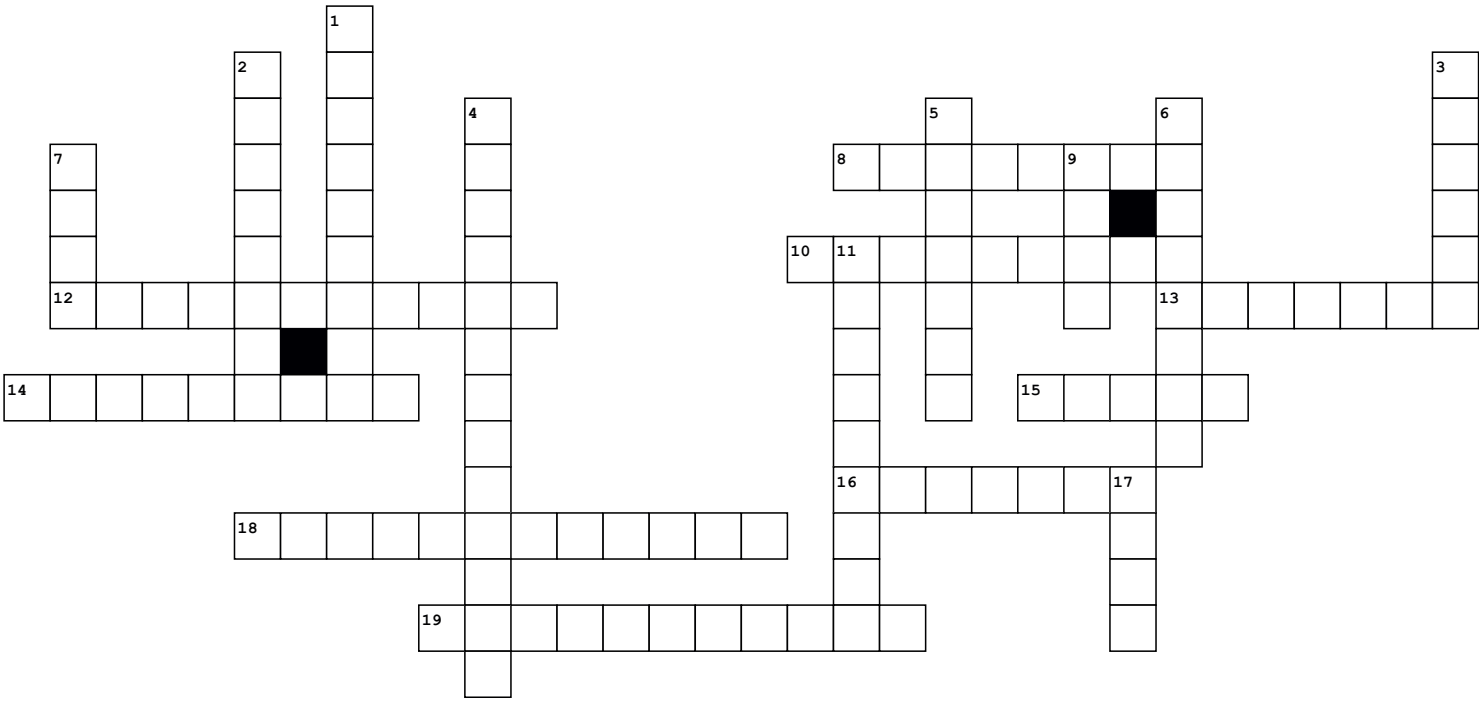
Clinical Trial Terminology



Across

8. This document describes every step of the study and contains answers to many study-related questions

10. Forgetting to complete a required procedure or conducting a visit out of window may result in one of these

12. Fill out one of these at every visit in which labs are collected. Make sure to include a copy with your samples!

13. This must be obtained prior to performing and study related procedures

14. The period in which medical history is reviewed and eligibility is determined

15. When data is missing, vague, or incomplete a CRA may issue one of these.

16. Subjects receive one of these after completing study visits

18. A cooler sounding name for the doctor overseeing the clinical trial

19. Medications are considered this term if they are being taken at the time of consent, or started after consenting

Down

1. All of these criteria must be met before a subject can enroll

2. The one-stop-spot for all your regulatory needs

3. Like an ICF, but for minors

4. The point at which a subject is officially enrolled into a study

5. Another name for a clinical research associate (CRA)

6. The service that Optimed sites use to compensate study participants

7. This phase of clinical research is also called “post market surveillance”

9. Complete study visits and answer internal queries on this site

11. None of these criteria can be met in order for a subject to enroll

17. This is collected by the coordinator and entered into the EDC by one of Optimed’s data managers