

Clinical Data Management Module Resources

Selected References

- Nahm, M., Shepherd, J., Buzenberg, A., Rostami, R., Corcoran, A., McCall, J., & Pietrobon, R. (2011). Design and implementation of an institutional case report form library. [Clinical Trials, 8, 94–102.](#)
- Reeves, D. & Walden, A. (2016). The Research Team. In A.D. Klimaszewski, M.A. Bacon, E.A. Ness, J.G. Westendorp, & K. Willingberg (Eds.), *Manual for Clinical Trials Nursing* (3rd edition, pp. 369-384). Pittsburgh, PA: [Oncology Nursing Society.](#)
- U.S. Food and Drug Administration. (2013, September). Guidance for industry electronic source data in clinical investigations. Retrieved from <http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM328691.pdf>
- Walther, B., Hossin, S., Townend, J., Abernethy, N., Parker, D., & Jeffries, D. (2011). Comparison of electronic data capture (EDC) with the standard data capture method for clinical trial data. [PLos One, 6\(9\).](#)

Selected Websites

- [Clinical Data Interchange Standards Consortium](#)
- [Common Data Element Resource Portal](#)
- [Critical Path Institute](#)
- [NCI-EVS](#) (National Cancer Institute – Enterprise Vocabulary Service)