

LUPUS LA MEET OUR LUPUS HEROES

Michelle & Alan Kaye: Advocates, Parents, Philanthropists

a lot of things to a lot of people. While the entire Lupus LA community is a family in and of itself, the Board of Directors is a small close-knit group who has been together since the early beginnings of Lupus LA. They are a team of truly dedicated people fighting to find the causes of and a cure for lupus. We are pleased to introduce you to our March Lupus Heroes, longtime Lupus LA board members Michelle and Alan Kaye. In 2005, Lupus LA founder Daniel J. Wallace, MD, diagnosed the Kayes'

It's a Family Affair! Family can mean

daughter Gillian with lupus. Since then they have committed themselves to doing all that they can to find a cure. "We saw how our daughter suffered from lupus and wanted to help in any way we could. Our focus then and to this day has been on raising money to fund innovative research with the expectation that it will result in new treatments and an eventual cure for this debilitating disease."

Alan's professional expertise has enabled him to lead the financial growth and management of Lupus LA as

ually raise more money as well as awareness for the

have common issues and goals!"

disease.

the board treasurer. Michelle's creative vision and artis-

tic flair have helped Lupus LA's events shine and contin-



March 2015

anyone with lupus, or family members of

those with lupus,

to find information

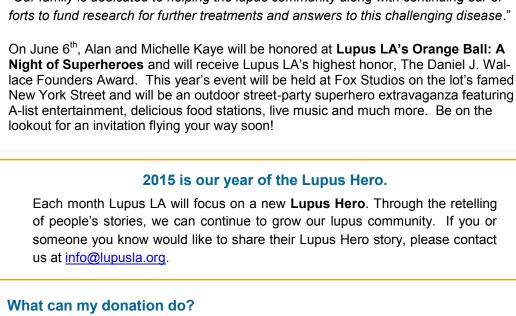
and support. - Michelle & Alan

Together, they are a force to be reckoned with. The Kayes have been active both in and out of the boardroom and look to Lupus LA for cutting-edge information on lupus treatments and education. They share, "We have attended some really informative patient conferences and been able to network with others to discuss what has worked for them. It helps to be around others who All that being said, what matters

> most to Michelle and Alan is their family. The Kayes have two beautiful girls, Gillian and

Danielle. While Gillian continues to face lupus-related challenges,

she is living life to the fullest and is currently working in New York City on a TV show. Her sister Danielle is graduating from The University of Arizona in May and returning to Los Angeles to pur-The Kaye Family sue a career in communications.



help support our programs and initiatives:

\$ 1,000 = One (1) week for a Rheumatology Fellow \$ 5,000 = Two (2) weeks of Scientific Research \$10,000 = One (1) day-long Patient Conference

DONATE HERE

Questions or comments? E-mail us at info@lupusla.org or call us at (310) 657-5667.

To remove your name from our mailing list, please click here.

Every dollar you donate counts at Lupus LA. Here are some of the ways your donation can

\$ 100 = One (1) monthly adult support group session \$ 250 = One (1) monthly pediatric support group session

\$ 500 = One (1) Emergency Grant for a lupus patient

camp for kids with lupus

\$ 750 = Sponsorship for one family to attend a special summer

