

## SOLVE M.E. LEADERSHIP



**Oved Amitay** is the Chief Executive Officer at Solve M.E. He has dedicated most of his professional career to the development of life-changing therapeutic options for people affected by rare genetic diseases. Throughout his career, Amitay has enjoyed extensive engagement with healthcare systems and patient advocacy organizations around the world, forging collaborations to address the needs of people affected by rare genetic diseases.

Amitay recognizes opportunities for Solve M.E. to evolve their work to include enabling public-private collaborations and expanding the connectivity within the ME/CFS community—both in the U.S. and globally. His overall vision for the organization includes a better understanding of ME/CFS as a disease, its underlying causes, personal variability, and the full scope of impact to ultimately arrive at the identification of targets for intervention and the availability of therapies that can improve quality of life.



**Emily Taylor** is the Vice President of Advocacy and Engagement for Solve M.E.. She brings to the organization over 15 years of policy, organization, and advocacy experience in both the non-profit and government sectors. With five years' experience on Capitol Hill, Emily contributes a keen understanding of both state and federal policy processes while cultivating strong grassroots organization and patient representation.

Prior to joining the Solve M.E., Emily served as the director of policy and advocacy for an award-winning autism organization. There she spearheaded major overhauls in disability, early intervention, and education policies. She is also a veteran of several successful electoral campaigns, where she trained others in effective social media and online organization. Emily draws inspiration from her mother who has battled ME/CFS as well as chronic autoimmune and thyroid conditions since 1999.

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## ABOUT SOLVE M.E.

Founded in 1987, Solve ME/CFS Initiative (Solve M.E.) is a non-profit organization that serves as a catalyst for critical research into diagnostics, treatments, and cures for myalgic encephalomyelitis/chronic fatigue syndrome (ME/CFS), Long Covid and other post-infection diseases.

Our work with the scientific, medical, and pharmaceutical communities, advocacy with government agencies, and alliances with patient groups around the world are laying the foundation for breakthroughs that can improve the lives of millions who suffer from various “long haul” diseases.