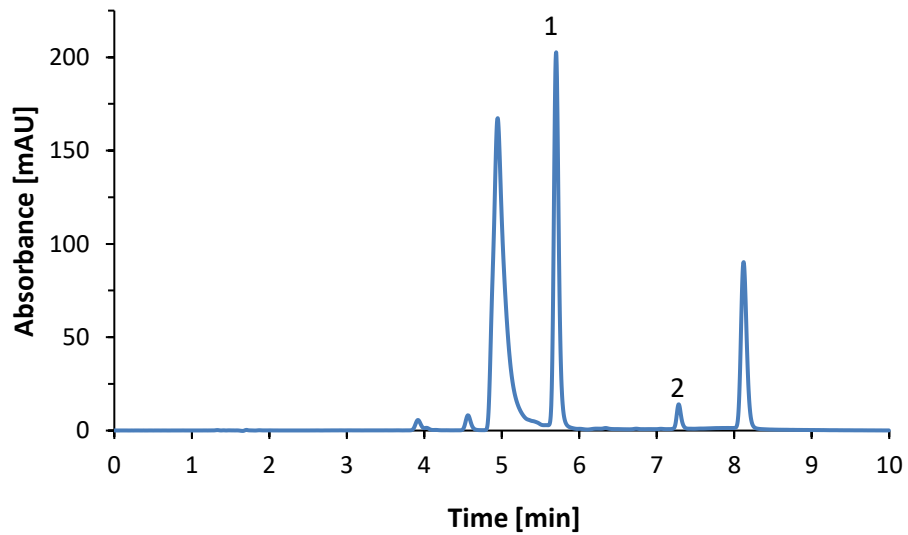


Figure S5. Representative chromatogram for glutathione and cysteine assay in the hippocampus (HIP) homogenate. Peaks: 1 - glutathione, 2 - cysteine.

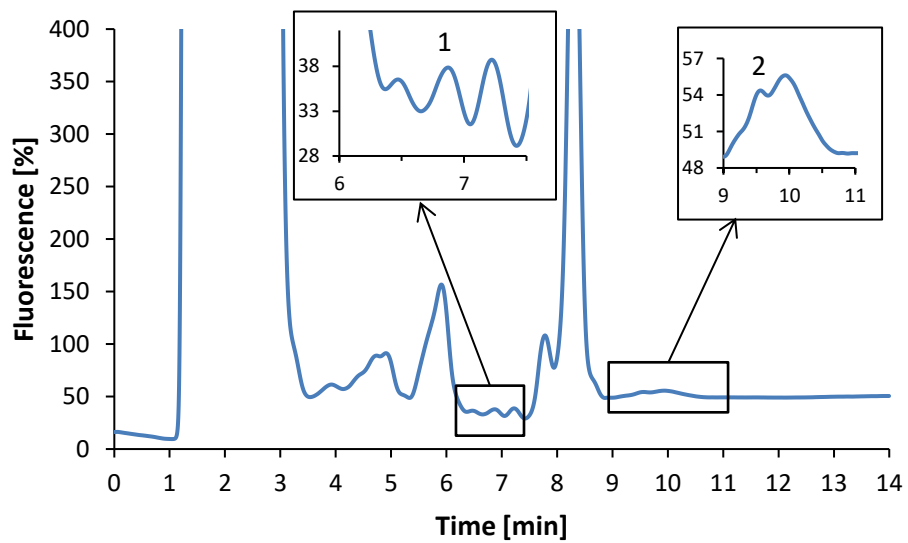


Figure S6. Representative chromatogram for methionine and homocysteine assay in the HIP homogenate. Peaks: 1 - methionine, 2 - homocysteine.