FOR ADDITIONAL INFORMATION REGARDING YOUR CONDITION OR DIAGNOSIS, YOU CAN VISIT THE FOLLOWING WEBSITES:

Condition	Additional Resources
Atopic Dermatitis	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/adult-dermatology** **Asthma and Allergy Foundation of America** The Asthma and Allergy Foundation of America (AAFA), a not-for-profit organization founded in 1953, is the leading patient organization for people with asthma and allergies, and the oldest asthma and allergy patient group in the world. **https://www.aafa.org/eczema/** **National Eczema Association** The National Eczema Association (NEA) is a non-profit, 501(c)(3) organization with a mission to improve the health and quality of life for individuals with eczema through research, support, and education. **https://nationaleczema.org/**
Crohn's Disease	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/digestive-and-liver-health** **Crohn's & Colitis Foundation of America** The Crohn's & Colitis Foundation of America (CCFA) is a non-profit, volunteer-driven organization dedicated to finding the cures for Crohn's Disease and ulcerative colitis. **http://www.ccfa.org/science-and-professionals/programs-materials/patient-brochures/** **Crohn's Online** This web page provides individuals who have Crohn's with education, support and treatment options. It gives Crohn's patients access to their Crohn's Patient Advocate Program which will provide them or their loved one with one-on-one education and free support. **http://www.crohnsonline.com/** **Crohn's Forum** Crohn's Forum is a support group and forum for people who are personally affected or who know someone who is affected by Crohn's Disease, Ulcerative Colitis and other IBDs. The forum is a community that offers support, friendship and understanding. Crohn's Forum also has a Twitter page where you can get instant updates and news articles about Crohn's. **http://www.crohnsforum.com/**
Cystic Fibrosis	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/pulmonary/cystic-fibrosis**

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	Cystic Fibrosis Foundation The Cystic Fibrosis Foundation is a nonprofit organization that provides information, assistance services, and support to those with cystic fibrosis. Their mission is to search for a cure for cystic fibrosis through extensive research that is donor-funded. **https://www.cff.org/** **Michigan Pulmonary Disease Community** The Cystic Fibrosis Support Network of Michigan is a non-profit Michigan Charity dedicated to providing direct and indirect services to enhance the lives of people with cystic fibrosis, their families, and the communities that support them. They work year-round providing support and financial assistance. **http://www.mpdci.org/**
Eosinophilic Asthma	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/pulmonary/asthma** **American Partnership for Eosinophilic Disorders** The American Partnership for Eosinophilic Disorders (APFED) is a 501c3 nonprofit organization founded in December 2001 by a group of mothers of young children living with eosinophil-associated diseases. We are a patient advocacy group dedicated to improving the lives of those living with eosinophilic disorders. **https://apfed.org/about-apfed/mission-and-objectives/** **Asthma and Allergy Foundation of America** The Asthma and Allergy Foundation of America (AAFA), a not-for-profit organization founded in 1953, is the leading patient organization for people with asthma and allergies, and the oldest asthma and allergy patient group in the world. **https://www.aafa.org/**
Hepatitis	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/digestive-and-liver-health** **American Liver Foundation** The American Liver Foundation was created in 1976 with a mission "To facilitate, advocate and promote education, support and research for the prevention, treatment and cure of liver disease." The American Liver Foundation has programs for education, Hepatitis C, Liver Wellness and Liver Disease. They also provide support services which include a help line, support guide and support groups and resources like Inspire, NeedyMeds and CaringBridge. **http://www.liverfoundation.org/** **Hepatitis Central**

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	Hepatitis Central is an organization that provides people with Hepatitis C information, news, treatments and support. Hepatitis Central can provide you with information about support groups in your area along with online support. Hepatitis Central also provides a free newsletter to keep you up to date with the latest news on hepatitis treatments, clinical trials, social issues and important breakthroughs. **http://www.hepatitis-central.com/** **Hepatitis B Foundation** The Hepatitis B Foundation is dedicated to finding a cure and improving the quality of life for those affected by hepatitis B worldwide. The Hepatitis B Foundation's commitment includes funding focused research, promoting disease awareness, supporting immunization and treatment initiatives, and serving as the primary source of information for patients and their families, the medical and scientific community, and the general public. **http://www.hepb.org/resources/printable_information.htm** **Centers of Disease Control and Prevention** CDC works 24/7 to protect America from health, safety and security threats, both foreign and in the U.S. The CDC fights disease and supports communities and citizens to do the same. • CDC's Hepatitis B Patient Education Resource http://www.cdc.gov/hepatitis/hbv/patienteduhbv.htm • CDC's Hepatitis C Patient Education Resource http://www.cdc.gov/hepatitis/hcv/patienteduhcv.htm
Hereditary Angioedema	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/allergy** **Hereditary Angioedema Association, Inc.** Founded and staffed by HAE patients and HAE patient caregivers, U.S. Hereditary Angioedema Association, Inc. (US HAEA) is a non-profit patient advocacy organization dedicated to serving persons with angioedema. The Association provides HAE patients and their families with a support network and a wide range of services including physician referrals, and individualized patient support. **https://www.haea.org/** **National Organization for Rare Disorders** The National Organization for Rare Disorders (NORD) is a patient advocacy organization dedicated to individuals with rare diseases and the organizations that serve them. NORD, along with its more than 300 patient organization members, is committed to the identification, treatment, and cure of rare disorders through programs of education, advocacy, research, and patient services. **https://rarediseases.org/**
Immune Deficiency	**Michigan Medicine**

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and Related Disorders	**https://www.mottchildren.org/conditions-treatments/peds-immunodeficiencies** **American Academy of Allergy Asthma & Immunology** Founded in 1943, the American Academy of Allergy, Asthma & Immunology is a professional medical membership organization of nearly 6,800 allergist/immunologists and related professionals around the world with advanced training and experience in allergy, asthma and other immunologic diseases. **https://www.aaaai.org/** **Immune Deficiency Foundation** The Immune Deficiency Foundation, founded in 1980, is the national non-profit patient organization dedicated to improving the diagnosis and treatment of patients with primary immunodeficiency diseases through research, education and advocacy **https://primaryimmune.org/** **Chronic Granulomatous Disease Association** The Chronic Granulomatous Disease Association (CGDA), founded in 1982, is a non-profit international support group for persons with chronic granulomatous disease (CGD), their families and physicians. The organization networks patients with similar CGD-related illnesses or infecting organisms. It provides research grants aimed at finding a cure for CGD. **http://www.cgdassociation.org/**
Multiple Sclerosis	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/brain-neurological-conditions/multiple-sclerosis-ms** **Multiple Sclerosis Association of America** The Multiple Sclerosis Association of America (MSAA) is a national, nonprofit organization founded in 1970 and is dedicated to improving lives today. MSAA provides ongoing support and direct services to individuals with MS, their families, and their care partners. **http://www.mymsaa.org/** **Multiple Sclerosis Foundation** The Multiple Sclerosis Foundation (MSF) helps individuals affected by multiple sclerosis maintain their health and well-being. MSF offers programs and support to keep those with multiple sclerosis self-sufficient and safe in their homes. They also provide educational programs to increase public awareness and understanding of their own disease. MSF offers support groups for people with MS and their families to share their experiences. **http://www.msfocus.org/** **National Multiple Sclerosis Society**

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	The National MS Society is a group of individuals whose mission is to "mobilize people and resources to drive research for a cure and to address the challenges of everyone affected by MS." The National Multiple Sclerosis Society has a chapter in all 50 states and an online community to help those who have the disease. The National MS Society also offers hundreds of special events a year to engage participants to support their mission. **http://www.nationalmssociety.org/**
Oncology	**Michigan Medicine** **https://www.mcancer.org/** **American Cancer Society** The American Cancer Society provides a wide range of programs in the areas of Research, Education, Support, and Advocacy. Their mission is to help improve the quality of life of cancer patients while working toward prevention and a cure. **https://www.cancer.org/** **Patient Advocate Foundation** At LIVESTRONG, support services are provided to anyone affected by cancer. They also have the ability to connect people and communities with specific services needed. **https://www.livestrong.org/we-can-help**
Psoriasis	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/adult-dermatology/psoriasis-eczema-other-inflammatory-skin-conditions** **National Psoriasis Foundation** The National Psoriasis Foundation (NPF) is "working to find a cure for psoriasis and psoriatic arthritis and to eliminate their devastating effects through research, advocacy, and education." The NPF can help those with psoriasis and psoriasis arthritis learn about types, treatments, living well, related conditions, accessing care and more. The NPF can also keep you up to date with different events in your area and has a network of support groups in different cities. **http://www.psoriasis.org/**
Rheumatoid Arthritis	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/cmc/arthritis/rheumatoid** **American College of Rheumatology** The American College of Rheumatology's (ACR) mission is advancing rheumatology. The organization represents over 9,400 rheumatologists and rheumatology health professionals around the world. The ACR offers its members the support they need to ensure that they are able to continue their innovative work by providing programs of education, research, advocacy, and practice support.

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	https://www.rheumatology.org/practice/clinical/patients/diseases_and_conditions/osteoporosis.asp **Arthritis Foundation** The Arthritis Foundation is the largest nonprofit organization that supports 100 different types of arthritis and related conditions and has service points located throughout the country. The Arthritis Foundation provides information and news on the different types of arthritis and can keep you up to date about different events and support programs related to arthritis. **http://www.arthritis.org/** **National Institute of Arthritis and Musculoskeletal and Skin Diseases** The mission of the National Institute of Arthritis and Musculoskeletal and Skin Diseases is to support research into the causes, treatment, and prevention of arthritis and musculoskeletal and skin diseases. **http://www.niams.nih.gov/** **Rheumatoid Arthritis Life Guide** Rheumatoid Arthritis Life Guide is a resource that can provide you with information about living with rheumatoid arthritis. They provide information about understanding, diagnosing, treating and living with rheumatoid arthritis. **http://www.rheumatoidarthritis.com/**
Systemic Lupus Erythematosus	**Michigan Medicine** **https://www.uofmhealth.org/conditions-treatments/rheumatology/lupus** **Centers of Disease Control and Prevention** CDC works 24/7 to protect America from health, safety and security threats, both foreign and in the U.S. The CDC fights disease and supports communities and citizens to do the same. **https://www.cdc.gov/lupus/index.htm** **Lupus Foundation of America** The mission of the Lupus Foundation of America is to improve the quality of life for all people affected by lupus through programs of research, education, support and advocacy. **https://www.lupus.org/** **The Lupus Initiative** The Lupus Initiative (TLI) is a national program of the American College of Rheumatology (ACR) dedicated to reducing health disparities in lupus. **https://thelupusinitiative.org/**

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