

Selective GSK3 β inhibition mediates an Nrf2-independent anti-inflammatory microglial response

Mohamed H. Yousef, Mohamed Salama, Hassan A. N. El-Fawal and Anwar Abdelnaser

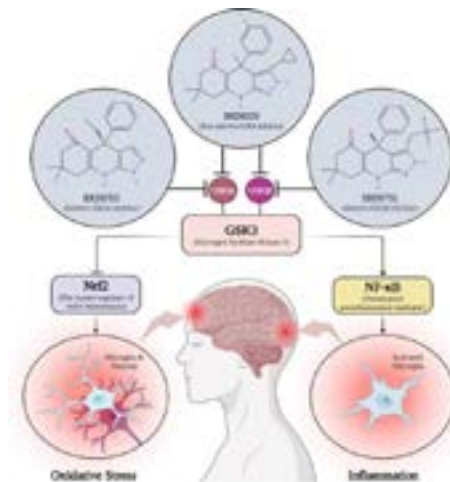
The American University in Cairo, Egypt

Glycogen Synthase Kinase 3 (GSK3) is associated with the proinflammatory phenotype of microglia and has been shown to act in concert with nuclear factor kappa B (NF- κ B). GSK3 is also a suppressor of nuclear factor erythroid 2-related factor 2 (Nrf2), the principal regulator of redox homeostasis. Agreeing with the oxidative paradigm of aging, Nrf2 is often deregulated in parainflammatory and neurodegenerative diseases. In this study, we aimed to explore a multimodal disease-modifying utility of GSK3 inhibition, beyond neuronal proteopathologies. Furthermore; we aimed to underscore the difference in therapeutic value between the two GSK3 paralogs by isoform-selective chemical inhibition.

The anti-inflammatory effects of paralog-selective GSK3 inhibitors were evaluated as a function of the reductive capacity of each to mitigate LPS-induced activation of SIM-A9 microglia. The Griess method was employed to detect the nitrate-lowering capacity of selective GSK3 inhibition. Real-time PCR was used to assess post-treatment expression levels of pro-inflammatory markers and antioxidant genes; pro-inflammatory cytokines were assayed by ELISA. Nuclear lysates of treated cells were examined for Nrf2 and NF- κ B accumulation by immunoblotting. Finally, to infer whether the counter-inflammatory activity of GSK3 inhibition was Nrf2-dependent, DsiRNA-mediated knockdown of Nrf2 was attempted.

Results from our experiments reveal a superior anti-inflammatory and anti-oxidative efficacy for GSK3 β -selective inhibition, compared to GSK3 α -selective and non-selective pan-inhibition; hence use of selective GSK3 β inhibitors is likely to be more propitious than non-selective dual inhibitors administered at comparable doses. Moreover, our results suggest that the anti-inflammatory effects of GSK3 inhibition is not Nrf2 dependent.

Keywords: GSK3; Paralog selectivity; Microglia; Neuroinflammation; Neurodegenerative diseases; Oxidative stress; Nrf2; NF- κ B.



References:

1. Nagamoto-Combs K, Kulas J, Combs CK. A novel cell line from spontaneously immortalized murine microglia. *J Neurosci Methods*. 2014 Aug; 233:187–98.
2. Beurel E, Jope RS, E B, RS J, Beurel E, Jope RS, et al. Lipopolysaccharide-induced interleukin-6 production is controlled by glycogen synthase kinase-3 and STAT3 in the brain. *J Neuroinflammation*. 2009 Mar 1; 6(1):1–11.
3. Yuskaitis CJ, Jope RS, CJ Y, RS J, Yuskaitis CJ, Jope RS. Glycogen synthase kinase-3 regulates microglial migration, inflammation, and inflammation-induced neurotoxicity. *Cell Signal*. 2009 Feb; 21(2).
4. Guzman-Martinez L, Maccioni RB, Andrade V, Navarrete LP, Pastor MG, Ramos-Escobar N. Neuroinflammation as a common feature of neurodegenerative disorders. *Front Pharmacol*. 2019; 10(SEP).

Biography

Mohamed H. Yousef currently working in the School of Sciences and Engineering, Biotechnology Graduate Program, The American University in Cairo, Cairo, Egypt.

Received: November 12, 2022; **Accepted:** November 15, 2022; **Published:** January 25, 2023