

STANFORD UNIVERSITY
Non-Clinical Survey Research Information Sheet

Protocol Director: Joseph Liao MD

IRB Use Only

Approval Date: June 25, 2026

Expiration Date: January 13, 2027

Protocol Title: Leveraging Artificial Neural Networks to Enhance Detection of Urothelial Carcinoma

IRB# 82508

FOR QUESTIONS ABOUT THE STUDY, CONTACT: Joseph Liao MD, 453
Quarry Rd, Palo Alto, CA 94304, (650) 852-3284

DESCRIPTION: You are invited to participate in **a research study** to evaluate the benefits of artificial intelligence (AI) models in the detection of urothelial cancer during endoscopy. You will be asked to complete two online surveys at different time points. You will be notified when you are eligible for each part of the survey. Each survey will contain cystoscopy images and/or video and you will be asked to give your impression of the media. Survey data is identifiable for the purposes of compensation. It will be deidentified for analysis and publication. Participation in this research is voluntary, and you are free to withdraw your consent at any time.

TIME INVOLVEMENT: Your participation will take approximately 1 hour total, spread across two 30 minute surveys.

RISKS AND BENEFITS: The risks associated with this study are none. The benefits which may reasonably be expected to result from this study are none. We cannot and do not guarantee or promise that you will receive any benefits from this study. If you are affiliated with Stanford, Your decision whether or not to participate in this study will not affect your employment.

PAYMENTS: You will receive \$100 in the form of gift cards as payment for your participation.

SPONSOR: The National Institutes of Health and Liao Lab/Stanford Department of Urology are providing financial support and/or material for this study.

PARTICIPANT'S RIGHTS: If you have read this form and have decided to participate in this project, please understand your participation is voluntary and you have the right to withdraw your consent or discontinue participation at any time without penalty or loss of benefits to which you are otherwise entitled. The alternative is not to participate. You have the right to refuse to answer particular questions. The results of this research study may be presented at scientific or professional meetings or published in scientific journals. Your individual privacy will be maintained in all published and written data resulting from the study.

Data collected on you to the point of withdrawal remains part of the study database and may not be removed per the Food and Drug Administration.

Your private information collected as part of the research, even if identifiers are removed, will not be used or distributed for future research studies.

CERTIFICATE OF CONFIDENTIALITY

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This research is covered by a Certificate of Confidentiality from the National Institutes of Health (NIH). The researchers with this Certificate may not disclose or use information, documents, or biospecimens that may identify you in any federal, state, or local civil, criminal, administrative, legislative, or other action, suit, or proceeding, or be used as evidence, for example, if there is a court subpoena, unless you have consented for this use. Information, documents, or biospecimens protected by this Certificate cannot be disclosed to anyone else who is not connected with the research except, if there is a federal, state, or local law that requires disclosure (such as to report child abuse or communicable diseases but not for federal, state, or local civil, criminal, administrative, legislative, or other proceedings, see below); if you have consented to the disclosure, including for your medical treatment; or if it is used for other scientific research, as allowed by federal regulations protecting research subjects

The Certificate cannot be used to refuse a request for information from personnel of the United States federal or state government agency sponsoring the project that is needed for auditing or program evaluation by National Institutes of Health which is funding this project or for information that must be disclosed in order to meet the requirements of the federal Food and Drug Administration (FDA). You should understand that a Certificate of Confidentiality does not prevent you from voluntarily releasing information about yourself or your involvement in this research. If you want your research information released to an insurer, medical care provider, or any other person not connected with the research, you must provide consent to allow the researchers to release it.

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CONTACT INFORMATION:

Questions: If you have any questions, concerns or complaints about this research, its procedures, risks and benefits, contact the Protocol Director, Joseph Liao MD, (650) 852-3284. You should also contact them at any time if you feel you have been hurt by being a part of this study.

Independent Contact: If you are not satisfied with how this study is being conducted, or if you have any concerns, complaints, or general questions about the research or your rights as a participant, please contact the Stanford Institutional Review Board (IRB) to speak to someone independent of the research team at 650-723-5244 or toll free at 1-866-680-2906. You can also write to the Stanford IRB at irbeducation@stanford.edu.

Please print a copy of this page for your records.

If you agree to participate in this research, please complete the provided electronic survey.