

Considerations on safety assurance for AI

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What is “assurance”?

- ISO/IEC/IEEE 15026-1:2019, “Systems and software engineering — Systems and software assurance”
- Assurance: grounds for justified confidence that a claim has been or will be achieved
- Assurance case: reasoned, auditable artefact created that supports the contention that its top-level claim (or set of claims) is satisfied, including systematic [arguments] and its underlying evidence and explicit assumptions that support the claim(s)
 - Note 1 to entry: An assurance case contains the following and their relationships:
 - one or more claims about properties
 - arguments that logically link the evidence and any assumptions to the claim(s)
 - a body of evidence and possibly assumptions supporting these arguments for the claim(s)
 - justification of the choice of top-level claim and the method of reasoning

What do standards require?

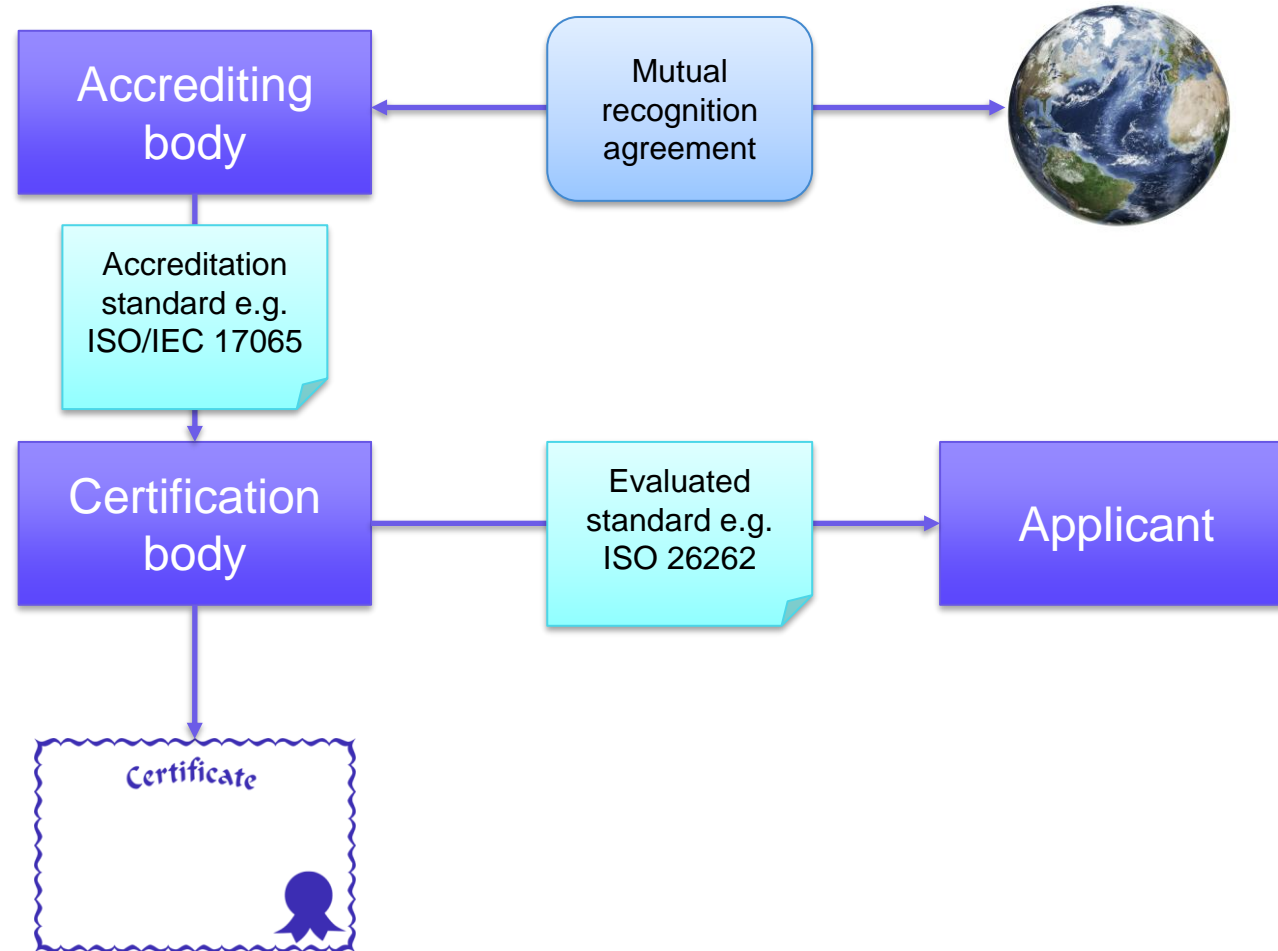
- Standards for functional safety and cybersecurity specify a number of activities to give assurance in the robustness of the product, including independent reviews
- These activities include the following (but precise application varies by standard)
 - Verification reviews
 - Confirmation reviews
 - Validation
 - Assurance case
 - Management system (engineering process) audit
 - Product audit
 - Product assessment
 - Release for production

- Clause 7 – AI safety management
 - Calls up ISO 26262 Part 2 including the confirmation measures
 - Confirmation review of safety plan, AI system validation report, AI safety analyses, AI safety assurance argument
 - AI safety audit and AI safety assessment
- Clause 8 – assurance arguments for AI systems
 - Seen as part of the ISO 26262 safety case
 - Objectives
 - Demonstrate the safety requirements allocated to the AI system are fulfilled
 - Evaluate whether the assurance argument reflects the actual residual risk of the AI system violating its safety requirements
 - Significant content around evidence and structure of the assurance argument

What about certification?

- Certification is a more formal activity of declaring that a product or process is in conformity with something, usually a standard
- Certification is not a requirement of ISO 26262 etc. and they are not “certification” standards that have to be met in the same formal way as e.g. DO-178C
- “Anyone” can issue a certificate, but the gold standard is for these to be accredited by a national body (e.g. UKAS in the UK, DAkkS in DE)
- Accreditation brings impartiality requirements that are much more onerous than “I3” independence in ISO 26262
 - In particular a certification body cannot provide conformity assessment on a product or process it (or its related bodies) has also provided consultancy on

Certification relationships



- ISO 26262, ISO/SAE 21434, etc. are engineering standards and certification against them is not mandatory
 - Any certification schemes are commercial products of the bodies offering them, are not “official” and do not follow a common approach
- Certification can be seen as attractive in the supply chain so that providers of components can demonstrate that a competent independent party has reviewed a product or process
- A certificate does not guarantee a product is correct for its application
- A certificate does not remove the need to follow a safety manual, integrator guidance, etc.
- A certificate does not guarantee a product is free from issues, nor grant immunity from marketplace actions

Questions to ask a Certification Body

- A certification body will have a “scheme document” that is required to be public and should be made available on request
- Who is the accrediting body and which accreditation standard has been used?
 - The appropriate standards are
 - ISO/IEC 17021 – for xSMS audit alone
 - ISO/IEC 17065 – for product certification (also covers process audit)
- How is impartiality managed?
- What are the surveillance requirements?
- What is the scope of a certificate? Also ask to see the associated report
 - What Parts or Clauses of a standard have been examined?
 - What process or product variants does it apply to?
 - What are the obligations on the end user? e.g. requirements in a safety manual



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