Resume of Yao CHEN

Basic Information



School : Gender: Date of Birth: Title: Education:

Tutor: Interest of research: Life Sciences and Health Engineering Female 198202 Lecturer Ph.D of Biochemistry and Molecular Biology Master degree Ptotein engineering , Virus molecular biology

Academic Background

From September 1998 to July 2002, Huazhong University of Science and Technology, Bachelor's degree in Biological technology;

From September 2004 to July 2012, Wuhan Institute of Virology, Chinese Academy of Sciences, Ph.D of Biochemistry and Molecular Biology.

Enrollment Information

- 1. Enrollment Discipline: Master of Bioengineering
- 2. Research direction: ptotein engineering, virus molecular biology
- 3. Enrollment Year: 2024-2025

Representative Projects

1.National Natural Science Foundation of China "Study on the function and mechanism of lef-9, a subunit of baculovirus polymerase, involved in the selection of transcription start sites and catalytic RNA synthesis of late viral genes", China, Project leader.

2. Hubei University of Technology Doctoral startup Fund Project "Structure and function of oncogene mdmx",Hubei Province, Project leader.

3.National Natural Science Foundation of China "Study on genes affecting hearNPV ODV embedding and their functions", China.

4.Major project of Hubei Province's special technology "innovation research and development of key technologies for the synthesis of anticancer drug carbinolone acetate based on phytosterol semibiological method", Hubei Province.

5.Wuhan Science and Technology Bureau, Wuhan Natural Science Foundation Key Project "Early diagnosis of liver cancer and targeted anticancer drug design", Wuhan City.

Representative Articles

1.Cysteines 128 and 250 are essential for the functions of the baculovirus core gene ac109,Virology,2023,587,109857.

2.Helicoverpa armaigera nucleopolyhedrovirus ORF50 is an early gene not essential for virus propagation in vitro and in vivo., Virus Genes, 2012, 45(1): 149-160.

3.H5N1 influenza virus-like particles produced by transient expression in mammalian cells induce humoral and cellular immune responses in mice., Canadian Journal of Microbiology, 2012, 58(4): 391-401.

4.Efficient reactivation of p53 in cancer cells by a dual MdmX/Mdm2 inhibitor., J Am Chem Soc, 2014, 136(52): 18023-18033.

5.Recombinant butelase-mediated cyclization of the p53-binding domain of the oncoprotein MdmX stabilized protein conformation as a promising

model for structural investigation, Biochemistry, 2019, 58(27): 3015.

6.A Fusion Protein of the p53 Transaction Domain and the p53-Binding Domain of the Oncoprotein MdmX as an Efficient System for High-Throughput Screening of MdmX Inhibitors, Biochemistry, 2017, 56(25): 3273-3282.