Swedish Match USA Inc.

ZYN 3 mg and ZYN 6 mg MRTPAs

Module 1: Administrative Information
1.2 Tables of Content - Index
1.2.2 - List of all Studies

Type of Studies	Module	Study Name - ID	Product Used	Study Description	Study Design	Study Outcomes
Biomarker Study	5.2	SM 22-03	Nicotine pouch users: One Swedish Match brand nicotine pouch product (usual brand) containing 3-16 mg nicotine per pouch Snus users: One Swedish pouched snus product (usual brand) containing 4-20 mg nicotine per pouch Combusted cigarette users: One commercially manufactured combusted cigarette (usual brand) Non-users of tobacco/nicotine products: None	SM 22-03 was a multi-center, cross-sectional, four-group, non-randomized study completed in Sweden, designed to assess BoEs and BoPHs in current, daily users of nicotine pouches, snus, or combusted cigarettes, relative to non-users of tobacco or nicotine products (TNPs).	Eligible subjects in the nicotine user groups participated in the study for 14 days, while non-users of TNPs participated for 1 day, excluding the up to 28-day screening period. Following a screening visit (Visit 1), all subjects attended Visit 2 at the study sites. After initial collection they exclusively used their product of choice ad libitum for 14 days, following their regular usage pattern and documenting. Addtional samples were collected followed by the analysis of BoEs and BoPHs and interviewed for adverse events (AEs).	(b)(4)
Observational Survey Study	6.2	SMNA 5240072	N/A	SMNA 5240072 was an observational survey study conducted to assess perceptions of and likelihood to use the proposed MRTPs with the reduced risk claim among 3400 U.S. adults (21+) in five cohorts	Data were obtained using responses from a customized web-based survey of invited consumers who met inclusion and exclusion criteria and who agreed to participate.	(b)(4)

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