

# The European Union Artificial Intelligence Act

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# The EU Artificial Intelligence Act

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The EU AI Act was adopted at the end of May 2024 and enters into force from 1 August 2024. Thereafter, the Act's tiered compliance obligations will progressively come into application over a phased timeline, with most requirements taking effect by mid-2027.

The Artificial Intelligence (AI) Act is a landmark in global AI regulation, reflecting the EU's objective to lead the way in promoting a comprehensive legislative approach to support the trustworthy and responsible use of AI systems. The AI Act follows other major EU digital legislation, such as the General Data Protection Regulation (GDPR), the Digital Services Act, the Digital Markets Act, the Data Act, and the Cyber Resilience Act.

This paper outlines key elements of the AI Act and provides an overview of the Act's tiered compliance obligations.

This paper does not constitute legal advice.

The AI Act will unify how AI is regulated across the single market of the 27 EU Member States. It also has important extraterritorial implications, as it covers all AI systems impacting people in the EU, regardless of where these systems are developed or deployed from.

Compliance obligations are significant, and largely determined by the level of risk the usage of an AI system poses to people's safety, security, or fundamental rights. Obligations apply along the AI value chain. The AI Act applies a tiered compliance framework: most requirements fall upon the developers and deployers of AI systems classified as "high-risk", and on general-purpose AI models (including foundation models and generative AI systems) posing "systemic risks".

Reflecting these tiers, the AI Act sets-out a phased application timeline, starting with requirements for prohibited AI systems in February 2025 and progressively extending to all AI systems by 2030. There are significant financial penalties for noncompliance, up to 7% of worldwide annual turnover (revenue).

It is important for business leaders, both in the EU and beyond, to consider the implications of this complex legislation without delay. This consideration includes understanding how the AI Act interacts with existing and emerging rules and regulations in other jurisdictions, as well as with voluntary AI codes and principles.

Businesses and other organizations should ensure they have an up-to-date inventory of the AI systems that they are developing or deploying. They will need to assess whether their systems are subject to new compliance obligations and, if so, under which classification. Developers and deployers of high-risk and general-purpose AI systems in particular, will also need to ensure that they have robust and effective AI governance frameworks and compliance systems in place.

## Key takeaways

### Who will the AI Act affect?

- ▶ The AI Act applies to all AI systems impacting people in the EU (whether these AI systems are built and operated from within the EU or from elsewhere). It applies across all sectors.
- ▶ The AI Act imposes different obligations across all actors in the AI value chain.
- ▶ In certain cases, the AI Act also applies to AI models and systems placed on the market *prior* to the Act coming into application, including:
  - ▶ If these are general purpose AI (GPAI) models or systems (see GPAI definition below)
  - ▶ If these are AI systems which fall into the "prohibited" category, or if these are "high-risk" AI systems that are intended to be used by public authorities.



- ▶ Moreover, if an existing AI system undergoes significant changes, it will be treated like the other systems in its “updated” risk category that are being placed on the market at the same time as when the update took place.

### What are the key features of the AI Act?

- ▶ **Definition of AI:** The AI Act applies a broad definition of an AI system derived from the Organisation for Economic Co-operation and Development definition (see relevant section below).
- ▶ **Risk-based approach focusing on use cases:** Obligations are primarily based on the level of risk posed by how an AI system is used (or could be used), not the technology on which it is based.
  - ▶ GPAI models and systems are treated separately due to the breadth of their potential use cases.
- ▶ **Risk classification system:** The AI Act establishes a tiered compliance framework consisting of different categories of risk, with different requirements for each category. All AI systems will need to be inventoried and assessed to determine their risk category and the ensuing responsibilities.
  - ▶ **Prohibited systems:** AI systems posing what legislators consider an unacceptable risk to people’s safety, security and fundamental rights will be banned from use in the EU.
  - ▶ **High-risk AI systems:** These systems will carry the majority of compliance obligations (alongside GPAI systems - see below), including the establishment of risk and quality management systems, data governance, human oversight, cybersecurity measures, post-market monitoring, and maintenance of the required technical documentation. (Further obligations may be specified in subsequent AI regulations for healthcare, financial services, automotive, aviation, and other sectors.)
  - ▶ **Minimal-risk AI systems:** Beyond the initial risk assessment and some transparency requirements for certain AI systems, the AI Act imposes no additional obligations on these systems, but invites companies to commit to codes of conduct on a voluntary basis.
- ▶ **Pre-market conformity assessments for high-risk AI systems:** High-risk systems will require a conformity assessment to evidence their compliance before being placed on the market:
  - ▶ The application of EU harmonized standards (currently under development, see below) will allow AI system providers to demonstrate compliance by self-assessment.
  - ▶ In limited cases, a third-party conformity assessment performed by an accredited independent assessor (“notified body”) will be required.
- ▶ **General purpose AI models (GPAI), including foundation models and generative AI:** These advanced models, and the systems built on these models, will be regulated through a separate tiered approach, with additional obligations for models deemed to pose a “systemic risk”.
- ▶ **Measures to support innovation:** Regulatory “sandboxes” will be put in place across the EU for operators (especially small and medium enterprises) to access voluntarily. Here they can innovate, experiment, test, and validate the compliance of their AI systems with the AI Act in a safe environment.
- ▶ **Interaction with other EU laws:** Obligations under the AI Act will need to be integrated into the compliance processes already established to implement existing EU laws, including, GDPR and data laws, product safety laws, copyright, cybersecurity, and sectorial regulations (e.g. financial services, energy, automotive, health).
- ▶ **Enforcement and penalties:** National competent authorities will have enforcement powers with the capacity to impose significant fines depending on the level of noncompliance.
  - ▶ For use of prohibited AI systems, fines may be up to 7% of worldwide annual turnover (revenue), while noncompliance with requirements for high-risk AI systems will be subject to fines of up to 3% of the same.

### When will the AI Act take effect?

- ▶ The Act enters into force from 1 August 2024, with different obligations then taking effect in stages. Some key dates are outlined below:

- ▶ AI Act prohibitions will start to be enforced six months after the Act enters into force (so, from 2 February 2025).
- ▶ GPAI obligations will take effect 12 months after entry into force (2 August 2025), but with one exception: GPAI models which have been placed on the market before this date will have an additional 24 months to comply (so from 2 August 2027).
- ▶ Most other obligations will take effect 24 months after the Act enters into force (so 2 August 2026).
- ▶ However:
  - ▶ Obligations for AI systems that are classified as high-risk because they are a safety component of a system that is subject to Union harmonization legislation (listed in Annex I), will only take effect 36 months after the Act enters in force (so from 2 August 2027).
  - ▶ Obligations for high-risk AI systems intended for use by public authorities that were on the market before the entry into force of the AI Act, will only take effect from 31 December 2030).

#### **What actions should companies and other organizations take from the outset?**

- 1) Inventory all AI systems you have (or potentially will have) developed or deployed and determine whether any of these systems falls within the scope of the AI Act.
- 2) Assess and categorize the in-scope AI systems to determine their risk classification and identify the applicable compliance requirements, including taking prohibited systems out of service ahead of the legal deadline (six months after the entry into force of the Act i.e., February 2025).
- 3) Understand your organization's position in relevant AI value chains, the associated compliance obligations and how these obligations will be met. Compliance will need to be embedded in all functions responsible for the AI systems along the value chain throughout their lifecycle.
- 4) Consider what other questions, risks (e.g., interaction with other EU or non-EU regulations, including on data privacy), and opportunities (e.g., access to AI Act sandboxes for innovators, small and medium enterprises, and others) the AI Act poses to your organization's operations and strategy.
- 5) Develop and execute a plan to ensure that the appropriate accountability and governance frameworks, risk management and control systems, quality management, monitoring, and documentation are in place when the Act comes into force considering the phased implementation timeline.

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## Context

The AI Act is intended to advance four key objectives:<sup>1</sup>

- (i) To ensure that AI systems placed on the EU market are safe and respect fundamental rights
- (ii) To ensure legal certainty to facilitate investment and innovation in AI
- (iii) To enhance governance and effective enforcement of EU law on fundamental rights and safety requirements applicable to AI systems
- (iv) To facilitate the development of a single market for lawful, safe and trustworthy AI applications, and prevent market fragmentation

## Who is affected?

The AI Act is broad in scope and comes with significant obligations along the value chain. It focuses on the impact of AI systems on people, specifically on their wellbeing and fundamental rights.

It also contains extraterritorial measures, affecting any business or organization that offers an AI system impacting people within the EU, regardless of where the organization is headquartered.

Under certain conditions the AI Act also applies to AI systems that were put on the market prior to the Act taking effect:

- ▶ If these are GPAI models
- ▶ If these are AI systems which fall into the “prohibited” category, or if these are “high-risk” AI systems that are intended to be used by public authorities
- ▶ Moreover, if an existing AI system undergoes significant changes, it will be treated like the other systems in its “updated” risk category that are being placed on the market at the same time

The AI Act will apply to (please see the appendix section below for full definitions of terms):

- ▶ Providers putting AI systems on the market within the EU, regardless of their location
- ▶ Providers and deployers of AI systems located in a non-EU country, where the output of the AI system is used within the EU
- ▶ Deployers of AI systems located in the EU
- ▶ Importers and distributors placing AI systems on the EU market
- ▶ Product manufacturers placing products with AI systems on the EU market under their own name or trademark

The AI Act will **not** apply to:

- ▶ Public authorities in non-EU countries and international organizations that have law enforcement and judicial cooperation agreements with the EU, provided that adequate safeguards are in place
- ▶ AI systems used for purposes outside the scope of EU law-making authority, such as military or defense
- ▶ AI systems specifically developed and used for the sole purpose of scientific research and discovery
- ▶ Research, testing and development activity regarding AI systems prior to placement on the market or into service
- ▶ Free and open-source software, unless their use would classify them as a prohibited or high-risk AI system, or their use would subject them to transparency obligations.

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<sup>1</sup> “EU AI Act Proposal, 2021 - Explanatory Memorandum”, European Commission, April 2021 <https://eur-lex.europa.eu/legal-content/EN/TXT/HTML/?uri=CELEX:52021PC0206>

## When will the AI Act be implemented?

The AI Act has been approved by the European Parliament and Council and was published in the Official Journal on 12 July 2024, entering into force on 1 August 2024. As an EU regulation (as opposed to a directive), it will be directly effective in Member States without the need for local enabling legislation.

The timeline for compliance with the provisions of the AI Act will be as follows:

Timeframe	Development
1 August 2024	AI Act enters into force.
Immediately after entry into force	The European Commission must begin work to establish the AI Office (EU oversight body) while Member States make provisions to establish AI regulatory sandboxes. (To note: the AI Office has already been formally established)
Six months after entry into force (2 February 2025)	AI Act prohibitions will come into effect.
12 months after entry into force (2 August 2025)	Requirements for GPAI models will come into effect. However, GPAI models that were already on the market before this date will have an additional 24 months to comply (see below).
24 months after entry into force (2 August 2026)	Requirements for high-risk AI systems (classified under uses listed in Annex III) will come into effect, alongside transparency requirements for certain other AI systems.
36 months after entry into force (2 August 2027)	Requirements for high-risk AI systems classified under EU harmonization laws contained in Annex I will come into effect. GPAI models that were already on the market before obligations began to apply twelve months after entry into force (see above), will now have to comply.
31 December 2030	High-risk AI systems intended for use by public authorities that were on the market before the entry into force of the AI Act should now be compliant.

## How does the EU define an AI system?

The AI Act's definition of an AI system is derived from the definition used by the Organisation for Economic Co-operation and Development (OECD). The objective in using the OECD definition as a basis, is to encourage international alignment and continuity with other laws and codes. The AI Act defines an AI system as follows:

*"An AI system is a machine-based system designed to operate with varying levels of autonomy and that may exhibit adaptiveness after deployment and that, for explicit or implicit objectives, infers, from the input it receives, how to generate outputs such as predictions, content, recommendations, or decisions that can influence physical or virtual environments."*

The AI Act emphasizes that a key characteristic that differentiates AI systems from simpler and more traditional software systems is their capability to infer. It states that the techniques that enable inference while building an AI system include machine learning approaches that learn from data how to achieve a certain objective, and logic- and knowledge-based approaches that infer from encoded knowledge or symbolic representation of the task to be solved. The capacity of an AI system to infer goes beyond basic data processing, enable learning, reasoning, or modelling.

## How are AI systems classified?

The AI Act sets compliance obligations based on the inherent risks that arise from the application for which AI systems are used.



General-purpose AI models (GPAI), including foundation models and generative AI systems, follow a separate classification framework. Please see the relevant section below.

AI systems are classified as follows in the Act:

Classification (Risk-based tier)	Description	Compliance level	Use case examples (see sections below for fuller details)
<b>Prohibited AI systems</b>	<b>Prohibited</b> because uses pose an unacceptable risk to the safety, security, and fundamental rights of people.	Prohibition	Includes use of AI for <b>social scoring</b> which could lead to detrimental treatment, <b>emotional recognition</b> systems in the workplace, <b>biometric categorization</b> to infer sensitive data, and <b>predictive policing</b> of individuals, among other uses. Some exemptions will apply.
<b>High-risk AI systems</b>	<b>Permitted</b> , subject to compliance with the requirements of the AI Act (including conformity assessments before being placed on the market).	Significant	Includes use of AI in:  Employment, biometric identification surveillance systems, Safety components of systems covered by harmonised legislation (e.g., medical devices, automotive), access to essential private and public services (e.g., creditworthiness, benefits, health and life insurance), safety of critical infrastructure (e.g., energy, transport).
<b>Minimal risk AI systems</b>	<b>Permitted</b> , subject to specific transparency and disclosure obligations where uses pose a limited risk.	Limited	Certain AI systems that interact directly with people (e.g., <b>chatbots</b> ), and visual or audio <b>“deepfake”</b> content that has been manipulated by an AI system.
	<b>Permitted</b> , with no additional AI Act requirements where uses pose minimal risk.	Minimal	By default, all other AI systems that do not fall into the above categories (e.g., photo-editing software, product-recommender systems, spam filtering software, scheduling software)

## Prohibited systems: which use cases pose an unacceptable risk?

The AI Act prohibits AI systems that pose unacceptable risks and that can be used to undermine a person’s fundamental rights, or that may subject them to physical or psychological harm. These prohibitions include:

- ▶ AI systems that exploit vulnerabilities, or deploy subliminal techniques, to manipulate a person or a specific group (e.g., children, the elderly, or people with disabilities), circumventing the users’ free will in a manner likely to cause harm.
- ▶ AI systems used for the social scoring, evaluation, or classification of people based on their social behavior, inferred, or predicted, or personal characteristics, leading to detrimental treatment.
- ▶ AI systems used to infer emotions of people in the workplace (such as human resource functions) and educational institutions. Exemptions apply for some safety systems (e.g., detection of the drowsiness of pilots).
- ▶ Biometric categorization to infer sensitive data, such as race, sexual orientation, or religious beliefs.
- ▶ Indiscriminate and untargeted scraping of facial images from the internet or CCTV to populate facial recognition databases.

- ▶ Predictive policing of individuals based on the profiling or assessment of their personality traits and characteristics to predict the risk of a natural person committing a criminal offense. This prohibition does not apply to AI systems used in assessment of involvement in occurred criminal activity which is already based on objective and verifiable facts.
- ▶ Law enforcement use of real-time remote biometric identification (RBI) systems in publicly accessible spaces (certain exceptions apply subject to prior judicial authorization and for strictly defined lists of criminal offenses).

## High-risk systems: which use cases are subject to conformity assessments and obligations?

The AI Act identifies high-risk uses in Annex I and Annex III. The European Commission is empowered to update these annexes as new uses and risks are identified. The following high-risk uses are currently listed:

- ▶ AI systems used as a safety component of a product covered by EU harmonization legislation, including but not limited to:<sup>2</sup>
  - ▶ Medical devices
  - ▶ Motor vehicles
  - ▶ Machinery
  - ▶ Civil aviation
  - ▶ Radio equipment
  - ▶ Pressure equipment
  - ▶ Marine equipment
  - ▶ Agricultural vehicles
  - ▶ Railway interoperability
  - ▶ Toys
  - ▶ Watercraft
  - ▶ Lifts
- ▶ AI systems applied in uses that pose a significant risk of harm to health, safety, or fundamental rights:<sup>3</sup>
  - ▶ Biometric identification and categorization of people (including those emotional recognition systems and remote biometric identification systems that are not prohibited)
  - ▶ Management and operation of critical infrastructure (specifically, safety components of traffic, water, gas, heating, and electricity infrastructure)
  - ▶ Education and vocational training (specifically, systems determining access to education and assessment of students)
  - ▶ Employment, worker management and access to self-employment (including recruitment and performance monitoring)
  - ▶ Access to and enjoyment of essential private and public services and benefits (including eligibility for benefits, evaluating creditworthiness, and pricing of life and health insurance, although those used for purposes of detecting financial fraud are specifically not included)
  - ▶ Law enforcement uses such as data analytics systems to assess evidence of criminal activity
  - ▶ Migration, asylum, and border control management (including monitoring of migration trends, border surveillance, verification of travel documents, and examination of applications for visas, asylum, and residence permits)
  - ▶ Administration of justice and democratic processes (including researching and interpreting the law, and those used for influencing the outcome of an election)

The Commission has powers to add or modify these high-risk use-cases.

### Exceptions to high-risk classification:

However, an AI system that's use is referred to in Annex III will not be considered high-risk if it does not pose a significant risk of harm to the health, safety or fundamental rights of natural persons because the system:

<sup>2</sup> Annex I, List of Union harmonisation legislation, EU Artificial Intelligence Act, [Regulation - EU - 2024/1689 - EN - EUR-Lex \(europa.eu\)](#), 12 July 2024

<sup>3</sup> Annex III, High-risk AI systems referred to in Article 6(2), EU Artificial Intelligence Act [Regulation - EU - 2024/1689 - EN - EUR-Lex \(europa.eu\)](#), 12 July 2024

- ▶ Performs a narrow procedural task
- ▶ Is intended to improve the result of a previously completed human activity
- ▶ Is used to detect decision-making patterns or deviations from existing patterns to flag inconsistencies and is not meant to replace or influence the previously completed human assessment, without proper human review (e.g., fraud detection)
- ▶ Is intended to perform a preparatory task to an assessment relevant to the use cases in Annex III

Providers that determine that their AI system is not high-risk for one of these reasons (despite being referred to in Annex III) should document their assessment in the event that this documentation is requested by a national competent authority.

## What are the obligations for providers of high-risk AI systems?

### General obligations

Requirements for high-risk AI systems include:

- ▶ Establishing and maintaining appropriate AI **risk** and **quality management systems**
- ▶ Effective **data governance**
- ▶ Maintaining appropriate **technical documentation** and **record-keeping**
- ▶ **Transparency** and provision of information to users
- ▶ Enabling and conducting **human oversight**
- ▶ Compliance with standards for **accuracy, robustness, and cybersecurity** for the intended purpose
- ▶ **Registering high-risk AI systems on the EU database** before placing them on the market; systems used for law enforcement, migration, asylum and border control, and critical infrastructure will be registered in a non-public section of the database

### Pre-market conformity assessment for high-risk systems

Providers must perform a conformity assessment on the high-risk AI system before placing it on the market:

- ▶ The conformity assessment should examine whether the requirements laid out above have been met

In most cases, **providers can self-assess** and will benefit from a presumption of conformity if:

- ▶ They apply procedures and methodologies that follow EU approved technical standards (harmonized standards)

A **third-party conformity assessment** by an accredited body (a “notified body”) is required if any of the following criteria apply:

- ▶ The AI system is part of a safety component subject to third-party assessment under Union harmonized regulations (see above)
- ▶ The AI system is part of a biometric identification system and harmonized standards are not used, or are only partly applied

### Post-market obligations

Once a high-risk AI system has been placed on the market, providers continue to have obligations to ensure ongoing safe performance and conformity over the system’s lifecycle. These include:

- ▶ **Maintaining logs** generated by high-risk systems, to the extent that they are under their control, for a period of at least six months
- ▶ **Immediately taking the necessary corrective actions** for nonconforming systems already on the market and informing other operators in the value chain of the nonconforming systems

- ▶ **Cooperating with the national competent authorities or the AI Office** (see relevant section below) by sharing all the information and documentation necessary to show conformity upon receiving a reasonable request
- ▶ **Monitoring performance and safety** of AI systems throughout their lifetime and actively evaluating continuous compliance with the AI Act
- ▶ **Reporting to the appropriate authorities, serious incidents** and malfunctions that lead to breaches of fundamental rights
- ▶ **Undergoing new conformity assessments for substantial modifications** (e.g., changes to a system's intended purpose or changes that affect how it meets regulations):
  - ▶ This applies whether the changes are made by the original provider or any third party.
  - ▶ For AI systems that are considered to have limited or minimal risk, it will be important to check whether the original risk classification still applies after any changes.

## What are the obligations for deployers, importers and distributors of high-risk AI systems?

**Obligations of deployers** of high-risk AI systems include:

- ▶ Completing a fundamental rights impact assessment (FRIA) before putting the AI system in use, if the deployer:
  - ▶ Is a public body or private entity providing public services
  - ▶ Provides essential private service that cover creditworthiness evaluation of persons, and risk assessment and pricing in relation to life and health insurance
- ▶ Implementing human oversight by people with the appropriate training and competence
- ▶ Ensuring that input data is relevant to the use of the system
- ▶ Suspending the use of the system if it poses a risk at a national level
- ▶ Informing the AI system provider of any serious incidents
  - ▶ Retaining the automatically-generated system logs
  - ▶ Complying with the relevant registration requirements when the user is a public authority
- ▶ Complying with GDPR obligations to perform a data protection impact assessment
- ▶ Verifying the AI system is compliant with the AI Act and that all relevant documentation is evidenced
- ▶ Informing people, they might be subject to the use of high-risk AI

**Before placing a high-risk AI system on the market, it is the responsibility of importers and distributors to:**

- ▶ Verify that the system complies with the AI Act, ensure that all relevant documentation is evidenced, and communicate with the provider and market surveillance authorities accordingly

## Minimal-risk systems: what obligations apply?

For some specific AI systems, limited transparency obligations apply.

Providers must:

- ▶ Design and develop systems in a way to make certain that people understand that they are interacting with an AI system from the outset (e.g., chatbots)

Deployers must:

- ▶ Inform and obtain the consent of people exposed to permitted emotion recognition or biometric categorization systems (e.g., safety systems monitoring driver attentiveness)
- ▶ Disclose and clearly label where visual or audio “deep fake” content has been manipulated by AI.

## How will general-purpose AI be regulated?

The definition in the AI Act of general-purpose AI (GPAI) models is:

*“General-purpose AI model” means an AI model, including when trained with a large amount of data using self-supervision at scale, that displays significant generality and is capable to competently perform a wide range of distinct tasks regardless of the way the model is placed on the market and that can be integrated into a variety of downstream systems or applications. This does not cover AI models that are used before release on the market for research, development and prototyping activities.”*

The AI Act adopts a tiered approach to compliance obligations, **differentiating between high-impact GPAI models with systemic risk, and other GPAI models**. The AI Act defines “systemic risk at Union level” as:

*“A risk that is specific to the high-impact capabilities of general-purpose AI models, having a significant impact on the internal market due to its reach, and with actual or reasonably foreseeable negative effects on public health, safety, public security, fundamental rights, or the society as a whole, that can be propagated at scale across the value chain.”*

The GPAI tiers are as follows:

Tier	Description	Compliance level
Base-level tier	Models meeting the GPAI definition	Limited transparency obligations
Systemic risk tier	<p>High-impact GPAI models posing a systemic risk are provisionally identified based on cumulative amount of computing power used for training (with power greater than <math>10^{25}</math> floating point operations [FLOPs]).</p> <p>A model can also be classified in this tier based on a decision of the Commission that a general-purpose AI model has capabilities or impact equivalent to those above.</p>	Significant obligations

**Providers of all GPAI models** will be required to:

- ▶ Keep and maintain up-to-date technical documentation.
- ▶ Make information available to downstream providers who intend to integrate the GPAI model into their AI systems.
- ▶ Put in place a policy to respect EU copyright law.
- ▶ Disseminate detailed summaries about the content used for training.

**Exceptions to base-level GPAI transparency obligations (first two bullet points directly above only):**

- ▶ Unless the GPAI models present systemic risks, these obligations shall not apply to providers of GPAI models that are made accessible to the public under a free and open-source license, and whose parameters are made publicly available.

**In addition, providers of high-impact GPAI models posing a systemic risk must:**

- ▶ Perform model evaluations.



- ▶ Assess and mitigate systemic risks.
- ▶ Document and report to the AI Office any serious incidents and the corrective action taken.
- ▶ Conduct adversarial training of the model (i.e., “red-teaming”).
- ▶ Ensure that an adequate level of both cybersecurity and physical protections are in place.
- ▶ Document and report the estimated energy consumption of the model.

**To provide agility for adapting to rapid GPAI technology developments, the AI Office (see relevant section below) will:**

- ▶ Update the designation criteria for high-impact GPAI, with possible inclusion of criteria related to the number of model parameters, quality or size of datasets, number of registered business or end users.
- ▶ Facilitate the formulation of codes of practice to support the application of the compliance requirements. Providers may rely on these codes of practice to demonstrate compliance until a harmonized standard is published (see below).

## How will the AI Act interact with existing legislation and standards?

- ▶ AI providers must continue to adhere to all relevant EU laws while incorporating requirements of the AI Act.
- ▶ Providers can combine AI Act compliance with existing procedures to avoid duplication and ease the compliance workload.
- ▶ Where applicable, the AI Act should be embedded into relevant EU laws (e.g., financial services regulations). Sectoral regulators will be designated as the relevant competent authorities to supervise the enforcement of the AI Act for their sector.

## How will new standards be developed and when will they be ready?

To reduce compliance burdens and speed up time-to-market, the AI Act allows for compliance self-assessment, provided the obligations are met using European Commission-approved industry best practices as formalized in “harmonized standards”.

- ▶ The European Commission has issued a “standardization request” to the European standards bodies (CEN and CENELEC), listing a series of topics for which new harmonized standards are required to cover the compliance obligations in the AI Act (see section on pre-market obligations of high-risk AI systems above).
- ▶ The European standardization bodies aim to have standards available in time for implementation of the AI Act in accordance with the agreed timelines (see above), but their readiness is not guaranteed.
- ▶ Where possible the European standardization bodies will seek to adopt standards created by the international standards bodies (ISO and IEC), with minimal modification.

## Codes of Practice to guide and support compliance

Providers of high-impact GPAI models posing a systemic risk may rely on codes of practice (see above) to demonstrate compliance until a harmonized standard is published.

The EU’s new AI Office (see below) shall facilitate the drawing up of codes of practice at Union level to guide the effective implementation of the obligations regarding the detection and labelling of artificially generated or manipulated content. Codes of practice should be prepared within nine months after the entry into force of the AI Act, to allow providers sufficient time to demonstrate compliance. The Commission is empowered to adopt implementing acts to approve these codes of practice.

The AI Office is expected to continue monitoring these codes to ensure that they are updated to sufficiently meet technological developments.

## How does the AI Act aim to support AI innovation in the EU?

### AI regulatory sandboxes

The AI Act mandates the establishment of AI regulatory sandboxes to offer innovation support across the EU.

- ▶ These regulatory sandboxes are controlled environments in which providers and deployers (e.g., small and medium enterprises) can voluntarily experiment, test, train, and validate their systems under regulatory supervision before placing them on the market.
- ▶ Each Member State will be expected to create a sandbox with common rules for consistent use across the EU to be operational within 24 months of the AI Act's entry into force (i.e., 2 August 2026).
- ▶ AI system providers will be able to receive a written report about their sandbox activities as evidence that they have met AI Act requirements. This is intended to speed up the approval process to take AI systems to market.

### Real-world testing

Testing of AI systems in real-world conditions outside of AI regulatory sandboxes may be conducted by providers or prospective providers of the high-risk AI systems listed in Annex III of the AI Act (see above), at any time before being placed on the market, if the following conditions are met:

- ▶ A testing plan has been submitted to, and approved by the market surveillance authorities
- ▶ The provider is established in the EU
- ▶ Data protection rules are observed
- ▶ Testing does not last longer than necessary and no more than six months (with the option to extend by an additional six months)
- ▶ End users have been informed, given their consent and have been provided with relevant instructions
- ▶ The predictions, recommendations and decisions of the AI system can be effectively reversed or disregarded

## What will the regulatory oversight model for the AI Act look like?

National competent authorities will be given oversight powers in Member States. These are likely to take different forms depending on the Member State.

At an EU level, the AI Act governance framework also establishes the:

- ▶ **AI Office** within the EU Commission, but with functional independence
  - ▶ This new body will have oversight responsibilities for GPAI models. It will contribute to the development of standards and testing practices, coordinate with the national competent authorities and help enforce the rules in Member States
- ▶ **AI Board** representing the Member States to provide strategic oversight for the AI Office
  - ▶ The Board will support the implementation of the AI Act and regulations promulgated pursuant to it, including the design of codes of practice for GPAI models
- ▶ **Scientific panel of independent experts** to support the activities of the AI Office
  - ▶ The panel will contribute to the development of methodologies for evaluating the capabilities of GPAI models and their subsequent classification, while also monitoring possible safety risks
- ▶ **Advisory forum** with representatives of industry and civil society
  - ▶ Will provide technical expertise to the AI Board

## What are the penalties for noncompliance?

The AI Act sets out a strict enforcement regime for noncompliance.

There are three notional levels of noncompliance, each with significant financial penalties. Depending on the level of violation (in line with the risk-based approach), the Act applies the following penalties:

Noncompliance case	Proposed fine
Breach of AI Act prohibitions	Fines up to €35 million or 7% of total worldwide annual turnover (revenue), whichever is higher
Noncompliance with the obligations set out for providers of high-risk AI systems or GPAI models, authorized representatives, importers, distributors, users or notified bodies	Fines up to €15 million or 3% of total worldwide annual turnover (revenue), whichever is higher
Supply of incorrect or misleading information to the notified bodies or national competent authorities in reply to a request	Fines up to €7.5 million or 1% of total worldwide annual turnover (revenue), whichever is higher

In the case of small and medium enterprises, fines will be as described above, but whichever amount is lower.

National competent authorities will determine the fines in line with the guidance provided above.

## What are the next steps around and beyond the AI Act?

### International alignment

At an international level, the European Commission and other EU institutions will continue to work with multi-national organizations including the OECD, the G7, The G20, the Council of Europe, the U.S.- EU Trade and Technology Council (TTC), and the UN to promote the development and adoption of rules beyond the EU that are compatible with the requirements of the AI Act.

### The EU AI Pact

A voluntary initiative led by the Commission to support the implementation of the AI Act, and referred to as the [AI Pact](#), will provide a forum for organizations to learn more about the AI Act and how to prepare for compliance. Participants may share and exchange good practices through engagement with the Commission and one another during the implementation period.

Furthermore, interested organizations also have the opportunity to make voluntary public 'pledges' reflecting steps they are taking to prepare for compliance with aspects of the Act.

### Delegated and Implementing Acts

The Commission is empowered to draft a number of delegated and implementing acts to clarify, complement or update the AI Act as appropriate. Some of these include:

- ▶ Adding conditions whereby a high-risk AI system listed in Annex III does not pose risk to health, safety or fundamental rights.
- ▶ Adding or removing high-risk use-cases from Annex III, within one of the existing categories.
- ▶ Amending thresholds for the classification of GPAI models posing a systemic risk.

## Appendix

AI Act term	AI Act definition
Provider	A natural or legal person, public authority, agency, or other body that is or has <b>developed</b> an AI system to place on the market, or to put into service under its own name or trademark whether for payment, or free of charge.
Deployer	A natural or legal person, public authority, agency, or other body <b>using</b> an AI system under its authority.
Authorized representative	Any natural or legal person located or established in the EU who has received and accepted a written mandate from a provider to <b>carry out its obligations on its behalf</b> .
Importer	Any natural or legal person located or established in the EU that places on the market an AI system that bears the name or trademark of a natural or legal person established outside the EU.
Distributor	Any natural or legal person in the supply chain, <b>not being the provider or importer, who makes an AI system available in the EU market</b> .
Product manufacturer	A manufacturer of an AI system that is put on the market or a manufacturer that puts into service an AI system <b>together with its product</b> and under its own name or trademark.
Operator	A general term referring to all the terms above (provider, deployer, authorized representative, importer, distributor, or product manufacturer).