

- 3) Prospective collection of biological specimens for research purposes by noninvasive means.
- 4) Collection of data through noninvasive procedures (not involving general anesthesia or sedation) routinely employed in clinical practice, excluding procedures involving x-rays or microwaves.
- 5) Research involving materials (data, documents, records, or specimens) that have been collected or will be collected solely for nonresearch purposes (such as medical treatment or diagnosis).
- 6) Collection of data from voice, video, digital, or image recordings made for research purposes.
- 7) Research on individual or group characteristics or behavior (including, but not limited to, research on perception, cognition, motivation, identity, language, communication, cultural beliefs or practices, and social behavior) or research employing survey, interview, oral history, focus group, program evaluation, human factors evaluation, or quality assurance methodologies.
- 8) Continuing review of research previously approved by the convened IRB as follows:
 - (a) the research is permanently closed to the enrollment of new subjects; **and** all subjects have completed all research-related interventions; **and** the research remains active only for long-term follow-up of subjects; **or**
 - b) Where no subjects have been enrolled and no additional risks have been identified; **or**
 - (c) Where the remaining research activities are limited to data analysis
- 9) Continuing review of research, not conducted under an investigational new drug application or investigational device exemption where categories 2 through 8 do not apply but the IRB has determined and documented at a convened meeting that the research involves no greater than minimal risk and no additional risks have been identified.

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Comments

Does this study meet expedited review criteria?

No Yes

IRB Administrator

Date

IRB Reviewer

Date