



Mccowen and Scurlock's life has been turned upside down. We have seen little Brennan sick and in pain from the chemo and cancer. We have seen him cry from the daily injections. We cry because there is nothing we can do to take it away. We have also seen the love and happiness when he feels good. We have seen all the good people and foundations that have prayed for him and helped our family in this fight for his life. It has opened our eyes to the horror that is childhood cancer!

The following pages will tell the story of Brennan's treatment

April 9 2011

Brennan officially diagnosed with Stage IV High Risk neuroblastoma.

He began his first of 5 rounds of chemotherapy.

May 26 2011 [Brennan's Birthday](#)

Brennan spent his 1st birthday in surgery to remove the tumor on his left kidney, adrenal gland & the diseased lymph nodes surrounding the abdominal aorta.

June 2011

Stem cell harvesting

August 2011

15 rounds of radiation treatment totaling 3 weeks

September 12 2011

In-patient @ Texas children's until about Oct. 20 for high dose chemo. And stem cell replacement. This will be the hardest on his little body, and some of the family because he can't have visitors due to his immune system being wiped out.

October 5 2011

Some not so good news.

A new MRI compared to August's shows the tumor on his cerebellum which had shrunk from the first series of chemo has since grown. Not sure what it will mean for his

treatment plan yet. Some extra

prayers will sure help

October 10 2011

The plan is back into radiation next week for the tumor on Brennan's skull. We don't know yet how many treatments will be needed

October 14 2011

[Brennan comes home today!](#)

His new MIBG scan showed no findings of new or progressive metastatic disease! Which is VERY good news. Now just a week more of radiation and then Brennan's last phase of treatment. By this time next year he will be in REMISSION!

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Our Brennan's Buddies have moved [HERE](#). Be sure to check them out!

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Brennan has always been a happy baby. Always laughing and playing. He loves helicopters, balloons, bubbles, Yo Gabba Gabba and playing in your hair. he loves his moms hair. As he got a little older he really liked watching zombie movies. He liked to pretend to be a zombie and growl and play bite. He loves his sister Katelyn a lot , she can always make him laugh.

- November 3 2011
- December 19 2011

Last day of radiation!

Brennan's scans are good! Except for a new spot showing up to left side of his spine. It lit up on MIBG scan, but doesn't show on any other scans. He also has a bony calcification growth on his spine, which is most likely his cause of pain every night for the past 3 weeks. He'll be scanned again early before 3 months to keep an eye on the spine and the possible lesion that has shown up. If bone marrow is clear he'll begin last phase on Jan. Also hopefully we'll be making plans to remove his port.

January 3 2012

Brennan Began his first round of Acutane treatments.

Here is a link about **Accutane** its mainly for treatment of severe acne but is also used to treat some cancers. It has some nasty side effects, but then so do all the drugs used for cancer.

Brennan was also to start Antibody therapy in this phase but slightly low kidney functions disqualifies him for that treatment here. Cindy is contacting Sloan-Kettering Cancer center in NYC to see if he would qualify for their clinical trials in their antibody therapy Cindy really feels he

January 22 2012

Brennan was admitted to TCH today. He hasn't been eating and has been vomiting all day. Unknown at this time

January 23 2012

A CT scan today found a fairly large mass in his abdomen, around his bowel causing what looks to be an obstruction in his small bowel, near stomach. Which is causing his bile to back up, and go now where but up and out of his

January 24 2012

We are very sad to say the least. Brennan will be in surgery today to biopsy the new tumor in his abdomen which is blocking the bowels and causing his constant vomiting, and also repair the connection from stomach to bowels so he can eat. pathology of the Neuroblastoma has changed, meaning a slightly different form all together. This form is very aggressive, and should this tumor be that. There's not much, if anything they can do for him. They will not be removing the tumor. Pray for little Brennan.

I sit here at this computer lost as what to do or say. This was completely unexpected, he was doing so well in his treatment. Just Saturday he was happy as always making his usual zombie growls. As Cindy's father I don't know what to do to take my little girls pain away and there seems

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Check our [Brennan's Buddies](#) page. There are many children that need help and prayers. Some are still fighting, some are Angels in Heaven. The one thing they all need is for people like you to share their story and raise awareness so changes can be made.

How fast things change. Just a couple of weeks ago we thought the worst was behind us. Little did we know what lay ahead.

• January 24 2012

Brennan's surgery was 2.5 hrs. It went well. They were able to remove the obstruction in his bowels. Hope to start some more chemo in next couple of days

• January 25 2012

Brennan is pale, elevated heart rate staying at 160, and NO urine since 1:30 pm today even after they've increased fluid intake greatly. And turns out he has a fever now. Brennan's right side is swollen and he's still in a lot of pain even with increasing morphine pain pump twice.

• January 25 2012

Brennan will also begin chemotherapy tomorrow evening. From my understanding the 2 drugs he will be given are 2 which he has not had. And typically can be done as an out patient but given Brennan's current status he will remain inpatient.. No clue as to when we will return home. Our hopes are to make this tumor respond to treatment.

Palliative care are words I wish I never knew.

Brennan is classified as Relapsed.

Things change so fast, not even a month ago we thought Brennan was doing so much better in his treatment.

• January 26 2012

No chemo today. The Dr's need to identify the new cancer first in order to find the right drugs to treat.

• January 27 2012

The Pathology Dept. @ TCH is working the weekend to try and identify Brennan's cancer. Without a type they can not treat.

• January 28 2012

Well its sure Brennan DOESN'T have neuroblastoma anymore. Apparently no one knows what Brennan has, but that its a high grade malignant sarcoma.. they have no clue what cell this came from, in turn have no plan in place to effectively treat him right now. So we sit and wait for a plan. They did say, Brennan went from NB to a different aggressive cancer. =(

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January 30 2012

Dr's are leaning towards an "undifferentiated Sarcoma" basically meaning they don't know where it came from but its not the obvious sarcoma cancers. A Dr. at St. Judes who is speaking with Brennan's Dr. claims to never have seen a child with NB switch to this cancer.. so Brennan is a rare case.. Cindy has contacted the Sarcoma dept at Sloan-Kettering.. She just doesn't believe Brennan is the only child to deal with this.

February 1 2012

After 9 painful days of waiting Brennan now has a diagnosis. This tumor is now diagnosed as Highly Aggressive secondary tumor of Undifferentiated Sarcoma. .

Today 46 children will be diagnosed with some form of cancer, Today 7 children will DIE from cancer!

The two Chemo drugs Brennan will start sometime in the middle of the night are Ifosfamide and Doxorubicin.. At high doses.. much higher than his previous induction cycle for Neuroblastoma. His Ifosfamide previously was given at 300mg.. It is now 1,470mg.. his Dox was 4mg, and is now 18mg.. both of these are very harsh, especially at high doses. Brennan will have a heart ultrasound before each cycle is to be given because Dox is very damaging to the heart. Ifosfamide can at high doses cause acute neurological problems like seizures, extreme sedation and some others..

February 2 2012

Ch t t d t d



February 4 2012

Since his blood pressure dropped it can either be chemo effect, which is very soon or just that they have diluted his blood with all the hydration he's been getting last several days. He's going to get blood and as long as he doesn't worsen he will be ok.. they say his bowel isn't obstructed all the way so an NG tube right now won't really help him much. So its a wait and see, hope he gets better type of thing. Until then he's on stronger doses of morphine and we're going to start our last day of chemo in a couple of hrs.

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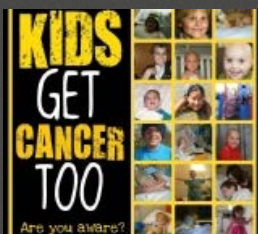
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February 4 2012

Brennan has been moved to ICU because of high heart rate, difficulty breathing and needing more observation than regular floor nurses can handle.. of course he's throwing up again. Not sure what will transpire being here now. He's also on oxygen now.

Chemo stopped for now. [Keep praying for our little one](#)

February 5 2012

Not sure today is going to bring us good news.. Brennan's vomiting has increased quite a bit, and now surgery team is going to come back and look at him.. I have a huge fear Brennan's partial bowel obstruction is worse. =(the night hasn't been good to Brennan. Several x-rays, and an arm IV as well early this morning, on top of getting sick. =(

There are so many ways to help, Donate to one of the great foundations you see throughout our site. Raise money to shave your head in a [St.Baldrick's](#) event near you. If it's something you've considered Register for [bone marrow donation](#), there are many children and adults that would benefit from your gift.

Brennan on his way to ct scan. Checking blockage again. Also there is a concern about the fluid in his lungs and threat of pneumonia when his counts drop from chemo.

February 5 2012

Trying to get Brennan on clear liquids and going to try and start formula. NG tube going back in. We hope he can come home for a bit before his next round of chemo Next Thursday.

CT scan shows 3 small liver spots, most likely tumor too.. it wasn't there before surgery, so its unsure if it just popped up in the 9 days waiting for diagnosis/chemo or if he's progressing thru chemo. I guess that'll be one more thing to pray for.. we can't have him not respond to this chemo!

So that makes abdominal tumor from left kidney, lung spot and 3 liver spots. =(and there's an abdominal abscess which is small but may need to be handled if it grows more

February 6 2012

Not much new today. He still has fever, heart rate still high. I did get to see him today. He was asleep when Me and his aunt Shanna got there, but he woke up and got a bit fussy. I sat next to him was holding his little hand and he looked at me and made the loudest zombie growl..If you don't know him we love to watch Walking Dead and other zombie movies. He always thought he was a zombie. He would come after you like he was going to eat you and growl. It made me so very happy to see him do that. Even when he must feel like crap he still tries to make you laugh. I love him and miss him being home.

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Treatment for childhood cancer can last years. Being a child, in most cases one parent must leave work to care for their child. It is 24 hour non-stop care. It leaves many families devastated financially and emotionally. The help of other family, friends, community and other foundations is essential. THANK YOU to everyone helping these families.

February 7 2012

Brennan's Attending Oncologist says Brennan remains critically ill, but is showing small progress. He says Brennan remains this way because things can always change. Brennan finally passed out after being up since 3:30 am this morning. =)

February 8 2012

Well not much new today, he's had a rash and they are ever so slowly lowering his blood pressure medication. He cant leave ICU until they can get him off it. Next round of chemo coming this month sometime. Everyday we pray. And we pray for the others, RJ, Emily, Rachel, Allison, Emma, Liam, these are just a few I have at the top of my head, there are so so many more.

February 11 2012

Brennan has been out of ICU for a day now! He's doing well, managed to somehow have his NG tube "slip/pulled" out. so now he is multiple-tubes free! =) Still not allowed to eat, hopefully soon that changes. And I (grandpa) got to spend the night in his room watching him. Had the best time EVER!

February 17 2012

Brennan is home! We had some setbacks once we came home, ended up back at ER for a central line repair(again) but all is well.

Back to clinic on Tuesday and next round of chemo on Thursday. He's so happy to be home. And we are so

February 19 2012

Tonight I get to watch "The Walking Dead" with the best zombie impersonator

February 23 2012

Brennan is back to Texas Children's. He should be admitted today for his next round of Chemo. Hope to have him back home Sunday or Monday!!

February 27 2012

Brennan is home again after 2 days of chemo.

So happy he's here. Should be getting scanned soon. Home for a few weeks before the next round of chemo. Keep

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Only about 3% of NCI's budget is allocated to ALL childhood cancers. That's only about \$197,000,000 it sounds like a lot but it's not really. Children still get adult chemotherapy drugs just scaled down. They should have their own, a child's body is not just a small adult. I believe better drugs and treatment protocols need to be made available. TOO MANY

March 2 2012

Brennan is back to TCH ER this morning. Throwing up a lot and not himself. Lets hope and pray it's nothing.

March 2 2012

Brennan was admitted from ER to PICU today for high heart rate and persistent fevers, from chest X-ray it looks as though he either has partial collapsed left lung or pneumonia.. leaning towards pneumonia. he's been pretty sick today, and hemoglobin at admission was 6.7 and platelets around 17..both very very low. he had blood and platelet transfusion already. just waiting to see if next X-ray will show improvement if so then its collapsed lung, if not then pneumonia.

March 6 2012

Brennan's second opinion from St. Jude's came back (head of their solid tumor board was head of tumor board here at TCH previously) and they are NOT comfortable calling Brennan's new tumors Undifferentiated Sarcoma, nor Neuroblastoma anymore. Although when they requested Brennan's original tumor slides of NB there were tumor markers one being ALK and another one (I cant remember) that showed on the NB tumor, which continue to show now.. they said its very rare to be neuroblastoma and lose the Mnyc amplified gene (which made his NB considered High risk) So literally they are calling it a tumor of Undifferentiated "something". Because they really aren't sure how/why this happened. These tumors do NOT pick up on the Neuroblastoma scan MIBG. So where this leaves us is in another "unknown" about Brennan's cancer, but potentially leaves us with a Phase 1 clinical trial should be be accepted,

even though he's not technically NB anymore.. Results for that acceptance are pending a "special" lab testing which takes 4-6 weeks.. meanwhile we continue, scans begin tomorrow and finish next week right before chemo again.. potentially Brennan could be looking at surgery before, or after the chemo.. it all depends on how this tumor has responded and if the surgery team feel it can be removed now.

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Something I've noticed in this fight with my grandson, Brennan. These kids are the toughest, bravest I have ever met. I've seen the warnings on the drugs that are pumped into their veins. Seen first hand the side effects. They take it all in stride. Hardly ever complaining. I wish I could have half the strength they do. They are my HEROES!

March 14 2012

Brennan did come home @ about 1am back to TCH today for chemo, but Chemo wont start today, and he's going to have x-ray/possible ultrasound to find out why he's throwing up so much more now. And depending what those show he may be seen by gastroenterologist. It's a never ending problem. So much changing, good news, stable news, bad news. It's been a roller coaster ride we want off of!

The chemo he is on will treat neuroblastoma, but we don't know why after 2 high dose rounds its showing still. Which is scary. His kidney tumor shrank slightly, but not as much as we had hoped. Liver tumor appears stable in size if not slightly smaller.

March 15 2012

Today waiting on Brennan to get his blood transfusion started. Being here since Thursday his Hemoglobin (carries oxygen through red blood cells) has been steadily dropping from almost 10 to now today 7.6. Not sure why over 3 weeks from last chemo he's dropping now. But I'm sure no one is concerned with it. Still waiting to hear from the Doctors/surgery team as far as their plan for Brennan, since we are no better with the bile vomiting. Which clearly means there's something going on with this tumor/stomach. He's also been switched to a continuous morphine pain pump for his pain. I'm hoping today brings us the answers we are looking for and some resolve for Brennan discomfort and vomiting.

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March 9 2012

Brennan is home! He's doing well Always so happy to be home. Brennan has scans next week then Thursday or Friday back to TCH for next round of chemo.

Be sure to check our Brennan's Buddies page.

March 13 2012

Brennan is back at Texas Children's ER. Has a fever of 102. Chemo is supposed to start tomorrow. Let's pray there is no delay and nothing to the fever.

March 15 2012

The lesion that picked up on last MIBG scan (looks for neuroblastoma) and they weren't sure then if it was true tumor, still shows now and apparently its active neuroblastoma. The other tumors on lung, left kidney and liver are the new cancer. So they say he most likely has 2 cancers right now. Neuroblastoma and this "not sure what it is" cancer.

St. Baldrick's Childhood cancer takes hope for a cure. StBaldricks.org



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March 19 2012

Well it just never ends...Brennan's oncologist failed to mention to Cindy last Friday when she gave her the results of the MRI's and other scans that Breanan also has a .5 cm lesion about in the center of his brain. She had to find out this terrible news by asking for the reports and reading it herself! The whole family needs some prayers especially now. So Neuroblastoma has returned along with the abdomen tumor. lung and liver tumor are stable, now a brain lesion. We just want our baby well. I get my updates from Cindy most times before she updates Brennan's Caringbridges site but she goes into much more detail, so check there also.

BRENNAN WAS DIAGNOSED ON APRIL 8TH 2011 WITH HIGH RISK STAGE IV NEUROBLASTOMA. JAN. 22, 2012 BRENNAN RELAPSED WITH A NEW DIFFERENT CANCER. APRIL 9TH 2012 BRENNAN IS DONE WITH

March 20 2012

Brennan is in for upper GI and small bowel test today.

March 21 2012

Brennan is having his Med's switched to treat Pneumonia caused by aspiration from throwing up. Which is most likely his cause of fevers, and steadily rising WBC. They're also going to begin chemo (hopefully sometime tonight) Hopefully Brennan can complete his 10 days of treatment before counts drop, otherwise there's a big risk of his pneumonia being able to do whatever it wants. Tumor boards met today about Brennan's recent scans. No one seems to be convinced this brain lesion is NEW spread from his secondary cancer.. apparently from his DR's words.. Neuroblastoma & Sarcoma don't "commonly" do this where the spot is, its too small to biopsy.

So all we can do is proceed forward and hope its no longer there, or not bigger when he is re-scanned. Good thing is Brennan can still receive radiation to his hipbone for the NB tumor and possibly (if needed) his liver lesion and lung lesion after surgical removal of abdominal tumor. It's also suspected Brennan has Gastric paresis.. basically paralysis of his stomach, which doesn't allow or move enough to empty enough. So after they get a handle on the pneumonia they will begin meds to stimulate his stomach to move more, and hopefully this will stop his throwing up.

March 26 2012

Brennan is home and doing well.

He's always happy at home. He has been

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April 8 2012

His mouth sores since being here got worse. He ended up waking up Friday with at least 5 visible white sores near the tip of his tongue. He's on 3 different med's to help get rid of or treat his pain for his mouth. He had to be switched from Morphine to Dilaudid to control his intense itching from the IV morphine. He seems constantly agitated. Although he's more awake now being switched from the morphine, he is in no mood to cuddle with his momma or look at any toys/gifts he's received. We're supposed to be admitted again Thursday for chemo.. I'm hoping and praying this last cycle isn't delayed a week like our previous one was. With his cancer being so aggressive, you fear just waiting 7 days from the normal 21 is just giving his tumors more time untreated, which can give them opportunity to grow.

the 5mm lesion we found 2.5 weeks ago has grown and there is a second lesion on his brain. They have grown and has caused Brennan's brain to bleed and cause pressure. Every one of his sarcoma tumors have grown. The kidney one, his liver and lung tumors. Brennan CANNOT continue chemo anymore because its too risky with his hemorrhage. This was not supposed to happen to my son. My heart is shattered. we have chosen to not place a drain because Brennan would be unable to move and his comfort would be compromised. ICU is not a place we want Brennan right now. He is comfortable and shows no signs of discomfort. Please pray for my love.

March 30 2012

At clinic all day. Had to have blood and platelets. Counts are low but all is good when Brennan is home. I think another 2 weeks at home then round 4 of chemo. Keep our little guy in your prayers, and all the other little fighters. And don't forget prayers for the little angels and their families. RJ, Rachel we think about you everyday.

April 9 2012

Hospice is coming today to talk about plans when Brennan is home. Please pray that Cindy and Blake feel Gods hand over their hearts.

April 5 2012

Well all had been quiet at home, until last night Brennan has been cranky and not a happy boy. Not sleeping, crying and throwing up a lot. Today was clinic day and they decided to keep them(don't know for how long yet) he's dehydrated and possibly an ulcer. He also has some mouth sores from the chemo.

Brennan should be home by the weekend. I'm not sure if Cindy & Blake will want many visitors at first. It's a very sad time right now for everyone.

April 9 2012

This afternoon Brennan gets re scanned with Ct/MRI then sometime soon MIBG to check the interpolations. His pain is very bad. We need some answers the scans may provide.

April 16 2012

Brennan, Blake, Cindy and Kate should be home sometime today(Monday). I'm both happy and very very sad at the same time. I cant wait to see Brennan but I don't want him to ever leave.

Easter Sunday will mark 1 year since his official diagnoses, and it seems his battle is far from over. so please send him some

Brennan's scans were rushed because of the fears the Dr's had.

Brennan's scans showed what we feared. Brennan's sarcoma has

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April 21 2012

Had some family and close friends come visit the past few days. Brennan sleeps all day but will respond when asked questions with yes/no head shakes.

I truly hate cancer and what it does to the children



April 17 2012

Brennan knows he's home. he keeps making noises and comfortable sighs, has even opened his eyes and been responding to us.. we haven't had him acknowledge us in 24 hrs!



April 19 2012

Not much new. Brennan is still in pain, med's have been increased a couple of times now. We are just trying to keep him comfortable and enjoy the time with him.

April 22 2012

Brennan is still a little fighter. He's very weak but still with us. He drinks a little water now and then. Sleeps all day. Continue to pray for our baby



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April 25, 2012

Brennan Became the sweetest angel this morning @ 5:15, with mom and dad, sister, aunt, uncle and grandfather by his side. Our life will never be the same. We love you so much little Brennan

The song playing "I have a photo of you" was written and performed by Katelyn, Brennan's big sister.



Saturday April 28 2012

I never expected this day to come. The day we have to say our final good bye to Brennan. It wasn't supposed to be this way. Today will be a hard day for me, I can only imagine how Blake and Cindy feel. We will all miss you Brennan. But we will never forget.

Brennan's service will be private for family and a few friends.



By: Matthew McCowen
(Brennan's 12 yr.old uncle)

As the soul departed young Brennan's dying heart the sky opened for him and down came the Angel's singing in harmony. As they flew down to take him away to heaven. Just after he past I was informed he had gone away but yet I still felt as if he was sitting next to me biting my arm like the zombie he would pretend to be. And as I looked over he smiled and laughed as he flew away and left us down alone and sad wondering how he's doing with God, past family members, and the Angel's calling to him from above. I loved him so much. We all did.



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