

Jacy's Wagons

Turning a Test into a Testimony One Wagon at a Time

Note from Jacy:

October has been an awesome month! We have hit and surpassed our goal of 150 and we keep growing everyday! We took many trips to speak at different engagements and are so thankful for donations that were received. Our group also got to make two trips to Memphis to deliver wagons to the families. It is such a blessing to be able to meet some of these amazing warriors and talk to them about their experiences. On a HUGE note, Jacy's Wagons is nw a member of the McNairy County Chamber of Commerce and we are so grateful to have been asked to join and can't wait to see the growth that the Commerce and Mrs. Jessica Huff will bring to our mission!





Jacy's Wagons

Michie, Tennessee www.mcnairy.com

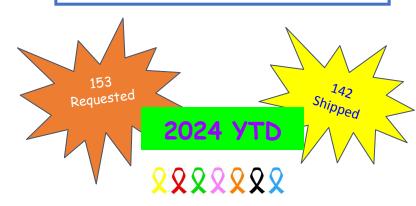
UPCOMING EVENTS

1 Year Celebration Ribbon Cutting Jan 18, 2025 at Acton Church of Christ Ministry Building 2-4

Save the Date: Valentine's Day Dinner February 15, 2025 at Acton Church of Christ Ministry Building

Executive Board Members:

Jennifer Raburn, *President*, (662)664-2547 Jacy Raburn, *Vice President* (731) 610-9406 Tasha Barnett, *Secretary* (662) 664-3305 Kelly Broadway, *Treasurer* (731) 267-1470







Mail donations to: Jacy's Wagons PO Box 355 Michie, TN 38357



venmo



2nd Annual Jacy's Wagons Music Festival

Save the Date: April 12, 2025



Where: Michie City Park, Michie, TN

Time: loam-7pm

What's Happening: MUSIC, Food Trucks, Vendors, Kids Corner, Fellowship, and Fundraising









Lydia was born with a genetic condition called Saethre Chotzen Syndrome (SCS). This caused one of her soft spots to close too early and other soft spots to be too large/open. At two weeks old, Lydia was sent to LeBonheur for failure to thrive. They started the work up for SCS while we there. On scans, they found Lydia had two strokes while in utero. We were then told her was blind. She did not respond to light when ophthalmologist assessed her. Her pupils also got larger instead of smaller when light was shined. We were terrified. Shortly after we got back home from that 9 day stay, she started having seizures. We had multiple trips, EEGs, and started on meds. Seizures stopped and she was able to stop meds. Then at 4 months old, she was diagnosed with AnoRectal Malformation (ARM). She was born without an anal opening. Lydia started therapy- OT, PT, ST, and vision therapy at about 8 weeks. We had so many questions about what would her future look like. Would she continue to have deficits from the strokes. Would she be completely blind her whole life? When she was almost 5 months old, she for the first time looked AT me and not through me. It is a time I will never forget! she was suppose to have cranial surgery at 3 months old but the surgeons were not comfortable doing it yet. They wanted her to grow a little more. They plan to do it the first part of 2025. She just underwent surgery to reconstruct her bottom and create an anus. She did amazingly well with the surgery. We are having some complications with her incision but she's handling everything so well. You wouldn't even know she had that major surgery.

Lydia likes to watch cocomelon, bluey, and chip and potato

We are now able to start eating baby foods, since her reconstructive surgery. That has been very fun

