http://dx.doi.org/10.2106/JBJS.19.00004

Question Text	Author Response	Custom Submission Question ID
Please complete the following questions in accordance with the 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 de-identified data set will be available.	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.)	The investigators will create a complete, cleaned, de-identified copy of the final data set that will include the patient survey data and key variables (e.g. knowledge scores) underlying the main results.	55
What other documents will be available?	A description of the study dataset, including a code book and syntax of the code used for creating the final study variables will be available. The study protocol paper includes detailed plans for conducting the outcomes analyses.	56
When will data be available (start and end dates)?	The investigators plan to make the data available within 3 months of the end of funding (anticipated release May 2020).	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 may be required to provide evidence of IRB approval (or exemption) and/or complete a data sharing agreement.	58
For what types of analyses?	The analyses will need to follow those outlined in the approved IRB protocol.	59
By what mechanism will data be made available?	Those interested in accessing the data should contact the corresponding author.	60