Investigator Financial Interests and Disclosure Statement Form



In compliance with the U.S. Code of Federal Regulations 21CFR54, clinical Investigators are required to disclose to the Study Sponsor their financial interests for the period of time he or she participated in the study, from the date the investigator entered into agreement with the sponsor, and for one year following the end of his or her participation in the study.

Protocol Study Number:	J2N-MC-JZNX	Study Sponsor:	Eli Lilly and Company	
Study Site No.: (if unknown leave blank)	89971	Other Study Sponsor/Co- Development Partner:	Specify	
Investigator Name (First, Middle, Last Name as applicable):	Martin Špaček			
Information collected at study time- point:	Initial DisclosureUpdated Disclosure	Update if legal name or financial interests and arrangements changes from the information provided during the clinical study or within 1 year post clinical study close/end of study participation. (*4)		
TO BE COMPLETED AND SIGNED BY EA	ACH PARTICIPATING INVESTIGATOR			
	n below, retain in your records and provide to the Study Sponsoi study after the site initiation date, complete and sign this form be		98	
Please indicate by marking YES or NO b	elow if any of the financial interests or arrangements applies to y	ou, your spouse, dependent children	, or any combination.	
Are you, your spouse or any of your dependent children an employee of the Study Sponsor(s)/Co-Development Partner(s)?				
2. Have you, your spouse or any of your dependent children entered into a financial arrangement with the Study Sponsor(s)/Co-Development Partner(s) whereby the value of the compensation could be influenced by the outcome of the trial, such as a bonus, royalty or other financial incentive (i.e., compensation that could be higher for a favorable outcome than for an unfavorable outcome)? This could be compensation that is explicitly greater for a favorable result, compensation in the form of an equity interest in Study Sponsor(s) or compensation tied to sales of the product, such as a royalty interest.				ℂ Yes 僑 No
3. Do you, your spouse or any of your dependent children have a proprietary interest in the tested product, including, but not limited to, a patent, trademark, copyright or licensing agreement?				
4. Do you, your spouse, any of your dependent children, or any combination hold any significant equity interest in Study Sponsor(s)/ Co-Development Partner(s) (stock, stock options, or other financial interest) that exceeds \$50,000.00 U.S. dollars? Equity interest includes any options, puts, calls, straddles and other privileges in addition to an equity ownership position in Study Sponsor(s). This does not include ownership interest, stock options or other financial interest over which you have no direct control or input as to the quantities or amounts, e.g., a 401k, IRA, Mutual Fund.				
5. Have you, your spouse, any of your dependent children, or any combination received significant payments of other sorts (SPOOS) having total value in excess of \$25,000.00 from Study Sponsor(s)/Co-Development Partner other than payments for conducting this clinical study or other clinical studies. Examples of such significant payment, include, but are not limited to, grants or funding for ongoing research, compensation in the form of equipment, retains for ongoing consultation or honoraria that are (A) paid directly to me or the institution with which I am affiliated, and (B) paid in support of my activities (i.e., payment paid directly or indirectly to me by Study Sponsor(s)?				✓ Yes ⑥ No
For each YES response above, please prov Indicate the number of attached pages	ide detailed information disclosing the nature of the financial arrangen $\underline{\ \ \ }$).	nent, including total value amounts. (If a	dditional space is needed, ple	ease attach to this document.
I also confirm that to the extent I have provided any I consent to the disclosure, collection and further us regulatory authorities such as United States Food a recipients may be based in countries whose laws de I agree to promptly update the above information	is form is, to the best of my knowledge and belief, true, complete and correct information about other individuals, I have appropriate permission to provide the e of the relevant financial information outside of my country/region to employees, and Drug Administration regulation as required by Title 21 of the Code of Federal I on to provide equivalent protection for personal data to those in the country in whi if my legal name or financial interests and arrangements, or those of my sponsor to conduct the clinical study and within 1 year post clinical study close.	agents and contractors of Study Sponsor(s), it Regulations Part 54, Financial Disclosure by Cl ch I reside. Douse and dependent children, changes fro	s representatives, and business p inical Investigators. I further unde	rstand and agree that such
Signature of investigator:	×	Date:	v 69-	SEP-2024