Neuroprotective Potential of Seaweed Extracts and Naringenin
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Outline
Introduction
Seaweeds' neuroprotective potential set in vitro on a human cellular stress model
Naringenin alleviates 6-hydroxydopamine (6-OHDA) induced Parkinsonism in SH- SY5Y cells and zebrafish model
Conclusion
Abstract
Parkinson's disease (PD) is a progressive neurodegenerative disorder predominantly acting the motor system. It is characterized by the loss of dopamine-producing neurons the substantia nigra region of the midbrain. Seaweeds are known to contain bioactive lecules with high antioxidant capacity. Naringenin possesses anti-inflammatory perties, high antioxidant capacity, and free radical scavenging abilities, and it can etrate the blood-brain barrier. Some studies suggest that antioxidants may have roprotective activities. Therefore, this report investigates the potential roprotective effects of seaweed extracts and naringenin on SH-SY5Y cells. The ills indicate that seaweed extracts can enhance the viability of SH-SY5Y cells and ibit dopamine-induced neurotoxicity. Under the influence of dopamine, cell viability apartially restored by 20-30% when treated with certain seaweed extracts. Moreover, pase-3 activity was effectively inhibited, returning to levels similar to those before amine induction. In terms of mitochondrial membrane potential, it could be restored up to 60% compared to the depolarization induced by dopamine. Principal Component alysis (PCA) suggested that caspase-3 activity was related to neuroprotective activity not directly associated with mitochondrial membrane potential, indicating that the tective mechanism of seaweed extracts may not be directly linked to mitochondrial mbrane potential. Naringenin effectively reduced the increased levels of ROS induced 6-OHDA, restored antioxidant enzyme levels (CAT, GSH, SOD), and mitochondrial mbrane potential in SH-SY5Y cells. It also altered the mRNA expressions of $LRRK2$ , $IK1$ , $CASP9$ , $POLG$ which are related to Parkinson's disease, and improved behavioral