Date		//
Patient	Participant ID	
Practitioner completing form		🗆 Dentist
		Dental Hygienist
	on Criteria	
	te: Any 'no' response in this section disqualifies the patient from stu	
	estionnaire is ended if a participant answers 'no' in this section. Plea	
1.	Potential participant is at least 40 years old; <i>Practitioner: please</i>	□ Yes
	confirm participant's age. You will be asked to enter their date of	🗆 No
	birth before randomization to ensure the participant is 40 or older.	
2.	Potential participant has 20 or more permanent teeth excluding	□ Yes
2.	3 rd molars.	
2		
3.	Potential participant is in good general health as evidenced by medical history (ASA Class I or II) per the practitioner.	□ Yes
		□ No
4.	Potential participant needs periodontal care for Generalized	□ Yes
	Stage II-III, Grade A-C periodontitis (previously classified as moderate to severe periodontitis), and a minimum of two	🗆 No
	quadrants of SRP (CDT code 4341) in practices participating in	
	the National Dental PBRN PAAS study, with current need for	
	NSPT.	
5.	Potential participant is willing to comply with all study visits and	□ Yes
	be available for the duration of the study (12-15 months).	□ No
6.	Potential participant is willing to provide contact information for	□ Yes
	self, including a cellular phone number for study text, and one	□ No
	to two emergency contacts to be reached for the follow-up visits	
	and any other study-related matters for the duration of the	
	study.	
	on Criteria	
	Any 'yes' response in this section disqualifies the patient from stud	
-	onnaire is ended if a participant answers 'yes' in this section. Please	
/.	Potential participant has a known drug allergy to antibiotics or	□ Yes
	anesthetics.	□ No
8.	Potential participant has used systemic antibiotics within the last	🗆 Yes
	3 months prior to enrollment.	□ No
9.	Potential participant has a medical condition which requires	🗆 Yes
	antibiotic prophylaxis prior to receiving dental treatments/visits.	🗆 No
10.	Potential participant currently uses medications that may cause	🗆 Yes
	adverse effects with Amoxicillin/Metronidazole (AMXM) such as	🗆 No
	disulfiram, warfarin, oral contraceptives.	
11.	Potential participant has a history of receiving periodontal	🗆 Yes
	therapy (including SRP D4341, D4342) within the last 6 months	🗆 No
1	prior to enrollment.	

12. Potential participant is currently pregnant or lactating per self-	□ Yes
report.	□ No
13. Potential participant is considered immunocompromised, in the opinion of the practitioner (including diseases and conditions such as HIV/AIDS, immunosuppressive drug therapy and/or radiation), or has chronic mucosal lesions (e.g. pemphigus vulgaris) affecting the gingiva.	□ Yes □ No
 Potential participant has Diabetes Mellitus with an HBA1c score	□ Yes
of >/= 10% within the past 3-months as per self-report.	□ No
15. Potential participant is planning to receive another type of adjunct periodontal therapy during the course of the study (the next 12 months). This includes but is not limited to laser therapy, local antibiotic, or periodontal surgery.	□ Yes □ No

Eligible: ____

This patient is eligible for the PAAS study. Please proceed to the consent form.

Pass the tablet to the participant after you click the Submit button below.

Not Eligible: ____

This patient does not qualify for the PAAS study. Please click submit below.

Date		//
Patient Pa	rticipant ID	
	hich practitioner is completing the form?	Dentist
		Dental Hygienist
1. Ha	as OHIP-5 form been completed by the participant?	🗆 Yes
		🗆 No
2. Ha	as patient participant been randomized?	🗆 Yes
		□ No
3. Ha	as medication been distributed and administered, to include	□ Yes
	edication instructions? <i>Recommended at least 30-60 minutes</i>	
	ior to the procedure	
	- F	
4. Ha	as plaque sample been collected? only if patient participant	□ Yes
	onsented to collection	
		□ N/A
	a laad an athatic been administered to do. 2	
5. Ha	as local anesthetic been administered today?	□ Yes
		□ No
	"yes" to Question 5,	
5a	a. Which type of anesthetic was administered	\Box Local anesthetic
		Topical anesthetic
		(e.g., Oraqix)
		🗆 None
		□ Other
		specify:
		. ,
6. W	'hat quadrants were treatment planned to undergo SRP?	Upper Right (UR)
		□ Yes
		□ No
		Upper Left (UL)
		\Box Yes
		□ No
		Lower Left (LL)
		□ Yes
		Lower Right (LR)
		□ Yes
		🗆 No
7. W	'hat quadrants received SRP today?	Upper Right (UR)
		□ Yes

	🗆 No
	Upper Left (UL)
	□ Yes
	🗆 No
	Lower Left (LL)
	🗆 Yes
	🗆 No
	Lower Right (LR)
	□ Yes
	🗆 No
8. Has the treatment area been verified for completion of residual	□ Yes
calculus with an ODU 11/12 explorer (included in patient	🗆 No
participant package)?	
9. Has patient participant reported an Adverse Event?	🗆 Yes
	🗆 No
10. Has practitioner reviewed and provided patient participant with	🗆 Yes
the post-op oral hygiene instructions at this visit?	🗆 No
11. Does a second appointment for scaling and root planing need to	🗆 Yes
be made to complete this procedure?	🗆 No
11a. If yes, enter date	
11b. If no, please answer question 12.	
12. Has practitioner uploaded the periodontal chart image?	□ Yes
Please ensure PD, BoP, and GR are captured	🗆 No

Date	//
Patient Participant ID	
Which Practitioner is completing the form?	□ Dentist
	Dental Hygienist
Days since first Baseline visit	
1. Has local anesthetic been administered today?	□ Yes
	□ No
If "yes" to Question 1,	
1a. Which type of anesthetic was administered	□ Local anesthetic
	□ Topical anesthetic
	(e.g., Oraqix)
	□ Other
	specify:
	specify.
2. What quadrants received SRP today?	Upper Right (UR)
	🗆 Yes
	🗆 No
	Upper Left (UL)
	□ Yes
	□ No
	Lower Left (LL)
	□ Yes
	Lower Right (LR)
	□ Yes
3. Has the treatment area been verified for completion of residual	□ Yes
calculus with an ODU 11/12 explorer (included in patient	
participant package)?	
4. Has patient participant been provided oral hygiene instructions	□ Yes
at this visit?	🗆 No
5. Has patient participant reported an adverse event?	□ Yes
	□ No
6. Has practitioner uploaded the periodontal chart image?	□ Yes
Please ensure PD, BoP and GR have been captured	

PAAS Medication Compliance Form

Date: ___/___/ ____/ _____

Patient Participant ID: ___ __ __

1. Please count the number of capsules you have left in the vials of medications that were given to you at your research study visit and record below.

Vial A: ____ capsules

Vial B: ____ capsules

Date		//
Patien	t Participant ID	
	Which practitioner is completing the form?	🗆 Dentist
		Dental Hygienist
1.	Has OHIP-5 form been completed by the patient participant?	🗆 Yes
		□ No
2.	Has plaque sample been collected? only if patient participant	□ Yes
	consented to collection	🗆 No
		□ N/A
3.	Did patient participant previously report any Adverse Events?	□ Yes
		□ No
	If "yes" to Question 3,	
	3a. Were all open AEs reviewed with patient participant?	□ Yes
		□ No
4.	Did patient participant report any new adverse events?	□ Yes
		□ No
	If "yes" to Question 4,	
	4a. Were the new AEs recorded on the AE log?	□ Yes
	Ű	□ No
5.	Has patient participant received any subsequent antibiotics for a	□ Yes
	newly diagnosed condition?	
6.	Has the periodontal maintenance interval been established?	□ Yes
		□ No
	If "yes" to Question 5,	
	5a. How frequently should the patient participant return for	Every 3 months
	periodontal maintenance?	Every 4 months
7.	Has patient participant been provided oral hygiene instructions	□ Yes
	at this visit?	🗆 No
8.	Has practitioner uploaded the periodontal chart image?	□ Yes
	Please ensure PD, BoP and GR have been captured	□ No

Date		//
Patien	t Participant ID	
	Which practitioner is completing the form?	🗆 Dentist
		Dental Hygienist
1.	Has OHIP-5 form been completed by the patient participant?	□ Yes
		□ No
2.	Has plaque sample been collected? only if patient participant	□ Yes
	consented to collection	🗆 No
		□ N/A
3.	Did patient participant previously report any Adverse Events?	□ Yes
		□ No
	If "yes" to Question 3,	
	3a. Were all open AEs reviewed with patient participant?	□ Yes
		□ No
4.	Did patient participant report any new adverse events?	□ Yes
		□ No
	If "yes" to Question 4,	
	4a. Were the new AEs recorded on the AE log?	□ Yes
	Ű	□ No
5.	Has patient participant received any subsequent antibiotics for a	□ Yes
	newly diagnosed condition?	
6.	Has the periodontal maintenance interval been established?	□ Yes
		□ No
	If "yes" to Question 5,	
	5a. How frequently should the patient participant return for	Every 3 months
	periodontal maintenance?	Every 4 months
7.	Has patient participant been provided oral hygiene instructions	□ Yes
	at this visit?	🗆 No
8.	Has practitioner uploaded the periodontal chart image?	□ Yes
	Please ensure PD, BoP and GR have been captured	□ No

PAAS Adverse Events Form

Instructions for adverse event follow up: Below are the unresolved adverse events previously reported by the participant. Review these adverse events with the participant and update the instance if anything has changed. The data is populated with the answers you entered the last time you submitted a report on this adverse event. Be sure to update the report type, Visit # and current status at a minimum.

If the event continues to be unresolved, you will follow up with the participant at the next visit.

After reviewing all unresolved adverse events, answer the question about any new adverse events the participant needs to report. If new adverse events are reported, you will complete a new adverse event instance for each new symptom.

Question at Baseline visit: Has the participant reported any adverse events at this visit? Yes/No

Question at all other visits: Has the participant experienced any NEW adverse events since their last visit? Yes/No

Participant ID: Report date: This is the date you are filling out the report •	Visit #: Select the visit number below. This should be the visit where this event is being reported OR the closest visit that has already occurred if reporting between visits. Baseline (V1) Optional Baseline (V1b) Re-evaluation (V2) Final (V3)
1. Description of event <i>Describe the</i> symptom/event in detail. Only include one symptom per report.	Between visits
 Date of event This is the date the event happened. The date may be today or may be a date in the past. 	
 3. According to study personnel, was this event expected or unexpected? Expected events are as follows (you will be asked to select the appropriate expected symptom in the next question if expected is marked): Post procedural tooth sensitivity Post procedural pain Nausea Allergic reaction: rash or itching Allergic reaction: difficulties breathing Periodontal abscess Diarrhea 	 Expected Unexpected
 a. If expected, please select the expected symptom: 	 Post procedural tooth sensitivity Post procedural pain Nausea

4. Did the event result in any of the following actions or could it be classified as: <i>Checking any option except "None" makes this</i> <i>a serious adverse event. If the event is classified</i> <i>as serious, your Node Coordinator will reach out</i> <i>to collect additional information about the</i> <i>event.</i>	 Allergic reaction: rash or itching Allergic reaction: difficulties breathing Periodontal abscess Diarrhea A life-threatening event A hospitalization (initial or prolonged) that required at least one night in the hospital A disability or permanent damage that resulted in a substantial disruption to their ability to conduct normal life functions A pregnancy resulting in a congenital abnormality or birth defect Other serious important medical event that may require medical or surgical intervention to prevent one of the other outcomes listed above None of the above
5. How would you rate the severity of this event? In the opinion of the practitioner	MildModerateSevere
6. Was this event study drug related? <i>In the opinion of the practitioner</i>	 Definitely Probably Possibly Remotely Definitely not Unknown
 7. What is the current status of the event? Additional reporting will be required at future study visits for conditions that are improving, present and unchanged and deteriorating. If a death is being reported, additional reporting is required as a serious adverse event. Your Node Coordinator will reach out to gather additional details 	 Completely recovered Recovered with some residual problems Condition improving Condition present and unchanged Condition deteriorating Death (skip to 7b)
a. On what date was this event resolved?	MM/DD/YYYY
b. Date of death? Per the protocol, the study PI should review the report and confirm the following information:	MM/DD/YYYY
8. <i>PI complete this field only:</i> In the PI's view, was this event expected or unexpected? <i>Expected events are as follows (you will be asked to select the appropriate expected symptom in the next question)</i>	 Expected Unexpected

Post procedural tooth sensitivity	
Post procedural pain	
Nausea	
Allergic reaction: rash or itching	
Allergic reaction: difficulties breathing	
Periodontal abscess	
Diarrhea	
a.If expected, please select the expected	 Post procedural tooth sensitivity
symptom:	Post procedural pain
	Nausea
	Allergic reaction: rash or itching
	Allergic reaction: difficulties breathing
	Periodontal abscess
	Diarrhea
9. PI complete this field only: In the PI's view,	Definitely
was this event study drug related?	 Probably
	 Possibly
	Unlikely
10. Di semulata this field entre Dess the Di serves	Not related
10. PI complete this field only: Does the PI agree	A life-threatening event
with the classification of serious or not serious	• A hospitalization (initial or prolonged) that
in question 4 above?	required at least one night in the hospital
	A disability or permanent damage that
	resulted in a substantial disruption to their
	ability to conduct normal life functions
	 A pregnancy resulting in a congenital
	abnormality or birth defect
	Other serious important medical event that
	may require medical or surgical
	intervention to prevent one of the other
	outcomes listed above
	None of the above
If no to question 10: You indicated you do not	
agree with the practitioner's assessment of	
seriousness. Contact the NCC as soon as possible	
to resolve this issue.	
11. PI Complete this field only: In the PI's view,	• Yes
does this event suggest that research places	• No
participants or others at a greater risk of harm	
(including physical, psychological, economic, or	
social harm) than was previously known or	
recognized).	
	• Yes
PI attests they have reviewed this adverse event,	• No
submitted their evaluation and verified that all	

information is complete	
	MM/DD/YYYY

Date	//
Patient Participant ID	
Practitioner completing form	□ Dentist
	Dental Hygienist
Inclusion Criteria	
Note: Any 'no' response in this section disqualifies the patient from stu	
questionnaire is ended if a participant answers 'no' in this section. Plea	-
1. Potential participant is at least 35 years old; <i>Practitioner: please</i>	□ Yes
confirm participant's age. You will be asked to enter their date of	□ No
birth before randomization to ensure the participant is 35 or older.	
2. Potential participant has 15 or more permanent teeth excluding	
3 rd molars.	□ Yes
 Potential participant is in good general health as evidenced by medical history (ASA Class I or II) per the practitioner. 	□ Yes
4. Potential participant needs periodontal care for Generalized	□ Yes
periodontitis), and a minimum of two quadrants of SRP (CDT	□ No
code 4341) in practices participating in the National Dental PBRN PAAS study, with current need for NSPT.	
5. Potential participant is willing to comply with all study visits and	□ Yes
be available for the duration of the study (12-15 months).	
 6. Potential participant is willing to provide contact information for 	□ No □ Yes
self, including a cellular phone number for study text, and one	
to two emergency contacts to be reached for the follow-up visits	□ No
and any other study-related matters for the duration of the	
study.	
Exclusion Criteria	
Note: Any 'yes' response in this section disqualifies the patient from stud	y participation. The
questionnaire is ended if a participant answers 'yes' in this section. Please	note as screen fail.
7. Potential participant has a known drug allergy to antibiotics or	🗆 Yes
anesthetics.	🗆 No
8. Potential participant has used systemic antibiotics within the last	🗆 Yes
3 months prior to enrollment.	□ No
9. Potential participant has a medical condition which requires	□ Yes
antibiotic prophylaxis prior to receiving dental treatments/visits.	□ No
10. Potential participant currently uses medications that may cause	□ Yes
adverse effects with Amoxicillin/Metronidazole (AMXM) such as	□ No
disulfiram, warfarin, oral contraceptives.	
11. Potential participant has a history of receiving periodontal	🗆 Yes
therapy (including SRP D4341, D4342) within the last 6 months	🗆 No
prior to enrollment.	
12. Potential participant is currently pregnant or lactating per self-	□ Yes
report.	🗆 No

 Potential participant is considered immunocompromised, in the opinion of the practitioner (including diseases and conditions such as HIV/AIDS, immunosuppressive drug therapy and/or radiation), or has chronic mucosal lesions (e.g. pemphigus vulgaris) affecting the gingiva. 	□ Yes □ No
14. Potential participant has Diabetes Mellitus with an HBA1c score of >/= 10% within the past 3-months as per self-report.	□ Yes □ No
15. Potential participant is planning to receive another type of adjunct periodontal therapy during the course of the study (the next 12 months). This includes but is not limited to laser therapy, local antibiotic, or periodontal surgery.	□ Yes □ No

Eligible: ____

This patient is eligible for the PAAS study. Please proceed to the consent form.

Pass the tablet to the participant after you click the Submit button below.

Not Eligible: ____

This patient does not qualify for the PAAS study. Please click submit below.

Date	/
Patient Participant ID	
Which practitioner is completing the form?	 Dentist Dental Hygienist
1. Has OHIP-5 form been completed by the participation of the participat	
2. What is the Periodontitis Stage?	
3. What is the Periodontitis Grade?	□ A □ B □ C
4. Has patient participant been randomized?	□ Yes □ No
5. Has medication been distributed and administere medication instructions?	ed, to include
6. Has plaque sample been collected? <i>Mark N/A if p</i> consent to plaque sample OR if your office is not p plaque sampling.	
7. Has local anesthetic been administered today?	☐ Yes □ No
If "yes" to Question 5,	
7a. Which type of anesthetic was administered	□ Local anesthetic □ Topical anesthetic (e.g., Oraqix) □ None □ Other specify:
8. What quadrants were treatment planned to unde	ergo SRP? Upper Right (UR) Yes No Upper Left (UL) Yes No Lower Left (LL) Yes No Lower Right (LR) Yes No
9. What quadrants received SRP today?	Upper Right (UR)

	□ No
	Upper Left (UL)
	□ Yes
	□ No
	Lower Left (LL)
	🗆 Yes
	□ No
	Lower Right (LR)
	🗆 Yes
	□ No
10. Who completed the SRP treatment on the patient participant	Dentist
today?	🗆 Hygienist
11. Has the treatment area been verified for completion of residual	□ Yes
calculus with an ODU 11/12 explorer (included in patient	□ No
participant package)?	
12. Has patient participant reported an Adverse Event?	🗆 Yes
	🗆 No
If yes: You indicated the participant reported an adverse event. Be sure	
to complete an Adverse Event Form for each reported event.	
13. Has practitioner reviewed and provided patient participant with	🗆 Yes
the post-op oral hygiene instructions at this visit?	🗆 No
14. Is scaling and root planing treatment for this participant	□ Yes
complete?	□ No
14a. If SRP treatment is complete, enter the treatment	//
completion date:	
14b. If SRP treatment is not complete, enter date of next	//
treatment visit:	
If no to 14: The periodontal chart image cannot be uploaded	
until the SRP treatment is complete. You will upload the image	
after the participant's next visit.	
15. If yes to 14: Has practitioner uploaded or is planning to upload the periodontal chart image?	□ Yes
Please ensure PD, BoP, and GR are captured	□ No
rieuse ensuie rD, DOr, una On ale cuptalea	