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## Areas of expertise

Biodiversity | Sustainability pathways | Characterization of nanostructure | Analytical chemistry | Semi-solid formulation | Supply chain | Molecular modeling | Pharmaceutical technology | Regulatory Affairs and GMP Compliance

## **Selected Achievement**

**7 Patents**: 3 Granted USPTO/WIPO + 4 Applications PCT **40 Scientific communications** in conference outside Canada in the last 10 years. **500 products** (NHP OTC, Rx, C/T, N) launched to the market worldwide since 2006

2025 - Lead Auditor ISO 19011:2018 EU GMP / US FDA CFR 211

2024 - Executive Vice President, Biovantek - Labomar Canada

2023 - Senior Regulatory Affairs Officer, Medelys International Labs

2022 - Health Canada – Narcotic Exemption for Psilocybin – Award

2021 - SVP Innovation and Compliance at CYBIN<sup>™</sup>, Public company

2020 - Co-Chair ISCRE - Chemical Reaction Engineering - Award

2019 - CQIB Grant - Biotech Start-Up of the Year - Award

2018 - Chief Innovation Officer at WeedMD™, Public company

2017 - Head of Science − Earth Technology and Science Trust<sup>TM</sup> - Award

2015 - Henry Shibata Fellowship, Cedar Cancer Foundation - Award

2010 - International Year of Biodiversity™ Award, Colombia – Award

Dr. Domenico Fuoco is a certified lead auditor for ISO 19011:2018, with over 100 audits conducted in Pharmaceutical Quality Management Systems across the European Union, Turkey, and North America. Since the early 2000s, he has been directly involved in product and process development finalized to the licensing and market approval of over than 500 pharmaceutical products worldwide.

In late 2022, Dr. Fuoco returned to academia as an Associate Professor, bringing extensive industry expertise to his research and teaching. Unlike traditional academics, his authority in the field is not solely based on peer-reviewed publications, but on his 24 years of experience in the pharmaceutical industry, where he has contributed to hundreds of medical claims and patents.

Dr. Fuoco unique background bridges the gap between scientific innovation and regulatory application, making him a key figure in the translational development of pharmaceutical technologies.