

Unifying the common concepts shared by neurodegenerative diseases

Editorial

Neurodegenerative diseases (NDs) such as Alzheimer's disease^{1,2} and Parkinson disease³ are among the big challenges facing the researches and the clinical professionals. NDs have complex mechanism including pathways that are yet to be elucidated. Each of these diseases is often studied separately and investigated towards explaining the mechanisms and finding potential therapies of one disease. Changing such approach might be a way to improve the data we obtain.

Indeed, many common features exist between the NDs. Therefore, focusing on such common features and mechanisms will allow us to extrapolate the obtained results and explain a pathway involved in more than one neurodegenerative disease and thus find out potential therapies for more than on neurodegenerative disease. Such methodologies still require further elucidation of the common features linking the divers' neurodegenerative disease and whether they are such links are the results of common underlying pathways or only similar symptoms or phenotypes. These needs collaborations between experts in different fields including the concerned NDs and always study each neurodegenerative disease within the context of the common features shared between more than a neurodegenerative disease and complete the data by cell culture studies results,⁴ pharmacology⁵⁻¹⁰ and toxicology^{11,12} to reach the final goal which is identify efficient therapies.

Such approaches will optimize the efforts aiming to understand the NDs and treat them by reducing the research cost, efforts and time. For building an animal model of a neurodegenerative disease such as Alzheimer's disease¹³ expressing a specific feature will contribute to study all the NDs that include that specific feature within its pathogenesis. Importantly, this concept is further strengthened by several facts such as the physiology of the brain that constitutes of a network within which the neurotransmitters are in continuous interactions¹⁴ and the common molecular basics¹⁵ related to the G protein coupled receptors¹⁶ that are of a great importance in both neurophysiology¹⁷ and pharmacotherapy.¹⁸⁻²⁰ In addition, other non-degenerative diseases share also similar mechanisms or pathways with some NDs which means that the range of extrapolation of the common features shared by some neurodegenerative disease might also include some non-NDs.

Acknowledgements

Abdelaziz Ghanemi is a recipient of a 2013 CAS-TWAS President's Postgraduate Fellowship.

Conflicts of interest

None.

References

1. Vangavaragu JR, Vissavajhala P, Reddy CD. Alzheimer's disease: *Current perspectives – animal models, drugs under development, and*

Volume 2 Issue 5 - 2015

Abdelaziz Ghanemi,^{1,2,3}

¹Key Laboratory of Animal Models and Human Disease Mechanisms of the Chinese Academy of Sciences & Yunnan Province, Kunming Institute of Zoology, Chinese Academy of Sciences, Kunming, Yunnan 0, PR China

²Kunming College of Life Science, University of Chinese Academy of Sciences, Kunming 00, China

³University of Chinese Academy of Sciences, Beijing, China

Correspondence: Abdelaziz Ghanemi, Key Laboratory of Animal Models and Human Disease Mechanisms, Kunming Institute of Zoology Chinese Academy of Sciences, No. 32 Jiao chang dong lu, Kunming 650223, China, Email ghanemiabdelaziz@hotmail.com

Received: August 03, 2015 | **Published:** August 05, 2015

potential nutritional intervention. In: Watson RR (Ed.), *Foods and dietary supplements in the prevention and treatment of disease in older adults*. Academic Press, San Diego, California, USA. 2015;pp.13–28.

- Ghanemi A. Alzheimer's disease therapies: Selected advances and future perspectives. *Alexandria Journal of Medicine*. 2015;51(1):1–3.
- Ghanemi A. Schizophrenia and Parkinson's disease: Selected therapeutic advances beyond the dopaminergic etiologies. *Alexandria Journal of Medicine*. 2013;49(4):287–291.
- Ghanemi A. Cell cultures in drug development: Applications, challenges and limitations. *Saudi Pharm J*. 2014;doi:10.1016/j.jsps.2014.04.002.
- Ghanemi A. How to map the bridges between zoology and pharmacology? *The Journal of Basic & Applied Zoology*. 2015;doi:10.1016/j.jobaz.2014.12.003.
- Ghanemi A. For dentists and doctors: The neglected concepts about the factors influencing the effects of drugs. *The Saudi Dental Journal*. 2015;doi:10.1016/j.sdentj.2015.05.001.
- Ghanemi A, Boubertakh B. Shorter and sturdier bridges between traditional Chinese medicines and modern pharmacology. *Saudi Pharm J*. 2015;23(3):330–332.
- Ghanemi A. How important is pharmacognosy for doctors and dentists? *Saudi Dent J*. 2015;27(1):1–2.
- Boubertakh B, Liu XG, Cheng XL, et al. A Spotlight on Chemical Constituents and Pharmacological Activities of *Nigella glandulifera* Freyn et Sint Seeds. *Journal of Chemistry*. 2013;Article ID.820183.
- Lang P, Irnich D. *Systemic pharmacotherapy*. In: Irnich D (Ed.), *Myofascial Trigger Points*. Churchill Livingstone, Oxford. 2013;pp.253–259.
- Ghanemi A. How to define a pharmacological or a toxic food? *Alexandria Journal of Medicine*. 2014;doi:10.1016/j.ajme.2014.06.004.

12. Ghanemi A. Is mapping borders between pharmacology and toxicology a necessity? *Saudi Pharm J.* 2014;22(6):489–490.
13. Ghanemi A. *Animal models of Alzheimer's disease: Limits and challenges.* NPG Neurologie – Psychiatrie – Gériatrie. 2014;14(84):303–305.
14. Ghanemi A. Psychiatric neural networks and neuropharmacology: Selected advances and novel implications. *Saudi Pharm J.* 2014;22(2):95–100.
15. Ghanemi A. Biological properties and perspective applications of “Bio–neuter” chemicals? *Saudi Pharm J.* 2014;22(1):1–2.
16. Ghanemi A, He L, Yan M. New factors influencing G protein coupled receptors' system functions. *Alexandria Journal of Medicine.* 2013;49(1):1–5.
17. Ghanemi A, Hu X. Elements toward novel therapeutic targeting of the adrenergic system. *Neuropeptides.* 2015;49:25–35.
18. Ghanemi A. Targeting G protein coupled receptor–related pathways as emerging molecular therapies. *Saudi Pharm J.* 2015;23(2):115–129.
19. Ghanemi A, Hu X. Targeting the orexinergic system: Mainly but not only for sleep–wakefulness therapies. *Alexandria Journal of Medicine.* 2014;doi:10.1016/j.ajme.2014.07.002.
20. Ferre S. The GPCR heterotetramer: challenging classical pharmacology. *Trends Pharmacol Sci.* 2015;36(3):145–152.