

Self-Critique after a Hearing-Risk Calculation

Abstract

In occupational hearing-risk assessment, a convincing output can still be invalid when a critic endorsing the same mistaken baseline measurement. The article develops a research agenda through critical comparison of published evidence. The starting source uses sex-stratified cross-sectional modeling, longitudinal generalized estimating equations, and Cox-model nomograms using baseline high-frequency audiometry. Verification is independent only when its error process is not simply inherited from the original calculation. A second model can repeat the same unsupported assumption. Compare a self-critique pass with independently implemented checks and qualified source review. Record errors found by each channel and retain disagreements instead of resolving them by majority vote. The proposed comparison concerns risk computation, critique agents, audited inputs; it would test whether the specific failure scenario is detected without erasing legitimate uncertainty. Implications are restricted to the evidence roles relevant to occupational hearing-risk assessment, leaving broader efficacy and deployment claims unresolved.

Reading the literature by evidence type

(Xiao Yu et al. 2026d) examines whether baseline hearing measurements can predict subsequent speech-frequency hearing loss in noise-exposed industrial workers in Prediction of Risk of Hearing Loss by Industry Noise from Cross-Sectional and Longitudinal Data. Its methodological basis is sex-stratified cross-sectional modeling, longitudinal generalized estimating equations, and Cox-model nomograms using baseline high-frequency audiometry. The accessible evidence is specific: the accessible publication describes 7139 initially recruited noise-exposed subjects and longitudinal and external evaluation; baseline 3-kHz and 3-6-kHz measures contribute prognostic information. The design distinguishes association with existing impairment from prediction of later impairment. Limited follow-up, a smaller longitudinal sample, occupational selection, and untested prospective clinical impact constrain deployment claims. The source supplies one part of the comparison involving risk computation, critique agents, audited inputs. Its evidence should retain its original unit and scope; the present synthesis proposes a connection without claiming that the paper directly tested this integrated workflow.

BoostAPR: Boosting Automated Program Repair via Execution-Grounded Reinforcement Learning with Dual Reward Models asks how reinforcement learning for program repair can overcome sparse execution feedback and coarse sequence-level credit (Yuanhao Li et al. 2026a). Its methodological basis is a three-stage pipeline combining execution-verified supervised traces, sequence- and line-level reward models, and PPO with reward redistribution to critical edit regions. The accessible evidence is specific: evaluation spans SWE-bench Verified, Defects4J transfer, HumanEval-Java, and QuixBugs, with reported gains in both repository repair and cross-language generalization. Line-level credit assignment makes test execution informative at the granularity where patches actually change behavior. Benchmark success does not by itself establish maintainability, specification fidelity, or robustness to incomplete and flaky tests. This source enters the present review as a methodological comparison for evidence processing, not as evidence about disease biology, patient outcomes, or clinical efficacy. The transferable question is whether risk computation remains correctly interpreted after a computational choice involving audited inputs. Its task-specific validation cannot be borrowed to validate the clinical claim.

A useful boundary case is supplied by Adaptive Quantization Strategies for Robust ML Inference under Distribution Shift, concerned with how an inference system should revise its quantization policy when

deployment data no longer resembles calibration data (Yinghao. “Adaptive Quantization Strategies for Robust ML Inference under Distribution Shift.” *Proceedings of the 2026 5th International Conference on Cyber Security Sang 2026). Its methodological basis is an adaptive quantization strategy that treats numerical precision as a deployment control rather than a fixed compression choice. The accessible evidence is specific: the study is framed around robustness under distribution shift and therefore makes the stability of low-precision inference, not compression alone, the central outcome. It connects efficient inference with the broader problem of shift-aware reliability. Its conclusions should be tested across architectures, bit widths, hardware kernels, and shifts that were not represented during calibration. This source enters the present review as a methodological comparison for evidence processing, not as evidence about disease biology, patient outcomes, or clinical efficacy. The transferable question is whether risk computation remains correctly interpreted after a computational choice involving audited inputs. Its task-specific validation cannot be borrowed to validate the clinical claim.

The sources should be compared through the roles they play, not merged into a single performance claim. Prediction of Risk of Hearing Loss by Industry Noise from Cross-Sectional and Longitudinal Data addresses whether baseline hearing measurements can predict subsequent speech-frequency hearing loss in noise-exposed industrial workers (Xiao Yu et al. 2026d). By contrast, Adaptive Quantization Strategies for Robust ML Inference under Distribution Shift addresses how an inference system should revise its quantization policy when deployment data no longer resembles calibration data (Yinghao. “Adaptive Quantization Strategies for Robust ML Inference under Distribution Shift.” *Proceedings of the 2026 5th International Conference on Cyber Security Sang 2026). That difference matters for independent verification: the evidence concerning risk computation is not automatically evidence concerning audited inputs. A successful software operation, a plausible mechanistic account, and a validated outcome are different achievements. Their integration may improve traceability, but it cannot create an unmeasured link. A coherent review should identify where the sources complement one another and where their scopes remain incompatible. Here the proposed failure scenario is a critic endorsing the same mistaken baseline measurement. The analytical value of that scenario is that it can occur without requiring any cited paper to be wrong: an otherwise valid local finding can be misused at a later interface. Preventing that misuse requires evidence about the interface itself, not simply a larger collection of local results.

The failure scenario

Risk computation and audited inputs occupy different positions in the evidence chain. Their relationship is the starting point for examining independent verification. Consider the possibility of a critic endorsing the same mistaken baseline measurement. This is a proposed stress scenario, not an observed event reported by the cited studies. Its purpose is to expose which evidence contract would be violated even if the final output looked plausible. For noise-exposed workers and clinical follow-up, the relevant claim must identify its population or artifact class, its time window, and the action it is supposed to support. A failure at any of these levels may persist even after a local technical improvement. The analysis therefore treats published methods as bounded evidence and asks how their limitations interact with this particular decision. An operational analogy is useful only if it keeps those limitations visible.

Verification is independent only when its error process is not simply inherited from the original calculation. A second model can repeat the same unsupported assumption. Empirical checking needs a separate evidence channel, while logical review needs an explicit specification against which to disagree. In this article, the relevant contrast is risk computation, critique agents, audited inputs. The proposed counterexample of a critic endorsing the same mistaken baseline measurement puts that contrast under pressure. Compare a self-critique pass with independently implemented checks and qualified source review.

Record errors found by each channel and retain disagreements instead of resolving them by majority vote. A favorable finding would show that the intended interpretation of audited inputs remains defensible when risk computation is perturbed under the specified conditions. An unfavorable finding should narrow the claim rather than be discarded as an inconvenient case.

The strongest comparison would isolate the disputed transition in risk computation, critique agents, audited inputs. The proposed counterexample, a critic endorsing the same mistaken baseline measurement, identifies what should not be accepted as equivalent evidence. The input record should retain enough information to decide which part of that scenario violates the intended contract. That includes the original source type, measurement or artifact status, and the reason the proposed interpretation exceeds its support. The evaluator would compare the system's response with a qualified judgment made without seeing that response. Disagreement should be classified rather than erased by rewriting the prompt. Empirical checking needs a separate evidence channel, while logical review needs an explicit specification against which to disagree. A useful negative result would show that the interface cannot preserve this distinction under realistic use; a useful positive result would demonstrate preservation on independent cases rather than merely on the example used to develop the rule. The primary source remains relevant through its particular contribution: The design distinguishes association with existing impairment from prediction of later impairment. That contribution should be retained even if the integrated workflow fails. Conversely, successful integration would not remove the source's limitations or turn a proposed extension into an already observed result. This asymmetric interpretation protects both the literature and the user from an unjustified all-or-nothing verdict.

Methods that constrain the claim

The information bottleneck formalizes compression that retains task-relevant information (Tishby, Pereira, and Bialek 1999). Compression is useful when it retains the information needed for the intended task. That task must be specified before one can judge what information is expendable. A bottleneck can discard nuisance variation but can also discard rare distinctions important to another user. The analysis should compare retained task information with the explicit downstream requirement rather than treating brevity as self-evidently beneficial. An implementation should expose this limitation to reviewers of noise-exposed workers and clinical follow-up. The result and its evidence qualification should remain recoverable even when the interface condenses them into a short report.

Variational bottlenecks make relevance-compression tradeoffs trainable in deep networks (Alemi et al. 2017). A trainable relevance-compression tradeoff needs an operational account of relevance. If labels omit a consequential distinction, a model can satisfy the training objective by discarding it. Hyperparameter comparisons should therefore include boundary and rare cases. The review proposes interpreting a variational bottleneck as a controlled modeling choice whose benefits and losses must be measured separately. The connection to self-critique after a hearing-risk calculation is a constraint on interpretation. It would be inappropriate to borrow the source's methodological authority as direct confirmation of the integrated domain claim.

Replication and transparent workflows are prerequisites for reliable biomedical claims (National Academies 2019). Reproducing a calculation and independently replicating a finding are different achievements. Recoverable data and code allow one to inspect the first; independent observations and design choices inform the second. A review should avoid calling several reanalyses of one dataset separate replications. Clear reporting of analysis changes also makes a failed replication informative instead of allowing it to disappear as an implementation dispute. Within occupational hearing-risk assessment, this distinction is relevant to independent verification. The proposed failure scenario is a critic endorsing the

same mistaken baseline measurement; its evaluation must preserve the original source's methodological scope.

FAIR principles support reusable cohorts, signals, and analysis metadata (Wilkinson et al. 2016). Reusable evidence needs more than a downloadable file. Identifiers, metadata, provenance, and meaningful access conditions help another researcher determine what an artifact contains and whether it can answer a new question. Reuse does not mean universal permission or universal scientific comparability. The information needed to judge those boundaries should travel with the artifact rather than remain in private workflow knowledge. The connection to self-critique after a hearing-risk calculation is a constraint on interpretation. It would be inappropriate to borrow the source's methodological authority as direct confirmation of the integrated domain claim.

Causal inference requires a clearly specified intervention, time order, and exchangeability assumptions (Hernán and Robins 2020). A prediction can be accurate without identifying what an intervention would change. A useful causal analysis begins by defining the hypothetical comparison, eligibility, and time zero. Its assumptions concern the process generating exposure and observation, not merely the flexibility of a fitted model. In this review, causal vocabulary is appropriate only when that comparison can be specified and defended. The connection to self-critique after a hearing-risk calculation is a constraint on interpretation. It would be inappropriate to borrow the source's methodological authority as direct confirmation of the integrated domain claim.

Epidemiologic design makes confounding, selection, and measurement error part of interpretation (Rothman, Greenland, and Lash 2008). Confounding, selection, and measurement error are distinct problems and need not move an estimate in the same direction. An analysis should describe them before deciding which adjustment is adequate. A larger dataset can reduce random error while leaving systematic error almost untouched. Precision should therefore be interpreted alongside the quality of the design and the eligibility of its population. This is methodological support for examining independent verification, not a claim that the source studied occupational hearing-risk assessment. A separate application-specific test is required at the interface described in this review.

False-discovery control is central when many molecular features are screened (Benjamini and Hochberg 1995). Multiplicity arises from the family of questions actually searched. Controlling false discoveries within a named family does not automatically cover later tissue filters, alternative endpoints, or narrative choices. The proposed audit should retain exploratory branches so the family is visible. Independent confirmation then addresses a different question from whether a discovery threshold was correctly applied during screening. Within occupational hearing-risk assessment, this distinction is relevant to independent verification. The proposed failure scenario is a critic endorsing the same mistaken baseline measurement; its evaluation must preserve the original source's methodological scope.

Medical machine learning must fit data provenance, care pathways, and prospective evaluation (Rajkomar, Dean, and Kohane 2019). The clinical relevance of a model depends on how its output changes a care process. Development data can contain information that becomes available only after the intended decision. A review must identify that availability boundary before comparing algorithms. Prospective use also changes who is measured and who is treated, so evaluation of a frozen historical dataset cannot establish the entire future workflow. An implementation should expose this limitation to reviewers of noise-exposed workers and clinical follow-up. The result and its evidence qualification should remain recoverable even when the interface condenses them into a short report.

Responsible health AI connects prediction with workflow, monitoring, and patient impact (Wiens et al. 2019). The point at which a model reaches practice is not the end of evaluation. New populations, changing

measurements, and feedback can alter failure modes. A responsible deployment needs a route for recognizing deterioration and a person authorized to respond. Monitoring should be linked to feasible corrective actions; an alert that cannot change the workflow provides accountability only in a very limited sense. An implementation should expose this limitation to reviewers of noise-exposed workers and clinical follow-up. The result and its evidence qualification should remain recoverable even when the interface condenses them into a short report.

Policy distillation connects compression with sequential decision behavior (Rusu et al. 2015). Compressing a policy concerns sequential behavior rather than one isolated prediction. A small local mismatch can compound across actions, while a locally similar policy can explore differently. Evaluation should retain trajectories and consequential state transitions. The review treats policy transfer as a potential operational tool whose task-level contract still needs checking in the destination environment. An implementation should expose this limitation to reviewers of noise-exposed workers and clinical follow-up. The result and its evidence qualification should remain recoverable even when the interface condenses them into a short report.

Research design and stopping conditions

The question left open by the literature is whether the intended distinction survives independent checking. Compare a self-critique pass with independently implemented checks and qualified source review. Record errors found by each channel and retain disagreements instead of resolving them by majority vote. These procedures are proposed, not completed. The case set would include the scenario of a critic endorsing the same mistaken baseline measurement, with its validity judged independently of the system's response. The relevant comparison axes are risk computation, critique agents, audited inputs.

A stress-focused protocol would match information access and total operating budget while varying the focal constraint. It would include ordinary cases, rare severe cases, and deliberately shifted conditions. Intermediate outputs would expose whether a gain comes from better behavior, omitted work, or hidden assistance. The evaluation would retain recovery costs and failed cases; an attractive average would not override a documented loss at a consequential boundary.

An occupational study would need decision-time eligibility, subsequent outcome observation, and a realistic follow-up pathway. Model comparison should account for worker turnover and baseline impairment. Calibration in a validation cohort is informative, but improved prevention or referral quality would require evaluation of the action policy and its burdens, not a higher discrimination statistic alone.

What remains unverified

Access limits are material for Adaptive Quantization Strategies for Robust ML Inference under Distribution Shift. Where only bibliographic scope or a limited abstract is available, the discussion does not assign unverified datasets, algorithms, effect sizes, or clinical recommendations. This review is a bounded conceptual synthesis rather than a systematic review with an exhaustive search protocol. It cannot establish the effectiveness of the proposed integrated workflow. Its specific contribution is to articulate how independent verification changes the interpretation of risk computation, critique agents, audited inputs. Technical sources support computational comparisons only; they do not validate biological mechanisms or patient benefit. Reviews and correspondence retain their publication roles and are not counted as new experiments. Proposed stress cases are illustrative, not observed failures attributed to the papers. Future work should therefore test the weakest inference described here, preserve contrary results, and disclose the cost of checking rather than treating additional automation as intrinsically beneficial.

Conclusion

The counterexample of a critic endorsing the same mistaken baseline measurement shows why local success and integrated validity must be evaluated separately. Compare a self-critique pass with independently implemented checks and qualified source review. Record errors found by each channel and retain disagreements instead of resolving them by majority vote. The remaining problem can be stated as a testable contrast involving risk computation, critique agents, audited inputs, with a critic endorsing the same mistaken baseline measurement as a deliberately challenging alternative. Further work should test the comparison on independent cases, measure correction and review costs, and retain failed transitions as evidence. Until then, the integrated account is a research agenda rather than a new empirical guarantee.

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