Marine For Health Regulatory Seminar

10. December 2019	
10:00-10:45	Registration

10. December 2019	
10:45-11:00	**Wenche Uksnøy, Welcome and introduction**
11:00- 12:00	**Mari Lyyra,Regulatory Affairs Manager Food and Nutrition, Medfiles**, **Finland** Novel food and health claim dossiers-A well done plan guarantees the best outcome
12:00-13:00	**Lunch**
13:00-14:00	**Anders Mikael Sjödin, Professor, Division Nutrition, Exercise and Sports, University of Copenhagen, Denmark, EFSA Expert Panel** TBC
14:00-15:00	**Asim K. Duttaroy, Professor, Clinical Nutrition, University of Oslo, Norway** Functional food regulations in Europe: A case study
15:00-15:20	**Break**
15:20-16:20	**Jim Lund, Research Institutes of Sweden (RISE), Sweden** Marine research and business innovation platform-A pilot study
16:20-16:20	**Break**
16:40-17:40	**Maie Gall, CEO, Varanah Ventures, Switzerland** Disrupting a health paradigm; Setting a new way of thinking
18:15-22:30	**Dinner**

11. December 2019	
9:00-10:00	**Firoz Roshan, Pharmaceutical Regulatory Affairs Specialist, Sweden** Role of regulatory affairs in clinical drug development
10:00-10:30	**Break**
10:30-11:30	**Jan Petter Akselsen, Leader, The Norwegian Medicines Agency Innovation Office, Norway** Regulatory Requirements and Services in Drug Development
11:30-12:30	**Grethe Berven, Link Medical, Norway** Pharmaceutical regulatory affairs: A business perspective
12:30-13:30	**Lunch**

	Wenche Wenche has over fifteen years of operational and strategic experience from the international marine ingredients industry in various sales and marketing roles. She holds an international marketing degree from BI Norwegian School of Management Oslo / ESADE Barcelona. In her current position Wenche is the leader of the marine / biomarine cluster NCE Blue Legasea, which has been given the status of Norwegian Center of Expertise in 2018. The cluster gathers resources, experience and knowledge from all parts of the marine value chain, in close interaction with the maritime sector. The ambition of the cluster project is to be the driving force for innovation, value creation and knowledge building in the utilization of marine raw material together with its 50 partners.
	Mari Lyyra- Regulatory affairs manager, Food & Nutrition, Medfiles, Finland Mari is a nutritionist (M.Sc. in clinical nutrition) with over 20 years of experience working on food regulatory issues. She has been with Medfiles for the last five and a half years. Mari is specialized in novel food and health claim tasks but also ordinary foods, food supplements, food additives, foods for specific groups and animal feeds.
	Anders Mikael Sjödin
	Asim K. Duttaroy- Professor University of Oslo, Norway Dr. Duttaroy is a professor at the Department of Nutrition at the University of Oslo. His scientific work has been published in over 250 original contributions, reviews, books, book chapters, and editorials. Dr. Duttaroy is a holder of several international patents, including cardio-protective factors in tomatoes and kiwifruits. Fruitflow™ is a patented, scientifically-proven tomato extract which inhibits platelet aggregation, a known cause of heart attack, stroke and venous thrombosis. Fruitflow™ became the first product in Europe to obtain an approved positive health claim under Article 13.5 of the European Food Safety Authority regulations. Dr. Duttaroy has successfully supervised 25 PhD/MD students and has several international patents, that led to the establishment of two international companies Provexis Ltd. UK, and IDIA AS, Norway. Dr Duttaroy is also Editor in Chief of Food and Nutrition Research (previously known as Scandinavian Journal of Nutrition).

	Jim Lund- Research Institutes of Sweden (RISE) Jim works as a researcher and project leader for RISE. Jim focuses on coordination and participation of cross disciplinary R&D in collaboration with industry, academy and healthcare.
	Maie Gall – CEO Viskumed- Next Generation Healthcare Investor and Accelerator Maie is a healthcare entrepreneur, with a focus on investing in disruptive innovation which delivers sustainable healthcare innovation to society. She has over 30 years' experience in the healthcare sector and has led on the start-up of several successful companies. She is currently CEO of 9 disruptive healthcare start-ups and is passionate about corporate strategy, M&A, forecasting, projections, and identifying future trends.
	Firoz Roshan – Regulatory affairs specialist Firoz works for the CRO, Pharmaceutical Product Development (PPD). His work involves supporting client pharma companies in obtaining regulatory approvals especially in Phase III clinical trials. Previously Roshan has worked in the field of regulatory labeling and publishing at AstraZeneca. Roshan has a PhD in Neurobiology of Alzheimer's Disease and Masters in Medical Biosciences.