NCORP Electronic Medical Record (EMR) Utilization Survey 2021

OMB No.: 0925-0642
Expiration Date: 03/31/2023

1. NCORP Name

Enter your answer

2. Site/Institution Name

Enter your answer

3. Do you currently use Electronic Medical Record (EMR) Software?

- ◯ Yes
- ◯ No
- ◯ Don't Know

4. If not, why?

Enter your answer

5. If yes, does your site/institution use EMR software to prescreen or identify potential study participants?

- ◯ Yes
- ◯ No
- ◯ Don't Know

6. If applicable, has this feature helped prescreen or identify study participants?

- ◯ Yes
- ◯ No
- ◯ Other

7. How do you screen, recruit participants if not using EMR?

Enter your answer

8. Are protocol specific documents/templates (i.e. protocol order set, adverse event assessment, etc.) available via the EMR?

- ◯ Yes
- ◯ No
- ◯ Don't Know
- ◯ N/A

9. If applicable, please describe which protocol specific documents/templates (i.e. protocol order set, adverse event assessment, etc.) are available via the EMR?

Enter your answer

10. Are documents such as nursing notes toxicity checks/grids able to be routed to a physician/APP for signature?

- ◯ Yes
- ◯ No
- ◯ Don't Know
- ◯ N/A

11. Does your site/institution use standardized, retrievable text to document clinical research activities in the EMR (i.e. Smart Text or Smart Phones for consent, eligibility determination, etc.)?

- Yes
- No
- Don't Know
- N/A

12. If applicable, please describe the standardized, retrievable text that your site/institution uses to document clinical research activities in the EMR (i.e. Smart Text or Smart Phones for consent, eligibility determination, etc.)?

Enter your answer

13. Can your site/institution grant remote access to the EMR (for auditing or monitoring purposes)?

- Yes
- No
- Don't Know
- N/A

14. Does your site have an EMR with capability to redact documents?

- Yes
- No
- Don't Know
- N/A

15. Additional comments

Enter your answer

Submit