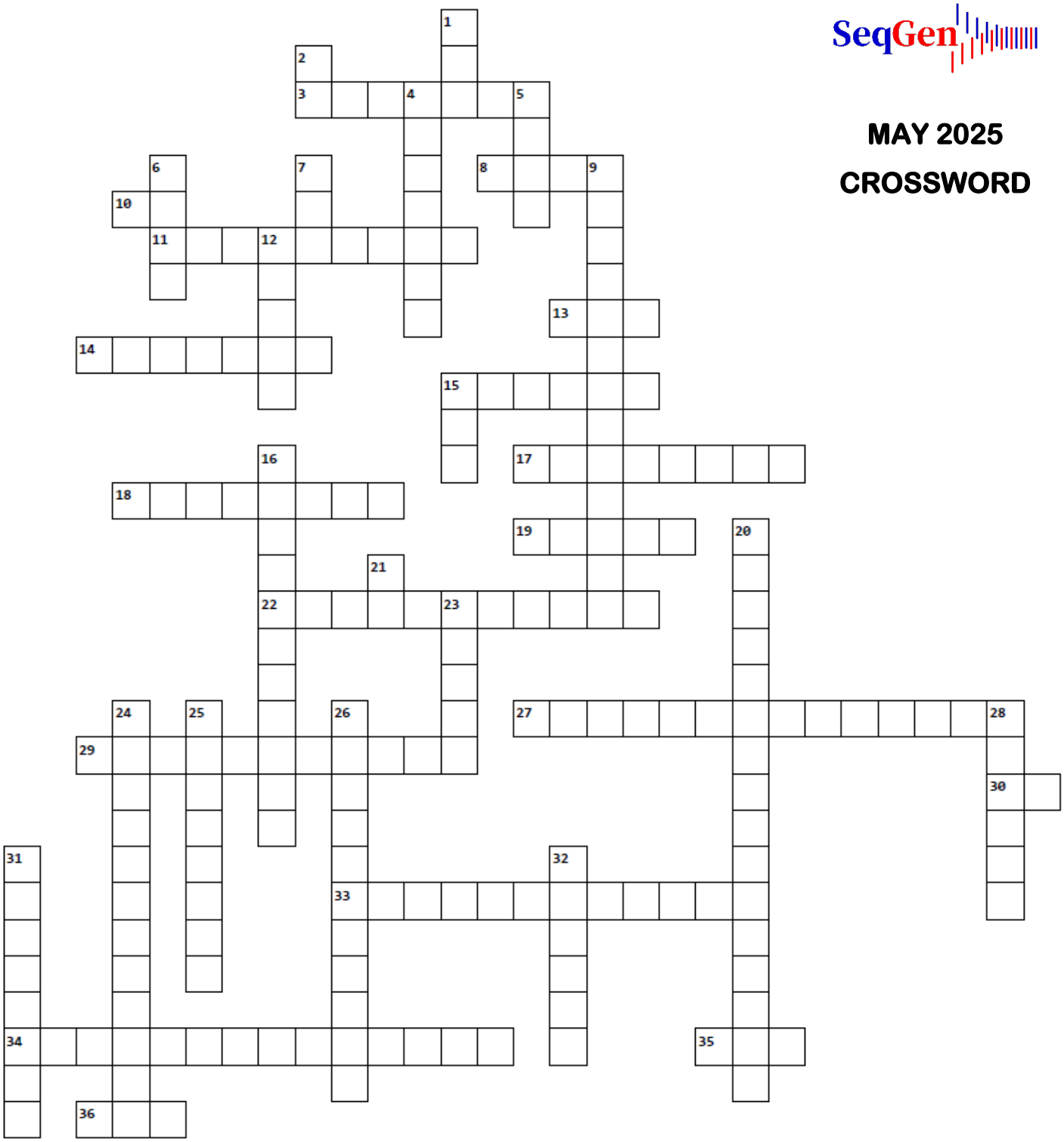
SeqGen

MAY 2025 CROSSWORD



Across

3. Control used to simulate treatment

8. ___ review – assessment by peers

10. Primary doctor overseeing a trial (abbr.)

11. Essential code for ethical research (1947)

13. Documented test procedures (abbr.)

Down

1. Phase ___ – first stage of human testing

2. Long-term study: follow-___

4. Subject group not receiving treatment

5. ___-label – all treatment details visible

6. James ___ – pioneer of clinical trials

14. Informed ___ (ethical requirement)

15. Location for testing new therapies

17. Common reference point in clinical comparisons

18. Predefined outcome used to assess a treatment

19. Type of trial where treatment is concealed

22. Study conducted without bias

27. U.S. database of registered trials

29. Consent must be given ___ (freely)

30. Common unit in dose measurement

33. Type of clinical trial that tests a treatment

34. A request for authorization from FDA to administer an investigational drug or biological product to humans

35. Acronym for Randomized Controlled Trial (abbr.)

36. Organization that regulates drugs (abbr.)

7. A group or treatment pathway in a clinical trial

9. Type of assignment that removes bias

12. Adverse ___ – undesirable effect

15. Form used for data entry (abbr.)

16. Clinical research ___ (supports investigators)

20. Study of how the body processes drugs

21. Investigator's Brochure (abbr.)

23. Trial halted before completion: ___ stop

24. Ideal trial design: randomized, double-blind, placebo-controlled

25. Participants are often called ___

26. Pre-trial testing phase (animal/in vitro)

28. Small group selected to represent a larger population

31. Favorable outcome or improvement seen in response to treatment

32. IRB stands for Institutional ___ Board