

General Information

Where is the venue? Do you need an invitation letter? Find out general information about ESPGHAN 2023 on this page.

The Austria Center Vienna

Bruno-Kreisky-Platz 1, 1220 Wien, Austria

<https://www.acv.at/en>The Annual Meeting will officially run on Central European Time CET – Vienna TimeThe official language of the Annual Meeting will be English. All abstract submissions and presentations should be in English.An official personalised letter of invitation will be provided to registered participants as part of the registration procedure. This invitation is prepared solely for the purpose of visa applications and is not a commitment on the part of the organisers to provide any financial support or confirmation of inclusion in the Scientific Programme.Participants are advised to wear business casual for all Annual Meeting functions and programs.All event activities (including educational sessions, meal functions, exhibit hall, etc.) are exclusively reserved for registered attendees. Non-registered guests (including children, family members, colleagues, etc.) are not allowed in any of the event areas. Badges provided at registration are required for entrance into all functions and will be strictly enforced.Please do not leave bags or suitcases unattended at any time, whether inside or outside the session halls. Hotels strongly recommend that you use their safety deposit boxes for your valuables.

The Congress Secretariat and Organisers cannot accept liability for personal accidents or loss of or damage to

private property of participants. Participants are advised to take out their own personal travel and health insurance for their trip. An exhibition will be held. See the Sponsorship & Exhibition page, by clicking [here](#), for more details regarding support. Support from pharmaceutical partners helps make this event possible. As such, it is important that we comply with the various codes and directives relating to the manufacturing, distribution, and marketing of pharmaceutical products. The European Union directive (2001/83/EC, article 86) provides guidelines relating to medicinal products for human use. Within these guidelines is the restriction that access to pharmaceutical product marketing activities must be limited to only those participants licensed/authorized to prescribe or supply medicinal products. All participants, with the exception of the exhibitors, are asked the following question during registration: Are you licensed/authorized to prescribe or supply prescription-only medicines?

Congress Organisers



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