MTS Sickle Cell Foundation, Inc. closed out its first year leading and being of service to the sickle cell community in Georgia and around the world. From sponsoring 26 kids suffering from SCD to sleep-away camp to raising funds for a family going through bone marrow transplant, we used our voice and platform to not only raise awareness about sickle cell disease, but provide much needed support and services to patients and their families.

The following pages highlight our 2016 efforts and set the stage for what’s to come.

OUR MISSION

MTS Sickle Cell Foundation, Inc. spreads awareness of sickle cell disease and provides moral and financial support to families affected by SCD.
This foundation was born out of a real need to see more support services provided to sickle cell patients and their families, while continuing to use our life to spread much needed awareness about sickle cell disease.

As a mother of three daughters who all suffer from this disease, I did not want families like ours, who struggle so much in and out of the hospital, to have to struggle for basic support services.

For the last year, it is my belief that our success has been a combination of several factors. The first, of course, is the quality and appropriateness of the services that we have made available to the families we serve. Other factors include the ability to empathize and connect with our families on a truly personal level, a genuine zeal to work hard to provide much needed support services, and a spirit of togetherness in working with others in the SCD community.

Ultimately, it is those who donate of their time, talents and finances who are our most valued resource. Quite simply, it is our donors who define our capabilities and our donors of whom I am most proud. It is solely because of their generosity that we have been able to set ourselves apart from other organizations in such a short amount of time. I believe this is why MTS Sickle Cell Foundation is one of the most go-to organizations for sickle cell families in Georgia.

"I want all sickle cell patients to live a full life of dignity and respect, while having access to the support services they deserve. To not be stigmatized, bullied, or discriminated against because of their illness. Until we can educate the public, that will never happen, and we have been doing just that. But our mission is not complete."

Mapillar Dahn
FOUNDER/CEO
MTS SICKLE CELL FOUNDATION, INC.
"MTS Sickle Cell Foundation, Inc. provides an invaluable service to our sickle cell families in providing resources, support, and promoting the best interest of the sickle cell community. Sickle cell communities all over the world need more organizations like MTS."

PATRICIA WATERS
HEMATOLOGY NURSE, CHILDREN’S HEALTHCARE OF ATLANTA

HOSPITAL VISITS
Visited patients in the hospital and took care packages.

PARKING PASSES
Provided parking passes for families visiting CHOA*.

FAMILY SUPPORT
Raised $549.03 for Jahzara Thompson’s transplant fund.

FAMILY PHOTOS
Hosted a photo shoot for family photos for SCD families.

CAMP SPONSORSHIP
Sponsored 26 campers and printing of t-shirts at Camp New Hope.

TOYS GIVEAWAY
Hosted Holiday Breakfast/Toys Giveaway in partnership with Toys4Tots and Sickle Cell Matters.

* CHOA - Children’s Healthcare of Atlanta
FEB 1 - FOX5 ATLANTA FEATURE
Fox5 Atlanta shares the story of Amatullah, Khadeejah, and Hajar, three sisters who all battle sickle cell disease. - https://bit.ly/2NogZd8

FEB 14 - PHOTO SHOOT

JULY 17-22 - CAMP!

APRIL 10 - BALLIN DOLLZ DONATION
MTSSCF was presented a $300 donation by The Ballin’ Dollz Kickball League at their opening day ceremony. https://bit.ly/2KKClh2

SEPT 3 - KICKBALL!
https://bit.ly/2z8z9Ne

SEPTEMBER 8 - WELLS FARGO COMMUNITY GRANT

SEPTEMBER 10 - WOMAN OF INFLUENCE AWARD

SEPTEMBER 15 - LIBERIAN OBSERVER FEATURE

DEC 24 - BREAKFAST AND TOYS GIVEAWAY

MARCH 25 - STEPATHON

AUG 17 - RED CROSS
MTSSCF founder speaks at the U.S. Citizenship and Immigration Services luncheon about the importance of blood donation. - https://bit.ly/2lUlcZx
ORGANIZATIONAL FINANCIAL OVERVIEW

TOTAL EXPENSES FOR 2016
- 62% Programs
- 27% Management & General
- 11% Fundraising

HOW WE USED OUR RESOURCES
- $7693.57 Total Expenses
- 59% Programs
- 27% Management & General
- 3% Program Support
- 11% Fundraising

WHERE WE USED OUR RESOURCES
- 48% Camp Sponsorship
- 47% Family Support
- 5% Events
behind every great organization...

OUR OFFICERS

MAPILLAR DAHN, FOUNDER/CEO
WILLIAMENA DAHN, TREASURER
QIANA L. CRESSMAN, STRATEGIC PARTNERSHIPS
TERRY R. CRESSMAN, CHAPLAIN

WE ENVISION A WORLD WHERE ALL SICKLE CELL
PATIENTS CAN LIVE A FULL LIFE OF DIGNITY
AND RESPECT, WHILE HAVING ACCESS TO
THE SUPPORT SERVICES THEY DESERVE.

△

BUT WE CANNOT

do it

HELP US IMPACT
THE LIVES OF
HUNDREDS OF
THOUSANDS OF
SICKLE
CELL PATIENTS
AND THEIR FAMILIES

LEARN ALL ABOUT
HOW YOU CAN GET
INVOLVED AND MAKE
YOUR GENEROUS
CONTRIBUTION
TODAY AT

MYTHREESICKLERS.ORG

11030 GENOVA TERRACE | HAMPTON, GA 30228
404.925.4369 | INFO@MYTHREESICKLERS.ORG
MTS Sickle Cell Foundation was proud to have been nominated by the wonderful staff of the Lovejoy Branch of Well's Fargo to receive a $1,000 grant during the Well's Fargo Day of Giving on September 8th at the Chick-fil-A College Football Hall of Fame. Pictured above: Founder Mapillar Dahn and Branch Manager Adrian Moore.

**PLATINUM - $1,000 +**

Mapillar Dahn  
Larry J. Stephens  
Williamena Dahn  
Wells Fargo  
Sample Czar

**GOLD - $500 - $999**

Sheri Crawley, Pretty Brown Girl, LLC  
Terry and Qiana Cressman
MTS Sickle Cell Foundation, Inc. was presented a $300.00 donation by The Ballin’ Dollz Kickball League at their opening day ceremony on April 10, 2016 at Rum Creek Park.

**SILVER - $100 - $499**

Antonio and Anton Ford  
Spencer Clarke  
Ballin’ Dollz  
Christopher Davis, Sr  
Erica Williams  
Nia Colton  
Robert Garguah, Jr., ME1C2/LILFOC Foundation, Inc.  
Desmond Thomas  
Ebony Rice  
William and Bernice Dahn  
Susannah Tweh-Tarr  
C.T.C Events, LLC  
Fernanda Loconsolo  
Juanita Perry  
Edward and Kema Gaye  
Kirk Faust  
Jacob and Adriana Ortega  
Sandra Vanessa
WE LOVE OUR DONORS!

BRONZE - $50 - $99

Checago Bright-Sawo, The Checago Bright Foundation
   Joseph Bolay
   Samantha Barnum
   Whitney DeBoe
   Anna Echols Carlton
Charles and Charlene Dunbar, suakoko betty
   Clarewen Best
   Daniel Tor
   Deane Fraser
   Deye Gonyon
   Gus Kormah II
   Kafuba Donzon
   Kevin Strakal
   Kunneh Dickson
   Mr. Blitch
   Nicole Jenkins
   Tammie Keith, Red Cross
   Tarseh Manthan
   Zhane Hall

SUPPORTER - BELOW $50

   Benetta Vincent      KaTrina Woodson
   Lincoln Ward         Lynn Michelle Vittitoe
   Aretta Baldon       Shirley Jones
   Bridget Smith       Stacy Patillo
   Danielle Williams   Tish Sutton
   Melissa Ferguson    Antar Austin
   Mira Evans          Antoinette Faison
   Crissinda Ponder    Brandy Lee
   Amelia B. Wheeler   Cherish Abu
   Calvin Lea          Dana Williams
   William N. Dahn, Jr  Eullanda Broome
   Bartum Kulah        Felecia Holmes
   Carole Wooten       Jahannah Christian
   Clarice Ford-kulah  Linda Phillips
   Desiree Roberson    Missey Bowens
   Karen Koukou Twaglee Nikita Lee
WE LOVE OUR DONORS!

**SUPPORTER - BELOW $50**

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MTS Sickle Cell Foundation, Inc. is grateful for all of our supporters, donors, sponsors, volunteers, and friends who made our 2016 programs, initiatives and events a success. **THANK YOU SO MUCH.** A special thank you to the compassionate individuals, businesses, and foundations listed here who gave generously from October 22, 2015 to December 31, 2016. You all seriously rock and we look forward to working together in 2017 and beyond. We have made every effort to ensure the accuracy of our lists. If we have misspelled or inadvertently omitted a name, please accept our apologies. Call the MTS Sickle Cell Foundation at 404-925-4369 with corrections.