

# Data Protection Impact Assessment

## **Purpose of policy**

Lowara Distribution Ireland is fully committed to protecting the personal data of its customers, employees, suppliers and other stakeholders in accordance with the requirements of the European Union General Data Protection Regulation 2016. We take the privacy of personal data very seriously and have initiated a variety of methods and controls to ensure we know what data we collect and hold and that we protect that data appropriately.

As part of this commitment, Lowara Distribution Ireland ensures that all business activities and projects that involve the use of personal data are subject to a data protection impact assessment. The purpose of this assessment is to ensure that our use of personal data is

Idireland.ie



IFS Template Reference: 1553318 Revision: 1

Name of Policy	Data Protection Impact Assessment Procedure			
Lowara Site	50 Broomhill Close, Tallaght, Dublin 24, D24 APP8, Ireland			
Applicable Policies	Data Protection Policy			
	<u>Breach Procedure</u>			
	Data Subject Request Procedure			
Last Updated	23/06/2025			

# Lowara Distribution Ireland Impact Assessment

## Procedure

Tricel (Baldonnell) Ltd, trading as Lowara Distribution Ireland (hereinafter referred to as "Lowara Distribution Ireland" or "the Company"), is a company registered in Ireland under registration number IE497442, with its registered office at 50 Broomhill Close, Tallaght, Dublin 24, D24 APP8, Ireland.

This document sets out our process for carrying out a data protection impact assessment and, in conjunction with the associated forms and guidance, should be used to ensure that our obligations and policies in this area are met.

# 1. SCOPE

This policy outlines the procedures and guidelines for conducting data protection impact assessments (DPIAs) at Lowara Distribution Ireland. The scope includes:

- **Commitment to GDPR Compliance**: Ensures Lowara Distribution Ireland's commitment to protecting personal data as per EU GDPR requirements.
- **Assessment Criteria**: Specifies conditions under which a DPIA is required, such as systematic evaluations, large-scale data processing, or new technology use.
- **Risk Management Process**: Details steps to identify, analyse, and evaluate risks to personal data, and establish risk treatment plans.



- **Documentation Requirements**: Provides guidelines for documenting the use of personal data, including how data is obtained, processed, stored, and accessed.
- **Management Approval**: Describes the need for obtaining management approval for residual risks and the process for prior consultation with supervisory authorities if high risks are identified.
- Implementation and Monitoring: Covers the implementation of risk treatment actions, monitoring key performance indicators, and conducting regular reviews.
- **Roles and Responsibilities**: Defines roles and responsibilities using a RACI chart to ensure effective risk assessment and management.

This comprehensive approach ensures that all personal data processing activities are thoroughly assessed and that appropriate measures are taken to protect data throughout its lifecycle.

# 2. OUT OF SCOPE

- General Data Protection Compliance: This document focuses specifically on the data protection impact assessment (DPIA) process and does not cover the broader aspects of GDPR compliance.
- Non-Personal Data: Procedures related to data that does not qualify as personal data under GDPR are not addressed in this document.
- Detailed IT Security Measures: While the document touches on risk assessment related to personal data, it does not provide detailed IT security protocols or technical measures for safeguarding data.
- Operational Data Processing: Regular data processing activities and operational procedures that do not require a DPIA are excluded.
- Incident Response: The document does not cover the procedures for responding to data breaches or incidents, which are likely covered by separate incident response policies.



• Legal and Regulatory Advice: The document does not provide specific legal or regulatory advice beyond what is necessary for the DPIA process.

## 3. ROLES & RESPONSIBILITIES

Within the process of risk assessment there are several key roles that play a part in ensuring that all risks are identified, addressed and managed. These roles are shown in the RACI table below, together with their relative responsibilities at each stage of the process.

#### RACI Chart

The table below clarifies the responsibilities at each step using the RACI model, i.e. R= Responsible A= Accountable C= Consulted I= Informed

ROLE:	ASSESSMENT	RISK	ТОР
STEP	LEAD		MANAGEMENT
		OWNERS	
ESTABLISH THE NEED AND	R	С	Α
CONTEXT			
DOCUMENT THE USE OF PERSONAL	R	С	А
DATA			
IDENTIFY THE RISKS	С	R	Α
ANALYSE THE RISKS	С	R	А
EVALUATE THE RISKS	С	R	А
DEFINE RISK TREATMENT PLAN	R	С	А
MANAGEMENT APPROVAL FOR	С	С	A/R
RESIDUAL RISKS			
IMPLEMENT RISK TREATMENT PLAN	R	R	A/R
MONITOR AND REPORT	R	I	А
REGULAR REVIEW	R	С	А

TABLE 3 - RACI CHART



Further roles and responsibilities may be added to the above table as the data protection impact assessment process matures within Lowara Distribution Ireland .

# 4. POLICY

#### **Data Protection Impact Assessment Process**

#### **Process Diagram**



#### FIGURE 1 - DATA PROTECTION IMPACT ASSESSMENT PROCESS DIAGRAM

## 4.1 Establish The Need and Context

There are a number of criteria that determine when a data protection impact assessment should be carried out within Lowara Distribution Ireland . The General Data Protection Regulation (Article 35) specifies that an impact assessment *shall be required* where the proposed processing involves:

> IFS 1554054 Rev 1 Page **4** of **17**



- a systematic and extensive evaluation of personal aspects relating to natural persons which is based on automated processing, including profiling, and on which decisions are based that produce legal effects concerning the natural person or similarly significantly affect the natural person;
- processing on a large scale of special categories of data referred to in Article 9(1), or of personal data relating to criminal convictions and offences referred to in Article 10; or
- a systematic monitoring of a publicly accessible area on a large scale

Note: Article 9(1) refers to processing of personal data revealing racial or ethnic origin, political opinions, religious or philosophical beliefs, or trade union membership, and the processing of genetic data, biometric data for the purpose of uniquely identifying a natural person, data concerning health or data concerning a natural person's sex life or sexual orientation.

In general, Lowara Distribution Ireland specifies that data protection impact assessments are appropriate for projects where one or more of the following applies:

- a) information about living individuals will be collected and processed for the first time
- b) information about living individuals will be shared with people or organizations that previously did not have access to it.
- c) change of use of existing personal data.
- d) the use of new technology that collects or uses data of a personal nature e.g. biometrics.
- e) existing personal data will be used to reach decisions as part of an automated process.
- f) it might reasonably be expected that an individual may find any aspect of the project intrusive or the data involved private.

If there is uncertainty regarding whether it is appropriate to carry out a data protection impact assessment for a specific project, by default the project team



should stay on the side of caution and ensure that one is performed. The Data Protection Officer may be consulted for clarification and further guidance may have been issued by the supervisory authority representing the EU within the country or countries in which the processing will be carried out, in which case this should be consulted also.

The overall environment in which the data protection impact assessment is carried out should be described and the reasons for it explained. This should include a description of the internal and external context of the project and its overall objectives.

The scope of the assessment should also be clearly defined. This may be expressed in terms of the defined scope of the project itself and may include factors such as:

- geographical location e.g. countries, offices, data centres
- organizational units e.g. specific departments
- business process(es)
- IT services, systems and networks
- Customers, products or services

Any specific exclusions to the scope should be stated with reasons.

## 4.2 Document The Use Of Personal Data

An appropriate level of detail should be gathered and documented regarding the personal data that is relevant to the project, including:

- Definitions of the specific data items to be stored and processed
- How the data will be obtained
- How the data will be processed
- Retention timescales of the data
- How the data will be stored
- Possible future uses of the data
- Where the data may be transferred to and under what circumstances

IFS 1554054 Rev 1 Page **6** of **17** 



• Who will have access to the data and how

This information may be collected and represented in an appropriate combination of information asset registers, flowcharts and via the use of relevant data mapping tools, if available.

## 4.3 Identify The Risks

The process of identifying risks to the personal data we collect, process and hold will consist of the following steps.

## 4.3.1 Identify Risk Scenarios

The identification of risks to the identified personal data will be performed by a combination of group discussion and interview with interested parties.

Such interested parties will normally include (where possible):

- Manager(s) responsible for each business-critical activity.
- Representatives of the people that normally carry out each aspect of the activity.
- Providers of the inputs to the activity.
- Recipients of the outputs of the activity.
- Appropriate third parties with relevant knowledge.
- Representatives of those providing supporting services and resources to the activity.
- Any other party that is felt to provide useful input to the risk identification process.

Identified risks will be recorded with as full a description as possible that allows the likelihood and impact of the risk to be assessed. Each risk should also be allocated an owner.



### 4.4 Analyse The Risks

Risk analysis within this process involves assigning a numerical value to the a) likelihood and b) impact of a risk. These values are then multiplied to arrive at a classification level of high, medium or low for the risk.

## 4.4.1 Assess The Likelihood -

An estimate of the likelihood of a risk occurring must be made. This should take into account whether it has happened before either to this organization or similar organizations in the same industry or location and whether there exists sufficient motive, opportunity and capability for a threat to be realized.

The likelihood of each risk should be graded on a numerical scale of 1 (low) to 5 (high). General guidance for the meaning of each grade is given in table 1. When assessing the likelihood of a risk, existing controls should be taken into account. This may require an assessment to be made as to the effectiveness of existing controls.

More detailed guidance may be decided for each grade of likelihood, depending on the subject of the risk assessment.

CLINANAADV

GRADE	DESCRIPTION	SUMMARY		
1	Improbable	Has never happened before and there is no reason to think it		
		is any more likely now		
2	Unlikely	There is a possibility that it could happen, but it probably		
		won't		
3	Likely	On balance, the risk is more likely to happen than not		
4	Very Likely	It would be a surprise if the risk did not occur either based on		
		past frequency or current circumstances		
5	Almost	Either already happens regularly or there is some reason to		
	certain	believe it is virtually imminent		





The rationale for allocating the grade given should be recorded to aid understanding and allow repeatability in future assessments.

## 4.4.2 Assess The Impact

An estimate of the impact that the risk could have on the organization should be given. This should take into account existing controls that lessen the impact, as long as these controls are seen to be effective.

Consideration should be given to the impact in the following areas:

- Customers
- Finance
- Health and Safety
- Reputation
- Knock-on impact within the organization
- Legal, contractual or organizational obligations

The impact of each risk should be graded on a numerical scale of 1 (low) to 5 (high). General guidance for the meaning of each grade is given in table 2.

More detailed guidance may be defined for each grade of impact, depending on the subject of the risk assessment.

The rationale for allocating the grade given should be recorded to aid understanding and allow repeatability in future assessments.

GRADE	DESCRIPTION	CUSTOMER	FINANCIAL	HEALTH AND SAFETY	IMPACT ON REPUTATION	LEGAL IMPACT
		IMPACT	ІМРАСТ			
1	Negligible	No effect	Very little or	Very small	Negligible	No
			none	additional		implication
				risk		S
2	Slight	Some local	Some	Within	Slight	Small risk
		disturbanc		acceptabl		of not
		e to normal		e limits		meeting

IFS 1554054 Rev 1 Page **9** of **17** 



		business operations				complianc e
3	Moderate	Can still deliver product/se rvice with some difficulty	Unwelcom e but could be borne	Elevated risk requiring immediat e attention	Moderate	In definite danger of operating illegally
4	High	Business is crippled in key areas	Severe on effect on income and/or profit	Significant danger to life	High	Operating illegally in some areas
5	Very High	Out of business; no service to customers	Crippling; the organisatio n will go out of business	Real or strong potential loss of life	Very High	Severe fines and possible imprisonm ent of staff

#### TABLE 2 – RISK IMPACT GUIDANCE

## 4.4.3 Risk Classification

Based on the assessment of the grade of likelihood and impact, a score is calculated for each risk by multiplying the two numbers. This resulting score is then used to decide the classification of the risk based on the matrix shown in figure 2.

Each risk will be allocated a classification based on its score as follows:

- HIGH 12 or more
- MEDIUM 5 to 10 inclusive
- LOW 1 to 4 inclusive





FIGURE 2 - RISK MATRIX CHART

The classification of each risk will be recorded as input to the risk evaluation stage of the process.

## 4.5 Evaluate The Risks

The purpose of risk evaluation is to decide which risks can be accepted and which ones need to be treated. This should take into account the risk acceptance criteria established for this specific risk assessment (*see Risk Acceptance Criteria*, above).

The matrix in *Figure 2* shows the classifications of risk, where green indicates that the risk is below the acceptable threshold. The orange and red areas generally indicate that a risk does not meet the acceptance criteria and so is a candidate for treatment.

Risks will be prioritized for treatment according to their score and classification so that very high scoring risks are recommended to be addressed before those with lower levels of exposure for the organization.

> IFS 1554054 Rev 1 Page **11** of **17**



## 4.6 Define Risk Treatment Plan

For those risks that are agreed to be above the threshold for acceptance by Lowara Distribution Ireland , the options for treatment will then be explored.

The overall intention of risk treatment is to reduce the classification of a risk to an acceptable level. This is not always possible as sometimes although the score is reduced, it remains in the same classification e.g. reducing the score from 8 to 6 means it still remains a medium level risk. The organization may decide to accept these risks even though they remain at a medium rating. Such decisions should be recorded with a suitable explanation.

## 4.6.1 Risk Treatment Options -

The following options may be applied to the treatment of the risks that have been agreed to be unacceptable:

- 1. *Modify* the risk apply appropriate controls to lessen the likelihood and/or impact of the risk
- 2. Avoid the risk by taking action that means it no longer applies
- 3. Share the risk with another party e.g. insurer or supplier

Judgement will be used in the decision as to which course of action to follow, based on a sound knowledge of the circumstances surrounding the risk e.g.

- Business strategy
- Regulatory and legislative considerations
- Technical issues
- Commercial and contractual issues

The Risk Manager will ensure that all parties who have an interest or bearing on the treatment of the risk are consulted, including the risk owner.



## 4.6.2 Selection Of Controls

Appropriate controls will then be identified to reduce either the likelihood or impact (or both) of each risk in order to bring it within acceptable bounds.

In accordance with Lowara Distribution Ireland 's adoption of the ISO/IEC 27001 standard, Annex A of that document will be used as the starting point for the identification of appropriate controls to address the risk treatment requirements identified as part of the risk assessment exercise. The controls set out in Annex A will be supplemented by the extended and additional guidance set out in the following codes of practice:

- ISO/IEC 27002 Code of practice for information security controls
- ISO/IEC 27017 Code of practice for information security controls based on ISO/IEC 27002 for cloud services
- ISO/IEC 27018 Code of practice for protection of personally identifiable information (PII) in public clouds acting as PII processors

The last two of these provide specific application of the Annex A controls to a cloud service provider scenario and address the area of the protection of PII more comprehensively than the ISO/IEC 27001 standard on its own.

## 4.6.3 Data Protection Impact Assessment Report

The evaluation of the treatment options will result in the production of the data protection impact assessment report which will detail:

- A description of the proposed processing operations and the personal data involved
- The purposes of the processing including, where applicable the legitimate interest of the controller of the personal data as defined by the GDPR
- An assessment of the necessity and proportionality of the processing



- The results of the assessment of the risks to the rights and freedoms of the data subjects
- Whether each risk is recommended for acceptance or treatment
- Priority of risks for treatment
- Risk owners
- Recommended treatment option
- Control(s) to be implemented
- Responsibility for the identified actions
- Timescales for actions
- Residual risk levels after the controls have been implemented

## 4.7 Obtain Management Approval For Residual Risks

At each stage of the data protection impact assessment process, management will be kept informed of progress and decisions made, including formal signoff of the proposed residual risks. Management will approve the data protection impact assessment report and will consider to what extent the report should be made public, either in full or in summarized form.

Signoff will be indicated according to Lowara Distribution Ireland 's documentation standards.

In addition to overall management approval, the acceptance or treatment of each risk should be signed off by the relevant risk owner.

Prior Consultation With Supervisory Authority

In the event that the results of the data protection impact assessment indicate a high level of risk prior to the identified controls being implemented, the GDPR requires that the supervisory authority is consulted before any processing takes place.



- The following information must be provided:
  - Details of the respective responsibilities of the controller, joint controllers and processors, where applicable
  - Purposes and means of the processing
  - The controls that will be implemented to protect the data
  - Contact details for the data protection officer (if applicable)
  - A copy of the impact assessment report

The supervisory authority has eight weeks (extendable by a further six weeks) to provide a judgement on the proposed processing and, if appropriate, give details of what must be done to make the processing acceptable under the GDPR.

## 4.8 Implement Risk Treatment Actions

Once the risk treatment plan has been approved, the necessary actions should be tracked and completed as part of the day to day control of the project. In the event that any actions are delayed or cannot be completed, the implications of this to the protection of the personal data involved must be assessed by management and a decision taken about what to do next. If the untreated risk is sufficiently serious, this may have a significant impact on the viability of the project from a compliance viewpoint and advice should be sought from the Data Protection Officer and/or the supervisory authority in the country or countries affected.

## 4.9 Risk Monitoring And Reporting

As part of the implementation of new controls and the maintenance of existing ones, key performance indicators will be identified which will allow the measurement of the success of the controls in addressing the relevant risks.

These indicators will be reported on a regular basis and trend information produced so that exception situations can be identified and dealt with as part of the management review process.

> IFS 1554054 Rev 1 Page **15** of **17**



#### 4.10 Regular Review

In addition to a full annual review, risk assessments will be evaluated on a regular basis to ensure that they remain current and the applied controls valid. The relevant risk assessments will also be reviewed upon major changes to the business such as office moves, mergers and acquisitions or introduction or new or changed IT services.

#### MONITORING AND REVIEW

#### 4.10.1 Risk Monitoring and Reporting:

- Implementation of new controls and maintenance of existing ones involves identifying key performance indicators (KPIs).
- These KPIs help measure the effectiveness of controls in mitigating identified risks.
- Regular reporting on these KPIs and trend analysis helps in identifying exceptions and addressing them through management review.

#### 4.10.2 Regular Review:

- Risk assessments undergo regular evaluations to ensure their relevance and the validity of applied controls.
- Major business changes such as office relocations, mergers, acquisitions, or significant IT service changes trigger additional reviews.
- An annual comprehensive review is also conducted to maintain up-to-date risk assessments.

#### 4.10.3 Roles and Responsibilities:

- The DPIA process assigns specific roles and responsibilities using a RACI chart, clarifying who is Responsible, Accountable, Consulted, and Informed at each step.
- Regular monitoring and review tasks are primarily assigned to those responsible (R) for overseeing the process and ensuring compliance with established controls and measures.



# 5. SUPPORTING DOCUMENTATION

- Data Protection Policy
- Breach Procedure
- Data Subject Request Procedure

# 6. ACKNOWLEDGMENT AND COMPLIANCE

The process of data protection impact assessment is fundamental to the implementation of a successful project that handles personal data and is a significant part of the GDPR legislation. Only by fully understanding its risks with regard to personal data can an organization hope to ensure that the controls it has in place are sufficient to provide an appropriate level of privacy and meet the high standard expected of it.

For Lowara Distribution Ireland , the regular assessment of risks to personal data and the application of comprehensive controls is vital to the continuing confidence of its customers and in meeting its obligations to protect personal data from all too common threats.

By following this process Lowara Distribution Ireland will go some way to ensuring that the risks that it faces in the day to day operation of its business are effectively managed and controlled.

IFS 1554054 Rev 1 Page **17** of **17**