Informed Consent Statement

All participants provided electronic informed consent before accessing the anonymous online questionnaire. Participation was voluntary, and respondents could decline participation or stop completing the survey at any time. Because the study was an anonymous, non-interventional online survey of physicians and did not involve patients, direct patient contact, or patient-level clinical data, signed written informed consent was not applicable. No personally identifiable information was collected, recorded, or reported at any stage of the study.