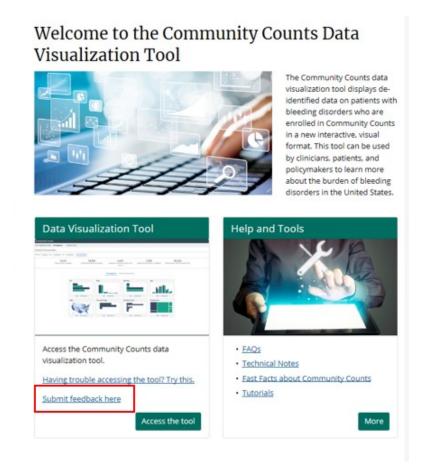
## Screenshots of Feedback Survey for Community Counts Data Viz Tool

View the developer webpage for the feedback survey here: https://wwwdev.cdc.gov/ncbddd/hemophilia/communitycounts/data-viz-feedback.html

The picture below is a screenshot of the survey link on the CC Data Viz webpage that users will click on to take the feedback survey.



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## Community Counts Data Visualization Tool Feedback

We welcome your feedback!

1. Thank you for using the Community Counts Data Visualization Tool. What is your primary role? (optional)
Healthcare provider
O Member of the media
O Scientist or researcher
O Educator
O General public
Bleeding disorder patient
Patient advocacy organization/chapter
Pharmaceutical company representative
Other - please specify below
Other Role: (optional)
2. How do you rate the tutorials? If you have not used or seen the tutorials, please select this option below. (optional)
○ I have not used the tutorials
O Excellent
○ Good
○ Average
O Poor
O Very poor
If "Poor" or "Very Poor," please describe how the tutorials can improve: (optional)

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3. Did you experience any technical issues with the Community Counts Data visualiza Tool? For example, a graph did not load or text not displaying properly.  (optional)	alization
O No	
○ Yes	
If yes, please describe the issues: (optional)	
4. How frequently do you access the Community Counts Data Visualization Tool? {opti	onal)
O This is my first time	,
O Daily	
O Weekly	
O Monthly	
O Infrequently	
5. How likely are you to revisit the Community Counts Data Visualization Tool? (optiona	d)
O Not likely at all	
Somewhat unlikely	
O Not sure	
Somewhat likely	
Extremely likely	

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6. Please briefly explain how you are using this data. This helps us learn how and why people are using the Community Counts Data Visualization Tool. (optional)

"I'm interested in using this data for research" or "I have a bleeding disorder and I want to learn more about it."

7. What would enhance your experience with the Community Counts Data Visualizati Tool? $(optional)$	
8. If you have any additional comments, please write them below. (optional)	
protected by reCAPTCHA	



Privacy - Terms

## Submit

CDC estimates the average public reporting burden for this collection of information as 3 minutes per response, including the time for reviewing instructions, searching existing data/information sources, gathering and maintaining the data/information needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0920-1154).