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1st Roundtable
What You Need to Know
About Lupus Live
Webcast

Speaker
Dr. Megan Krause,
MD

Let's Get the conversation Started



Mission

Marlene's Kaleidoscope is dedicated to improving the lives of people affected by all forms of Lupus. We strive to empower people by providing information about Lupus and offering support, so that their voices can be heard in service of creating a better quality of life through research, public awareness, advocacy and education.

Vision

A world where people with lupus can lead a full and active life.

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Ultimately find a cure.



Did You Know that Lupus is more prevalent than AIDS, sickle cell anemia, muscular dystrophy, cerebral palsy, multiple sclerosis, and cystic fibrosis **COMBINED**. It is estimated that over 1.5 million Americans have been diagnosed with Lupus; 16,000 people are diagnosed each year. That's why we need to **Get The Conversation Started** KC!

Sources <http://www.lupusinternational.com/Living-With-Lupus/Newly-Diagnosed-1-1/Lupus-FAQ-s.aspx>
<https://www.lupus.org/heartland>

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Lunch with Doctors

This component requires a partnership with a rheumatologist who can provide continuing education to primary care physicians (internists, family practices, etc. Once the presentation was developed, it would need to be an approving body for physician Continuing Medical Education (CME) so that physicians who attend can get credit toward their required education units.

Coping with Lupus Awareness and Education (CLAE) Round Table

The purpose of the CLAE round table is to provide information and awareness regarding living with Lupus. This quarterly webinar covers topics like: Getting Diagnosed, what you need to Know about Lupus, Wellness and Lifestyle Changes, Psychological Factors in Coping with Lupus, etc.

RoundTable

Promote awareness across our community

Monthly Lupus Support Group

This group is designed for individuals living with Lupus who can benefit from ongoing social support and encouragement from others who are coping with the disease.

Online group

Once quarter group outing

Lupus Love Gift – Hospital / Home

Scholarship

Legacy Scholarship

Established in 2019, it aims to support KC area high school students, living with Lupus.

Applications are available online Wednesday, December 1st each year

The application deadline is April 31st each year

Grant Amount is \$1,000

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LET'S GET THE CONVERSATION

AWARENESS

STARTED ABOUT LUPUS

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We are please to have Dr. Megan Krause

Dr. Krause received her undergraduate degree from Northwestern University in Evanston, IL. She went to medical school at the University of Kansas. She completed her Internal Medicine Residency, Chief Residency, and Rheumatology fellowship at Mayo Clinic in Rochester, MN. She joined the staff at the University of Kansas in 2016 as an assistant professor and is the Associate Program Director for the Rheumatology fellowship. She is active in clinical practice and teaching at the medical school, resident, and fellowship level.

Agenda

- 10.00 am: Introduction By La Tanya Lipprand with Marlene's
- 10.15 am: What You Need to Know About Lupus by Dr. Megan Krause, Kansas University Hospital
- 11:15 am: Q &A
- 11:45 am: Final thoughts and call to action
- 12.00 pm: End of Session



**TO LEARN MORE ABOUT GETTING THE CONVERSATION STARTED OR
MARLENE'S KALEIDOSCOPE**

GOT TO

WWW.MK4LUPUS.COM

OR

EMAIL: KBUTTERFLIES2017@GMAIL.COM

573-738-8648

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[HTTPS://WWW.MK4LUPUS.COM/DONATE](https://WWW.MK4LUPUS.COM/DONATE)



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