

worth the ride

A journey with pachyostosis, muscular dystrophy,



Josh Winheld

www.cuppress.com/joshwinheld

Worth The Ride My Journey With Duchenne Muscular Dystrophy

Cyndie Degnan



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Worth the Ride Little Treasure Books,2008-03 **Migraine Expressions , Eco-Teen Films** Robin L.

Murray,Joseph K. Heumann,2024-09-06 Illuminating the impacts of environmental disasters and climate crises globally this book examines the experiences of teens grappling with eco disasters and issues in films of the twenty first century With an emphasis on teen activism international settings and filmmakers and marginalized perspectives this book showcases teens on film that are struggling with present and future everyday eco disasters amplified by climate change By highlighting and interrogating diverse genres of teen films in which young adults encounter address and battle environmental issues and calamities while also struggling with adolescent development this book acknowledges the young adult point of view missing from most critical ecocinema research and underlines connections between the more complex coming of age themes found in teen films with ecocinema themes and approaches The films examined navigate increasingly realistic conditions even in fantastical settings as they showcase teens relationships with and responses to environmental issues and eco disasters Emphasizing teen activism and under represented intersectional perspectives outside Hollywood it establishes the eco teen film as a notable subgenre This book will be of interest to students and scholars of film studies ecocriticism and environmental studies especially those with a particular interest in ecocinema and or ecocritical readings of films **The**

Oxford Compendium of Hope Anthony Scioli,Steven C. van den Heuvel,2025-12-22 The Oxford Compendium of Hope presents an authoritative and comprehensive guide to hope with five main aims provide a historical context for the interdisciplinary study of hope condense and organize seminal writings on hope document the state of research on hope across 12 scholarly areas prioritize scholarship with generative power and cross disciplinary potential and anchor itself in a unifying philosophical meta perspective on hope Central to the volume s purpose is the editor s third culture approach to the study of hope encompassing both a scientific and a humanities based examination Chapter authors hailing from psychology philosophy health studies anthropology and other disciplines examine hope from their unique vantage points **Be the**

Change Rita Verma,2010 At a time when teachers are the scapegoats for all that is wrong with education Rita Verma and colleagues push back by illuminating the critical and creative roles that teachers and youth are playing to make education impactful The examples in this book model the possibilities for anti oppressive activism through education and inspire Kevin Kumashiro author of The Seduction of Common Sense How the Right has Framed the Debate on America s Schools

Worth the Ride Josh Winheld,2009-05-01 Enter the world and mind of a young man who lives with Duchenne muscular dystrophy a fatal genetic neuromuscular disorder that he battles with his incredible sense of humor warm heart and awe inspiring passion for life Calling this the master s thesis he never wrote Josh Winheld s self reflection and candor teaches us all valuable life lessons Find out why Worth the Ride is well worth a read All proceeds from this book will go directly to Parent Project Muscular Dystrophy which works to improve the treatment quality of life and long term outlook for all

individuals affected by Duchenne muscular dystrophy through research advocacy education and compassion Health Humanities Reader Therese Jones, Delese Wear, Lester D. Friedman, 2014-08-28 Over the past forty years the health humanities previously called the medical humanities has emerged as one of the most exciting fields for interdisciplinary scholarship advancing humanistic inquiry into bioethics human rights health care and the uses of technology It has also helped inspire medical practitioners to engage in deeper reflection about the human elements of their practice In Health Humanities Reader editors Therese Jones Delese Wear and Lester D Friedman have assembled fifty four leading scholars educators artists and clinicians to survey the rich body of work that has already emerged from the field and to imagine fresh approaches to the health humanities in these original essays The collection s contributors reflect the extraordinary diversity of the field including scholars from the disciplines of disability studies history literature nursing religion narrative medicine philosophy bioethics medicine and the social sciences With warmth and humor critical acumen and ethical insight Health Humanities Reader truly humanizes the field of medicine Its accessible language and broad scope offers something for everyone from the experienced medical professional to a reader interested in health and illness Video Source Book ,2006 A guide to programs currently available on video in the areas of movies entertainment general interest education sports recreation fine arts health science business industry children juvenile how to instruction *Our Journey with Duchenne Muscular Dystrophy* Vicki Spence, Stacy Spence, 2010 *Fueled by Misery: My Journey Through Life with Muscular Dystrophy* Peter Li, 2017-07 When I first met Peter Li he greeted me from his wheelchair with a smile and hearty Nice to meet you Despite the fact that MD has taken away most of his mobility and that he is 100% dependent on his care takers at Angel s View he maintains a pretty cheerful spirit and a smile that fills the room Behind that spirit though is a story of surviving and navigating life events that would leave most people without any reason to smile Peter is a Chinese immigrant who sailed his way to American with his family His childhood MD diagnosis came with devastating implications for a boy who just wanted to fit in and for a family unfamiliar with the disease And as MD began to take over his body Peter s own bout with depression had him wondering if medical care was even worth it As we sat and talked that sunny day in California Peter shared the tragedy of his life but he also shared the thankfulness and gratitude that has been a part of his experience He receives incredible care from the nurses who serve the home he and his brother and several other friends share all battling MD He has met celebrities He has been the keynote at fund raising galas and has helped spread the message that life after MD can still be richly fulfilling Spend a few hours with Peter as I did and you ll walk away with a smile on your face and boost to your own spirit That s Peter s hope for this book that you will be encouraged to live life to its fullest and embrace all parts of it the good parts and the hard parts as aspects of the journey So go ahead meet Peter by reading his book and see if you can find a smile and some encouragement for your life Joe Castaneda Publisher **Journey of Love** ,1998 *Heart Shaped Rocks, Don't Die Without Me* Misty Vanderweele, 2014-05-23 Heart Shaped Rocks is about a sister named Jenna as she lives

knowing her brother could die at any time from Duchenne Muscular Dystrophy What has brought Jenna extreme hardship and struggle over and over she is anything but broken Jenna s story will not only leave a mark on your heart you ll start seeing hearts everywhere In nature and maybe even finding them living on your windowsill The Diagnosis Luke Dalien,2018-12-05 *The Explanation: Epic Battle with Muscular Dystrophy* Luke Dalien,2019-03-25 It s a fact that children understand difficult situations at a much different level than grown ups Mark The Mighty Muskrat is a series of children s books that tackles this exact reality The Explanation is a real life story of how the author and his wife explained Duchenne Muscular Dystrophy to their newly diagnosed four year old son and his six year old sister Each page in this incredibly personal family centric story is filled with beautifully illustrated characters and rhyme filled poetry The second in a series intent on battling the stigma and fear so often associated with tough things this children s book describes 1 Just how unique each person is in this world 2 How each person learns to adapt to their reality 3 The importance of understanding and respecting each other The final page in the book describes it all So the next time you see someone who looks or acts differently than you Remember their body was made in a unique way and they re special too Believe God & Me: His Story. My Journey with Muscular Dystrophy Peter E. Brown,2018-10-17 He wasn t supposed to live The odds were against him and he beat them He saw victory in things impossible He sees the silver lining instead of the darkness He sees hope when there is none He speaks inspiration when others delve into the negative Nothing in his life was simple and easy but he persevered This is his story **Duchenne Muscular Dystrophy** Josephine M. Green,Frances E. Murton,1994

Moonrise Penny Wolfson,2014-04-22 Moonrise is Penny Wolfson s first person account of her family her son Ansel and his progressive disability caused by the genetic disease Duchenne muscular dystrophy The journey begins when he is born and deemed a particularly beautiful and magical baby continues with the alarming possibility at the age of two of wrongness takes us through the diagnosis of disease and prognosis of early death and brings us to his adolescence where his parents are never sure if the moon is rising or setting over his life As she traces her son s development and the impact of his disability on her worldview she embarks on a quest to understand scientific advances and their implications The gene was isolated at approximately the time Ansel was diagnosed She also explores special education giftedness prenatal testing and the genetic links she shares with her mother sisters and son Questions about the disease causing mutation persist What does knowledge of the self on a molecular level mean Is genetic self knowledge our goal now much as knowledge of the psyche was in the last century Moonrise is an essential contribution to the dialogue about genetics as well as a deeply human story about a remarkable child and his family **The Logan Letters** Logan Shannon,2017-06-05 Logan is the oldest son of Michael and Tyra Shannon and the second of four children When he was a toddler his parents noticed his physical development began to deteriorate Doctors in Cincinnati diagnosed him with Duchenne Muscular Dystrophy Read the chronicles of how others shared his journey from walking with assistance to being confined to a wheelchair you ll feel a part of his perspective in the

value of family friends and faith in God Back cover His Strength Made Perfect Cyndie Degnan,2012-02 At the age of four Michael Degnan was diagnosed with Duchenne muscular dystrophy His parents were told he had only ten years of quality life ahead after that he would steadily deteriorate Through journal entries Michael s mother shares the family s journey In Your Face Duchenne Muscular Dystrophy All Pain All Glory Misty VanderWeele,2010 A mother s story of life with her son who has Duchenne Muscular Dystrophy

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