



Laina Shamblin: Advocate, Leader, and Inspiration for the Voiceless

Laina Shamblin is the President of the Board for Alliance League Matching Services, Inc. (ALMS). She is a woman with Down's Syndrome, born to a mother with Down's Syndrome, making her a rare and genetically unique individual. She is also a member of the Shoshone Indian tribe in the Pacific Northwest region of the United States.

From an early age, Laina demonstrated resilience—she underwent open-heart surgery at just six months old. Despite challenges, she leads an active and engaged life. Laina enjoys swimming, singing karaoke, all things Disney, and helping to care for her family.

Laina is not only a guiding force within ALMS but also an inspiration for those without a voice in society—individuals who refuse to comply with mRNA-based technologies encoding synthetic DNA ledger systems as a requirement to buy, sell, or receive funding and services in the future. With her unique DNA and history of heart-related surgery, the safety and efficacy of such substances have never been tested or proven for her. Her personal reality underscores the ethical and medical concerns surrounding forced compliance with unproven bio-cybernetic systems.

Her perspective on the world's trajectory regarding healthcare, biotechnology, and environmental neglect is summed up powerfully in her words: "Not good." She serves as a beacon of integrity, standing firm against coerced participation in synthetic identity systems and inspiring others to reject technologies that compromise human autonomy and dignity.