



*5<sup>th</sup> Annual*



**TIJERAS CREEK GOLF CLUB**

**Friday, June 6, 2025**

# Reception Schedule

4:00PM

COCKTAIL HOUR, CASINO TABLES OPEN,  
& SILENT AUCTION PREVIEW

4:30PM

HELICOPTER BALL DROP

5:00PM

DINNER SEATING

5:10PM

NATIONAL ANTHEM  
BY ANTHONY BERNASCONI

5:15PM

WELCOME REMARKS

JOHN PORRELLO & MARK HERSHEY  
GOLF CLASSIC EVENT CO-CHAIRS

5:20PM

LIVE AUCTION

5:35PM

DR. NAMITA GOYAL, MD &  
DR. TAHSEEN MOZAFFAR, MD  
UCI ALS NEUROMUSCULAR CENTER

5:40PM

FUND-A-NEED

5:50PM

AWARDS

*Checkout*

5:55PM

SILENT AUCTION CLOSES

6:00PM

COLLECT ITEMS  
EVENT CLOSES

# Golf Committee

JOHN PORRELLO &  
MARK HERSHEY

Co-Chairs

NATALIE VILLEGAS

Executive Director  
ALS United Orange County

DENISE GREEK

Development & Operations Manager  
ALS United Orange County

MARK CHAVEZ

Project Coordinator  
ALS United Orange County

LIZ DORAM

Procurement Manager  
ALS United Orange County

MICHAEL BASSETT

Consultant

JACK EVANS

Consultant

CAROLYN PLASCENCIA

Director of Client Development  
and Retention  
PriorityWorkforce

CLARA LOPEZ

Client Concierge Coordinator  
PriorityWorkforce

CHRISTINA VOGT

Assistant Director, Marketing & Sales  
PriorityWorkforce

**THANK YOU TO ALL**  
*the hands that helped make  
this golf tournament  
the best yet!*



ALS United Orange County is committed to supporting individuals and families impacted by Amyotrophic Lateral Sclerosis (ALS). Through collaboration with the community, medical experts, and researchers, we aim to improve quality of life, raise awareness, and advance effective treatments toward finding a cure for this devastating disease. Our dedication remains steadfast as we stand united in the fight against ALS.

## *Thank You!*

Thank you for participating in our 5th Annual PriorityWorkforce, Oak Laguna Beach, PEO Exchange and ALS United Orange County Golf Tournament at Tijeras Golf Creek.

This golf tournament, dinner, and auction event contribute to ALS United Orange County by funding research, equipment loan programs, patient and family support initiatives, case management, patient advocacy and education, transportation, and respite care for families affected by ALS (Lou Gehrig's Disease), an illness that currently has no a cure.

## *Save the Date!*

**ALS GOLF TOURNAMENT**  
**Friday, June 5, 2026**

*Reserve your spot now for only \$75!*



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\* DEPOSITS ARE TAX-DEDUCTIBLE & NON-REFUNDABLE.  
BALANCE DUE ON OR BEFORE APRIL 26, 2026.

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**2021 • \$161,750**

**2022 • \$167,466**

**2023 • \$205,107**

**2024 • \$225,628**

**TO REACH \$1 MILLION**  
**WE NEED TO RAISE**  
**THIS YEAR**

**2025 • \$240,049**

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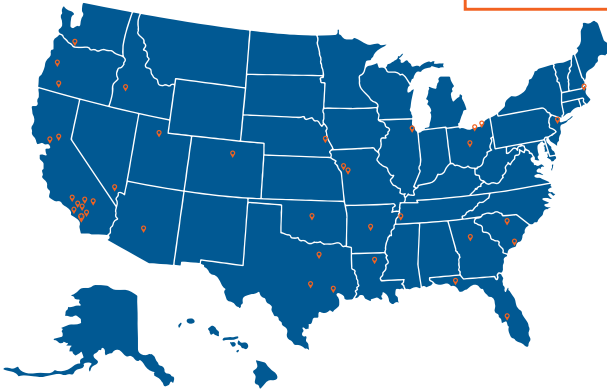


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# UCI Health ALS & Neuromuscular Center

At the UCI Health ALS & Neuromuscular Center, we provide comprehensive care for patients facing ALS and other neuromuscular conditions, including peripheral neuropathy, myasthenia gravis, and muscular dystrophies. Our expert team integrates leading-edge research, diagnostic excellence and personalized treatment plans to address each patient's unique needs. As part of Orange County's only academic health system, we are committed to advancing clinical care, offering access to clinical trials and providing holistic support for patients and their families throughout their care journey.



**Namita A. Goyal, MD**  
Neuromuscular Medicine, Neurology  
ALS United OC Board of Directors



**Tahseen Mozaffar, MD**  
Neuromuscular Medicine, Neurology

# UCI Health

# *In Loving Honor of* ISABEL PORRELLO

Isabel Marie Porrello was born March 29th, 1945 and passed away on July 9th, 2003 from ALS. She was diagnosed 13 months prior to her death but it was suspected that the disease was taking over well before that. My mother was born in New York City and moved to California with my father and started their life in Los Angeles. She was a mother of three ... kind, caring and loved with a big heart. She faced this horrible disease head on, never felt sorry for herself and always remained positive. She laughed, smiled and joked often, even up until her final days. She was loved by all and touched the hearts of so many during and after she passed.

When our mother was diagnosed with ALS, it changed our lives forever. She wasn't only a mother, wife and grandparent, but she was the Matriarch of our family. We quickly had to make mom comfortable and learn to start living without the strongest, most loving person in our entire family. We found a way to get through it, but needless to say, it was very challenging.

After mom passed, we felt it was important to keep her legacy alive by supporting our local and National ALS Associations in her honor. We were blessed to have access to resources that many may not always be able to obtain. Therefore, we are doing our best to give back and help those who are diagnosed with this horrible disease, along with their families and caregivers.

Supporting this cause is and will always be very important to us.

*Written by John Porrello*

Isabel's son &  
ALS Golf Classic Co-Chair



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# Veteran JOSEPH SCRIVNER

Born and raised in Southern California, Joe joined the United States Marine Corps in 2012 at the age of 18 and served 5 years. He deployed 3 times, rose to the rank of Corporal and became a subject matter expert on urban combat and amphibious raids.

After his service he followed in his family's footsteps of working in the construction trade and started his own company alongside his girlfriend, Cat, providing home improvement services throughout Orange County. Joe now lives in Anaheim with his girlfriend and their 3 kids: 6-year-old Isabella, 4-year-old Cody, and 2-year-old Sophia. Together they are learning to navigate work, kids and life with a terminal incurable illness, ALS.



## 2025 LOU GEHRIG DAY

Lou Gehrig Day, held annually by MLB on June 2nd, marks the day Gehrig became the starting first baseman for the New York Yankees, the beginning of his incredible streak of consecutive games played.



# *In Honor of* JODY BLACK

My sister, Jody Black, was a focused and intelligent woman who cared deeply for her family and had a great passion for education. Her idea of a great vacation was sitting in her pajamas with her dog and a big stack of books. She was a loving mother and wife, and an amazing daughter, appreciated the time she spent with other PALS at lunches, holiday parties, walks, and the National ALS meeting sister, aunt, and friend. She was never okay with any of us settling for second best. Her diagnosis with ALS in 2013 forever altered our lives. We had no way of knowing how to wage a fight against this ruthless disease – a disease that has no cure and robs you of hope.

ALS United Orange County provided us with the light we desperately needed to create the best possible life for Jody and to accept the tough reality we faced. Their support came in many forms – healthcare, education, equipment, camaraderie, and most importantly, emotional support. Jody deeply in Washington, D.C., where she somehow managed to pull off a Dukes of Hazzard styled two-wheel maneuver on her three-wheel scooter. Our champion lost her battle in 2015 and we are forever grateful to ALS United Orange County.



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