



2025

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Dear MPO Summit Attendee,

We are pleased to welcome you to the MPO Summit in Silicon Valley, home to fantastic innovation and healthcare technology development. The region serves as a reflection of the entire state regarding that continued effort, as the California Life Sciences Association (CLSA) asserts investment in biotech and medtech was \$4.5 billion in 2023. The region's medical device segment includes companies such as Abbott, Medtronic, J&J, Intuitive, Stryker, and BD, which all maintain facilities in the area. We were pleased with the event that took place here in 2019 and are happy to have the opportunity to return in 2025 to continue to learn more about the technologies being supported and the companies involved.

According to CLSA, life science companies in the Bay Area employ more than 143,682 people—the most of all California's major life sciences hubs. Of that figure, 17% is directly attributed to organizations in the medical devices and equipment sector. The average annual life sciences industry wage here is \$191,213. Overall, the life sciences industry supports more than 254,000 jobs in the region, according to BIOCOM (per its Economic Impact Report). In addition, there are more than 3,800 life science establishments in the Bay Area.

Innovation is alive and well in the Bay area. In addition to the aforementioned \$4.5 billion in VC investment in medtech (and biotech), there were 50 medical device venture raises in 2023, resulting in a total of \$1.4 billion. In addition, real estate firm JLL shared midsize and larger requirements of space of more than 30,000 square feet are creating demand for lab space in the region. These mid- to large-sized labs accounted for about 45% of demand in the fourth quarter (2023).

Finally, the sector enjoys support from the dedicated organizations that aid life sciences companies in California, including the aforementioned CLSA as well as Biocom California and OCTANe. Additionally, talent for the sector originates in part from a wide variety of educational institutions in the area.

With every MPO Summit, the primary focus is on facilitating education and networking for all attendees; we kept this mission at the top of our minds for this year's event. Maintaining a focus on medical device outsourcing alongside related critical concerns, the MPO Summit draws from having a close eye on significant issues relevant to members of the industry. We've assembled speakers to address these vital concerns with an educational agenda we feel addresses your needs perfectly.

Take advantage of the networking breaks between sessions to visit with sponsor companies, supporting organizations, speakers, and, of course, the staff of MPO. All are excited to have an opportunity to hear your specific challenges and concerns.

We hope you enjoy your time in California and find the 2025 MPO Summit to be a valuable experience that caters to your professional needs. If there's anything we can do to make your time here more beneficial, please let us know.

Sincerely,



Howard A. Revitch
Group Publisher
Medical Product Outsourcing
Orthopedic Design & Technology



Sean Fenske
Editor-in-Chief
Medical Product Outsourcing
Orthopedic Design & Technology



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THURSDAY, SEPTEMBER 18

7:00-8:55 a.m.

Registration & Continental Breakfast

Sponsored by Evolution Free Zone

8:55-9:00 a.m.

Welcome Address

Speakers:

- Howard Revitch, MPO/ODT Group Publisher
- Sean Fenske, MPO/ODT Editor-in-Chief

9:00-9:45 a.m.

State of the Industry

Speaker: Bryan Hughes, P&M Corporate Finance (PMCF)

9:45-10:30 a.m.

Beyond Reshoring: What Companies Are Doing Now

Speaker: Rosemary Coates, Reshoring Institute

10:30-11:15 a.m.

Networking Break

Sponsored by BMP Medical

11:15 a.m.-12:15 p.m.

Make vs. Buy: An OEM/CM Supply Chain Discussion

Moderator: Barry Schnur, David Schnur Associates

Panelists:

- Jim Fitzgerald, ATL Technology
- Brent Hahn, Miniaturization and Precision Manufacturing SME
- Ryan Hazelton, Dexcom
- John Schneider, Edwards Lifesciences
- Eric Steuben, Calyxo

12:15-1:45 p.m.

Networking Luncheon

Sponsored by Vance Street Capital

1:45-2:30 p.m.

Washington Update

Speaker: Brendan Benner, MDMA

2:30-3:15 p.m.

Lean Isn't Dead—It's Digital: Rethinking Inventory in the Age of Smart Manufacturing

Speaker: Matt Stekier, Plante Moran

3:15-4:00 p.m.

Networking Break

Sponsored by MTD Micro Molding

4:00-5:00 p.m.

Face to Face: Industry Execs Address

Medtech's Most Pressing Issues

Moderator: Julie Schulte, MBA, Chamfr

Panelists:

- Nora Crivello, WESTPAK Inc.
- James W. LaVersa, Jr., Harmac Medical Products Inc.
- John Nino, Life Science Outsourcing
- Sarah Ptach, Canyon Labs
- David Robson, Iantrek

5:15-7:00 p.m.

Networking Reception

Sponsored by Life Science Outsourcing

FRIDAY, SEPTEMBER 19

7:30-9:00 a.m.

Registration & Continental Breakfast

Sponsored by Evolution Free Zone

9:00-9:45 a.m.

FDA's New QMSR Is on the Horizon—

Will Your Company Be Ready?

Speaker: Christopher Joseph Devine,
Devine Guidance International Inc.

9:45-10:30 a.m.

Scientific AI for the Medical Device Industry

An Open Conversation Featuring:

- Ethan Mirsky, Ph.D., NobleAI
- Paul Orlando, Schivo Medical
- Matt Rudow, NobleAI
- Jeff Wheeler, NobleAI

10:30-11:15 a.m.

Networking Break

Sponsored by MPO

11:15 a.m.-12:15 p.m.

The Sterilization Road Less Traveled: A Look at Alternatives

Moderator: Darci Diage, TPL Consulting

Panelists:

- Megan C. Frost, Ph.D., Sterile State
- Derek Prince, Ph.D., Prince Sterilization Services
- Michelle Kennedy Scott, Gulf Sterilization

12:15 p.m.

End of Conference

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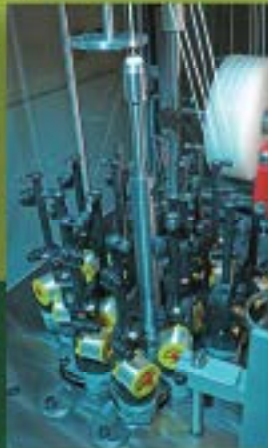


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Brendan Benner

Executive Vice President of Public Affairs for MDMA

Brendan Benner is the executive vice president of public affairs for MDMA, where he is in charge of communications, strategic outreach, and grassroots development. Benner has extensive experience in the private and public sectors, as well as working with nonprofits. He spent several years working in Congress, serving both as a Communications Director and District Director for former Congresswoman Melissa Hart (PA) and Congressman Jim Gerlach (PA), respectively. He oversaw and implemented aggressive media and outreach plans in these highly competitive congressional districts. Benner is also a veteran of numerous Congressional and Presidential campaigns. Benner is a member of the Pennsylvania Bar and a graduate of Georgetown University and Temple University's Beasley School of Law.



Rosemary Coates

Founder and Executive Director of the Reshoring Institute

Rosemary Coates is the founder and executive director of the Reshoring Institute, a 501c3 nonprofit and non-partisan organization focused on expanding U.S. manufacturing. She is also the president of Blue Silk Consulting, a global supply-chain management consulting firm. Coates also works as an expert witness on legal cases involving global supply chain disputes and has worked on over 60 legal cases. As a management consultant for 35+ years, She has helped over 80 global supply chain clients worldwide. She is an Amazon.com best-selling author with five supply chain management books, including *The Reshoring Guidebook*, *42 Rules for Sourcing and Manufacturing in China*, and *Legal Blacksmith: How to Avoid and Defend Supply Chain Disputes*.



Nora Crivello

President & CEO of WESTPAK Inc.

Nora Crivello is the president and CEO of WESTPAK Inc., an independent testing laboratory network in California. Through her 20+ years at WESTPAK, she has been "boots on the ground" as a Test engineer, quality manager, and vice president, making her uniquely qualified to now serve as president and CEO. Crivello is also the Chair of the ISTA Global Board of Directors and sits on numerous industry committees including Kilmer Innovations, Cal Poly Consortium, and ISTA Technical Committees. In her free time, she serves as Chair of the Board for Navigator Schools and raises guide dog puppies for Guide Dogs for the Blind with her family. Crivello holds a BS from California Lutheran University and an MBA from Pepperdine University.



Christopher Joseph Devine

Founder and President of Devine Guidance International Inc.

Dr. Christopher J. Devine is the founder and president of Devine Guidance International Inc. (DGII), a medtech establishment that provides consulting services to the medical device and pharma industries. Dr. Devine has over 45 years of combined experience in the fields of quality assurance, regulatory affairs, and design and development of medical devices. He is a senior member of the American Society of Quality (ASQ), a member of the Regulatory Affairs Professionals Society (RAPS), a member of the Society of Manufacturing Engineers (SME), and a member of the Project Management Institute (PMI). Dr. Devine received his doctorate from Northcentral University, with his doctoral dissertation entitled, "Exploring the Effectiveness of Defensive-Receiving Inspection for Medical Device Manufacturers: A Mixed-Method Study." Prior to launching his commercial career, Dr. Devine served honorably in the United States Marine Corps. During his downtime, Dr. Devine is active in his community, volunteering as a deputy sheriff supporting the Search and Rescue Division of the Lyon County Sheriff's Office.



Darci Diage

Owner, Founder, and CEO of TPL Consulting

Darci Diage, founder and CEO of TPL Consulting, provides consulting services for all facets of medical device development, manufacturing, and market approval, from concept to commercial. After more than 20 years of running complex laboratories and quality departments for medical device manufacturers, Diage decided to apply her unique contributions to the "total product lifecycle" to accelerate development timelines and launch products. Her BS in Molecular Biology from Sonoma State and MS in Biomedical Laboratory Science from San Francisco State springboarded her career in combination products, cardiovascular, peripheral vasculature, spinal, and software devices. Diage's knowledge and expertise in feasibility testing, design control, risk management, and quality system implementation has further expanded her regulatory role, resulting in the successful submission of numerous device approvals in the U.S., EU, Canada, and Australia. With a portfolio of diverse device developments, Diage has firsthand experience in overcoming challenges within the design control process, given the continually evolving regulations and guidance documents. Having owned and operated TPL Consulting for seven successful years, Diage comprehends the intricate relationship between suppliers, service providers, and contract manufacturers to get devices to the market and in the hands of users to improve patient lives.

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Jim Fitzgerald
CEO at ATL Technology

Jim Fitzgerald has joined ATL Technology as its new chief executive officer. He brings an extensive track record of executive leadership in the medical device contract manufacturing industry, with a career spanning multiple globally recognized organizations. Fitzgerald brings a wealth of experience with him, having held CEO and senior executive leadership positions at Flexan, MRP Solutions, ITxM, Vesta, and Cardinal Health. His professional journey in the life sciences sector began at Baxter Healthcare, where he developed a deep understanding of the industry's evolving landscape. Throughout his career, he has demonstrated a steadfast commitment to operational excellence, customer-centric innovation, and strategic growth, making him exceptionally well-suited to lead ATL into its next phase of advancement.



Megan C. Frost, Ph.D.
Founder and Chief Technology Officer for Sterile State

Dr. Megan C. Frost is the founder and chief technology officer for Sterile State. She has worked on the development of medical polymers and their applications to devices for over 25 years. She earned a BS in Biological Sciences from the University of Notre Dame, a BS in Chemistry and an MS in Analytical Chemistry from Purdue University—Indianapolis, and a Ph.D. in Chemistry from the University of Michigan - Ann Arbor. Dr. Frost was a post-doctoral researcher in the Department of Surgery at the University of Michigan Medical School.



Brent Hahn
Miniaturization and Precision Manufacturing SME

For over 25 years, Brent Hahn has been recognized as a leading authority in the micro molding and materials industry, specializing in precision medical devices and innovative drug delivery systems. Throughout his career, he has led miniaturization strategies for groundbreaking micro advancements. His industry achievements include developing solutions for the thinnest micro catheter, leadless pacing, minimally invasive eye surgery, and micro drug-eluting devices. As a seasoned speaker and recognized subject matter expert, Hahn has educated audiences worldwide with his expertise in micro molding and miniaturization. He has been regularly featured in industry articles, conference expert panels, and as a keynote speaker. He also dedicates his time to mentoring peers, making a lasting impact on the plastics community.



Ryan Hazelton
Sr. Manager of Direct Procurement for Dexcom

Ryan Hazelton is a Sr. Manager of Direct Procurement at Dexcom. He leads global procurement categories of Capital Equipment, Sterilization, and Chemicals globally. Ryan has over 20 years of experience in supply chain, procurement, supplier management, and new product development. Ryan's team is responsible for delivering exceptional value by sourcing quality products to enhance patient outcomes while driving innovation and sustainability within Dexcom's supply chain. He has driven growth and global expansion within medtech, defense, and retail industries throughout his career. Ryan holds a B.S. in Economics from the United States Military Academy at West Point and a M.B.A. from Price College of Business at the University of Oklahoma.



Bryan Hughes
Managing Director at P&M Corporate Finance (PMCF)

Bryan Hughes is a director of PMCF and leads the firm's medical technology team. His practice focuses on assisting clients with mergers and acquisitions, leveraged buyouts, private placements, financings, valuation, and strategic consulting. His clients have ranged from global medical technology companies to small, privately held businesses.



James W. LaVersa, Jr.
Vice President, Business Development, for Harmac Medical Products Inc.

James W. LaVersa, Jr., serves as vice president of business development at Harmac Medical Products Inc., a global contract design and manufacturing organization specializing in single-use medical devices. Based at the company's headquarters in Buffalo, NY, he oversees global business development efforts and contributes to corporate strategy across engineering, operations, quality, supply chain, and finance. With more than 35 years of experience in sales leadership and strategic growth, LaVersa has helped guide Harmac's expansion across four manufacturing plants located in the United States, Ireland, and Mexico. He has worked extensively across five continents, forming long-term partnerships with some of the world's leading medical device OEMs as well as well-funded startups seeking scalable, high-quality manufacturing solutions. LaVersa is recognized for his cross-functional leadership, international business expertise, and commitment to delivering customer-focused solutions in highly regulated markets.

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**Ethan Mirsky, Ph.D.****Chief Product and Strategy Officer at NobleAI**

Ethan Mirsky currently serves as the chief product and strategy officer at NobleAI, focusing on the advancement of sustainable and high-performance chemical and material products. His educational credentials include a Ph.D. in Biophysics from the University of California, San Francisco, a Master's in electrical engineering and computer science, and a Bachelor's in computer science and engineering, both from the Massachusetts Institute of Technology.

**John Nino****CEO at Life Science Outsourcing**

John Nino brings over 35 years of leadership and innovation in manufacturing, product development, and operations across the aerospace, medical device, and automotive industries. He began his career at Rockwell International's Rocketdyne Division, contributing to the development of propulsion systems for the Space Shuttle and the International Space Station. As CEO of Aveox and ECA Medical Instruments, Nino drove groundbreaking advancements in motion control systems and surgical technologies. He has also held senior operations positions for industry-leading BDC, American Capital Strategies (Nasdaq: ACAS), and Simulation Software & Services Leader MSC Software (Nasdaq: MSCS). Currently, as CEO of Life Science Outsourcing, he applies his vast expertise to support the evolving needs of global medical device OEMs worldwide. A University of California graduate with a BS in Electrical Engineering, Nino is Lean Manufacturing trained and holds several patents associated with pioneering surgical instruments and technologies.

**Paul Orlando****Business Development and Operations Executive with Schivo Medical**

Paul Orlando is currently working in a business development role at Schivo Medical since April 2025. Before joining Schivo Medical, he was with Olympus Surgical for 10 years, where he held an operations learnership role in support of M&A strategies and also served as vice president of procurement, where he oversaw the discovery, assessment, selection, and management of external solution providers, including finished devices, components, and design services. Prior to Olympus, Orlando held various operations, plant, engineering, business unit, and product development leadership positions with Motorola, Johnson & Johnson, and Vention Medical. He is a Boston native with more than 40 years of manufacturing, operations, supply chain, and engineering experience. Orlando has an Industrial Engineering degree from Northeastern University.

**Derek Prince, Ph.D.****President of Prince Sterilization Services**

Derek Prince began his professional career working at Gibraltar Laboratories, where he became an expert across various compendial tests on sterile and non-sterile pharmaceutical and health care related products. He has experience performing and developing procedures and test methods for a variety of microbiology, virology, and molecular biology-related test services, including rapid micro methods. In 2018, Prince formally departed Gibraltar to lead Prince Sterilization Services, a leader in environmentally friendly and safe outsourced pharmaceutical and medical device sterilization, wash & pack services, cleanroom assembly and kitting, and RTU products. The company is FDA registered, cGMP, and ISO 13485 certified. As president, Prince leads a high-performing team and fosters a quality-minded, performance-focused, safe, and friendly work environment. Prince is a subject matter expert in microbiology and sterilization science. He is a member of two Kilmer Collaboration sterilization work-stream groups focused on Parametric Release and VH2O2 sterilization. Prince has a bachelor's degree in biology, a Master's degree in microbiology, and a Ph.D. in microbiology from Rutgers University. He lives in New Jersey and enjoys playing hockey, hiking with his dog, and spending time with his family.

**Sarah Ptach****President and CEO of Canyon Labs**

Sarah Ptach is the president and CEO of Canyon Labs, a leading provider of medical device and pharmaceutical testing, consulting, and sterilization services. She joined the company during a pivotal ownership transition with a clear mission to elevate industry standards and build a true end-to-end solutions partner. Leveraging her background in packaging engineering and testing, Ptach has expanded Canyon's focus beyond packaging to deliver a more integrated, accessible, and expertise-driven experience for clients. She began her career in professional sports and advertising. A personal turning point came when her father was diagnosed with Parkinson's disease, which inspired her to pursue work that could create a meaningful impact in healthcare. She went on to help grow and successfully exit a packaging firm before bringing her vision and leadership to Canyon Labs. In addition to her role at Canyon, Ptach co-leads Kilmer Innovations in Packaging (KiIP) and serves on the board of the Medical Device Packaging Technical Committee (MDPTC) of the Institute of Packaging Professionals (IoPP). She remains deeply committed to advancing healthcare through innovation, expertise, and strong partnerships.

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**David Robson**

Founding COO and VP Quality & Regulatory Affairs at Iantrek

David Robson has spent his career concentrating on medical device development, including roles in R&D, project management, and regulatory compliance. In addition to multiple stints in startups and larger-scale OEMs, Robson spent 17 years at Ximedica (now Veranex), a contract medical device and IVD instruments design and development firm. In 2019, he was instrumental in converting an eye surgeon's concept for micro-interventional glaucoma surgery instruments (MIGS) into a feasible and documented design suitable for human use. He joined the resulting company, Iantrek Inc., as the founding COO and was the primary architect for establishing the facility as well as the quality management system and the development of the company's portfolio of products. As a result of Robson's 35+ years in the medical device industry, he has accumulated an uncommon set of experiences with respect to outsourcing services, both as a customer and provider. He has experienced the many benefits (and some of the pitfalls) of strategically balancing permanent employees and vertically integrated manufacturing with outsourced expertise and services such as contract manufacturing.

**Matt Rudow**

Sales Director for NobleAI

Matt Rudow is a sales and engineering professional based in San Francisco, CA. He has a background in material science, failure analysis, material characterization, physics-based simulation, material informatics, and enterprise strategy. Rudow has helped global enterprises evaluate, adopt, and strategically deploy AI to improve their products and performance. He is passionate about demystifying Practical AI for leadership in the product, chemical, material, and energy sectors. He believes medical devices could see a revolution by embracing and leveraging AI for their product development needs.

**John Schneider**

Senior Manager, Global Supplier Management, at Edwards Lifesciences

John Schneider is a senior manager in the Global Supplier Management organization for Edwards Lifesciences and leads the commercial supply chain strategy for the packaging, advanced catheters, and extrusion categories. He has over 25 years of increasing responsibility and complexity in the fields of sourcing, procurement, and supply chain. His expansive global experience includes the medtech, semiconductor, and aerospace industries. Schneider received his B.A. in Business Administration from Cal State Fullerton and his MBA from the University of Redlands. He is also APICS CPIM certified. Among his extensive endeavors, Schneider is deeply involved in various medtech and packaging conferences and leads the annual Edwards' Supplier Partner Awards Forum.

**Barry Schnur**

President and CEO at David Schnur Associates

Barry Schnur brings more than 30 years of medical device industry experience and is well-versed in the design, development, and manufacturing of medical components and devices. Under his leadership, David Schnur Associates (DSA) has become a leading global provider of outsourced technical sales and marketing services for the medical device industry. Leveraging a global network of partners, DSA consults with customers to objectively identify the right resources for their project, including materials, components, and commercial manufacturing strategy to launch devices efficiently and cost effectively.

**Julie Schulte, MBA**

Co-Founder and CEO of Chamfr

Julie Schulte brings over 20 years of experience in the medical device industry. She got her start in the industry with a small contract manufacturer focused on implantable and single-use devices that was acquired several times and eventually became part of Greatbatch (now Integer). Schulte led sales, marketing, and corporate sustainability efforts at Vention before it was sold to Nordson MEDICAL and MedPlast (now Viant). She co-founded Meraqi Medical, a device design, development, and manufacturing services provider, for which she served as president for two years prior to its acquisition by Viant where she led its design and development business as the vice president of commercial. Additionally, Schulte co-founded Chamfr, a medical device component marketplace, for which she currently serves as CEO. She holds a BS in Business Administration and Organizational Management as well as an MBA with an emphasis in Marketing.



Michelle Kennedy Scott

President and CEO of Gulf Sterilization

Before founding Gulf Sterilization, Michelle Scott built a well-respected speech pathology private practice, which continues to thrive today, 15 years later. During and after the pandemic, she became increasingly aware of medical product supply chain challenges through conversations with colleagues and family in the healthcare industry. Her research revealed a critical issue: limited sterilization capacity, increased turnaround times, and growing concerns over the environmental and safety risks associated with traditional sterilization methods. Determined to provide a safer, more sustainable solution, Scott established Gulf Sterilization—a state-of-the-art facility specializing in chlorine dioxide (CD) gas sterilization. As CEO, she led the company through the rigorous ISO 13485 certification process and continues to drive innovation in CD technology, including process optimization and packaging development. Most recently, she was named one of Alabama's top 25 women in Bio Tech for 2025. An active leader in the sterilization industry, Scott serves as a member of Bio Alabama, Florida Medical Manufacturers Consortium, Parenteral Drug Association, Women in Bio, Kilmer Modalities Team, the Biotechnician Task Force for ACCS Innovation Center and the Advancing Sight Network Innovation committee. She also contributes to the Industrial Sterilization and

Sterile Processing Working Groups and is currently drafting TIR124: Guidance Document on CD Sterilization for the Association for the Advancement of Medical Instrumentation (AAMI).



Matt Stekier

Principal—Supply Chain & Operations Consulting at Plante Moran

Matt Stekier serves clients throughout the supply chain continuum by quickly identifying improvement opportunities that deliver tangible results and lower costs in a variety of industries, including the medical device, food and beverage, footwear and apparel, military vehicle, and automotive manufacturing sectors. He earned his Bachelor's in supply chain management from Central Michigan University and a Master's in business administration from Wayne State University. Stekier also has obtained certifications as a Six Sigma Green Belt and in Theory of Constraints. A 10-year veteran of Plante Moran, he previously served in supply chain roles at Mercedes-Benz Technology and Ford Motor Company.

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Eric Steuben

Senior Vice President, Operations, at Calyxo

Eric Steuben is an energetic leader with 30 years of progressive achievements from hands-on engineering to executive management. He has broad corporate experience ranging from high-growth start-ups and world-leading corporations. Steuben has successfully transferred manufacturing operations offshore to China, Mexico, Pakistan, Dubai, and Costa Rica. Prior to joining Calyxo, he held executive leadership roles in manufacturing, operations, regulatory affairs, and quality assurance with PROCEPT BioRobotics, Clarity Medical Systems, Asante Solutions, Align Technology, and MicroDental Laboratories. He started his career at Varian Medical. Steuben holds both a BS degree in Mechanical Engineering and a Master's degree in Business Administration from Santa Clara University.



Jeff Wheeler

Director of Sales at NobleAI

Jeff Wheeler is a technical sales executive based in Boston, MA, with a focus on enterprise analytics, advanced manufacturing, and Practical AI. He enjoys working with global organizations to unlock the full value of their scientific and product data—accelerating innovation, improving margins, and reducing regulatory risk. Wheeler has helped enterprises evaluate, adopt, and deploy science-based AI to streamline formulation, optimize product performance, and modernize decision-making across the product development lifecycle.

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www.itek.net



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727-570-2293

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www.ndhmedical.com



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Wayne, PA

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www.tekni-plex.com/healthcare



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Wakefield, MA

978-573-2500

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www.amt-mat.com

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boulderiq.com



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www.cadenceinc.com

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Tracy, CA

209-532-5146

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www.centplasticmfg.com

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315-255-1779

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currierplastics.com/markets/healthcare/

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www.eptam.com

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www.forefrontmedical.com

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www.hahnautomation.group

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impactplastics.co

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www.indo-mim.com



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www.irpmedical.com

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kmmgrp.com

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801-972-6150

KP provides marketing services and print supply chain solutions for medical device and pharmaceutical companies. This includes IFUs, inserts, packaging, labels, training kits, promotional materials, direct mail, non-sterile kitting, inventory management, and distribution, all available via online ordering tools. KP operates five locations, distributing throughout the U.S., Mexico, and internationally. KP is also ISO-certified.

www.kpcorp.com

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The Woodlands, TX

346-327-2412

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www.punch-us.com

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www.roechling.com/medical

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www.sigmatronintl.com

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www.ableelectropolishing.com

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argonmedicalcps.com

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617-314-3950

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www.covllc.com

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www.dschnur.com

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Buffalo, NY

716-897-4500

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MICRO

Somerset, NJ

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www.micro-co.com

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Minneapolis, MN

Resolution Medical is a privately held contract medical-device design and manufacturing firm. They specialize in complex catheter delivery systems, active implantable devices, and component production—supporting neuromodulation, electrophysiology, structural-heart, cardiology & vascular, and heart-failure markets. Their integrated, ISO-certified and FDA-registered capabilities span concept through commercial finished goods manufacturing.

resolutionmedical.com

Resonetics

Nashua, NH

603-886-6772

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ADDITIONAL INFORMATION TO COME!

Any questions, please contact Howard Revitch
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Supporting Organizations

BMP Medical

Sterling, MA

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BMP Medical is a U.S.-based, 80,000-sq.-ft. plastic manufacturing facility for medical devices and components, with 45 years of experience in helping customers bridge their capability gaps and get products to market quickly. We partner with OEMs, CMOs, Engineers, and Designers to develop high-quality products that enhance patient care.

www.bmpmedical.com

Evolution Free Zone

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Evolution Free Zone is the fastest-growing business ecosystem in Costa Rica. Offers state-of-the-art infrastructure, strategic geographical location in the center of the Americas, and the best complementary services for innovative medtech companies.

www.evolutionfz.com

Life Science Outsourcing Inc.

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Life Science Outsourcing is an FDA-registered, ISO 13485-certified CMO specializing in assembly, packaging, sterilization, and specialized capabilities in diagnostics packaging & design. We offer end-to-end services with deep regulatory expertise, enabling agile market launches while standardizing supply chains and mitigating risk.

www.iso-inc.com

MDMA - Medical Device Manufacturers Association

Washington, DC

202-354-7171

MDMA is a national trade association based in Washington, DC. We provide educational and advocacy assistance to innovative and entrepreneurial medical technology companies. Since 1992, MDMA has been the voice for medical technology innovation, leading the industry's role to help shape policies that impact and improve the ecosystem for patients and innovators. This is accomplished by working with key Members of Congress, senior staff at FDA, CMS, and other agencies, and through the grassroots support of our members.

www.medicaldevices.org

The MedTech Conference

Washington, D.C.

202-434-7213

Hosted by AdvaMed, The MedTech Conference brings the global medtech community to San Diego, October 5–8, 2025, for four days of critical education, hands-on innovation, and valuable networking that drives insight, collaboration, and progress across the industry. Register now!

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MedWorld Advisors

Andover, MA

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MedWorld Advisors (MWA) is a leading international boutique mergers and acquisition advisory firm based in Andover, Mass., U.S. MWA specializes in helping small- to middle-market companies in medical device, medtech, biotech, dental, life sciences, digital health, medical, and dental practice, and anything healthcare to achieve their stakeholder objectives.

www.medworldadvisors.com

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MTD Micro Molding is your single-source partner for micro medical manufacturing, from material selection to assembly and packaging. With our exclusive focus on the medical market, we offer unparalleled bioabsorbable expertise, breakthrough drug delivery molding capabilities, and advanced overmolding services. Bring us your most challenging micro medical designs and our team of engineers will get your products to market faster with improved manufacturability solutions. ISO 13485; ISO 15378; ISO 9001; FDA Registered; ISO Class 8 Cleanrooms.

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The Future of E-beam and X-ray



SteriTek is a high-volume E-beam/X-ray contract sterilizer and R&D innovation center serving the medical device, biotech, pharmaceutical, and other industries.

Particularly with sensitive materials and complex devices, SteriTek has developed a proprietary system for optimizing E-beam/X-ray for radiation sensitive materials such as drugs/biologics, combination devices, implantables, bioresorbables, and other complex products.

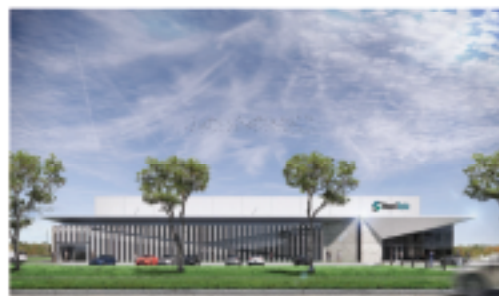
Established in 2016, SteriTek has emerged a well-recognized and trusted E-beam/X-ray partner for startups and global companies alike and we are expanding to serve growing demand.



SteriTek's flagship facility in Silicon Valley (Fremont, CA) boasts two state-of-the-art 10 MeV 20KW linear accelerators, using simultaneous beam processing that allows for high volume production, providing uniform dose to the product without having to rotate the customer's boxes.

This DualBeam™ configuration significantly increases efficiencies, expands product options, serves as an effective back-up for the accelerators sterilization needs.

Our newest facility in Dallas (Lewisville, TX) with operations having started in May 2023 will ultimately house three separate lines, two E-beam/X-ray 10MeV, 30KW DualBeam™ lines and a dedicated X-ray 7 MeV, 560 KW line which together will increase total throughput fivefold.



With this operational format, SteriTek offers medical device, biotech, and pharmaceutical companies with:

- Expertise in optimizing E-beam/X-ray for complex devices and sensitive materials (combination devices, drugs, biologics, allograft tissue)
- Fastest standard turnaround in the industry (Routine 6 business days, RUSH 24 hours and 4 hours)
- Highest up time in the industry
- Direct involvement in ISO 11137 and AAMI

SteriTek can help with the R&D testing and determination of E-beam or X-ray as the most optimal sterilization method for your new product.

Please contact us today at (510) 933-9700 to inquire how we can help solve your complex sterilization needs.

(510) 933 - 9700 | www.steri-tek.com | [@ebeamxray](https://twitter.com/ebeamxray)
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