



VOLUNTEER

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JANUARY 2024



HAPPY NEW YEAR!!

Welcome to 2024 and our 28th year as a leading and influential Sarcoma organization within the U.S. focusing on anyone affected by Sarcoma in the Pacific Northwest. Our history is rich, especially as the oldest Sarcoma organization in the U.S., we continue to strive to add programs and outreach within our Sarcoma communities.

As we launch into this new year, and you may begin to think about your resolutions please consider the Northwest Sarcoma Foundation as a place for your time and energy. We are

welcoming 4 new board members in 2024, along with new committee members, plus we are always searching for new committee members. Maybe this is the year you join us in our efforts.

We are excited to be bringing back programs that were on hold until we could meet in person again. We had our 1st Community Fun event in Seattle in Oct. 2023, but are launching them further in 2024 to include hosting locations in Oregon. Another return and expansion this year is the Pediatric Days. We bring fun to any cancer patient within certain medical centers in Washington and Oregon. The hope is to add more states as our funding continues to increase.

A few pages down you will see what we have currently in store for 2024! Mark your calendars.



Volunteer with us!

Upcoming Committee meetings

Programs Committee 1.15.2024 usually 2nd Mon. ea. month 1 p.m. PT Events Committee 1.11.2024 2nd Thurs. ea. month 6 p.m. PT

Marketing-Communications Committee 1.10.2024 1st Wed. ea. month 3 p.m.PT *Moved in Jan. 2024 to accommodate a schedule change.

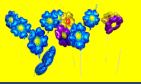
Development Committee 1.9.2024 2nd Tues. ea. month 6 p.m. PT

YOUR BOARD OF DIRECTORS

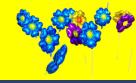


During 2023 we welcomed 2 new board members Elana Emlen and Dr. Kenny Gundle. At the start of 2024 we are
welcoming 4 new board members -

Mike Bell, Nick Casillas, Mike Lillie, and Bill O'Connell!



DRAGONSLAYER TRIBUTE FUNDS DECEMBER NEW TRIBUTE FUNDS





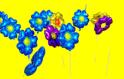




MATT EMLEN

Matt Emlen was someone who people would meet, and immediately feel comfortable and happy to be talking with him. He knew so much about history, politics, and music; and he was always curious about people and the world. He was not defined by his job but approached his career in resource conservation strategies with integrity and steadfastness.

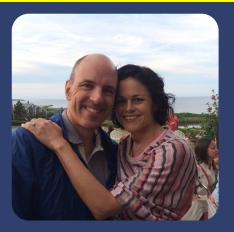
Matt's full story HERE.



DRAGONSLAYER TRIBUTE FUNDS JANUARY'S NEW TRIBUTE FUNDS









MARIA ANDERSON

In thinking about Maria's journey, I always go back to those years between ages 27 and 43, when Maria suffered through surgeries, procedures, and pain management on her right arm. Ultimately, I wonder what could have happened if Maria's doctors had been exposed to some of Chappie's expertise here in Seattle. In this way, the work of the Northwest Sarcoma Foundation is very real for me.

TRIBUTE FUNDS

Maria's full story **HERE**

Dragonslayer Tribute Funds are established by friends and/or families to honor the lives of specific Sarcoma patients, survivors, or loved ones lost to Sarcoma. These funds assist the Northwest Sarcoma Foundation in its mission and ensures that

No One Faces Sarcoma Alone You helping to create *thri-vors* in 2024!

2024 NWSF EVENTS



IANUARY

Patient Education Night

25 Topic: Chemo! Radiation! A Port! OH MY! RN Panel Discussion

FFBRUARY

Researcher Update Night

21 Topic: Tenosynovial giant cell tumor

Dr. Elizabeth Loggers

MARCH

O6 Dragonslayer Days
Seattle Children's Hospital

TBD Sarcoma Journal Club

Dr. Lee Cranmer

APRIL

Dragonslayer Days

TBD Hope Grows Here

Spokane

27 Hope Grows Here

MAY

Hope Grows Here

Dragonslayer Days

Providence Sacred Heart

Sarcoma Journal Club **TBD**

Patient Education Night

TBD Topic: Radiation Therapy and what it means for me

Dr. Art Hung

JUNE

Researcher Update Night

Topic: PEComas and new

Dr. Lee Cranmer

treatments

Adult Patients and Survivors Casual Chat - 2nd Wed. each month Caregivers Casual Chat - 3rd Tues. each month Sarcoma Bereavement Group - 4th Mon. each month





Casual Chats are free events, register once, the same Zoom link each time. Adult patients & survivors: Second Wednesday of every month.

Next meeting, 1/10/2024 6 - 7 p.m.

Led by either Jo McNeal or Chelsea Winn

<u>Caregivers:</u> Third Tuesday of every month.

Next meeting, 1/16/2024 6 - 7 p.m.

Volunteer peer-led by Anita Nelson

Sarcoma Bereavement: Fourth Monday of every month.

Next meeting, 1/22/2024 6 - 7 p.m.

Volunteer peer-led by Elana Emlen

PATIENT EDUCATION NIGHT

Jan. 25 @6:30 PM PST VIA ZOOM

Chemo! Radiation! A Port! OH MY!



2024 NWSF EVENTS



Register!

JULY

Researcher Update Night

23 Topic: Chondrosarcoma and chonDragon tx Dr. Lara Davis

AUGUST

Researcher Update Night

o6 Topic: Gastrointestinal stromal tumor (GIST)
Dr. Limin Zhu

SEPTEMBER

O7 Dragonslayer Walk

Portland

Dragonslayer Walk

Boise

Dragonslayer Walk

Spokane

Dragonslayer Walk

Seattle

OCTOBER

Patient Education Night

16 Topic: Palliative care

Dr. Elizabeth Loggers

NOVEMBER

06 Sarcoma Journal Club

Dr. Jeremy Sharib

DECEMBER

Patient Education Night

Topic: Pathology - The hidden heroes

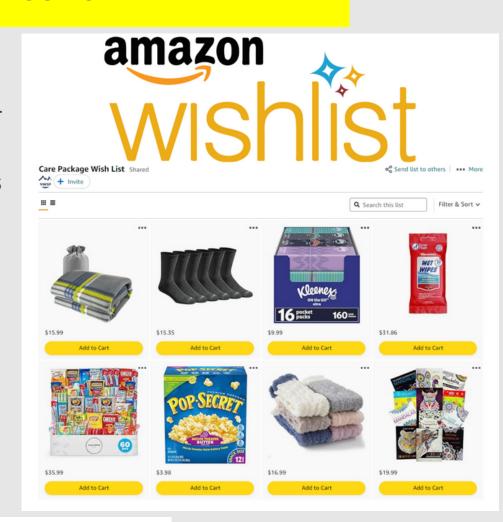
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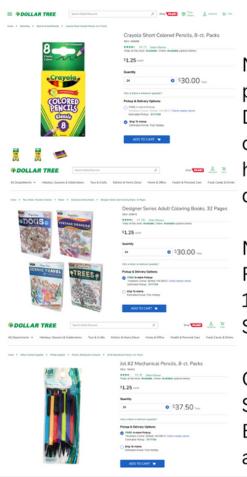


CARE PACKAGES NEED YOUR CARE

In 2023, we provided a record number of care packages. Our supplies on specific items are getting low, and we are always in need of additional supplies. Please consider helping restock our shelves.

Items in most need:
Blankets & men's warm
socks.





POLLAR TREE

Not an Amazon purchaser, consider the Dollar Tree. You can order items in bulk and have them shipped to us directly.

Northwest Sarcoma Foundation 117 E. Louisa St., #443 Seattle, WA 98102

Or if you live in the Seattle, Portland, or Boise areas, we can arrange to pick up.





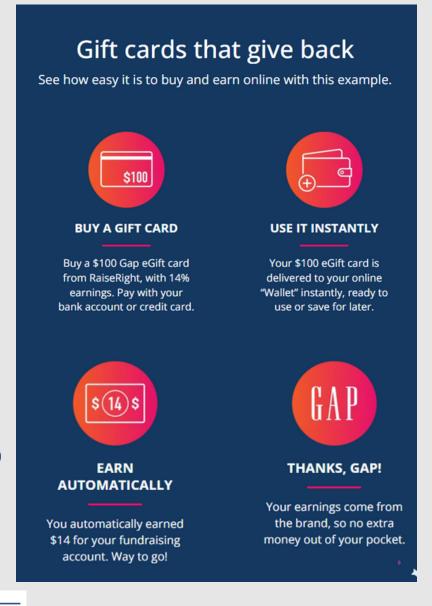
OTHER DONATION OPTIONS

Fundraise with Raise Right. You can purchase gift cards (eCards) through our Raise Right account and a percentage of your purchase comes back to us. Plus you can donate gift cards to grocery stores, Starbucks, etc. to the care package program. Just choose to have them physically mailed to NWSF.

During the holidays, Exec. Dir. Jo registered, purchased, and donated \$18 back to NWSF!

Register and get started today. Create a free account on the RaiseRight mobile app or at RaiseRight.com.

Enrollment Code: TJ4HZXHS2PA6 Have questions contact: info@nwsarcoma.org





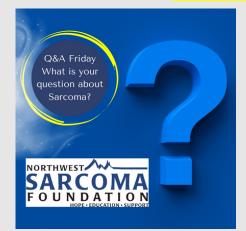
Link your Fred Meyer rewards card to donate back to the NW Sarcoma Foundation!

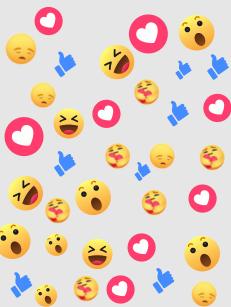
Northwest Sarcoma Foundation - Seattle (HN504)



DECEMBER'S SOCIAL MEDIA BESTS







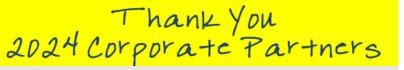
Is there a national hub where all information is being compiled and sorted?

There are some efforts to establish databases to collect information on sarcoma patient outcomes on a large scale, but there are many complicating factors like making sure that all of the data is consistently and reliably entered and regulatory issues to ensure protection of patient data and patient privacy. On a national (and international) level, many sarcoma centers collaborate on clinical trials, database work, and basic science to establish sarcoma treatment guidelines and to find new (and hopefully better!) ways to treat sarcomas in the future. Even with the high level of collaboration between sarcoma specialists, there are sometimes differences of opinion and it never hurts to get a second opinion to discuss a treatment plan.

-Michael Wagner M.D.



Each month we host Q&A Friday. I'm thrilled that our question and answer in December were so well received. Thank you, Dr. Michael Wagner for providing the answer!















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