

TOURNAL

2024

FAMILIAL DYSAUTONOMIA FOUNDATION, INC



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JORNAL 2024

Familial Dysautonomia Foundation 315 West 39th Street, Suite 701 New York, New York 10018

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Dear FD Families, Friends and Allies -

It's always a pleasure to put our annual Journal together; it helps us see how much love is out there for our beloved FD community. This past year has been challenging, to say the least; the situation in Israel/Gaza continues to be of enormous concern. We send not only our thoughts and prayers to our FD families in Israel but have also provided support at their request to be sure that they have access to mental health counseling during this very worrisome time.

Given all that has been going on, we are especially grateful to everyone who took the time to offer beautiful messages and send in wonderful photos for this year's FD Journal. Between the covers of this book, it's possible to literally see the FD world materialize, with enormous care, heartfelt wishes, as well as rich memories that mark the present while also helping us build the future. The contributions that underwrite each page provide crucial support to our wide-ranging efforts to help those with FD have the best lives possible. This Journal – along with our many other fundraising activities -- mean that as a Foundation, we have been able to:

- continue to support the expert care provided by our outstanding medical team at the Dysautonomia Treatment Center in New York City;
- provide virtual meet-ups and individualized mental health support for our community;
- offer group meetings for FD family members who are eager to connect with others who understand the complexities of managing FD;
- organize and run our annual virtual FD Day in cooperation with the Dysautonomia Treatment Center, bringing hundreds of people together to learn about the latest FD research and most effective treatments, and to connect people across the globe; and
- collaborate with Rick Guidotti and his team at Positive Exposure for an incredible February 2024 solo exhibition of FD artist Peter Sonenshein at the PE109 Gallery.

The recent past has also been marked by sadness for those who have lost loved ones who lived with FD. As some of you know, my husband Fred and I lost our beloved daughter Samantha, z"I, on Sunday, September 24/Elul 29, the last day of 5782, at the age of 33. Along with others, we have had to find our way through this difficult time when it feels like there is a hole in the universe. For me, my memories of Sam's efforts to build awareness of FD are particularly comforting, inspiring me to continue in my 12th year as President of the Familial Dysautonomia Foundation. I'd like to give you a sense of Sam's commitments and her voice by sharing a short excerpt of her Dvar Torah from her Bat Mitzvah.



My Torah portion is Devarim. Moses and the Israelites are at the River Jordan. They have been wandering in the desert for forty years since they left Egypt and are about to cross into Israel. That was when God told Moses that he could not enter the Promised Land. My connection with the Torah portion is very personal. Due to my familial dysautonomia, I have a lot of challenges in my life and deal with many medical issues every day. For me, that is like wandering in the desert. I also have a dream of my own Promised Land, a world with a cure or at least ways to make life easier for me and everyone with FD. That helps me get through all the tough times, just like the idea of the Promised Land did for Moses and the Jews. I think that almost everyone has difficult times, but if they have dreams about how to make the world better, it makes a big difference. And I think that even though Moses didn't get to the Promised Land, his dream was fulfilled because he helped the next generation. To help make my dream come true, I have dedicated my bat mitzvah to raising awareness and money for the Familial Dysautonomia Foundation to support FD research and treatments. No matter what happens, it makes me feel really great that I can make a difference in the lives of people with FD, even if I don't get to the Promised Land.

Sam never got to her version of the Promised Land, but her life WAS made so much longer, better, and remarkable because of the superb care that she received from the Dysautonomia Treatment Center staff. They were always there to help her through many medical emergencies, while also serving the whole FD community, whether providing direct care or by working with families at a distance, consulting with their local medical team to be sure all involved are aware of the most effective protocols. For a condition as rare as FD, this kind of sharing of the most up-to-date knowledge is essential.

In addition to our medical staff, we have an incredible network of FD researchers who make up our Scientific Advisory Board, working on everything from new treatments for crisis, to breakthrough genetic interventions that have the potential to alter the effects of the FD mutation.

On behalf of the Foundation and its dedicated and talented staff, I want to express profound gratitude to each and every one of you for your contributions to this year's Journal. I hope we can all continue to live up to Sam's dream, one shared by so many in the FD world: to carry forward the mission of our small but mighty Foundation, founded over 70 years ago by a small group of extraordinary parents dedicated to improving the lives of those with FD.

We're all in this together.

With deep appreciation,

Talg Censon

Faye Ginsburg President.

Familial Dysautonomia Foundation





WHAT IS FD

Familial Dysautonomia (FD) – pronounced "dys-auto-NO-mia" – is a devastating neurological disease that occurs almost exclusively in people with Ashkenazi Jewish heritage. FD is inherited in an autosomal recessive manner, when both otherwise healthy parents pass on their copy of a mutated IKAP gene to their unborn child. The misread DNA "blueprint" profoundly disrupts the normal development of the sensory and autonomic nervous systems. Infants are born unable to sense information coming from inside their own bodies.

All the normal bodily functions we take for granted are gone awry in people with FD. As people with FD are insensitive to pain, they lack an essential protective mechanism and often severely burn or injure themselves. Blood pressure swings from dangerously high to extremely low. People with FD also face problems with feeding, breathing and walking and confront an uncertain future with kidney failure and blindness looming at a young age.

Crying without tears is a signature feature of FD, hence the teardrop in our "FD" logo. A world without pain and tears exists, but it is far from a carefree existence.

People with FD face unimaginable daily struggles. They must survive frequent bouts of pneumonia, uncontrollable vomiting attacks triggered by their own emotions, and there are times when the drive to take another breath fails while sleeping.

FD is a devastating disorder that causes severe physical, emotional and social problems. Nearly every major bodily function is impaired. Living with FD is a daily challenge for those who are affected as well as their families.

Features of FD include:

- Infants born with poor suck & feeding problems
- Malnutrition & failure to thrive
- Poor muscle tone (hypotonia)
- Aspiration pneumonias
- Delays in milestones like walking & talking
- Unexplained high fevers & very low temperatures
- Almost no reaction to pain

- · Inability to taste
- Episodic vomiting
- Drenching sweats
- · Uncontrollable flushing of the skin
- · Abnormal spine curvature
- Learning disabilities
- Kidney failure
- Sleep apnea
- Walking & coordination problems
- Blindness



OUR MISSION & HISTORY

The Familial Dysautonomia Foundation is a non-profit organization that was founded in 1951 by parents of children with FD who had nowhere else to turn for help and support. More than seven decades later it has grown to include chapters throughout the USA, Israel, and Canada. The Foundation is the leading source of funding for FD treatment and research and maintains the world's only FD treatment centers in NY and Israel. The Foundation raises funds and operates programs to pursue the best possible medical treatment, scientific research, public education, and social services for the benefit of those afflicted with, or at risk for familial dysautonomia.

We have established the world's only clinical research lab for the investigation of new FD therapies. Through expert medical care, patients who once had a 50% chance of reaching age 5, now have a 75% chance of living to 40 years old and beyond. While improvements in treatment and research have increased life expectancy for patients, FD remains a life threatening and heartbreaking disease with no cure.

The Foundation has funded scientific and medical research around the world. Through drug trials, animal models, and studies on blood pressure, gait, eyesight and the underlying genetics of FD, our researchers aim to eliminate the daily struggles faced by people with the disease. Additionally, one of the goals of the Foundation is to provide a continual flow of information to FD families and offer a variety of programs that include: an annual symposium for FD families; medical conferences for doctors and scientists; oxygen concentrators for air travel; and community building programs including virtual hang-out groups.

The Familial Dysautonomia Foundation is proud to have funded the research that led to the 2001 discovery of the gene that causes FD. This breakthrough enabled scientists to provide prenatal and carrier screening for FD for the first time. As a result, general population screening has become available to Jewish couples, and countless healthy babies have been born to FD affected families. In addition, we have successfully petitioned the American College of Obstetrics and Gynecologists (ACOG) to recommend genetic screening for FD, so that, as with Tay-Sachs, physicians have an obligation to offer carrier screening to at-risk patients.

Discovery of the FD gene has also opened the door to a better understanding of FD and to the development of genetic therapies that will hopefully alleviate the symptoms, slow the progression of the disease, and provide a better quality of life for those affected. It is only through the support of our generous donors that we have been able to accomplish so much. Among rare disease groups, ours is an amazing success story, but our mission will not be fulfilled until we achieve out ultimate goal: a cure for FD.



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THE FD FOUNDATION THANKS

ITS WONDERFUL STAFF

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Lareina Wang

Your Efforts Make All
Our Good Work Possible.
Thank You!



Dear FD Community,

We commence 2024 with great anticipation and are thrilled to share the latest advancements and innovations happening at the Dysautonomia Center. Our commitment to improving the lives of those affected by FD remains unwavering, and we are excited to introduce groundbreaking developments that redefine patient care.

We have embraced cutting-edge technologies to assess blood pressure in the comfort of your own home for extended periods. This innovative approach ensures a more relaxed testing environment and provides a richer dataset as patients continue with their regular activities while wearing the device.

Our aspiration to enhance the quality of life takes a significant step forward with a new modality designed to treat autonomic crises at home, reducing the need for hospitalizations. We are embarking on a clinical trial that brings the hospital to your homes, pioneering a patient-centric approach to care. Ensuring every patient receives the best medical care remains our top priority. We diligently update our care guidelines, incorporating recommendations that aid in preventing complications, and making certain that you receive the highest standard of healthcare. After decades of dedicated work in collaboration with basic scientists, we are on the verge of testing disease-modifying treatments.

Recognizing the diverse needs of our community, the Dysautonomia Center continues to offer both face-to-face and telehealth options. Whether you reside far away or face difficulties in coming to the center, we are delighted to welcome you and your families. We aim to foster a continuous connection with you, ensuring that you always feel engaged with us.

We have expanded our support groups for family and friends, fostering a stronger sense of community. Our annual art competition, a tradition during FD day, has seen an exceptional number of submissions. Evolving from a photography contest, it now embraces all forms of artistic expression and media, capturing the full range of creativity within the FD brain.

We express our heartfelt gratitude for your ongoing support. Your contributions enable the Dysautonomia Center to drive forward these innovative projects and make a tangible difference in the lives of those affected by dysautonomia.

Together, let's redefine possibilities and empower lives.

Warm regards,

Horacio Kaufmann, MD Director, NYU Dysautonomia Center Alejandra Gonzalez-Duarte, MD Associate Director, NYU Dysautonomia Center



The Team at NYU's Dysautonomia Center



Horacio Kaufmann, MD



Maria Alejandra Gonzalez-Duarte Briseno, MD



Andreana Barnett, FNP-BC



Kaia Dalamo, DNP, FNP-BC



Jose Martinez, MS



Patricio Millar, MD



Mecky Kuijpers



Lee-Ann Lugg

Exceptional personnel dedicated to the well-being of the FD community



GENETIC RESEARCH

The Familial Dysautonomia Foundation has always supported biomedical research including the quest to find the FD gene mutation, believing from the outset this would reduce the number of FD births, help us better understand the disease and open the door to possible genetic therapies.

Our first breakthrough came in the 1990's, when researchers identified a DNA marker that enabled prenatal testing for families that already had a child with FD. In 2001, after a decade of searching, the FD gene mutation was finally discovered as a splice mutation in the gene IKBKAP (also called ELP1) on chromosome 9q. This finding not only sparked a flurry of interest into how the gene functions, but also allowed general population carrier testing.

In 2004, the American College of Obstetricians and Gynecologists formally recommended that FD testing be offered to at-risk families. As a result of this push, the birth rate of FD newborns plummeted worldwide. This was the starting line in our race to end FD; but we did not stop there.

After more than 15 years of continued efforts in the laboratory and in the clinic, 2020 marked a turning point for FD research. Under the new leadership of Dr. Frances Lefcort and Dr. Adrian Gilbert, the Scientific Advisory Board (SAB) added outstanding talent from the scientific, pharmaceutical and medical fields from the US, Israel and worldwide. The re-energized SAB is now taking a more active role in stewarding the progress of prospective FD therapies through the development pipeline.

Today, the FD community of researchers are pursuing a number of promising treatments including small molecules, anti-sense oligonucleotides and gene therapy, and they are optimistic that one or more may soon be ready for testing in the clinic. We are especially encouraged that several niche organizations have shown interest in supporting our efforts to take some of these initiatives to the next level.

We are grateful for the time, talent and dedication of our SAB. With the ongoing support of our community, together we will continue to prevent, treat and hopefully one day cure FD.



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WE ARE GRATEFUL FOR THE LONGSTANDING AND LOYAL SUPPORT OF

OUR CHAPTERS

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Michael,

We adore you and love spending time with you! You are so kind, smart, funny, and brave.

All our love, Mom, Dad, Rob and Rikki, Joe, Nate, Prissy and Ricky, Elly, and Emmy





IN HONOR OF OUR THREE BEAUTIFUL GIRLS

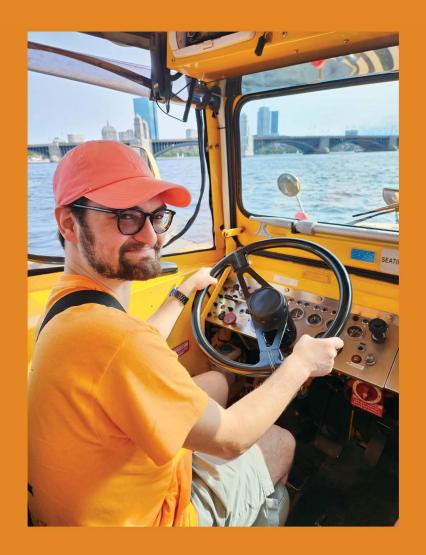
PERRY

SYDNEY

CODY

LOVE, MOM and DAD





Sammy, I cannot wait to see where you are going but please remember to keep your eyes on the road!

> Love, Grandma Soso



In loving memory of our dear mother and grandmother,

Balfura Lemer

The Byrne Family





Samantha Myers, Z″l, 1989-2022 You brought so much joy to everyone. We love you always.

Faye Ginsburg and Fred Myers



STEVEN WEXLER



Steven,

President John F. Kennedy won the Pulitzer Prize when he wrote "Profiles in Courage"

Had he Known you,

The first chapter would have been about you!

You inspire everyone you meet with your upbeat inquiring personality.

We are so proud of you and love you very much,

Mom and Dad



Wishing

Steven Wexler

a happy and healthy 2024.

Love,

Mark and Melissa Halperin



Photo Credits to Rick Guidotti of Positive Exposure

We dedicate this year's ad to our two beloved daughters, Julia and Rebecca, and to the Nurse Practitioners for working tirelessly to help the FD community.

Lisa and Jeff Newman

In Loving Memory of **Lauren Jamie Adler**

'And wherever you've gone
And wherever we might go
It doesn't seem fair
Today just disappeared
Your light's reflected now
Reflected from afar
We were but stones
Your light made us stars'

and
In honor of our amazing children
and grandchildren

Joseph, Eliana, Perry, and Sammy Adler

We love you so much!!

Vivian and Gerry Adler

MICHAEL BRENNER



THE DISNEY CRUISE WAS ANOTHER ITEM CROSSED OFF OF YOUR BUCKET LIST!

YOUR SMILE LIGHTS UP OUR WORLD.

WE LOVE YOU,

MOM AND SARAH

IN LOVING MEMORY OF

DAVID BRENNER

MAY HIS MEMORY BE A BLESSING

THE STEINER FAMILY

Best wishes to the

Famílial Dysautonomía Foundation

From

Ken Sonnenfeld

And

Peg Brivanlou

IN MEMORY



ANDREA LYNNE HAUBEN

FEBRUARY 2, 1984 - MARCH 21, 2009

FOREVER IN OUR HEARTS

The Hauben Family

IN HONOR OF OUR CHILDREN

JOSHUA

ADAM + LIZ

* * * * *

AND WE WELCOME
WITH LOVE

MADELEINE ELLE
JORDAN ISAAC

WE LOVE YOU SO MUCH
MOM AND DAD

Josh Kietz

Amy & Joseph Perella

FRIENDS OF

JOSH KIETZ



IN LOVING MEMORY

of

MY PARENTS

JULIAN and PHYLLIS LINKER

and

MY BROTHER

STEPHEN LINKER

YOUR STRENGTH, HUMOR AND LOVE CONTINUES TO GUIDE ME

~FOREVER IN MY HEART SUSAN LINKER

Steven Schwartzberg



Stevie,

From the moment you were born you brought so much light, joy and happiness into our lives. You attracted people like a magnet with your beautiful smile and upbeat personality. The way you lived your life has inspired us to be better people, more genuine, selfless and loving. Stevie, your passing has left a hole in our lives and hearts that can only be filled with the wonderful memories of the beautiful person you were. We miss you every minute of every day.

Love, Jack, Rowena and Daryl

IN HONOR

OF

BRIAN SOLOMONS AND HIS FAMILY

BEST WISHES FOR HEALTH

AND

HAPPINESS.

Love,

Jill, Sandy, Johnny and Brittany Sirulnick

In Honor of Michael Baranoff



The Ades Family Alan, Carla, Renna, Louis & Nancy

Michael Brenner

our amazing nephew and cousin.

May all good things come your way in 2024!

With Love,

Aunt Robin, Uncle Brian, Cousins Alex and Maddie.

MICHAEL BRENNER

WE LOVE YOU! YOU SMILE BRIGHTER THAN ANYONE!

AUNT ADELE & UNCLE PAUL COUSINS DANIELLE, ETHAN, HENRY & REISS COUSINS ALYSON, CRAIG, ERIC & ABIGAIL

In honor of Kaia Dalamo & Zenith Khan thank you for keeping me healthy so I can enjoy life!



Dreams really do come true!

Love, Mara Clawson & family

In loving memory of

Rachael Leah Eisenson Z"L

my best friend ever. I miss you so much.

Veronica Segal



Scott



Even though it has been six years, there isn't a day that goes by in which we don't think about you and feel your presence.

You are always in our hearts and words cannot express how much we miss you.

Love, Mom and Dad



Scott,

You have the soul of an angel, a heart of gold and a spirit that will forever remain with those who love you. You exemplify grace, courage, compassion, sweetness, love and everything that is good. Your quiet wisdom, your capacity to love and your bravery is something we will always admire and strive for. You led your life with dignity and you truly enjoyed life to the fullest. We will miss that sweet smile, but will carry it with us, always and forever. We love you with all of our heart.

Love, Rachel and Mark

REMEMBERING

SCOTT FASS

From Pepperoni to Piledrivers, and everything in between, We miss laughing, playing, and learning with you.
Your impact on us is forever.
WE LOVE YOU

Aunt Harriette and Uncle Neal Stacey and Gary Marnie and Jeff Ally, Casey, Dani and Sam In Honor of all the fabulous FD patients and their incredible families.

You have enriched my life immeasurably and will forever be in my heart.

Dr. Felicia Axelrod

In Honor of All Those Who Work So Hard In the Fight Against FD

Howard and Tova Weiser

IN MEMORY OF

ADAM SCOTT GERSON

SUSAN AND JIM MICHELLE - ALEXIS SHARI AND MIKE

IN LOVING MEMORY OF

KATHERINE MERLE IRLEN 1964 – 2006

Harvey and Barbara Irlen Kevin and Anita Irlen Steve and Jennifer Irlen

In Honor of Our Gabi

Who has surpassed all our expectations. She is now studying to be a life coach and work with her peers. As her Grandma, I'm willing to be her guinea pig.

We love you Gabi, and are very PROUD of your accomplishments.

Gma, Aunt Roni, Alexa, Milo & Thor

Welcome

Kietz Twins!

Madeleine Elle

Jordan Isaac

October 3, 2023

Love, Uncle Josh

Dear Uncle Josh,

We can't control the love we have for you and our growing Family.

Love Forever,

Grandma & Grandpa Anita & Stanley

Josh Kietz

We constantly learn from you what the really important things are:

Love, kindness, caring, family.

You are truly special.

Your accomplishments inspire all of us.

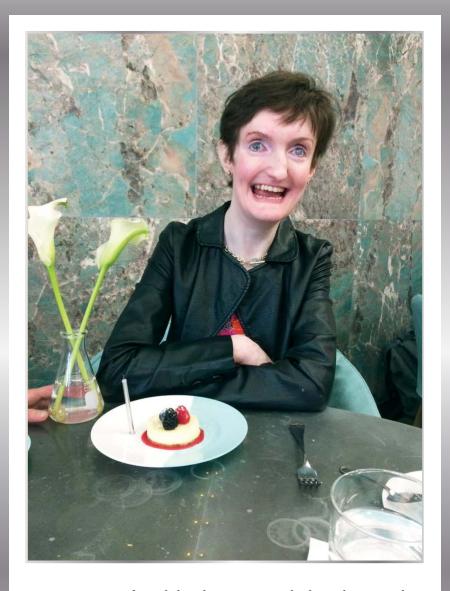
We Love you!!!!

Aunt Judi, Uncle Lenny, Shara, David, Kim, Jake



Sam,
I love seeing you with your cousins.
I am so proud of all of you!
Love,

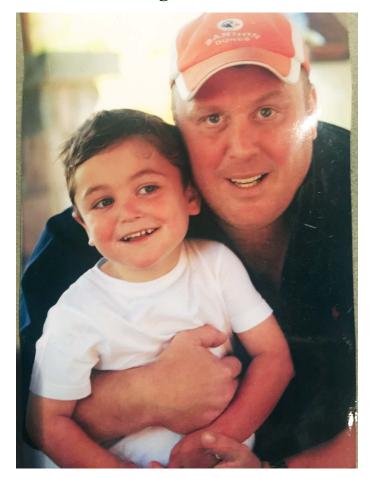
Aunt Ruth



In memory of my beloved niece, Sam, who brought joy and delight to all who knew her, and in honor of her extraordinary parents, Faye Ginsburg and Fred Myers, who do so much for the Foundation. I hold you all in love and gratitude.

Aunt Deb

Our Amazing Son & Grandson



In Memory of Adam
In Honor of Jack

Forever Loved

Gail & Stan Posnack

FOR A SPECIAL UNCLE, BROTHER AND FRIEND

BRIAN JAY SOLOMONS

WE LOVE YOU SO VERY MUCH! WE RESPECT AND ADMIRE YOUR COURAGE, STRENGTH AND CHARACTER.

Love,

Scott, Sheryl, Alex, Rachel, Josh & Ali Haberman

BRIAN SOLOMONS

AMY AND CLIFF GOLDMAN
AND FAMILY

In honor of those who fight this disease and the people who support them...

May there one day be a cure!

The Ginsberg Family
Stephanie, Ian, Alec, Wendy & Reed

IN HONOR OF

BRIAN SOLOMONS STEVEN WEXLER

and

IN MEMORY OF

SCOTT FASS

Your courage and strength inspire us. May a cure be found soon!

Adela and Mitchell Kahn

"SOMETIMES BEING A BROTHER IS EVEN BETTER THAN BEING A SUPERHERO." ~

MARC BROWN

STEVE, YOU'RE OUR SUPERHERO.

LOVE,

STEF, JAY, ELLA, & LJ BRYAN & NICOLE

Wishing Steven Wexler a great 2024

With Love and Admiration

All of the Greenlands

Steven Wexler

The most courageous and kind person we ever met.

He is truly a hero!

With love for 2024 and beyond

Carol Weissman Richard Kopelman

STEVEN WEXLER

"Strength does not come from winning. Your struggles develop your strengths. When you go through hardship and decide not to surrender, that is strength."

Arnold Schwarzenegger

Steven,

Though the hardships you have faced have certainly made you impressively strong, the positive attitude with which you approach the hardships is even more impressive.

We Love You,

The McAuliffe Family

Steven Wexler

And all his family and friends who fight this disease.

We all look forward to the day that there will be a cure!

With love,

The Zack Family

IN HONOR OF

Paul and Karen Wexler

FOR THE BOUNDLESS LOVE AND SUPPORT YOU HAVE GIVEN TO YOUR AMAZING SON STEVEN

CONTINUE THIS IMPORTANT WORK

DEBY & STEVE COHEN

Dear Lauren,

Your contagious smile, a radiant beam, In our hearts, it's an enduring dream. Everlasting love from a soul so pure, Gives us strength, helps us endure.

Life without you is never the same,

Only the memories, they do remain.

In every whisper of the wind, in every star's twinkle above,

We feel your presence, your endless love.

In our thoughts, you forever dwell,
In the beauty of stories we continue to tell.
Though time marches on, and the world spins anew,
Our love for you deepens, forever true.

With all our love,

Uncle Simon & Aunt Robin

Bracha & Oded, Chanoch & Ester, Yoel & Tali, Daniel & Yael

Elkana & Cherut, Bat Shachar, Eitan, Hodaya, Ayala, Yosef Eliyahu,

Sinai, Achinoam, Shai, Lavi, Harel, and Eden

Morgan Asinowski



Morgan's Story MMXXIII

I swear, someone moved the road!

Our understanding is that Morgan has been trying out professions for a while now. Karate, sailor, singer, and now bus driver. First it was karate but Morgan is a big guy and he kept yelling "Hulk Smash", and after a while nobody wanted to get on Morgan's bad side so, he couldn't do that for a living. Then there was sailing but the thrill in that was velling "Ramming Speed" and then aiming at the other boats, so that was not a great fit. Singing could

have worked, and Morgan still sings in a choir, but there is not much gelt in singing alongside 20 other people. No, Morgan needed something where people could see him the way we see him, as the leading man in a black and white picture show that turns to color as the show progresses, because that is his life. So, Morgan became a Bus Driver and I think the photo of him at the wheel says it all. That last turn was a little wet. But above all else Morgan is resilient so perhaps next year... Airplane Pilot!

Love from your not-so-secret admirers,

Auntie Carole, Uncle Bernie, Cousin(e)s Lauren, Sydnee, and Alex, Francis, James, and Lily

Michael Baranoff

My Dearest Michael,

You are my very special grandson. Your achievements are remarkable, including graphic designs, a great card and game player, and all around fun guy.

You're always an inspiration.

My love always,

Grandma Prissy

Michael Baranoff,

we love you!

Amy, Scott, Mia, Rachel and Jacob

In honor of our beautiful daughters



Frances Emily Natalie Rose

We are so proud of both of our daughters!!

You continue to awe us with your
determination, drive and tremendous
joy you bring to us each and every day!!

We love both of you so with all of our hearts!!

Mommy & Daddy

In honor of my two beautiful nieces and cousins



Frannie & Natalie

I'm so incredibly proud of both of you and your Extrordinary accomplishments!

Love you both!!

Uncle Ellis and Tessa

IN LOVING MEMORY OF

SCOTT FASS

YOU WILL CONTINUE TO INSPIRE US.
YOU WERE THE STRONGEST PERSON
WHO DEFIED ALL THE ODDS AND WILL
ALWAYS BE REMEMBERED.

LOVE,

BROOKE, JEREMY LUKE & BEN

Our dear Jamie,

Your life was a blessing, Your memory a treasure. You are loved beyond words And missed beyond measure.



All our love ... alwąys,

Mommy & Daddy Jessica, Matt, Leah & Emma

In loving memory of our precious Jonathan Michael Gordon

March 11, 1990 - October 3, 2008

Always smiling, feeling great and never complaining no matter what the situation was!

You are in our hearts forever!

Love,
Debbie, Daniel,
Benjamin, Shayla,
Mila, and Mason Gordon
and Edna Sydney

HONORING OUR AMAZING NURSE PRACTITIONERS,

ZENITH AND KAIA

THANK YOU FOR YOUR
HARD WORK AND DEDICATION
TO THE FD COMMUNITY.

WE APPRECIATE YOU!

THE BRENNER FAMILY



Josh!

What a year it's been! From our own private guided tour of the aeronautics museum where you work, to becoming an uncle for the very first time! So glad to have shared this year with you and can't wait to see what 2024 brings your way!

We love you to the moon and back!

xoxo aunt Amy, uncle Michael, Kayla, the Johnson family, the Walker family and the Reid family

IN HONOR OF

JOSH KIETZ

HARRY & ELIZABETH
DISIMONE

Jordan & Leslie Mayer

Honoring

Joshua Kietz

To Our Wonderful Grandson

Ezra Kress

Keep up your sunny disposition. You are the most courageous person we know.

May 2024 bring you better health and much happiness.

We love you very much,

Grandma & Grandpa

In Honor of

Ezra Kress

You have great taste in tv shows, movies, and cousins.

We love you!

Lisa, Brian, Shayna and Ari

TO SAM LANDAU

WE LOVE YOU

SUSAN AND ARNOLD SCHARF

IN EVERLASTING MEMORY OF OUR BELOVED DAUGHTER

AMY JILL LEHRER

AND HER BELOVED FATHER & MY DEAREST HUSBAND

DONALD LEHRER

IN LOVING MEMORY OF

PHYLLIS, JERRY, AND STEPHEN LINKER

JEAN AND STEVE ANREDER

In Loving Memory of

Evan Rosenthal

Your bright smile lit up our lives.

Sweet memories of you continue to be a blessing.

You are always in our hearts.



With love,



Debbie and Joan and Stephanie and Susan and our families

IN MEMORY OF:

THE ROSENTHAL MEN

RICHARD DAVIS ROSENTHAL EVAN ROBERT ROSENTHAL SETH ÅNDREW ROSENTHAL

ALWAYS IN MY HEART,

LOREN ROSENTHAL

IN HONOR OF

BRIAN SOLOMONS

YOUR COURAGE AND STRENGTH
ARE AN INSPIRATION

LYNDA AND HAL KATZ

In Honor of

SHERYL AND SCOTT HABERMAN

RANDY & JAY RAUBVOGEL

IN HONOR OF

BRIAN SOLOMONS

AND

STEVEN WEXLER

WISHING YOU THE BEST OF HEALTH AND HAPPINESS ALWAYS!!!

JOANIE & DON FISHER

In support of our friends,

The Habermans

and

The Wexlers

it is our great pleasure to contribute to the amazing work and the dedicated people of the Dysautonomia Foundation.

Karen and Steven Hess

David and Peter Sonenshein



We are thrilled to be able to see so many of Peter's beautiful paintings at David and Jennifer's home and are delighted to have several of his paintings in our home. We are grateful that he has survived Familial Dysautonomia for so long and that he is very active despite his health problems. It is always fun to spend time with Peter, his parents Jennifer and David and his sister and her family (Emily, Judah and children Sam and Ari), although we live so far away that we can't do it often enough.

With love - Peter's Aunt Gail and Uncle Linc, Cousin Dina, Cousin Adam and his wife Jenna and their children Samantha and Jack

Peter Sonenshein

Congratulations
On your
Wonderful
Art Show



Leslie and Warren Noel and Bruce

In Honor and Memory of

Harvey Rosenzweig

and his Strong and Deep Love of

Steven Wexler.

The Brauns

We are Honored to Support

Steven Wexler

And The

Familial Dysautonomia Foundation

Mark & Kimberley Elie



K&E Travel Gives Back

IN HONOR OF

STEVEN WEXLER

FROM
MICHAEL & AMY
KOPELMAN

IN HONOR OF

STEVEN WEXLER

AND THE WEXLER FAMILY

LESLIE AND MICHAEL ROSENBERG

Steven-

We are so proud of you!

You are such an awesome person!

Love you-

The Segal Family

Charlie,

Words can not express the effect that you have on Steven.

You are a major reason that Steven is able to enjoy life as he does today.

Thank You from the bottom of our hearts.

Love, Karen and Paul

Joy,

Thank you so much for being Steven's night owl.

Without you we would not be able to re-energize and make it through the year.

You have been part of the Fire Department for as long as we have known you.

You have been saving our lives for just as long.

Love, Karen and Paul

In honor of the Adler Family

WE ARE PROUD TO SUPPORT THE FOUNDATION'S WORK

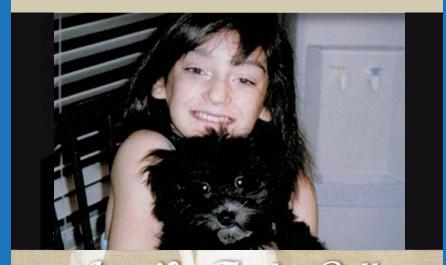
REYNA and PIERRE GENTIN

In Honor of Our Good Friends

Vivian & Gerry Adler

In recognition of all the work they do for the Foundation.

Bonnie and Russ Mannis Caron and Steve Gelles



Jennifer Taylor Bell
Time Goes on but Memories Live Forever
Always in our Hearts













Rachael, We miss you. You will always be with us and will always be remembered.

We love you.

Frannie, Gabi, Perry, Rebecca, and Sarah

In Loving Memory of

Scott Fass

Caring • Selfless • Smiling • Adorable
Thoughtful • Brave Warrior
Superhero • Angel

You are missed every day

Aunts Adele Fass and Barbara (Fass) Young



In Loving Memory of our wonderful cousin

Scott Fass

We miss you very much.

Cousins Sue Lipkowitz, Marcel Lipkowitz and Robert Fass In Loving Memory of

Scott Fass

Always Our Teacher

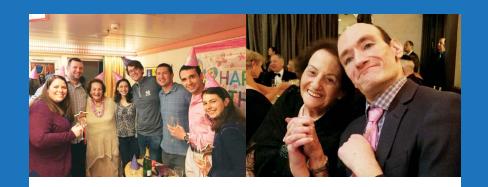
Sandy and Sandy Goodman

In Loving Memory of

Scott Fass,

a very special young man, who brought joy into the lives of all who knew him.

Sue and John Korn



In Loving Memory Of Maryon Weill Ea Scott Fass

Your courage and dignity in the face of adversity is an inspiration to all of us.

Forever you will remain in our hearts.

We miss you both.

Elena, Barry, Andrea, Steve Matt, Kelly, Joey, Marissa, Andy, Jon, and Charlotte



Joseph P. Navarra, R.Ph., FACA Owner joseph.navarra@towntotalcompound.com Town Total Compounding Center 415 Crossways Park Drive, Suite B Woodbury, NY 11797 t. 516.249.7436 x122/ f. 516.249.7437 www.towntotalcompound.com To honor the memory of

Benson and Pearl Ginsburg:

Truly a couple for the ages



Mark and Susan Hamilton

IN MEMORY OF OUR SON & BROTHER

ELLIOT LOUIS GOLDBERG

JUNE 1973 - JULY 2020

ALWAYS IN OUR HEARTS, FOREVER AN INSPIRATION, HE TURNED STRUGGLES INTO STRENGTH.

WITH LOVE,

THE GOLDBERG FAMILY

In loving memory of

Barbara A. Gould

Miriam K. Gould

Richard H. Gould

And with eternal gratitude

For all you were

Judith E. Gould

THE PAUL NORMAN GOULD MEMORIAL FUND

IN LOVING MEMORY

OF

SANA GOULD

and

DAVID GOULD

LOVING PARENTS, GRANDPARENTS and GREAT GRANDPARENTS

STEVEN GOULD and FAMILY

MARSHALL GOULD and FAMILY

THE PAUL NORMAN GOULD MEMORIAL FUND

IN LOVING MEMORY

OF

CAROL SUE GOULD

WE LOVE YOU AND MISS YOU

STEVEN GOULD and FAMILY

MARSHALL GOULD and FAMILY

THE PAUL NORMAN GOULD MEMORIAL FUND

IN LOVING MEMORY

OF

PAUL NORMAN GOULD

HE LIVES IN OUR HEARTS FOREVER

STEVEN GOULD and FAMILY

MARSHALL GOULD and FAMILY

Remembering Lisa & Bob Gross



M With Lots of Love M
M

Bobbi, Jason & Kevin

In Loving Memory of Our Son and Brother David Halle



After 63 years, we remember and miss you,

Ruth Halle, Esther Marcovici, and Linda Halle

And the rest of the family who wishes they knew you,

Richard, Rachel, Gabriel & Devin, Ben & Annie

Michael & Jack & Sloane

GABI,

30 years old! It feels like only yesterday that you were born. You've grown into an amazing person, a wonderful life coach, and we're just so proud of you. Not our own words, but this is the journey you've helped us to see as our own life coach:

"Behind You: all your memories
Before You: all your dreams
Around You: all who love you
Within You: all you need
Everything is possible, believe in yourself"

Mom & Dad







Mitchell, Always Loved Never forgotten!











In Honor of my Grandson

EVAN KAPLAN

Love,
Grandma Nancy
(and in memory of
Grandpa Bobby)

FOR MY GRANDSON,

JOSH KIETZ

TO THOSE WHO

WORK SO HARD

TO HELP US

DREAM THE POSSIBLE

DREAM...

FOR MY GRANDSON, JOSH

AND ALL THE FD KIDS...

THANK YOU SO VERY MUCH.

WITH LOVE AND HOPE,

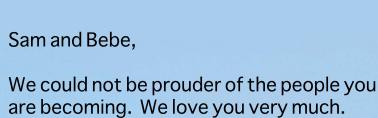
GRANDPA AL AND GLORIA

Dear Ezra,

You are the most amazing, sweetest, bravest, most epic young man we know.

We love you!

Aunt Sherry and Uncle Ira



Mom and Dad



IN MEMORY OF AUNT BESS

WE WILL ALWAYS LOVE YOU MARC, BRETT, AND BROOKE

WE ARE SAD THAT WE NEVER HAD THE OPPORTUNITY TO KNOW YOU.

LOVE YOUR GRAND NEPHEWS AND NIECE

ALEX, CARLY, JACKSON, AND RYDER



Rebecca,

You are our ray of sunshine! We love your optimistic attitude and adventurous spirit.

We love you so much, Rich & Sue

In Loving Memory of

Bennie & Zak



Richard, Joyce, & Samara Rosen







In Loving Memory of Evan Rosenthal January 27, 1983 - July 17, 2007

"There are stars up above
So far away we only see their light
Long, long after the star itself is gone.
And so it is, with people that we love,
Their memories keep shining, ever brightly
Though their time with us is done.
But the stars that light up the darkest night,
These are the lights that guide us...
As we live our days, these are the ways we remember ~
We remember."

Memories of our Evan shine brightly every day, and we pause to remember our wonderful friend and courageous young man!

You are Forever in Our Hearts

Ronnie and Bob Powers Lauren, Daniel, Fiona, and Aidan Powers Julie, Josh, Abigail, and Adam Lurie



THANK YOU, RENI
FOR BEING YOU.
MORE THAN ONE LIFE
HAS BREATHED EASIER
BECAUSE YOU HAVE LIVED.
WE TALK ABOUT YOU
REGULARLY AND YOU WILL
ALWAYS REMAIN A PART
OF OUR FAMILY.

renette mérose shafiq-berger

October 8, 1952

October 28, 1984

We are thankful for all the loving things Reni has given us, the memoires of innocence, sweetness and strength; her thoughtful ways, her gentleness, her caring and giving nature.

Reni was deeply devoted to those she loved. She added to the world these precious qualities – and they will be missed.

TAMAR JACOBS ADDY AND AURIANNE

In honor of **Brian Solomons,** an inspiration to us all.

Love, Mom



In loving memory of **David Solomons**

The Solomons and Haberman Families

In Honor of

Brian Solomons

Sheryl and Scott Haberman

We honor and support Your hard work, devotion And dedication to the FD Foundation.

With Love,

Deborah & Joel Brooks Sharon & Michael Tyner

We love you, Steven. You make us proud every day.

Love, The Diton Family and The Wealth Alliance



Eric Diton, CIMA*
President/Managing Director
105 Broadhollow Road
Melville, New York 11747
(631) 670-0702

Florida Office: 6501 Congress Ave. Suite 100 Boca Raton, FL 33487 (561) 910-8626

Experience the Future Today

In Honor Of

STEVEN WEXLER

"A Real Life Superhero"

Best wishes for a happy and healthy year.

THE ESTEROW FAMILY

Dear Steven:

We truly admire your courage and perseverance. May this year bring only good things... health, happiness and a bright smile each and every day!

> Love, Mark and Rachel

IN MEMORY OF

ERIC GOTTLIEB

LOVE, THE WEXLERS In Memory of

Eric Shapiro

Love, The Wexlers In honor of our wonderful daughter and sister

SARAH ZUCKER

Love, Mom, Dad, Josh, and Shelly (she's the furry one)



Many thanks to the entire FD team for all you do!

IN HONOR OF OUR GRANDDAUGHTER:

SARAH ZUCKER

How to describe her?

Amazing: what she has accomplished; her cheerful attitude (even when confronted with challenges); her loving disposition (family, friends, animals-especially dogs, chocolate chip cookies), her sense of humor, her thoughtfulness.

While we could come up with many more adjectives we won't (we couldn't stop - intelligent, compassionate, empathetic, considerate).

Best is she loves us almost as much as we love her.

Grandma and Grandpa

In memory of Zoe's and Sydney's brother

ZACHARY

With love, Steven, Naomi and Sam

IN MEMORY OF

IRENE WOHL BOCHNER

FROM THE BOCHNERS AND GOLDSMITHS

In Loving Memory of

Zachary Michael Bopp



We love and miss you
Grandma Ruth Halle,
Dad and Linda

In Honor of

Michael Brenner

Nicky, Priscilla and Harold

In Loving Memory

To our most beautiful wonderful girl **Falon Bromberg**

> From Mom, Dad & Phílíp

In Loving Memory of

Falon Bromberg

Soaring With The Angels Now

Love, Aunt Susan, Uncle Ira, Jason, Bridget, Rose & Caroline

IN HONOR OF

FRANNIE COHEN

OUR INSPIRATION

THE RAFFLER FAMILY

In Memory of

JILL COPLIN

Beloved Daughter and Sister 1966-1977

DIANE COPLIN LIPSITZ

Beloved Wife, Daughter, Sister and Aunt 1968-2010

The Coplin Family

IN LOVING MEMORY OF

JILL ANNE COPLIN

AND

DEDE COPLIN LIPSITZ

THE HOMER FAMILY

In Loving Memory of

Maximiliano and Lionel Donzis

Graciela, Carlos, Sebastian, Tracie, Sienna, Greyson, Ashton, Hernan, Melissa, Samuel & Reid Donzis

Forever Remembered, Forever Missed

Scott Fass

Geri, Harold, Jared, Mariel, Blake, Cori, Alex, Lexi & Bernie

In tribute to:

All those with FD and their loving families Our dedicated board, staff and volunteers The devoted NYU Treatment Center team Our brilliant Scientific Advisory Board

And with special recognition to:

Our Nurse Practitioners

For the Excellent Care they Provide 24/7

I am inspired by all of you and I'm proud to be part of the FD community

Lanie Etkind, Executive Director

In Memory of Scott Fass and his FD friends

To my friend and all-star neurologist **Dr. Alejandra Gonzalez Duarte**

I could not be more proud of your leadership and strength at the Center. I know the patients are in the best hands. I cannot think of a more thoughtful, thorough, and well trained doctor to provide care. I know you will take the research torch and do wonderful things. You are the very best!

Lucy Norcliffe-Kaufmann



There is a cast of **Super Heroes** walking through the streets of New York and they are not from the Marvel movies. Their "**Super suits**" tend to be **scrubs** or **lab coats**, not spandex or armor. They are larger than life, at least in our eyes. Individuals come and go but they become your friend, confidant, and crisis manager. The Nurse Practitioners at the NYU FD Center guide you through the maze of each yearly visit.

The most recent **Dynamic Duo** of Zenith Kahn & Kaia Dalamo have stood on the shoulders of their predecessors & have reached farther than ever to support the FD community. For this we are grateful & thankful! Wishing Zenith success in her new endeavor! Welcome to Andreana Barnett, the newest superhero on the team.

Florence Fried, Nelson & Morgan Asinowski

IN MEMORY OF

SY FEIT



SANDY Z''L AND LARRY GREEBEL

IN LOVING MEMORY OF MY NIECE

JAMIE GOLDBLAT

GAIL GOLDBLAT

In Honor of Evan Kaplan



EVAN
We are so proud of you and love you very much.
XOXO

In Honor of Tio Josh -

the Best Uncle in the World!!

Love, Maddy and Jordan Kietz Dear Josh...

To the finest "drum tech" in the world...

We love you!!

Aunt Ronni Uncle Phil Brian, Jacob, Claudia and the "Wiggler"

JOSH KIETZ IS:



WE LOVE YOU!
Aunt Pam, Uncle Jordan
& Cousin Lauren

In Honor of

JOSH KIETZ

Perri & Leo Klingbaum

We love, respect, and admire

Josh Kietz And his family

The Tortorellas

In Honor of my grandson

Mitchell Kofsky

and in recognition of

The Dysautonomia Foundation

Sandra Kofsky

With tremendous love for our grandson

EZRA KRESS

We celebrate your 26th birthday with joy, inspired by your spirit, love, and courage.

Saba and Savta, Bill & Beverly Lebeau

In Honor of the amazing and accomplished

Rebecca Newman

With love and admiration, Cousin Wendy

In Loving Memory of My Sister

Carly Allison Posner

In honor of our warrior princess,

Michelle Rappaport

and in gratitude for the

and in gratitude for the Dysautonomia Center for everything.



All our love, Mom, Dad, Jessica, & Aspen

DEBBIE SCHOENHOLZ

May 29, 1957 - January 24, 2024

The Dysautonomia Foundation really helped Debbie enjoy life.
Kaia Dalamo NP.

Dr. Gonzalez-Duarte, and Dr. Millar maintained communication with the Grandell Nursing Facility and did their best to help Debbie.

Debbie, you are a true star who made life as pleasurable as possible. We will miss you so much.

God bless you.

The Schoenholz Family



For his determination & courage

We honor

Brian Solomons

Mark & Joan Haberman

IN HONOR OF

BRIAN SOLOMONS

PERRY AND PENNY BERGER
AND FAMILY

In honor of **Brian Solomons and family**

with best wishes for a happy and healthy new year.

Love, The Siskind Family

In Honor of
Brian Solomons

Steven Wexler

Sheree & Jeffrey Markowitz Bette & Jeffrey Rubin

In Honor of

Peter Sonenshein

Friend, Artist, Inspiration

With love,

Marsha & Steve Dickstein

In honor and celebration of our incredible friend

PETER SONENSHEIN

With love from the Hoxter-Levine family

Debi and Ron Andrew and Heather Deanie, Gary, Sylvie and Teddy

IN HONOR OF A WONDERFUL FRIEND AND TALENTED ARTIST

PETER SONENSHEIN

WISHING YOU THE BEST ALWAYS...

FERN AND STEVE ROTFELD

In honor of our granddaughter

Símí Steiner

who fills our lives with music and love.

"If music be the food of love, play on."

Shakespeare, Twelfth Night

And the OSCAR goes to... **STEVEN A. WEXLER**

For BEST LIFE LIVED with PERSEVERANCE, KINDNESS, HUMOR and JOY!

With GRATITUDE from the ACADEMY & your #1 FAN

Charles Donovan

In Loving Memory of Lauren Adler Karyl and Asher Miller

In memory of our loving sister, **Deborah Begin**In memory of our loving brothers,

Dov and Yaakov Begin

All who suffered greatly from Familial Dysautonomia.

May their memories be a blessing to all who knew them.

They are greatly missed and forever in our hearts.

IN LOVING MEMORY OF GREGORY ASHER BRANDT

ALWAYS IN OUR HEARTS.
MOM, JASON & BENJAMIN



In Honor of

MICHAEL BRENNER

COUSINS LINDA & BOB



Judy Fettman Dreyfus

1970 - 2020

We miss her terribly.

Love, Mom & Dad

To my great friend Rachael Eisenson.
Think about you all the time.
Miss all our fun times together.

Love Always

Andrew Sigman

In loving memory of our dear cousin **Felicia Gail Gold**

And with gratitude to the devoted doctors and staff of the Dysautonomia Foundation

Joyce and Bern Saxe

In Honor of

Gabí Jassíe

With Love, Suzie

In Memory of Louis Kaley Roslyn Kaley Alisa Robin Kaley

Thank you to the staff of the FD Center for your care and support.

Stefanie Kaley Linakis

For Our Wonderful Cousin Josh Kietz!

Sending Big Love from Massachusetts!!

The Maimans and The Wanzers



Josh Kietz, You are our superhero! Love, Hal, Carole, Greg, Emily, and Maddy

and Frankie too!

Created by Mahmure Alp from Noun Project

Josh,
We wish you the best of health and joy in the upcoming year. Keep on rockin'!
The Glaser Family



IN HONOR OF JOSH KIETZ

LAW OFFICES OF HARRY WEINBERG, PLLC

ONE PENN PLAZA, SUITE 2110 NEW YORK, NY 10119 (212) 889-4100 LABOR AND EMPLOYMENT LAW

IN HONOR OF
EVAN WHITE
AND
EZRA KRESS

DAVID & BETTY ROTH

Rebecca Newman,

You are an inspiration to us all!

Sending lots of love your way, Eric & Julie, Josh, Lisa & Tony, Jeremy and Aliza

> Best Wishes to The Newman Family

> > Phyllis Honig

IN MEMORY OF ELLEN SUE NUGER

THE NUGER & WEINROTH FAMILIES

IN MEMORY OF

DR. ROBERT PORGES

EUGENE LIND



In Honor Of Jack Posnack

In Honor of Jack and Robin Posnack

You are both heroes and role models for the rest of the world. The time and dedication and love is endless. You both make this world a better place.

> With love and adoration always, Jessica and Mitchel Weiss



Chanie, we will always love you to the moon and back! You made us laugh, smile, and warmed our hearts with your kindness. We miss you everyday!

Love, Mommy, Daddy, Devorah & Gershie

SAM SERNOVITZ

We are so proud of how much you've grown and how you've approached the opportunities and challenges that come during the teenage years. We can't wait to see what is in store for you in the year to come and are grateful for your continued good health and the blessings that you bring to us and everyone who knows you.

Love, Mom, Dad, Daniella, Eden and Peaches

Ben Solomon,

you will always be our daily SUNSHINE!!!
1976-1999

With love, Mom and Adam

In Honor of

Brian Solomons

We are proud to support the FD Foundation

The Bochner Family

In Honor of

BRIAN SOLOMONS

SHERRIE AND MITCH MAYRSOHN

BRIAN SOLOMONS & STEVEN WEXLER

WITH FRIENDS SURROUNDING YOU, THERE IS ALWAYS HOPE!

ELLEN & BRUCE BELSKY

IN HONOR AND CELEBRATION OF PETE SONENSHEIN AND HIS GREAT FAMILY.

WITH LOVE, ELLEN AND STEVEN WOLF

In Honor of Steven Wexler

DRS. AJAY & SHASHI BERDIA

Wishing you great health and much love, Steven.

Love, The Gewirtz Family

In Honor of **Steven Wexler** and his Family.

Dayle and Carl

With continuing love and respect to the entire Wexler Family

-Ellen Kessler

In Honor of the Wexler Family Marvin & Wendy Natiss

LISA AND LON GOLDSTEIN



Peter Luftig

D: (516) 726-2635 C: (516) 476-7840 F: (516) 829-5857 pluftig@scsai.com

1981 Marcus Avenue Suite 125 Lake Success, NY 11042

Friends in Honor Of

Josh Kietz

Bauman + Krasnoff LLP
Matthew Bernardini
Meme and Art Peponis
Bonnie & Brian Roth
Marci and Dan Settle
Dawn Zappetti & Patrick Sullivan
Anonymous Friend

Rebecca Newman

Laury Hamilton Herbert Levitz Edith Rubin

Jack Posnack

Andrew & Jill Feldman

Brian Solomons

Richard Allerton & Catherine Stewart
Rona Cohen
Andrea & Jeffrey Lomasky
Pace Cares
Richard & Robin Reubenstone
Alex & Dawn Seaman

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Steven Wexler

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Lauren Adler

Laura and Eric Green

Mitchell Joseph

Carol & Richard Engelson

Philip Wexler

Mr. & Mrs. Monroe Jacobson

Boosters In Honor Of

Morgan Asinowski

Florence and Nelson Asinowski Grandma Goldie Asinowski Bubby Jenny Fried Jennie Gindin Sonia Elisha and Sol Shimshi

Michael Brenner

Alyson, Craig, Eric, and Abby Brener

Kaia Dalamo and Zenith Kahn

Elise & Robert Goldblat Robert Ross

Josh Kietz

Noel Ayala
Phil & Dore Handy
Jane Helfman
Staci and Jeff Kirschner
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Shelly and Wendy Nussbaum
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Ezra Kress

Renee & Hervey Sande

Rebecca Newman

Lucille Newman & Family Stacey & Peter Reichman

Sam Sernovitz

Betsy Newberry and Ken Friedrich
Debra & Arthur Skaroff

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Fern & Jeffrey Bernstein Michelle & Lennert Gruszecki Gladys Hymowitz Vanita & Jack Solomons

Peter Sonenshein

Ernest & Jane Dellheim

Steven Wexler and Family

Ira & Risa Bellach
Mr. David R. Bernstein
William & Ellen Enco
Mark & Gail Fialkov
Bruce Goldstein
Carol Gottesman
Marc & Suzie Harvey
Piper Lutbak
Stuart Plotkin
Debra & Michael Resnick
Jennifer Signet

Boosters In Memory Of

Florence Lewis Baker
Barbara & Peter Sperber

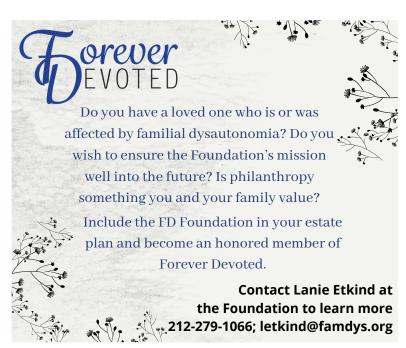
Jamie Goldblat
Rich & Karen Heller

Mitchell JosephRobert & Catherine Kushner

Samantha Myers
David and Susan Cohen

David RosenbergCarla & Jason Rosenberg

Naomi Walker Ben & Ruth Walker







RACHAEL EISENSON

7/1/88-8/6/23





JORDAN ERIC SCHERWIN

3/3/77 - 2/9/24





DEBBIE SCHOENHOLZ

5/29/57 - 1/24/24





ROXANNE STEIN

4/27/59 - 12/19/23





BENJAMIN TARYAN

4/14/93 - 12/14/23



FAMILY AND FRIENDS



DICK FRANKLIN
ALYCE JOAN FREEMAN
MARK HABERMAN
EILEEN GOLD HARRIS
TOVA ISRAELI
SYLVIA JASSIE
HARVEY ROSENZWEIG
MEL SACHS
SYLVIA SHANDLER
HARRY WEISER
JUDY WENER

To all who contributed to the 2024 Journal

THANK YOU!

Through your kindness and generosity, the FD Foundation will continue to provide vital medical care, scientific research, public education, and social services for the FD community.



IN HONOR OF STEVEN WEXLER

ALL OUR LOVE,

Rena, Josh, Elana & Jacob Kopelman