



# Jacy's Wagons

Oct 2024

*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 now 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!

## 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

153  
Requested

2024 YTD

142  
Shipped



*New* 2024 MEMBER

**Jacy's Wagons**

Michie, Tennessee  
www.mcnaairy.com

**WONDERFOLD**  
— WAGON —



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



venmo

**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*

**2nd Annual Jacy's  
Wagons Music Festival**

Save the Date :  
**April 12, 2025**



**Where: Michie City Park,  
Michie, TN**

**Time: 10am-7pm**

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





# Lydia



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 were there. On scans, they found Lydia had two strokes while in utero. We were then told she was blind. She did not respond to light when an 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 supposed 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

