

# What is EGPA?

### Eosinophilic

Granulomatosis with

Poly-

## Angiitis



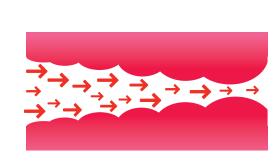




Formerly known as Churg-Strauss syndrome<sup>1</sup>, EGPA is a condition characterized by asthma, an increase in eosinophils, and inflammation of blood vessels (vasculitis).2

Vasculitis is inflammation of blood vessels such as capillaries, veins, or arteries.5

Inflamed blood vessels can reduce or stop blood flow, which may cause damage to tissue and organs.5



#### EGPA can result in damage to:





Heart





Gastrointestinal tract





and other parts of the body. It can even be life-threatening for some patients.3

#### **Eosinophil** ee-uh-sin-uh-fil

A type of white blood cell that is a normal part of the body's immune system.2

Based on global estimates, EGPA likely affects approximately



prevalence of 14.58 cases per million, based on a U.S. population of approximately 325 million in July 2017.

are affected by EGPA, and the average age of diagnosis is 48.8

Both men and women



#### The main characteristics of EGPA are:9

 The development of asthma as an adult



 Allergic rhinitis (hay fever)



Growths in the nose



An increase in

eosinophils



Of these, asthma symptoms are the most common.4

Not everyone experiences all three phases, and the phases may not occur in the same order:10

EGPA has 3 phases.

such as asthma, hay fever, and sinusitis. **Eosinophilic Phase:** Abnormally

**Allergic Phase:** Allergic reactions

high numbers of eosinophils in the blood or tissues. Vasculitic Phase: Blood vessel

inflammation.

present when a person has at least four of the following six:11 Asthma Sinus problems

EGPA can be difficult to diagnose because it is uncommon. EGPA is considered

- **Increased number of**
- **eosinophils** (higher than 10%) Nerve damage (numbness and
- pain in hands or feet)
- **Pulmonary infiltrates** (spots or
- lesions on a chest X-ray of the lungs) **Extravascular eosinophils**

(presence of eosinophils outside

of a blood vessel)

EGPA, patients may be referred to healthcare providers who specialize in treating vasculitis or eosinophilic conditions, such as allergists, immunologists, pulmonologists, and rheumatologists.12

To confirm the diagnosis of







inophilic-granulomatosis-with-polyangiitis#ref\_3233. Accessed March 2018.



Mayo Clinic. Churg-Strauss Syndrome: Symptoms & Causes. https://www.mayoclinic.org/diseases-conditions/churg-strauss-syndrome/symptoms-causes/syc-20353760. Accessed March 2018.

Masi A, et al. The American College of Rheumatology 1990 Criteria for the Classification of Churg-Strauss Syndrome (Allergic Granulomatosis and Angiitis). Arthritis Rhuem. 1990; 33(8): 1094-1100.

Seo P. Eosinophilic Granulomatosis with Polyangiitis: Challenges and Opportunities. J Allergy Clin Immunol Pract. 2016. 4(3): 520-521.

Jennette JC, et al. 2012 Revised International Chapel Hill Consensus Conference Nomenclature of Vasculitides. *Arthritis Rheum*. 2013; 65(1): 1-11. National Institutes of Health (NIH). Eosinophilic granulomatosis with polyangiitis. https://rarediseases.info.nih.gov/diseases/6111/eosinophilic-granul Baldini C, et al. Clinical Manifestations and Treatment of Churg-Strauss Syndrome. *Rheum Dis Clin North Am*. 36(2010): 527-543. Noth I, et al. Churg-Strauss syndrome. Lancet. 2003; 361: 587-94.
National Institutes of Health (NIH). Vasculitis. health-topics/vasculitis. https://www.nhlbi.nih.gov/health-topics/vasculitis. Accessed March 2018. Data on File DNG#2017N348628\_00. United States Census Bureau. U.S. and World Population Clock. www.census.gov/popclock. July 2017.

Mouthon L, et al. Diagnosis and classification of eosinophilic granulomatosis with polyangiitis (formerly named Churg-Strauss syndrome). J Autoimmun. 2014; 48-49: 99-103.

Giofreddi A, et al. Eosinophilic granulomatosis with polyangiitis: an overview. Front Immunol. Nov 2014; Vol. 5, Article 549; 1-7.