

Fiona is still imagining the possibilities of her wish

Fiona, 3
genetic disorder



Anyone who has had the pleasure of meeting three-year-old Fiona, instantly feels her warmth, calmness, and pure soul.

Fiona adores people and her favorite activity is to snuggle; which Mom and Dad are happy to comply with. Fiona enjoys the outdoors and loves feeling the wind in her hair and watching the sunshine glisten through the trees. She also loves to bust a move by turning up the tunes and dancing and twirling in her wheelchair or in someone's arms. Fiona was born with a rare genetic condition and spent the first month of her life in the NICU at the IWK. Overtime numerous health issues have arisen including epilepsy, feeding difficulties that require all her nutrition to be through G-Tube and Cortical Vision Impairment. She has seizures daily, does not have the ability to verbally communicate or walk. Fiona also has great difficulty when she is ill. Even common colds are difficult for her and she has had numerous hospital admissions that include being life-flighted and admitted to the PICU. Fiona is a warrior princess and continues to persevere with a smile on her face and joy in her heart. Her family, friends and community have been nothing short of amazing and Fiona is blessed to have so many people in her corner. She is a rare and beautiful unicorn and she and her family are thrilled she will be granted a wish.

“And though she be but little, she is fierce.” – William Shakespeare

Follow Fiona's journey through her personal blog on Facebook called “Friends of Fiona.”
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