

Bobby Jones Chiari & Syringomyelia Foundation

Fly me to the

MOON

CASINO NIGHT

HONORING

Kristin Means

SATURDAY, FEBRUARY 28, 2026

Dear Friends and Family,

Fly me to the moon... let us celebrate a night filled with purpose, possibility, and a little bit of luck.

Tonight, as the chips stack high and the cards are dealt, we gather not just for an evening of fun — but for a mission that reaches far beyond this room. Casino Night is about coming together as a community, united in our commitment to advancing research, raising awareness, and bringing answers to families affected by Chiari malformation, syringomyelia, and related disorders.

The spirit of “Fly Me to the Moon” reminds us to dream boldly. To reach higher. To believe that even the most complex medical challenges can be met with innovation, collaboration, and hope. Every event we host, every dollar raised, and every story shared moves us closer to breakthroughs that once felt out of reach.

This evening, we are proud to honor Kristin Means, whose dedication as a patient advocate and long-standing unite4answers Chair has helped strengthen and grow this community year after year. Her leadership reflects the very heart of our mission — compassion in action and hope in motion.

But tonight belongs to all of us. To our sponsors, our volunteers, our families, and our friends — thank you for believing in this cause and for helping us aim higher than ever before.

So high rollers and hopeful dreamers, it’s time to play your hand for a purpose. Let the chips fly, raise the stakes, and double down on doing good. Because when we gather like this — united by heart and driven by hope — we don’t just wish upon the moon...

We reach it! 🌙 ✨

Thank you for your generosity,

Dorothy J. Poppe





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Pam Fenner
Board of Directors Secretary

HONOREE REMARKS

Kristin Means
Honoree

GAMING

7:00 – 10:15pm

BASKET RAFFLE

7:00 - 9:30pm

SILENT AUCTION

Closes at 10:00pm



Thank you

WE ARE GRATEFUL TO OUR 2026 SPONSORS

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Kristin Means

*Bobby Jones CSF Patient Advocate
unite4answers Chairperson*



Kristin Means, a former schoolteacher and athlete, found a new purpose as a passionate advocate for families navigating complex medical diagnoses. After her 10-year fight for a diagnosis, suffering setback after setback, Kristin was diagnosed with Ehlers-Danlos Syndrome and Chiari malformation. Months later, both of her sons were diagnosed as well. Surviving 41 surgeries, Kristin knows firsthand the pain of being misunderstood by the medical system and the strength it takes to keep fighting. Today, she uses her voice to empower other families to trust their instincts, push for answers, and never apologize for advocating for their children.

Kristin discovered Bobby Jones CSF in 2010 and the organization quickly became a second family. Kristin has served as Walk Chair for a decade and has watched the event she leads grow and evolve with each passing year. She has also spent countless hours working to educate and guide other individuals and families suffering from similar circumstances. She is an active member in support groups and has worked tirelessly to try to provide support to anyone who needs it.

Kristin believes that while chronic illness may shape your story, it does not define your worth or limit your impact. With the unwavering support of her family, Kristin is able to keep a smile on her face and a positive attitude. She continues to navigate her complicated health as well as her two sons' and lives by the quote, "It's hard to beat a person who never gives up."

Congratulations

Kristin Means



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**ARLIA
& ASSOCIATES**

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"Your bridge to a positive future"

To our esteemed honoree

Kristin Means

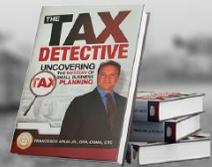
We admire and respect your dedication to
Bobby Jones Chiari & Syringomyelia.

May your family, friends, and the entire community follow
in your footsteps to help perpetuate the great work you
accomplished.

Also thank you to Dorothy Poppe, Kaitlyn Esposito,
Mary Poppe and Andrea Grosz for all your hard work and
dedication, always while wearing a smile!

We wish you continued success in all your endeavors.
Congratulations!

Frank Arlia & the staff at Arlia & Associates CPAs LLP



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Congratulations
to the
Bobby Jones
Chiari & Syringomyelia Foundation
with special Recognition to
Dorothy Poppe &
our Board Member, Kaitlyn Esposito



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**Congratulations to
this evening's honoree
Kristin Means**

**Your dedication to and support of
patients and Bobby Jones CSF is an
inspiration to us all.**

*Proud to support
Bobby Jones CSF
and this year's
honoree
Kristin Means
at the
2026 Casino Night*

*With love from
John Esposito*

Proud to support the
Bobby Jones Chiari &
Syringomyelia Foundation
Casino Night and to honor
Kristin Means



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*“Unless someone like you cares a whole
awful lot, nothing is going
to get better. It’s not.”*

—Dr. Seuss

We are proud to once again be a part of
this wonderful evening dedicated to
finding a cure for Chiari malformation and
syringomyelia.

We would like to congratulate Kristin
Means and thank you for your tireless
efforts in bringing awareness to these
diseases and your countless support to the
families who are affected by them.

Fondly,

The Farrell & Kosty Families





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KRISTIN MEANS

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**The Bobby Jones Chiari & Syringomyelia Foundation
and their research and educational efforts.**

Thank you for all you do.

God speed your efforts!

**Cesare Lucido
Grand Knight**

**Frank Tansey
Deputy Grand Knight**

**Matthew Leavy
Chancellor**

**Jack Lamantia
Brother Knight**



The Cleary's send our
love and support to our
cousin Paul and this
wonderful Foundation!

With all our admiration,

Jim and Lisa
Bob and Peggy
Ed and Kathy
John and Mai
Tom
Scott and Mary
Bill
Gary and Debbie



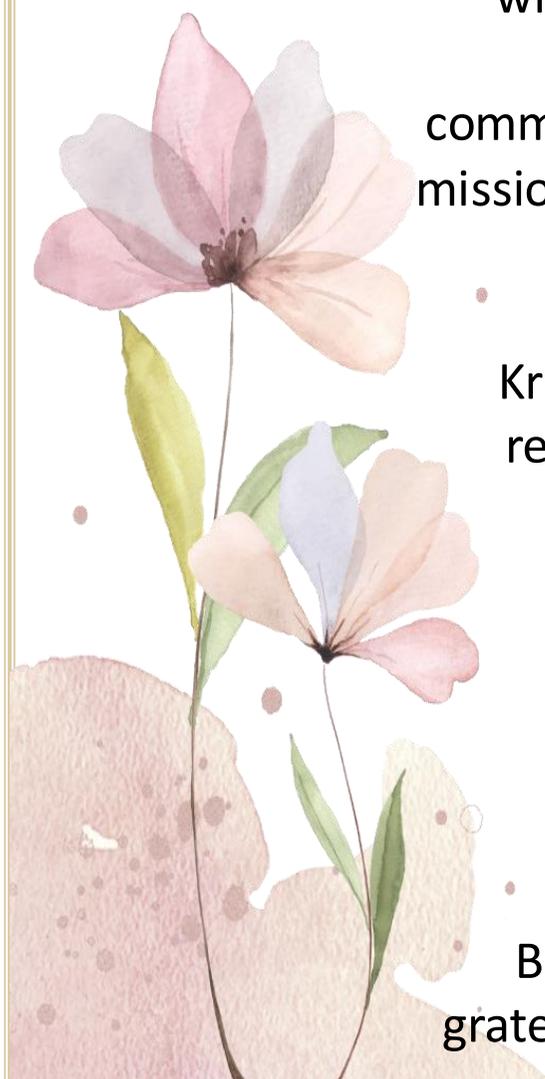
Congratulations to
Our Casino Night Honoree,
Kristin Means!

Tonight, we celebrate a
champion — someone
whose compassion,
generosity, and
commitment uplift our
mission and inspire our
community.

Kristin, your impact
reaches far beyond
this evening.

Thank you for
standing with us
and helping
change lives.

Bobby Jones CSF is
grateful to honor you!





Kristin,

Your strength and friendship mean the world to me! You've been more than a friend all of these years; you've been a source of comfort, courage, and understanding in a way only someone who truly "gets it" can be.

With love and appreciation always, Pam

Proud to Support the Bobby Jones Chiari & Syringomyelia Foundation

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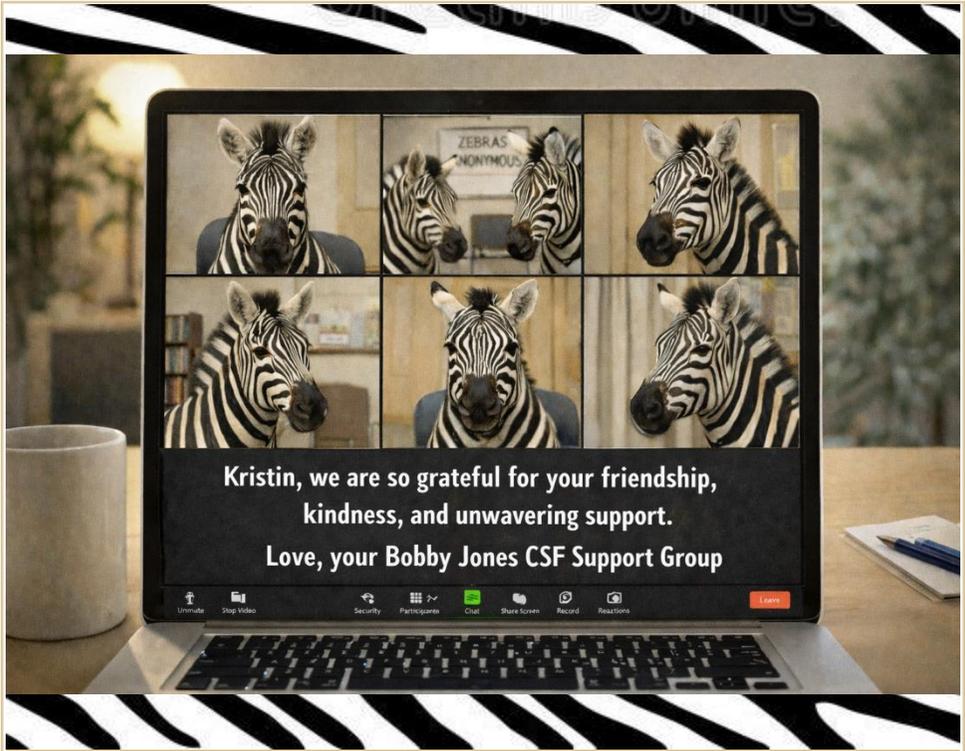
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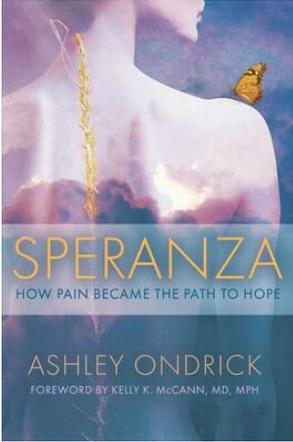
Daniel,
With all our love
to eternity and
beyond.

Love,
Nanny & Poppy



"The finest of the genre, intriguing and hard to put down, this book provides an extraordinary encouragement for persons suffering with chronic conditions. This is a must read, told through the culinary perspective of a young private chef with Ehlers Danlos Syndrome"

—Fraser C. Henderson Sr. MD,
Co-Author of Symptomatic: The Symptom-
Based Handbook for Ehlers Danlos Syndromes
and Hypermobility Spectrum Disorders



Speranza—Italian for hope—is more than a memoir. It's a feast for the senses and a testament to grace. For anyone living with chronic illness or aching for renewal, Ashley's story offers comfort, courage, and the reminder that even in our weakest moments, hope can rise.

Preorder HERE:

<https://www.mostlyhealthychef.com/speranza-book>



Thank you, Kristin for everything
that you do and always being YOU!

Tonya

"Play it as it Lies" Open
FOR BOBBY JONES CSF

OCTOBER 20, 2026

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