



Nia Tatsis, Ph.D.

Executive Vice President and Chief Regulatory and Quality Officer

Nia Tatsis, Ph.D., joined Vertex Pharmaceuticals in 2017 and is the company's Executive Vice President, Chief Regulatory and Quality Officer. In this role, she is responsible for managing all aspects of global regulatory affairs and quality assurance involving the research and development, manufacturing, and commercialization of Vertex's transformative medicines.

Prior to joining Vertex, Dr. Tatsis held positions of increasing responsibility at pharmaceutical companies including Sanofi, Pfizer and Wyeth. Most recently, she served as Vice President, Head of Global Regulatory Affairs, of the Sanofi Genzyme Business Unit.

Dr. Tatsis currently serves as a member of the board of directors for Verve Therapeutics. In 2024, she was named to the *Endpoints News* Women Leading Biopharma R&D list and the *PharmaVoice* 100. She was previously named to *Fierce Pharma's* Fiercest Women in Life Sciences list.

Dr. Tatsis received her Ph.D. in cell and molecular biology from the University of Vermont and also holds a B.S. in biology from Temple University. She worked as a staff scientist and research fellow in immunology and vaccine development at the Wistar Institute and completed a postdoctoral research fellowship in immunology at Thomas Jefferson University.

Contact:

Nia Tatsis, Ph.D.

Executive Vice President and Chief
Regulatory and Quality Officer
she/her/hers

Vertex Pharmaceuticals Incorporated
50 Northern Avenue
Boston, MA 02210
+1 (617) 341-6100
vrtx.com

Joined Vertex: 2017

Education:

Temple University
University of Vermont

Notable Recognitions:

PharmaVoice 100
Endpoints News Women Leading
Biopharma R&D
Fierce Pharma Fiercest Women in Life Sciences