

WeHaveAFace.org



Our Mission:

Founded in March 2011, WeHaveAFace.org is the first organization founded to increase international awareness of Huntington's disease through a cinematic form. The Documentary:

"The Faces of Huntington's Disease: I am No Longer a Faceless Face"

The Documentary focuses on the personal stories of the International Huntington's disease community, and incorporates education from Medical Professionals within its content. The aim of The Huntington's Disease Project is to produce an independent, full-length documentary. The film is the first of its kind, created by individuals within the JHD/HD community.

The Huntington's Disease Project is comprised of an incredible team of individuals within the Huntington's disease community. The [Team](#) has joined efforts to create the Documentary, and to continue to provide valuable support to others within the JHD/HD community.

Throughout the entire project, all members of the Film Team work selflessly with the understanding that there is no financial gain. This also applies to all individuals who will be interviewed (JHD/HD community participation and Medical Professionals). Feel free to [participate](#) in the Documentary.

The ultimate goal of this Documentary is to bring about international awareness of JHD/HD, and to provide a mechanism for individuals within the community to tell their stories. All donations to produce the Documentary are collected via Deshalamar CDC 501(c)3 of New York. Once the full-length documentary is released, any/all remaining donations, and future proceeds, will afford [Deshalamar CDC 501\(c\)3](#) the ability to create programs and services for individuals within the JHD/HD community. Advocacy for Huntington's disease is a key part of our daily lives, and our project is a legacy for future generations. If not now, when?

You Are Loved!

James Valvano

