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The Burden of Indolent Systemic Mastocytosis in Europe: Results From the PRISM Patient Survey

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PRISM: Introduction

- Systemic mastocytosis (SM) is a clonal mast cell disease primarily driven by the KIT D816V mutation¹
- The majority of patients with SM have indolent systemic mastocytosis (ISM), a subtype of non-advanced disease, characterized by unpredictable, severe, and debilitating skin, gastrointestinal, and systemic symptoms^{1,2}
- A substantial number of patients with ISM experience a significant reduction in their quality of life (QoL) due to chronic, debilitating symptoms such as fatigue, abdominal pain, and anaphylactic reactions. These symptoms often lead to frequent healthcare visits and interventions, significantly affecting patients' daily activities and overall well-being^{2,3}
- Previously, the US-based TouchStone survey reported that patients with advanced SM and ISM experienced high symptom burden, poor QoL, and significant impacts on patients' ability to work⁴
- The Perceptions, Realities & Insights on SM (PRISM) Patient Survey was administered in seven European countries to understand the experiences of patients with SM
- 540 patients with SM participated in PRISM; 237 patients reported having ISM
- This analysis reports PRISM survey results for patients with ISM



PRISM: Largest European survey to assess symptomology, use of medications, and impact of SM on QoL and overall functioning

Patient Survey

- 119-item questionnaire to assess:
 - SM diagnostic journey
 - Symptom assessment (ISM-SAF), QoL (SF-12), global health assessment (EQ-5D) and work/activity impairment measure (WPAI)
 - Use of medications and healthcare services
 - Satisfaction with treatments, physician communication

Methods

- Designed by an international steering committee composed of SM clinical experts and patient advocates
- Ethics committees for participating countries (Italy, the UK, Spain, Germany, France, Switzerland, and Austria) approved the study materials
- Eligible participants ≥18 years old with a self-reported SM diagnosis who provided informed consent were recruited to participate in this survey
- Descriptive statistics on survey responses from the ISM patient cohort are reported



EQ-5D, Euro-QoL Five-Dimension; ISM, indolent systemic mastocytosis; ISM-SAF, ISM Symptom Assessment Form; PRISM, Perceptions, Realities, and Insights on Systemic Mastocytosis; QoL, quality of life; SF-12, 12-item short form survey; SM, systemic mastocytosis; WPAI, Work Productivity and Activity Impairment Questionnaire.

The majority of these patients with ISM were female with a mean age of approximately 50 years and were diagnosed on average more than 9 years ago

	All patients with ISM (N=237)	Italy (n=62)	UK (n=50)	Spain (n=49)	Germany (n=43)	France (n=15)	Switzerland (n=13)	Austria (n=5)
Age, years, mean	49.1	45.6	51.0	50.5	56.6	37.1	46.6	36.8
(SD)	(13.8)	(12.7)	(14.3)	(10.0)	(10.8)	(8.2)	(9.1)	(9.7)
Sex, female,	168	42	43	38	30	3	10	2
n (%)	(71)	(68)	(86)	(78)	(70)	(20)	(77)	(40)
Time since ISM diagnosis, months, mean	114.7	80.0	95.8	152.0	170.5	68.9	111.1	37.6

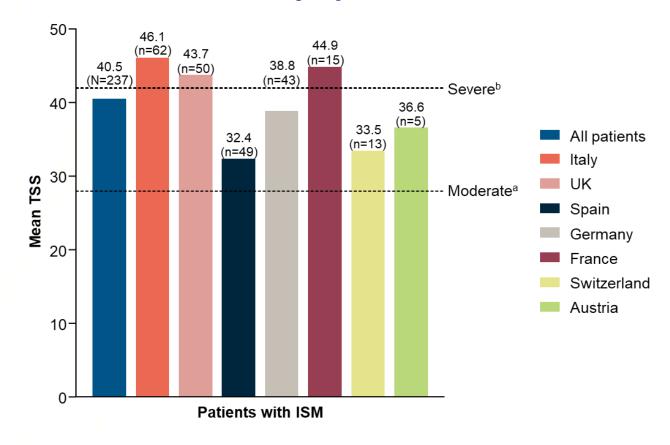


ISM, indolent systemic mastocytosis; SD, standard deviation.

Patients reported having moderate to severe ISM symptoms

Mean Total Symptom Score (TSS)

ISM Symptom Assessment Form (ISM-SAF)						
ISM Symptom	Scoring					
Abdominal pain						
Diarrhea	The ISM-SAF questionnaire calculates a					
Nausea	total symptom score (TSS) based on					
Spots	severity of 11 ISM symptoms					
Itching	Each symptom is scored 0–10;					
Flushing	where: 0 = no [symptom]					
Brain fog	10 = worst imaginable [symptom]					
Headache	Individual symptom severity scores are					
Dizziness	summed to generate a					
Bone pain	TSS (0–110)					
Fatigue						





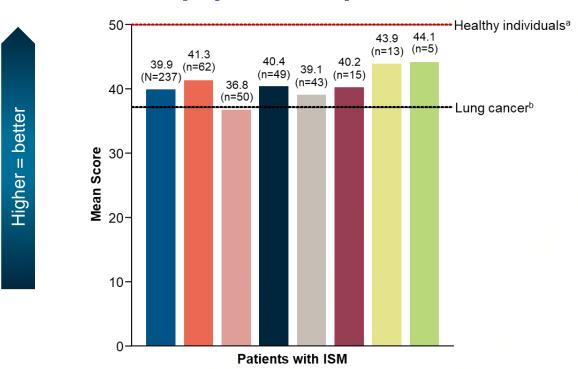
^aTSS ≥28 indicates moderate symptom profile.¹ ^bTSS >42 indicates severe symptom profile.²

ISM, indolent systemic mastocytosis; ISM-SAF, Indolent Systemic Mastocytosis Symptom Assessment Form.

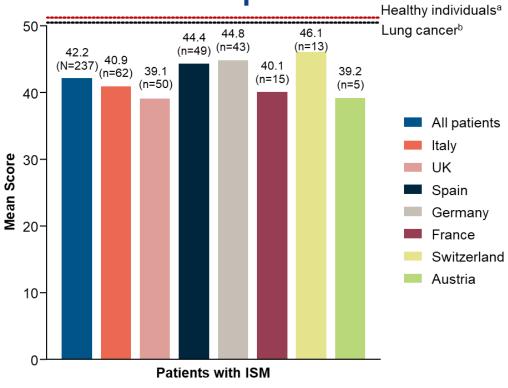
1. Shields AL, et al. *Orphanet J Rare Dis.* 2023;18(1):69; 2. Gotlib J, et al. *NEJM Evid.* 2023;2(6):EVIDoa2200339.

Patients with ISM reported substantially impaired physical and mental functioning

SF-12 physical component score



SF-12 mental component score





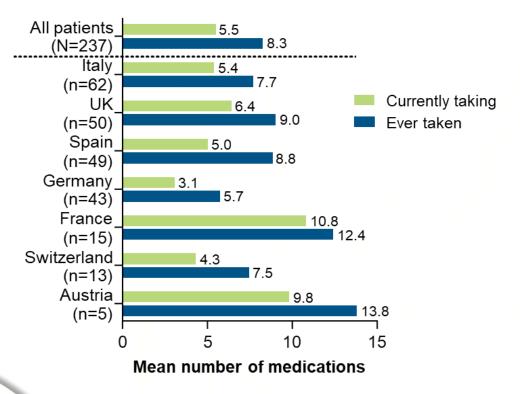
^bMean physical component score for patients with lung cancer is 37.2 and the mean mental component score is 50.1.² ISM, indolent systemic mastocytosis; SF-12, 12-item short form survey.

1. Ware JE, et al. SF-12: How to Score the SF-12 Physical and Mental Health Summary Scales. 4th ed. Quality Metric Inc..; Health Assessment Lab; 2002; 2. Kenzik KM et al. Cancer. 2015 Aug 15;121(16):2831-839.

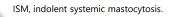


Patients reported high lifetime use of symptom-directed medications and rates of anaphylaxis

Mean number of ISM medications



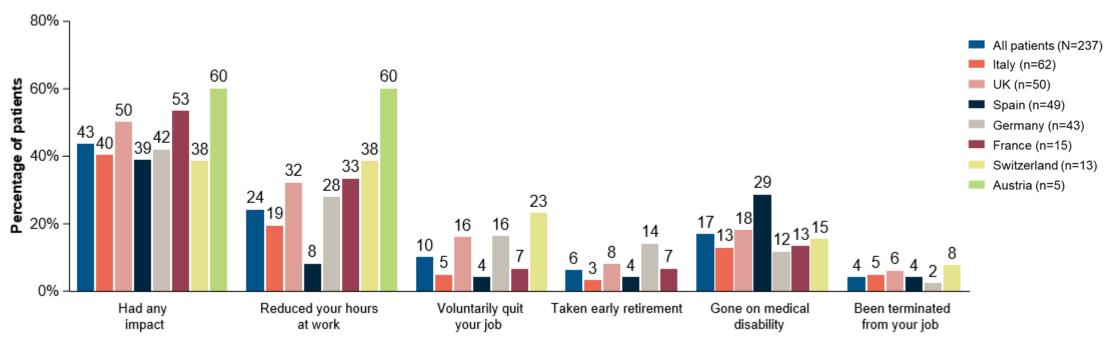
- Patients reported currently taking a mean number of 5.5 medications
- 38% of patients reported experiencing anaphylactic episodes during the prior year
 - 21% of patients experiencing anaphylaxis had 5 or more episodes
- Food (37%), stress (31%), medication (24%), and temperature changes (27%) were the most common anaphylaxis triggers for patients





Patients with ISM reported a significant workforce impact

Impact of ISM on work status



43% of patients reported that ISM had meaningful impact on their ability to work, with 24% reducing hours and 17% going on medical disability





Conclusions: PRISM survey findings in ISM show high disease burden and unmet need for patients despite use of polypharmacy

- To date, PRISM is the largest survey to evaluate the burden of ISM from a patient perspective
- These data show that patients with ISM report moderate to severe disease-related symptoms, impaired physical and mental functioning, and frequent anaphylaxis despite being treated with multiple symptom-targeted medications (current mean: 5.5)
- Additionally, ISM impacts patients' ability to work in approximately 43% of patients



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