## SM symptom checklist for HCPs

## **Could your patients have SM?**

Systemic mastocytosis (SM) is a **heterogeneous disorder** characterised by abnormal proliferation and activation of mast cells. SM has an estimated prevalence of  $\sim 1$  in 10,000 and affects at least one extracutaneous organ<sup>1,2</sup>

Prolonged timespan between symptom onset and diagnosis is common due to the **range of non-specific symptoms and signs**. Therefore, clinical suspicion of SM should start with recognising recurrent instances of the common symptoms<sup>1</sup>

For more information, visit www.systemicmastocytosis-hcp.com



## Complete this simple checklist to help you recognise the symptoms or features indicative of SM\*

Symptom/feature	Severity?	No. of episodes?
<ul> <li>Skin lesions<sup>3</sup></li> <li>Maculopapular lesions: urticaria pigmentosa; raised patches of brownish skin that becomes inflamed by rubbing (Darier's sign)<sup>4</sup></li> <li>Pruritus<sup>3</sup></li> </ul>		
Elevated tryptase levels <sup>3</sup> • Basal tryptase ≥20 ng/mL <sup>5</sup> • Basal tryptase 11.5–20 ng/mL <sup>5</sup>		
Anaphylaxis in the context of a hymenoptera venom allergy, or unexplained anaphylaxis <sup>3</sup> • Often coupled with hypotension/syncope <sup>1</sup>		
Osteopenia/osteoporosis (otherwise unexplained), <sup>3</sup> especially for young patients and young males <sup>5</sup>		
Histamine-induced symptoms (e.g. cramping, headache, hypotension, diarrhoea) <sup>3</sup>		
<ul> <li>Splenomegaly<sup>3</sup></li> <li>May be suspected by physical examination, and evaluated by a complete blood count or imaging studies (e.g. ultrasound and CT imaging). A spleen size of 12–20 cm in craniocaudal length indicates splenomegaly and</li> </ul>		

If your patient experiences episodic and recurrent instances of a combination of two or more symptoms, you should consider further diagnostic workup

>20 cm is definitive of massive splenomegaly6

<sup>\*</sup>Please note that this checklist is based on the 2022 ECNM-AIM User's Guide<sup>3</sup>; this is not an exhaustive list of all symptoms in SM, and symptoms may vary between individuals.

References: 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. Valent P, et al. J Allergy Clin Immunol. 2021;10(8):1999–2012.e6. 4. Hartmann K, et al. J Allergy Clin Immunol. 2016;137(1):35–45. 5. Theoharides TC, et al. N Engl J Med. 2015;373(2):163–172. 6. Chapman J, et al. Splenomegaly. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; August 2022.