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Law360's 2023 Life Sciences Editorial Advisory Board

Law360 (June 16, 2023, 3:00 PM EDT) -- Law360 is pleased to announce the formation of its 2023 Life Sciences Editorial Advisory Board.

The editorial advisory board provides feedback on Law360's coverage and expert insight on how best to shape future coverage.

The members of Law360's 2023 Life Sciences Editorial Advisory Board are:

Stacy Cline Amin - Morrison Foerster LLP
Stacy Cline Amin leads the FDA and health care regulatory and compliance practice in MoFo's global life sciences and health care group, and provides strategic regulatory and business advice to companies in the life sciences, health care and technology industries. Stacy previously served as chief counsel of the U.S. Food and Drug Administration.

Shireen Barday - Pallas Partners (US) LLP
Shireen Barday is a partner at New York litigation boutique Pallas Partners (US) LLP. She has been litigating complex commercial matters for the last 15 years. As a BigLaw alumna, Shireen offers advice that is practical, reasonable and commercially astute.

D'Lesli Davis - Norton Rose Fulbright
D'Lesli Davis is Norton Rose Fulbright's U.S. head of life sciences and health care. She represents pharmaceutical and medical device clients in high-stakes litigation in state and federal courts across the country, in MDLs and consolidated proceedings, governmental actions, commercial, economic loss and product liability matters.

Gary F. Giampetruzzi - Paul Hastings LLP
Gary Giampetruzzi is the global chair of Paul Hastings' life sciences department and one of the vice-chairs of the investigations and white collar department. He advises clients on day-to-day compliance matters and represents life sciences companies in high-profile investigations, including FCPA investigations, and other complex litigation.

Abeba Habtemariam - Arnold & Porter
Abeba Habtemariam advises life sciences companies on regulatory, compliance and legislative matters, particularly on compliance with the FD&C Act. She also routinely advises clients on the regulation of medical device software and health care IT, premarket approval and clearance strategies, promotional review matters, and cGMP compliance.

April Isaacson - Kilpatrick Townsend & Stockton LLP
April Isaacson is the managing partner for the firm's San Francisco office. She has over 25 years of experience as a trial lawyer and is a registered U.S. patent attorney. Her practice focuses on complex technical intellectual property litigation as well as Hatch-Waxman cases on behalf of drug innovators.

Marian Lee - Gibson Dunn & Crutcher LLP
Marian J. Lee is co-chair of the FDA and health care practice at Gibson Dunn. She brings over 18 years of experience advising on FDA regulatory, compliance and enforcement matters. Marian is listed in The Best Lawyers in America and Who's Who Legal's Thought Leaders. She graduated from Harvard Law.

Arman Oruc - Goodwin Procter LLP
Arman Oruc is co-chair of Goodwin's antitrust practice. Arman was most recently part of the founding team of a life sciences company helping grow the startup into a late-stage clinical company. Arman's experience combining private practice and in-house work gives him a unique perspective on the life sciences industry.

T. Reed Stephens - Winston & Strawn LLP
T. Reed Stephens, the co-chair of our health care and life sciences industry group, is a nationally recognized legal authority in the life sciences industry who focuses his practice on advising global life sciences companies on sensitive, high-risk corporate compliance matters often involving government investigations or congressional investigations.

Xin Tao - Baker McKenzie
Xin Tao is an FDA regulatory partner in Baker McKenzie's litigation and government enforcement practice in Washington, D.C. A former research biochemist, Xin understands the complex scientific issues related to the FDA's legal and regulatory requirements, enabling him to help clients with all phases of product development, manufacturing and marketing.

Eva Temkin - King & Spalding LLP
A partner at King & Spalding and former FDA policy director, Eva Temkin counsels clients on a variety of FDA-regulated products, with a particular focus on drugs, biologics and complex combination products. Eva advises companies across product development and lifecycle, from data generation and application submissions to exclusivity issues and dispute resolution.

Adam Yoffie - Bristol Myers Squibb
Adam G. Yoffie is senior corporate counsel, litigation and government investigations, at Bristol Myers Squibb. Prior to BMS, Adam served as a DOJ trial attorney on the health care fraud strike forces in Miami and Philadelphia. He previously worked as an associate at Williams & Connolly focusing on the life sciences industry.
