Inside: Your best guide to local The Wife The FAPPENINGS /p19

PRSRT STD U.S. Postage PAID Permit No. 5187 San Jose, CA

CampbellTimes

times@timesmediainc.com

nis

in

ng

nat

gic

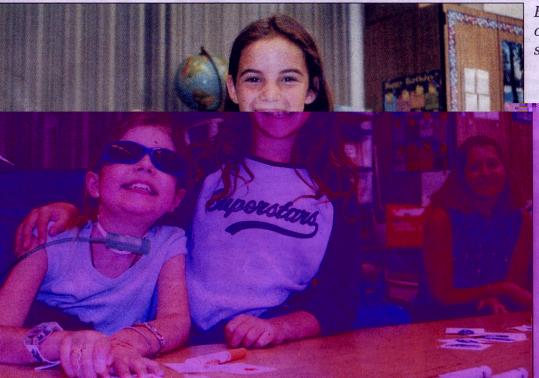
ns

om

17

October 2002 • Volume 12, Number 1

Kaitlyn returns to school



Eight-year-old struggles to overcome devastating effects of severe allergic reaction

By Julie Davis Berry Editor

t recess Kaitlyn Langstaff has alway been one of the most popular kids the Forest Hill Elementary School capus. And she still is, despite the fact that the fall she returned to school blind, mute and a wheelchair after surviving a devastati allergic reaction to Ibuprofen. The fact the she is alive at all is a testament to a vecourageous little girl, the strong love of hamily and the amazing compassion of a comunity touched by her health crisis.

Physical issues

While out of town last April, Kaitl became ill with a very rare and severe aller reaction to Ibuprofen, which is called Steve Johnson Syndrome. She was airlifted from the action to a Los Angeles hospital where a spent several weeks on life support.

See KAITLYN, page

Kaitlyn is especially glad to be sitting next to her b friend Arlana Gomez in Mrs. Dorsch's third gra class. Her mother Kerry watches in the background

Kaitlyn returns to school

KAITLYN, from page 1

Eventually she was flown back to northern California to Lucille Packard Children's Hospital at Stanford. These were very rough months for the eight-year-old and her family. She almost died several times, but each time she pulled through.

Dr. Benjamin Mandac is in charge of pediatric rehabilitation at Valley Medical Center. He heard about Kaitlyn's case and came to Packard after hearing that she almost died one night. Mandac took control of her

doctor at UC Davis recently told them that they should put off doing any surgery for a few years to benefit from any future technology advances. "He told us that he believes that at some point in Kaitlyn's life she'll be able to have some vision," said Kerry.

Due to the damage from Stevens Johnson Syndrome, Kaitlyn has no airway through her nose or mouth and she must breath with the assistance of oxygen through a tracheotomy tube in her throat. Because of the lung damage it



Kaitlyn returns to school

KATILYN, from page 17

But the family has not been able to access any financial help. "She doesn't qualify for social security benefits. She has a lot of problems but she doesn't qualify for government assistance because she doesn't have a developmental disability, just a physical disability," said Kerry. "A family of four cannot have assets of more than \$3,000 not including one car and a house. They said to call back when Kaitlyn is 18 or we have no income."

The ironic thing is that Kerry was scheduled to start a new job as a marketing director at Nuance the day that her daughter was airlifted to a Los Angeles hospital. Kerry feels that once Kaitlyn starts to feel comfortable being at school

without her in the very near future, she would like to return to consulting on a part time basis. Brad's specialty is putting on trade shows in the high tech sector and with that comes travel so it is unlikely that he will return to a similar job.

Kerry is looking to start up her market- given, given to the community. Now that everyone just wants to belouthe fare lading, ind corrections have been supported and outlewiness are translated one consistently in the constant of the community of the community.

more about Kerry's consulting business at www.langstaffcommunications.com

In the meantime the family says that the donations have been helping them to pay for things like the mortgage and food.

Community Support

The outpouring of support from family, friends and the community in general has been a real source of solace for the family during this extremely difficult time.

"It's impressive to see how nice people can be in this day and age," said Kerry. "All of these people, some who we barely know, are reaching out to help us. We are so grateful for this great community."

When Kaitlyn was first sick a family friend, Tammy Davies, flew down and

ended up staying for a week while Kaitlyn was transferred to Los Angeles. "It was a very awful week," said Davies. "I was just there to let Kerry vent because it was so tough. They are quite remarkable people. Kerry and Brad have always just given, given, given to the community. Now that

responding and they are reaping what they've sown."

Kerry has led Brownie troops, Destination Imagination groups and produced the school play last year for Rolling Hills Middle School. Brad was in charge of Indian Princesses at the Southwest YMCA and has coached Bobby Sox teams. They have touched many lives in the community and people have responded in droves.

Three times a week people bring dinner to the family. "Even the principal of Rolling Hills, Mrs. Atkinson, brought us dinner!" said Kerry. And, twice a month a group of moms the family met through school and brownies comes over to clean the family's house. These are all organized

by the Langstaff's neighbor Karen Pasos.
"It's pretty easy because I can send out
messages over the Internet and people just
sign up," said Pasos. "We have people
from Forest Hill, Rolling Hills, Indian
Princesses, Bobby Sox, Brownies and
everyone just wants to belothe fore: language.

"Keny's Gir "Scout troop is putting together all of the cards and letters in scrapbooks for the family. "There were over 814 get well cards sent to Kaitlyn," said Kerry. "The cards were just amazing. They were all so supportive."

Davies said that the other day she was thinking about the Langstaff's and she thought, "They could really use a miracle. But, you know they already have a miracle. Kaitlyn is still with us."

If you would like to donate money to Kaitlyn and her family send a check to Kaitlyn Langstaff, c/o Forest Hill Elementary School, 4450 McCoy Avenue, San Jose, 95130.



Kaitlyn is one of the top students in Mrs. Dorsch's third grade class. She constantly raises her

Nicole Zannini is her p

full time aide. She has heen and de with the district for nine years.

"I'm used to working with children who are born with disabilities. It's second nature to said Zannini. "When I found out that I would be working with Kaitlyn I knew it was going to be a challenge because of all of the adjustments that Kaitlyn was going to have to make. But, she is a great kid. And her family is really awesome. They don't baby her. They want her to be independent. They are verv upfront about her issues."

parents came in the first day and explained things to the class and told the kids about things they might see and hear which was very smart. They don't over accommodate her."

Kerry is overwhelmed by the support Kaitlyn gets from the other students. "The kids have been very supportive from the very first day. At recess the kids would take the bracelets off their arms and give them to Kaitlyn," said Kerry.

Her best friend Ariana Gomez sits next to her in class. They were on the same soccer team last year. This year Kaitlyn will have an honorary T-shirt to wear on the sidelines. "I saw her at Valley Med and she looked better than I thought she would," said Ariana. "I like Kaitlyn because she is really funny. Now I can understand

practice."

Since she was in the hospital Kaitlyn has had a dream of having a Golden Retriever puppy like her dad had when he was growing up. She would name the puppy Daisy. The family would like to make her dream come true soon.

Financial issues

The family is also struggling with devastating financial issues. Both Kerry and Brad were unemployed when Kaitlyn's illness occurred. They have been unable to work since then due to the 24-hour care that Kaitlyn's illness requires. "We have exhausted our unemployment. We have been living off of donations and our savings and they are almost gone," said a weary Kerry.

One would think that there would be help from social security or some other government