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Question Text	Author Response	Custom Submission Question ID
Please complete the following questions in accordance with International Committee of Medical Journal Editors' recommendations on data sharing in clinical trials (guidelines and examples are available here). Will individual participant data be available (including data dictionaries)?	A complete, cleaned, de-identified copy of the final data set will be available, include pre-visit, one week post visit and six month follow-up data.	54
What data in particular will be shared? (Examples include all individual participant data after deidentification, only participant data that underlies the results, or not available.)	All survey data from the three patient surveys and limited data from the chart review will be available.	55
What other documents will be available?	The following will be made available: a code book, a SAS file of the code used for creating the final study sample, the final study variables and plan for conducting the outcomes analyses outlined in the study protocol will be made available.	56
When will data be available (start and end dates)?	The data will be available March 2021 for at least 5 years.	57
With whom? (Examples include anyone who wishes to access the data, researchers who provide a proposal, or not applicable.)	The PI will share a de-identified data set with outside investigators according to the policies in the approved IRB protocol. Investigators will be required to provide evidence of IRB approval (or exemption) and/or complete a data sharing agreement.	58
For what types of analyses?	There are no restrictions.	59
By what mechanism will data be made available?	Interested investigators should contact the corresponding author and study PI.	60