

# Systemic mastocytosis (SM) is a rare form of mast cell disorder characterised by abnormal proliferation and activation of mast cells<sup>1,2</sup>

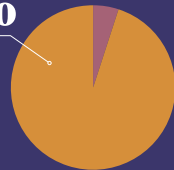
~1 in 10,000 people have SM<sup>3\*</sup>



Disease onset commonly occurs at 20–50 years of age<sup>1</sup>



SM accounts for >95% of adult mastocytosis cases<sup>4</sup>



◀ Mast cell mediator symptoms prominent

Organopathy prominent ▶

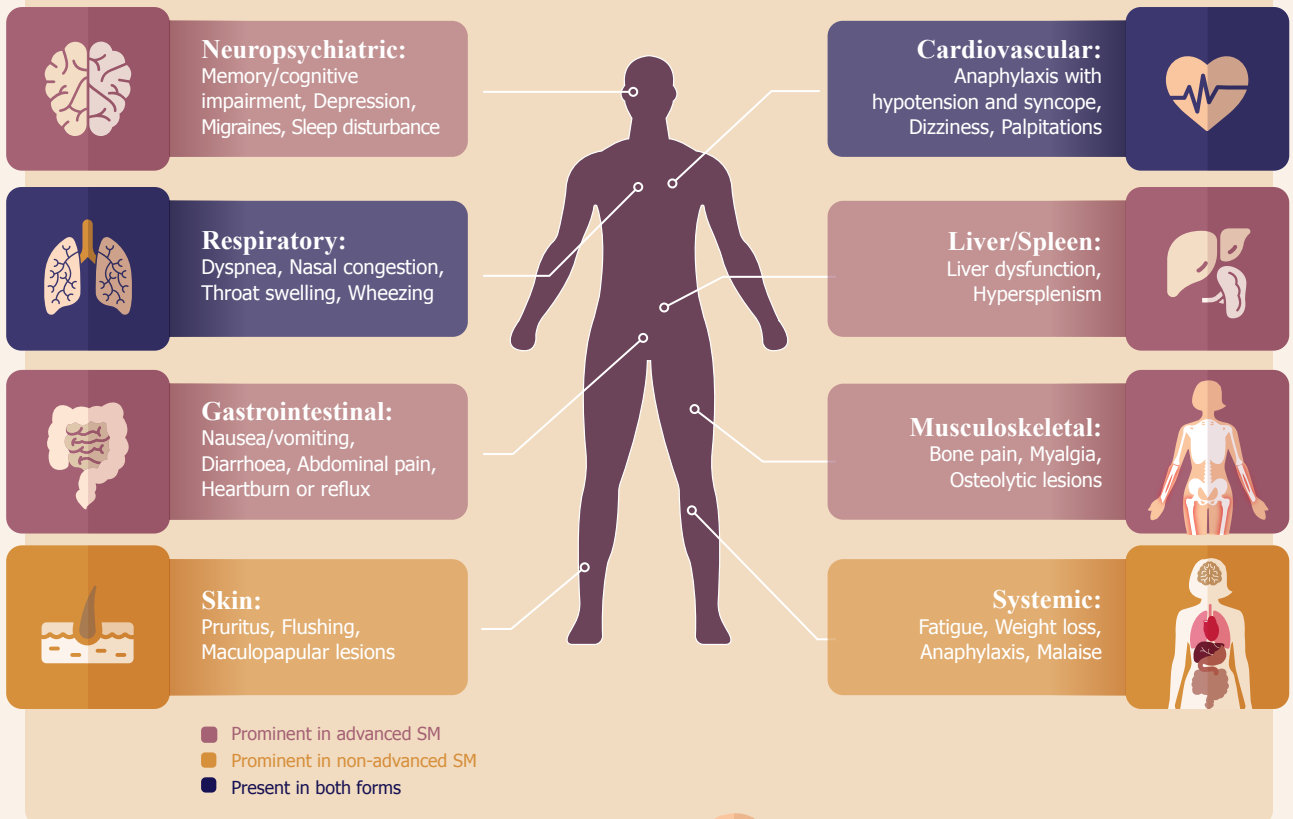
Non-Advanced SM accounts for ~90% of SM cases<sup>1,2</sup> predominantly indolent SM (ISM)

Advanced SM accounts for ~10% of SM cases<sup>1,2</sup> aggressive SM (ASM), mast cell leukaemia (MCL), associated haematological neoplasm (AHN)

## The clinical presentation often involves one or more extracutaneous sites (as well as skin involvement):<sup>1,2,5,6</sup>

Not an inclusive list of all symptoms.

The frequency and intensity of any given symptom may vary from person to person.

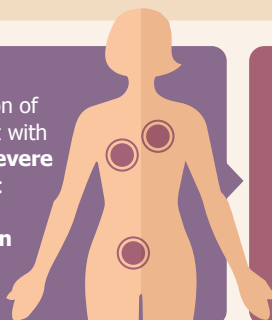


Due to the heterogeneity and unpredictability of SM symptoms, early diagnosis is challenging<sup>1</sup>

Median time from onset to diagnosis



Clinical suspicion of SM should start with recognising severe and recurrent instances of these common symptoms<sup>1</sup>



A combination of diagnostic tests, mast cell examination, high-sensitivity KIT D816V assay or serum tryptase test, is recommended to help confirm the SM suspicion<sup>7</sup>



To learn more about the real clinical impact of SM, visit: [www.systemicmastocytosis-hcp.com](http://www.systemicmastocytosis-hcp.com)

\*Based on Cohen 2014 study of 548 adults with SM diagnosed from 1997 to 2010 in linked Danish national health registries, with a 14-year limited-duration prevalence estimated at 9.59 per 100,000 as of 1 January 2011.<sup>3</sup> <sup>1</sup>Based on data from 149 patients with self-reported mastocytosis in Mast Cell Connect registry in Jennings 2018 study.<sup>9</sup>

1. Gülen T, et al. *J Intern Med.* 2016;279(3):211–228. 2. Pardanani A. *Am J Hematol.* 2021;96(4):508–525. 3. Cohen SS, et al. *Br J Haematol.* 2014;166(4):521–528. 4. Brockow K. *Immunol Allergy Clin North Am.* 2014;34(2):283–295. 5. Jennings SV, et al. *Immunol Allergy Clin North Am.* 2018;38(3):505–525. 6. Theoharides TC, et al. *N Engl J Med.* 2015;373(2):163–172. 7. Valent P, et al. *J Allergy Clin Immunol Pract.* 2022;10(8):1999–2012.