

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Almirall, S.A. works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organizations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Almirall, S.A. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Almirall, S.A. certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Almirall, S.A. certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Almirall, S.A. certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Almirall, S.A. certifies that its disclosure complies with the Data Privacy obligations.

Date: 25 June 2025

Name of signatory: Stacy Lockwood

Position in the Company: Global Head Compliance

Signature:

A handwritten signature in black ink, appearing to read 'Stacy Lockwood', written in a cursive style.



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom **Amgen** works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, **Amgen** hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Amgen certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Amgen certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Amgen certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Amgen certifies that its disclosure complies with the Data Privacy obligations.

Date: 4/14/2025

Name of signatory: Gilles Marrache

Position in the Company: SVP General Manager

Signature:

Signed by:
Gilles Marrache
 Signer Name: Gilles Marrache
Signing Reason: I approve this document
Signing Time: 4/14/2025 | 5:43:41 PM GMT
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EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Astellas Pharma Europe Ltd works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Astellas Pharma Europe Ltd hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2025 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Astellas Pharma Europe Ltd certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Astellas Pharma Europe Ltd certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Astellas Pharma Europe Ltd certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

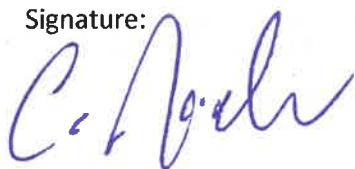
Astellas Pharma Europe Ltd certifies that its disclosure complies with the Data Privacy obligations.

Date: 5 JUNE 2025

Name of signatory: CLAUD ZIELER

Position in the Company: CHIEF COMMERCIAL & MEDICAL AFFAIRS OFFICER

Signature:



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom AstraZeneca works provide the pharmaceutical industry with valuable, independent, and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, AstraZeneca hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

AstraZeneca certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

AstraZeneca certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

AstraZeneca certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

AstraZeneca certifies that its disclosure complies with the Data Privacy obligations

Date:

Name of signatory: **Stefan Woxström**

Position in the Company: **SVP Europe & Canada**

Signature: 
Stefan Woxström (26-Jun-2025 12:36 GMT+2)



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Bayer AG works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Bayer AG hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Bayer AG certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Bayer AG certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



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Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Bayer AG certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Bayer AG certifies that its disclosure complies with the Data Privacy obligations.

Bayer Aktiengesellschaft

Berlin, 23/06/2025

A handwritten signature in blue ink, appearing to read "Stefan Oelrich", written over a horizontal line.

Stefan Oelrich
Member of the Board of Management
President Pharmaceuticals Division

Berlin, 23/06/2025

A handwritten signature in blue ink, appearing to read "ppa. Königer", written over a horizontal line.

Dr. Ursula Königer
Law, Patents and Compliance
Business Partner Pharmaceuticals



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Boehringer Ingelheim works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Boehringer Ingelheim hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Boehringer Ingelheim certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Boehringer Ingelheim certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Boehringer Ingelheim certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

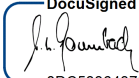
Boehringer Ingelheim certifies that its disclosure complies with the Data Privacy obligations.

Date: 26-Jun-2025

Name of signatory: Hubertus von Baumbach

Position in the Company: Chairman of the Board of Managing Directors

Signature:

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EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom **BIAL - Portela & C^a, S.A.** works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, **BIAL - Portela & C^a, S.A** hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles

Disclosure quality

BIAL - Portela & C^a, S.A certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

BIAL - Portela & C^a, S.A certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

BIAL - Portela & C^a. S.A certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

BIAL - Portela & C^a, S.A certifies that its disclosure complies with the Data Privacy obligations.

Date: 2025.05.13

Name of signatory: António Portela

Position in the Company: Chief Executive Officer

Signature:





EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom **Biogen International GmbH (Biogen)** works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, **Biogen** hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Biogen certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Biogen certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Biogen certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Biogen certifies that its disclosure complies with the Data Privacy obligations.

Date: 09-May-2025

Name of signatory: Dr. Wolfram Werner Schmidt

Position in the Company: President Region Europe

Signature:

DocuSigned by:
Wolfram Schmidt
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EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Bristol-Myers Squibb works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Bristol-Myers Squibb hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Bristol-Myers Squibb certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Bristol-Myers Squibb certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Bristol-Myers Squibb certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Bristol-Myers Squibb certifies that its disclosure complies with the Data Privacy obligations.

Date: 15th May 2025

Name of signatory: Emma Charles

Position in the Company: Senior Vice President Europe Region

Signature:



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Chiesi Farmaceutici S.p.A. works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Chiesi Farmaceutici S.p.A. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Chiesi Farmaceutici S.p.A. certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Chiesi Farmaceutici S.p.A. certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Chiesi Farmaceutici S.p.A. certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

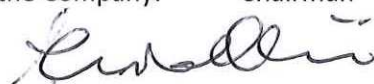
Chiesi Farmaceutici S.p.A. certifies that its disclosure complies with the Data Privacy obligations.

Date: June 11th, 2025

Name of signatory: Alessandro CHIESI

Position in the Company: Chairman

Signature:



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom CSL Behring works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients. In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, CSL Behring hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

CSL Behring certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

CSL Behring certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

CSL Behring certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

CSL Behring certifies that its disclosure complies with the Data Privacy obligations.

Date: 24.06.2025

Name of signatory: Debbie Drane

Position in the Company: SVP Global Strategy & Business Development

Signature: 8E07A715630B425...

Passion for Innovation.
Compassion for Patients.™



Daiichi Sankyo Europe GmbH
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www.daiichi-sankyo.eu

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Daiichi Sankyo Europe GmbH works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients. In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Daiichi Sankyo Europe GmbH hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Daiichi Sankyo Europe GmbH certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Passion for Innovation.
Compassion for Patients.™



Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Daiichi Sankyo Europe GmbH certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Daiichi Sankyo Europe GmbH certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Daiichi Sankyo Europe GmbH certifies that its disclosure complies with the Data Privacy obligations.

Date:	June 19, 2025	June 23, 2025
Name of signatory:	Curd Lejaegere	Charlotte Chui-Franz
Position in the Company:	VP Mid-Size Countries Efpi Representative Daiichi Sankyo Europe	Chief Compliance Officer Efpi Representative Daiichi Sankyo Europe

Signature:

DocuSigned by:
Curd Lejaegere
15BA0ABE3AC1406...

DocuSigned by:
Charlotte Chui-Franz
829C5C10A95648A...



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Eisai Europe Ltd works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Eisai Europe Ltd hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Eisai Europe Ltd certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Eisai Europe Ltd certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Eisai Europe Ltd certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

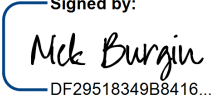
Ensuring compliance with Data Privacy Obligations

Eisai Europe Ltd certifies that its disclosure complies with the Data Privacy obligations.

Date: 27-Jun-2025 | 12:34:59 BST

Name of signatory: Nick Burgin

Position in the Company: President & COO EMEA & President GV&A

Signature:  Signed by:
DF29518349B8416...

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Gilead Sciences works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Gilead Sciences hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2023 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Gilead Sciences certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Gilead Sciences certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Gilead Sciences certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Gilead Sciences certifies that its disclosure complies with the Data Privacy obligations.

Date: May 26, 2025 | 7:08:45 AM PDT

Name of signatory: Josephine Comiskey

Position in the Company: Senior Vice President, ACE Region

Signature:

DocuSigned by:

2B2D2233141541D...



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom GSK works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, GSK hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

GSK certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

GSK certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

GSK certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

GSK certifies that its disclosure complies with the Data Privacy obligations.

Date: 16.05.25

Name of signatory: Lynn Baxter

Position in the Company: SVP, Head of Europe Commercial

Signature:

A handwritten signature in black ink, appearing to read 'Lynn Baxter', is written over a light grey circular stamp.



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Ipsen works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Ipsen hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Ipsen certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Ipsen certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Ipsen certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Ipsen certifies that its disclosure complies with the Data Privacy obligations.

Date: 20 June 2025

Name of signatory: David Loew

Position in the Company: Chief Executive Officer

Signature:

A handwritten signature in blue ink, consisting of a stylized 'D' and 'L' followed by a horizontal line.

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Jazz Pharmaceuticals works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients. In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Jazz Pharmaceuticals hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Jazz Pharmaceuticals certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Jazz Pharmaceuticals certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Jazz Pharmaceuticals certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

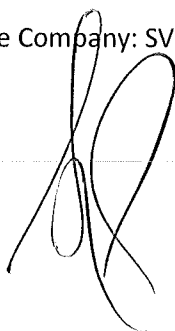
Jazz Pharmaceuticals certifies that its disclosure complies with the Data Privacy obligations.

Date: 29/7/25

Name of signatory: Amanda Flanagan

Position in the Company: SVP, Head of Europe & International

Signature:



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Janssen Pharmaceutica NV works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Janssen Pharmaceutica NV hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Janssen Pharmaceutica NV certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Janssen Pharmaceutica NV certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Janssen Pharmaceutica NV certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Janssen Pharmaceutica NV certifies that its disclosure complies with the Data Privacy obligations.

Date: May 27, 2025

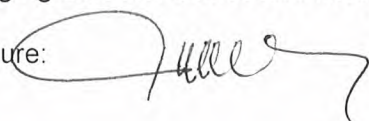
Name of signatory: Kris Sterkens

Position in the Company:

Company Group Chairman J&J Innovative Medicine EMEA

Managing Director Janssen Pharmaceutica N.V.

Signature:





EFPIA Code Disclosure 2025 Self-Certification Scheme

• **Dermatology
beyond the skin**

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom LEO Pharma works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organizations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

LEO Pharma A/S
Industriparken 55
2750 Ballerup
Denmark

Main +45 4494 5888

www.leo-pharma.com
CVR no.: 56 75 95 14

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, LEO Pharma hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2023 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

LEO Pharma certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

LEO Pharma certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



• Dermatology
beyond the skin

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

LEO Pharma certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

LEO Pharma certifies that its disclosure complies with the Data Privacy obligations.

Date:

Name of signatory: Christophe Bourdon

Position in the Company: CEO

Signature:



Eli Lilly and Company
Lilly Corporate Center
Indianapolis, Indiana 46285
U.S.A
+1 317 276 2000
www.lilly.com

EFPIA Disclosure Code 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Eli Lilly and Company (Lilly) works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted a code and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standard of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Lilly hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Lilly certifies that:

- Its disclosures are made in each country covered by the EFPIA code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Lilly certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs, HCOs and POs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Lilly certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO/PO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO/PO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Lilly certifies that its disclosure complies with the Data Privacy obligations.



Ilya Yuffa

Executive Vice-President and President of Lilly International

May 2025



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom H. Lundbeck A/S (“Lundbeck”) works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Lundbeck hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Lundbeck certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code’s requirements and applicable codes

Lundbeck certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs’ transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Lundbeck certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Lundbeck certifies that its disclosure complies with the Data Privacy obligations.

Date: 23 June 2025

Name of signatory: Charl van Zyl

Position in the Company: President and Chief Executive Officer (CEO)

Signature:

A handwritten signature in black ink, appearing to be "Charl van Zyl", written over a light blue circular stamp.



A. MENARINI
INDUSTRIE FARMACEUTICHE RIUNITE

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Menarini works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Menarini hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Menarini certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Menarini certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



A. MENARINI
INDUSTRIE FARMACEUTICHE RIUNITE

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Menarini certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Menarini certifies that its disclosure complies with the Data Privacy obligations.

Date: 26th May 2025

Name of signatory: Elcin Barker Ergun

Position in the Company: CEO and Member of the Board of Directors

Signature: _____



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Merck works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Merck hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Merck certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Merck certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Merck certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

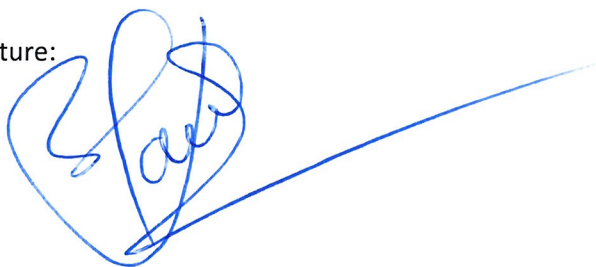
Merck certifies that its disclosure complies with the Data Privacy obligations.

Date: 22.05.25

Name of signatory: Belén Garijo

Position in the Company: Chair of the Executive Board and CEO of Merck

Signature:

A handwritten signature in blue ink, appearing to be 'Belén Garijo', with a long horizontal stroke extending to the right.

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom MSD works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, MSD hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

MSD certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

MSD certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

MSD certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

MSD certifies that its disclosure complies with the Data Privacy obligations.

Date: June 26, 2025

Name of signatory: Joseph Romanelli

Position in the Company: President, Human Health International, MSD

Signature:

A handwritten signature in black ink, appearing to read "J. Romanelli", with a stylized flourish at the end.



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom the Novartis Pharma AG works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, affiliates of Novartis Pharma AG below hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Novartis Pharma AG certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Novartis Pharma AG certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Novartis Pharma AG certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Novartis Pharma AG certifies that its disclosure complies with the Data Privacy obligations.

Date: 19 June 2025

Name of the Signatory: Patrick Horber

Position in the Company: Novartis President, International

Signature:

A handwritten signature in blue ink, appearing to be 'PH', written over a light blue horizontal line.



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Novo Nordisk A/S works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Novo Nordisk A/S hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Novo Nordisk A/S certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Novo Nordisk A/S certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Novo Nordisk A/S certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Novo Nordisk A/S certifies that its disclosure complies with the Data Privacy obligations.

Date: 23 June 2025

Name of signatory: Lars Fruergaard Jørgensen

Position in the Company: President & Chief Executive Officer

Signature:



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Otsuka Pharmaceutical Europe Ltd. works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Otsuka Pharmaceutical Europe Ltd. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Otsuka Pharmaceutical Europe Ltd. certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Otsuka Pharmaceutical Europe Ltd. certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Otsuka Pharmaceutical Europe Ltd. certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Otsuka Pharmaceutical Europe Ltd. certifies that its disclosure complies with the Data Privacy obligations.

Date: 09 May 2025

Name of signatory: Andy Hodge

Position in the Company: President & CEO, Europe

Signature:





EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Pfizer works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Pfizer hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Pfizer certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Pfizer certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Pfizer certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Pfizer certifies that its disclosure complies with the Data Privacy obligations.

Date: juin 20, 2025

Name of signatory: Alexandre de Germay

Position in the Company: Chief International Commercial Officer

Signature:

Signé par :

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EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Pierre Fabre Medical Care works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Pierre Fabre Medical Care hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Pierre Fabre Medical Care certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Pierre Fabre Medical Care certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Pierre Fabre Medical Care certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

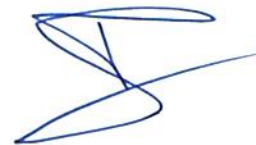
Pierre Fabre Medical Care certifies that its disclosure complies with the Data Privacy obligations.

Date: **21/05/2025**

Name of signatory: **Marie-Andrée GAMACHE**

Position in the Company: **Medical Care Business Unit Director**

Signature:





EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom **F. Hoffmann-La Roche AG** works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to build understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, **F. Hoffmann-La Roche AG** hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

F. Hoffmann-La Roche AG certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

F. Hoffmann-La Roche AG certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

F. Hoffmann-La Roche AG certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

F. Hoffmann-La Roche AG certifies that its disclosure complies with the Data Privacy obligations.

Padraic Ward
Head of Pharma International

Signed by:

Padraic Ward

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Date 28-May-2025 | 12:13 CEST

Teresa Graham
CEO Roche Pharmaceuticals

Signed by:

Teresa Graham

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Date 28-May-2025 | 01:07 CEST



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Laboratorios Farmaceuticos ROVI SA works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Laboratorios Farmaceuticos ROVI SA hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Laboratorios Farmaceuticos ROVI SA certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Laboratorios Farmaceuticos ROVI SA certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Laboratorios Farmaceuticos ROVI SA certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

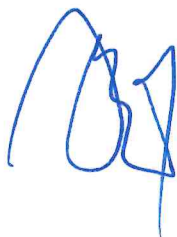
Laboratorios Farmaceuticos ROVI SA certifies that its disclosure complies with the Data Privacy obligations.

Date: 17/06/2025

Name of signatory: Juan López-Belmonte Encinas

Position in the Company: Chief Executive Officer

Signature:

A handwritten signature in blue ink, appearing to be 'JB', is written over a light blue horizontal line.

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Sanofi works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Sanofi hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Sanofi certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Sanofi certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Sanofi certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

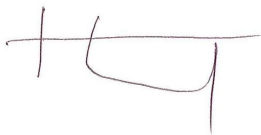
Sanofi certifies that its disclosure complies with the Data Privacy obligations.

Date: June 26, 2025

Name of signatory: OLIVIER CHARMEIL

Position in the Company: EVP, GENERAL MEDICINES

Signature:

A handwritten signature in blue ink, appearing to be 'O. Charmeil', written over a horizontal line.

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom SERVIER works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, SERVIER hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

SERVIER certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

SERVIER certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

4-



SERVIER certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

SERVIER certifies that its disclosure complies with the Data Privacy obligations.

Date: July 17th, 2025

Name of signatory: Mr. Olivier LAUREAU

Position in the Company: President

Signature:

EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom STALLERGENES GREER INTERNATIONAL AG works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, STALLERGENES GREER INTERNATIONAL AG hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

STALLERGENES GREER INTERNATIONAL AG certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

STALLERGENES GREER INTERNATIONAL AG certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

STALLERGENES GREER INTERNATIONAL AG certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

STALLERGENES GREER INTERNATIONAL AG certifies that its disclosure complies with the Data Privacy obligations.

Date: 22.09.2025

Name of signatory: Andreas AMREIN

Position in the Company: CEO

Signature: 



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Takeda Pharmaceuticals International AG works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients. In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Takeda Pharmaceuticals International AG hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Takeda Pharmaceuticals International AG certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Takeda Pharmaceuticals International AG certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Takeda Pharmaceuticals International AG certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

Takeda Pharmaceuticals International AG certifies that its disclosure complies with the Data Privacy obligations.

Date: 19th June 2025

Name of signatory: Jean-Luc Delay

Position in the Company: President Europe and Canada

Signature:

A handwritten signature in black ink, appearing to read 'Jean-Luc Delay', written over a light blue horizontal line.



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Teva Pharmaceuticals Europe B.V. works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, Teva Pharmaceuticals Europe B.V. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Teva Pharmaceuticals Europe B.V. certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

Teva Pharmaceuticals Europe B.V. certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).



Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

Teva Pharmaceuticals Europe B.V. certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

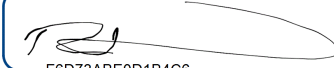
Teva Pharmaceuticals Europe B.V. certifies that its disclosure complies with the Data Privacy obligations.

Date: 13-Jun-2025 | 12:04 BST

Name of signatory: Richard Daniell

Position in the Company: Executive Vice President
Teva Pharmaceuticals Europe B.V.

Signature:

Signed by:

F6D73ABE0D1B4C6...



EFPIA Code Disclosure 2025 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom UCB works provide the pharmaceutical industry with valuable, independent and expert knowledge derived from their clinical and management experience. As owners of scientific knowledge and experts in medicinal products, pharmaceutical companies can be a unique resource to the healthcare systems and providers, which will ultimately benefit the patients.

In the same way, the pharmaceutical industry works with Patient Organisations (POs), this collaboration is essential in addressing patient needs. POs have a key role in helping to shape, develop and define the outcomes that make the most difference to patients.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs, HCOs and POs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs/POs aims to building understanding of the collaboration and recognition of its value to patient care.

Except in countries where disclosure is prescribed by laws, UCB hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2024 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

UCB certifies that:

- Its disclosures are made in each country covered by the EFPIA Code where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the Code and associated guidance issued by EFPIA;
- Its Methodological Note describes the process it has followed in order to compile the data hereby disclosed.

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Code's requirements and applicable codes

UCB certifies that:

- Data collection complies with the requirements of the EFPIA Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Code).

Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

UCB certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;
- If an HCP or HCO (where applicable) has provided consent to individual disclosure only in respect of part of the Transfers of Value he/she/it received, all Transfers of Value to such HCP or HCO (where applicable) are being disclosed in the aggregate.

Ensuring compliance with Data Privacy Obligations

UCB certifies that its disclosure complies with the Data Privacy obligations.

Date: 02-Jun-2025

Name of signatory:
Jean-Christophe Tellier

Position in the Company:
Head of UCB, CEO

Signature:  Signed by:
D8509A78164346B...

Certificate Of Completion

Envelope Id: E5835DBA-A4E1-4AFB-B136-CDFEAAC42B79	Status: Completed
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Contract:	
No	
Contract Title:	
Source Envelope:	
Document Pages: 2	Signatures: 1
Certificate Pages: 1	Initials: 0
AutoNav: Enabled	
Envelopeld Stamping: Enabled	
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Envelope Originator:	
Denise Michelle Jones	
Chemin du Foriest	
Braine l'Alleud, Belgium 1420	
DeniseMichelle.JONES@ucb.com	
IP Address: 71.204.91.96	

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Status: Original	Holder: Denise Michelle Jones	Location: DocuSign
6/2/2025 5:18:15 AM	DeniseMichelle.JONES@ucb.com	

Signer Events

Jean-Christophe Tellier
Jean-Christophe.Tellier@ucb.com
CEO

Security Level: Email, Account Authentication
(None)

Signature

Signed by:

D8509A78164346B...

Signature Adoption: Drawn on Device
Using IP Address: 194.187.114.162
Signed using mobile

Timestamp

Sent: 6/2/2025 5:24:57 AM
Viewed: 6/2/2025 7:49:16 AM
Signed: 6/2/2025 7:49:52 AM

Electronic Record and Signature Disclosure:
Not Offered via Docusign

In Person Signer Events

Signature

Timestamp

Editor Delivery Events

Status

Timestamp

Agent Delivery Events

Status

Timestamp

Intermediary Delivery Events

Status

Timestamp

Certified Delivery Events

Status

Timestamp

Carbon Copy Events

Status

Timestamp

Witness Events

Signature

Timestamp

Notary Events

Signature

Timestamp

Envelope Summary Events

Status

Timestamps

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Certified Delivered	Security Checked	6/2/2025 7:49:16 AM
Signing Complete	Security Checked	6/2/2025 7:49:52 AM
Completed	Security Checked	6/2/2025 7:49:52 AM

Payment Events

Status

Timestamps