

# HOW AI COULD IMPACT CANCER RESEARCH

The perspective of a clinical trialist,  
breast cancer screening researcher  
and the Chief Research Officer of the ACR

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GEOFFREY HINTON  
NOBEL LAUREATE IN PHYSICS 2024  
"GODFATHER OF AI"

"People should stop training  
radiologists now."

It is "completely obvious" that AI  
would outperform radiologists within  
5 years.

2016

# WHY AI HAS NOT REPLACED RADIOLOGISTS DESPITE EXCELLENT PEER-REVIEWED EVIDENCE

E.G. LOUIS ET AL. NATURE HEALTH.2025.[HTTPS://DOI.ORG/10.1038/S44360-025-00001-0](https://doi.org/10.1038/s44360-025-00001-0)



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Research

Original Investigation | LESS IS MORE

## Diagnostic Accuracy of Digital Screening Mammography With and Without Computer-Aided Detection

Constance D. Lehman, MD, PhD; Robert D. Wellman, MS; Diana S. M. Buist, PhD; Karla Kerlikowske, MD;  
Anna N. A. Tosteson, ScD; Diana L. Miglioretti, PhD; for the Breast Cancer Surveillance Consortium

Lehman et al. JAMA Internal Medicine 2015;175;(11):1828-37.doi 10.1001/jamainternmed.2015.5231

Laboratory Studies and even Human Reader Studies do not  
necessarily reflect real world performance

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Fee splitting with AI vendors?  
Sharing in Liability?  
Integration into hardware?  
New Capabilities - e.g. risk prediction?

Medicare and Private Insurers are unwilling to pay EXTRA for image interpretation using AI algorithms



THE  
TERMINATOR

# AUTONOMOUS AI IN IMAGE INTERPRETATION

- Adoption requires high level of trust in patient outcomes – by patients, clinicians, radiologists, which likely necessitates Real World Evidence without human-in-the-loop.
- Not permitted for breast cancer screening under the US Mammography Quality Standards Act.
- Removing cases from review by humans creates a more challenging case mix for humans. How will that impact performance and training?

# LESSONS FROM MY TIME AT ARPA-H LEADING THEIR CLINICAL TRIAL REFORM EFFORT

Too Expensive and  
Inefficient

Take Too Long

Not available to  
most Americans  
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Every Question cannot be answered through randomized trials.  
Pragmatic designs that use combined randomization and RW data will become  
more common and will use AI.

# WAYS AI COULD IMPROVE TRIAL EFFICIENCY

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- Identifying Eligible People
- Recruitment and Consenting
- Building Data Forms
- Extracting Data from EHRs
- Automatically identifying and flagging or correcting data errors
- Continuous engagement of trial participants to improve retention
- Flagging cross-over and other trial deviations and interrogating registries for those lost-to-follow-up
- Monitoring and Auditing

# ASSESS-AI

Algorithms drift. Staff and patients change

Monitoring performance in clinical practice by capturing clinical data

First national registry for monitoring AI performance

