



Our 2026 Impact Report is out!

Read about everything your continued support has allowed us to accomplish this year — from funding breakthrough lab research in institutions across the country; to answering an urgent need to step into the clinical trial space; to providing critical funds to families in financial crisis while receiving cancer treatments, and more.

Plus, you'll hear about remarkable young cancer survivors impacted by our work — now paying it forward to other kids facing cancer.

It's our biggest year of funding yet. And we couldn't do it without you. Together, we can do even bigger and bolder things next year!

Scan to read the report now:



Funding the next generation of scientific leaders!

The Morgan Adams Foundation is thrilled to support the Gates Summer Internship Program by sponsoring two students in this remarkable program as they work with our MAF funded researchers to help advance both their own development as well as critical kids' cancer research.



Meet Dina, a first-generation college student who immigrated from Russia to Atlanta with her musician parents.

Hello! My name is Dina and I am a sophomore Biology major from Georgia Tech! I am super passionate about

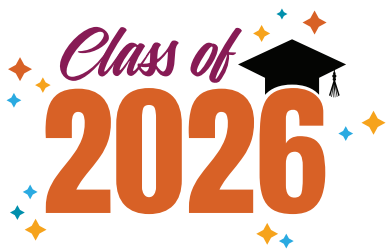
medicine, clinical research, oncology, and finding new ways to explore and integrate sciences into my life. I am super excited to get to work in the Franco Lab this summer, and I am very thankful for The Morgan Adams Foundation for funding this amazing opportunity!



Meet Stella, who has overcome tremendous health challenges to go on be rated #1 out of 70 interns in a previous position this past year.

My name is Stella, and I am so grateful to be supported by The Morgan Adams Foundation! I

am thrilled to be working with Dr. Sujatha Venkataraman on engineering and testing CAR T cells to treat solid tumors. I am passionate about finding better treatments and cures for childhood cancers and look forward to partnering with MAF to do so!



Congratulations to our Ambassador Graduates!



Cameron, age 18, diagnosed with Hypodiploid B Cell-Acute Lymphocytic Leukemia. Graduating from The Academy of Westminster and heading to Colorado School of Mines in the fall!



Priscilla, age 17, diagnosed with Hodgkin Lymphoma. Graduated early from High School in December, and heading off to college in the fall!

The end of May is almost upon us, marking a season of new beginnings! At The Morgan Adams Foundation, these milestones hold a special significance. We know that behind every cap and gown is a story of incredible strength — and often, a reliance on the breakthroughs made in pediatric cancer research.

Investment in research is what allows kids facing a cancer diagnosis to not only survive, but to truly thrive and reach their goals. Whether it was navigating treatments or balancing studies, our graduating Ambassadors have shown that a diagnosis does not define their potential. Because of advancements in treatments made possible by donors like you, these students can walk across their high school stages and step into their next chapter.

To Cameron and Priscilla: Your resilience and dedication have brought you to this moment. We are proud of everything you've accomplished! Congratulations on this monumental achievement — we'll be rooting for you in all your future chapters!

What's Up Doc? 2026: Research to Reality

Scan the QR code to watch the full event

Two of our esteemed researchers, Rajeev Vibhakar MD, PhD and Todd Hankinson, MD, shared updates on the innovative projects your gifts are helping to move forward. We were also deeply moved by MAF Ambassador Austin's story, as his parents spoke about his brain tumor diagnosis and journey—and how they credit Dr. Hankinson's research (and your donations) for his survival and quality of life.

Huge thanks to our sponsors Amber Page, Realtor and CHUBB North America.

WHAT'S UP DOC?



Your Legacy: Brighter Futures for Kids, Teens, and Young Adults with Cancer

When you make a planned gift to The Morgan Adams Foundation, you'll become a member of The Iris Society — a group of donors taking charge today of what their legacies mean tomorrow. It's perhaps the most meaningful gift you can make — from your family to families who need it the most.



The Iris Society

THE **Morgan Adams** FOUNDATION

A legacy gift matters

A planned gift is an incredible way to help secure the long-term stability of The Morgan Adams Foundation — no matter what the future holds. It's a gift that can help fund lifesaving treatments for generations to come.

“As we made our estate plans, we were determined to help ensure that the tireless efforts of The Morgan Adams Foundation had a shot at long-term stability. Our will was written so that MAF received a specified percentage of our estate before other distributions are made. It is one of our great wishes that this organization endures until there is a cure for all kids who, tragically, get cancer.”

— Dr. & Mrs. Neal Adams, Members of the Iris Society



Scan here more information



Expanding RadFlix Internationally!

We're thrilled to announce that RadFlix has launched at Princess Margaret Cancer Centre in Toronto — the first clinical site in Canada to offer this innovative technology.

RadFlix will allow the hundreds of kids receiving radiation each year at this institution to watch their favorite shows during treatments — reducing anxiety and often eliminating the need for anesthesia.

This is the power of donating to RadFlix: turning fear into comfort and innovation into compassion. Learn more at **RadFlix.Org**



Featured Project

A Promising Osteosarcoma Study linking human/canine tumor biology, and a path forward to treat both

Osteosarcoma is the most common bone cancer in children and teens. Treatment options have not meaningfully changed in decades, and survival rates remain unacceptably low.

Thanks to your generosity, The Morgan Adams Foundation is funding a new treatment strategy designed to make innovative immunotherapy treatments more effective ... by engaging and aiding “humans’ best friends” facing the same and biologically similar disease.

Dogs naturally develop osteosarcoma and are considered the most accurate non-human model of osteosarcoma. MAF funded researchers will combine innovative immunotherapies with readily available drugs to disrupt the cancer’s growth.

Ultimately, this study aims to generate critical insights that can translate to children with metastatic osteosarcoma, offering a promising new path toward more effective and less toxic treatments.



Janea

January 1, 2000 - April 28, 2014

Funding new hope for kids, teens, and young adults like Janea

Janea was diagnosed with Osteosarcoma at just 12 years old. By the time of diagnosis, the disease had spread to her lungs — making it much more difficult to cure.

For two very long years, Janea fought this cancer with courageous positivity and a happy personality as much as possible. She endured 10 months of chemo, three major surgeries (including a leg amputation), multiple minor surgeries, and countless other unbearable side effects ... which she took in stride.

Just a few short months after Janea’s initial treatments ended, the cancer relapsed. She and her family bravely faced what lay ahead. Devastatingly, the treatment options that remained for Janea failed.

But during this final battle, when she was only given two weeks to live, Janea had her last wish granted to attend Prom. Her brother escorted her, and she was crowned Prom Princess! Everyone who knew Janea, loved her. And she will be forever missed.





Links Fore Liam

Links Fore Liam joins forces with The Morgan Adams Foundation



Five years ago, in the face of unimaginable loss, Liam's family found a way to honor their child's memory while creating a rallying point for the community to come together in the fight against a devastating disease.

Today, the golf tournament series, *Links Fore Liam*, has expanded to multiple locations, raising close to \$1 million towards pediatric cancer research to date.

We couldn't be more excited to announce that we've partnered with Liam's Dad (and the organization's founder), Nick Bascle, to bring Links Fore Liam under our umbrella and help aid further growth, adding events next year in Vail, CO; Scottsdale, AZ; and beyond.

Together, we'll be able to invest the funds raised directly towards labs investigating the precise brain tumor (Ependymoma) that affected Liam, as well as other select childhood cancer research projects. We are incredibly honored to help extend Liam's legacy across the country.



Honoring Liam

Liam was a source of light and joy to all who knew him. Diagnosed with a brain tumor (ependymoma), at just ten months, his courage and resilience were an inspiration. Despite the challenges he faced, his spirit remained unbroken, and his ability to find joy in life's moments touched everyone around him. His seven-year battle was a reminder of the urgency of finding a cure for pediatric brain cancer and improving treatments for all affected.

Scan for more information on upcoming Links Fore Liam events, or visit: www.linksforeliam.org/



#PlatesForACure

4/7 is a date that carries heavy weight. Because every single day in the U.S., 47 children are diagnosed with cancer.



This year, as a sign of solidarity for these kids, we asked our community to post a pic of their Childhood Cancer License Plates — spreading awareness about the devastating disease. Proceeds from every license plate purchased benefit The Morgan Adams Foundation.

Learn more about how to get your plate by scanning the QR code.





Morgan Adams Foundation Experiences

Not your average fundraising events

Our organization is unique in our standards. So, it's no surprise that our events are either.

MAF Cultivated Experiences have garnered incredible followings and reputations for delivering spectacular ways to engage with like-minded individuals while enjoying events crafted to the attendees' tastes. All while raising critical funds for lifesaving pediatric cancer research.

Visit morganadamsfoundation.org/all-events/ or scan the QR Code for tickets and more info!



Serve up a difference in the fight against childhood cancer! This fun-filled pickleball tournament will raise funds to research how to prevent and better treat Graft versus Host Disease, in honor of Jack Paul. **All skill levels welcome!**

Saturday, June 20
Mile Hi Pickleball



An exhilarating morning of remote-control car racing on a massive racetrack outside Children's Hospital! Teams are made up of a pediatric cancer patient, local media or sports personality, local race car driver, and one of MAF's amazing community sponsors.

Thursday, August 6
Children's Hospital Colorado | Aurora





The longest-standing charity-run Concours event in the Western United States bringing together a collection that spans all collectible eras, representing an incredible array from the earliest significant air/auto/moto to breathtaking modern jets and supercars — alongside a live and silent auction with cocktails and fare from Denver's best restaurants.

It's a spectacular, three-day weekend with events for everyone including a Road Tour, Museum Tours, Whiskey and Watches at Stranahan's, Breakfast at Tiffany's event, and family-friendly Sunday Soirée at the Polo Club.

Friday, August 21 – Sunday, August 23
Centennial Airport



The Race Against Kids' Cancer is an exciting weekend of vintage car racing, hosted by Rocky Mountain Vintage Racing (RMVR). **The races are open to the public, free to attend and offer lunch time rides on the track!**

Friday, September 27 – Sunday, September 29
High Plains Raceway | Deer Trail, CO



This series of exclusive art events will be held bi-monthly on the first Tuesday of the month, hosted at galleries, museums and private collectors' homes. **It's a chance for our artma event lovers to engage further with the Colorado art community!**

**Tuesday, June 2
Tuesday, August 4**
Various Venues Across Denver



Framed in Hope

artma reaches an all-time high!

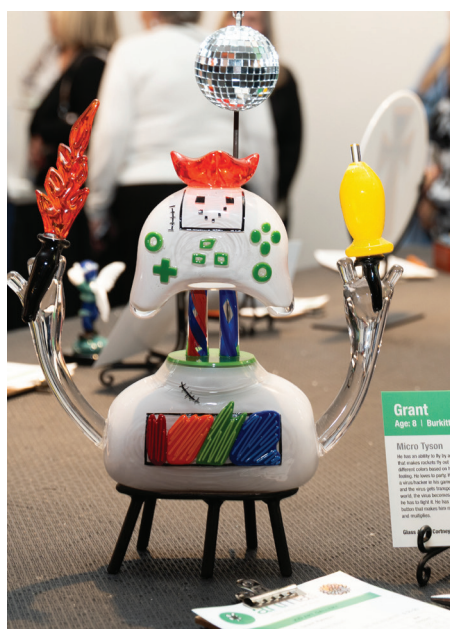
Our 25th “artma-versary” event on February 28th was nothing short of extraordinary — a vibrant celebration of creativity, community, and funding a cure for kids’ cancer that left a lasting impression on everyone who attended.

This milestone event carried special meaning. What began as a one-time, standalone gathering has grown into something far greater, helping launch The Morgan Adams Foundation and fueling a 25+ year commitment to finding a cure for pediatric cancer. That legacy was felt in every detail of the evening.

From the moment guests arrived, the energy was unmistakable. The space buzzed with conversation and curiosity as attendees explored more than 250 pieces of art. From bold contemporary works to elegantly crafted porcelain sculptures, the diversity and talent on display showcased the very best of our artistic community.



artma



Photography credits: John Flickinger – WOW Photography; Julie Kemerling – A Fine Photographer; Scott James Photography; Mike's Camera

The impact of the evening was just as powerful as the art itself.

- Thanks to the generosity of attendees and supporters, the event raised over \$400,000 for pediatric cancer research.
- In a particularly inspiring moment, the paddle raise reached record-breaking numbers, surpassing \$100,000, an incredible testament to the passion and commitment of this community.

In every sense, this year’s artma was a resounding success. It honored a remarkable 25-year legacy while continuing to push forward in the fight against pediatric cancer. We are grateful to everyone who made it possible — and we’re already looking forward to what’s ahead.

We extend our deepest gratitude to our sponsors, whose support made this unforgettable evening possible:

Minor Keele Parke, 5280 Magazine, Kitchen Distributors, Division One Construction, Mountain Safe Exteriors, Burr Law, Chubb Insurance, KOSI 101.1 Radio, Kylie Russell – LIV Sotheby’s, Children’s Hospital Colorado Foundation, Arapahoe Architects, BeSana Health & Wellness, and Jay’s Valet.