

EB Research Partnership was founded in 2010 by a group of parents set out to save their children's lives along with Jill and Eddie Vedder (Pearl Jam) with one bold audacious goal: cure the devastating and life-threatening genetic rare skin disease epidermolysis bullosa (EB) by 2030 and in the process pioneer an innovative business model to lead the way for 7,000 rare diseases that affect 400 million people.

The mission of EB Research Partnership is to advance life-saving treatments, and ultimately, find a cure for epidermolysis bullosa (EB) by the end of this decade. Secondly, EBRP's innovative model is leading the way for all rare diseases. While EB is a rare disease, there are 7,000 rare diseases that affect 1 in 10 people in the world, 95% which have no treatments, that can benefit from the research we fund and our innovative venture philanthropy - both of which are scalable to all rare diseases.