

# DIA CMC Workshop Exhibitor Directory

April 24-26 | Rockville, MD  
Hilton Washington DC/Rockville Hotel & Executive Meeting Center



## American Association of Pharmaceutical Scientists

**Phone:** +1.703.243.2800

**Email:** [aaps@aaps.org](mailto:aaps@aaps.org)

**Website:** [www.aaps.org](http://www.aaps.org)

A non-profit professional organization, AAPS aims to advance the capacity of pharmaceutical scientists to develop products and therapies to improve health world-wide. A global membership of over 9,000 pharmaceutical scientific leaders in industry, government, and academia discover, develop, and manufacture pharmaceutical products and therapies.

## FreeThink Technologies, Inc.

**Phone:** +1.860.237.5800

**Website:** [freethinktech.com](http://freethinktech.com)

FreeThink Technologies, Inc. is a thought leader in the science and technology of stability studies, with some of the most knowledgeable and experienced research scientists in the field. Our team is experienced in a range of product types, and routinely develop and execute studies for pharmaceuticals, biopharmaceuticals, generic drugs, OTC medicines, nutraceuticals, cosmetics and other household and consumer products.

## Intertek QTI

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Intertek's global network of laboratories is designed to exceed our customers' expectations with innovative services for their operations and supply chain. Our Whitehouse NJ laboratory is an FDA and DEA registered cGMP compliant facility with expertise in Method Development/Validation, Stability, Extractables & Leachables, and Elemental Impurities.

## MakroCare

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MakroCare is supporting clients in FSP&FTE capacities with quality, consistency & technical validity of CMC technical documentation. It's providing documentation hierarchy for regulatory submission for clinical trial applications (IMP/IND) or marketing authorization (NDA, BLA, ANDAs, MAA) in CTD Module3. Its CMCxtract tool helps retrieve & analyze CMC content.

## Mapi

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Mapi Group has 40+ years of experience supporting Life-Science companies in utilizing Strategic Regulatory Services, Patient Reported Outcomes measures, Value communications and Commercialization support, and gathering Real-World Evidence on Medical devices, Pharmaceuticals, and Biologicals.

## PharmaLex US

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