**MOvie Fest! Advanced ticketing required.**

Breaking down the silos in our health care system has a lot to do with aligning incentives of everyone involved—patients, entrepreneurs, hospitals, regulators, researchers, and industry. This year’s DIA Annual Meeting presents two film documentaries, shown Tuesday and Wednesday evening, brought to you by the Kauffman Foundation. Wine and light refreshments to be provided.

Seating is limited. Advanced ticketing is available on a first-come, first-served basis.

**Please note that the DIA Courtesy Shuttle will not be available after 7:30PM.**

### RARE
**June 25 | 5:30–7:30PM**

RARE tells the story of Donna Appell, an extraordinary mother, in a race against time to get a clinical trial going to treat her daughter’s rare genetic disease; the heartbreak when her daughter Ashley, whose disease is advancing with her age, is excluded from the trial; a surprising love story when Ashley falls in love with an earnest young man who also has Hermansky Pudlak Syndrome; and the intimate, intricate relationship between a research scientist and the patients to whom he inevitably becomes emotionally attached.

To obtain tickets for RARE, please visit the Attendee Registration Desk no later than Monday at 5:00PM.

### HERE.US.NOW.
**June 26 | 5:30–7:30PM**

The film, directed by Emmy award-winning filmmaker Rudy Poe, is a story of three realities in the US today. First, the patient advocate’s role in driving changes in policy and popular sentiment—we saw this in the AIDs movement and hope for the same in pediatric and rare diseases. Second, the Myelin Repair Foundation, Michael J Fox Foundation, Army of Women are depicted and are pursuing models of patient-driven R&D partnerships with academic and private sector scientists. These models, which are intent on utilizing a patient focus to accelerate research outcomes while also driving policy changes through their unique funding models, have grown significantly and are achieving critical outcomes. Last but not least, this documentary underscores the story of the Hempels, who represent a patient/parent-conducted R&D model. The intent of this documentary is to underscore the need for accelerating the alternative pathways from new discoveries to curing patients.

**This showing will be followed by an interactive panel discussion with Hugh Hempel, Lesa Mitchell, John Wilbanks, Marc Boutin.**

To obtain tickets for HERE.US.NOW, please visit the Attendee Registration Desk no later than Tuesday at 5:00PM.

### FEATURED SESSION
**Wednesday, 8:00AM | Room 154**

**#313 Using Legislation to Advance Regulatory Science: “I’m Just a Bill...”**

Several legislations have been introduced calling for regulatory reform for the drug approval process. FDA is under pressure from industry to enable a regulatory framework for translational medicine but at the same time preserve the public health. This session will discuss impacts to the pharmaceutical industry and investment community as well as the patient community.

**Session Chair:**
Karen E. Jaffe, MBA, MS, MSc, RAC
Regulatory Research, Alfred Mann Institute

**Panelists:**
Juergen Froehlich, MD, MBA, FFPM
Vice President, Global Regulatory Affairs
Vertex Pharmaceuticals

Marc M. Boutin, Esq, JD
Executive Vice President and Chief Operating Officer
National Health Council

Steven Walker
Co-founder, Abigail Alliance

Carla Mann Woods
CEO and Founder, Fight to Live