

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom AbbVie 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, AbbVie 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

AbbVie 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

AbbVie 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

AbbVie 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

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

Date: 5/6/2024

Name of signatory:
Jason SMITH

Position in the Company:
Senior Vice-President, Europe

Signature:

EFPIA Code Disclosure 2024 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 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, Almirall S.A. 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

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: 24 June 2024

Name of signatory: Stacy Lockwood

Position in the Company: Global Head Compliance

Signature:





EFPIA Code Disclosure 2024 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 2023 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: 5/29/2024

Name of signatory: Gilles Marrache

Position in the Company: SVP General Manager

Signature:

DocuSigned by:
Gilles Marrache
 Signer Name: Gilles Marrache
Signing Reason: I approve this document
Signing Time: 5/29/2024 | 6:19:52 PM GMT
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EFPIA Code Disclosure 2024 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 2023 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: 06 juin 2024

Name of signatory: Damien Bailly

Position in the Company: President - Established Markets

Signature: 4F5270BA99FC459...

EFPIA Code Disclosure 2024 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 2023 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: 27/06/2024

Name of signatory: **Stefan Woxström**

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

Signature: 
Stefan Woxström (Jun 27, 2024 20:27 GMT+2)



EFPIA Code Disclosure 2024 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 2023 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, 21.6.24

A blue ink signature of Stefan Oelrich, written in a cursive style, positioned above a horizontal line.

Stefan Oelrich
Member of the Board of Management
President Pharmaceuticals Division

Berlin, June 21, 2024

A blue ink signature of Dr. Ursula Königer, written in a cursive style, positioned above a horizontal line.

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

EFPIA Code Disclosure 2024 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom **Bial-Portela & C^a, 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, **Bial-Portela & C^a, SA** 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

Bial-Portela & C^a, 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

Bial-Portela & C^a, 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.

Bial-Portela & C^a, 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

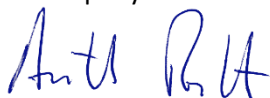
Bial-Portela & C^a, SA certifies that its disclosure complies with the Data Privacy obligations.

Date: 2024.06.20

Name of signatory: António Portela

Position in the Company: Chief Executive Officer

Signature:





EFPIA Code Disclosure 2024 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 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, **Biogen** hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2022 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

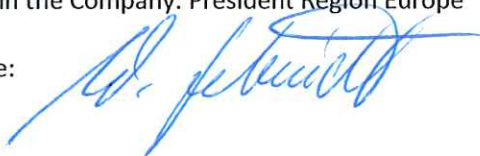
Biogen certifies that its disclosure complies with the Data Privacy obligations.

Date: 3rd June 2024

Name of signatory: Wolfram Werner Schmidt

Position in the Company: President Region Europe

Signature:



EFPIA Code Disclosure 2024 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 2023 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: 30 May 2024

Name of signatory: Monica Shaw

Position in the Company: SVP, Head of European Markets

Signature:

A handwritten signature in black ink, appearing to be 'MS' or similar, written in a cursive style.

EFPIA Code Disclosure 2024 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 2023 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.

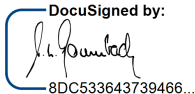
Ensuring compliance with Data Privacy Obligations

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

Date:

Name of signatory: Hubertus v. Baumbach

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

Signature: 8DC533643739466...

EFPIA Code Disclosure 2024 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 2023 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).



CHIESI	FARMACEUTICI	S.p.A.
Via	Palermo	26/A
43122,	Parma	(PR)
Tel.:	+39 0521	2791
Fax:	+39 0521	774468
Info@pec.chiesi.com		

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 21st, 2024

Name of signatory: Alessandro Chiesi

Position in the Company: President

Signature:

EFPIA Code Disclosure 2024 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Vifor 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, Vifor International AG 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

Vifor 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

Vifor 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.

Vifor 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

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

Date: 31 May 2024

Name of signatory: Hervé Gisserot

Position in the Company: General Manager, CSL Vifor

Signature:



EFPIA Code Disclosure 2024 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 2023 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.

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

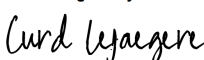
Daiichi Sankyo Europe GmbH certifies that its disclosure complies with the Data Privacy obligations.

Date:

Name of signatory: Curd Lejaegere ppa. Martin Fürle

Position in the Company: VP Mid-Size Countries GC & CCO Europe
Daiichi Sankyo Europe Efpia Representative

Signature:

DocuSigned by:

15BA0ABE3AC1406...

DocuSigned by:

AF72B0726EF9440...



EFPIA Code Disclosure 2024 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 2023 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-2024 | 17:29:50 BST

Name of signatory: Nick Burgin

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

Signature:  DF29518349B8416...

*Excludes Netherlands

EFPIA Code Disclosure 2024 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: 24th April 2024

Name of signatory: Josephine Comiskey

Position in the Company: Senior Vice President, ACE Region

Signature:





EFPIA Code Disclosure 2024 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 2023 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: 29 May 2024

Name of signatory: George Katzourakis

Position in the Company: SVP – Head of Europe



Signature:



EFPIA Code Disclosure 2024 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 2023 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

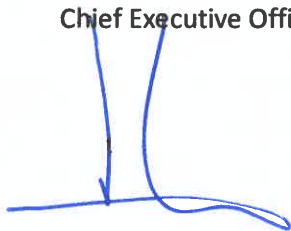
Ipsen certifies that its disclosure complies with the Data Privacy obligations.

Date: 19 June 2024

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 2024 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 2023 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: June 10, 2024

Name of signatory: Kris Sterkens

Position in the Company: Managing Director

Signature:



EFPIA Code Disclosure 2024 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 2024 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.

EFPIA and its members are fully committed to implement and apply the highest ethical standards.

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 2023 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 ToVs that cannot be disclosed on an individual basis for legal reasons

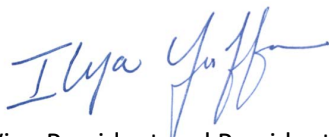
Lilly certifies that aggregate disclosure is limited to the following topics:

- 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 2024

EFPIA Code Disclosure 2024 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 2023 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:

Name of signatory: Charl van Zyl

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

Signature:





A. MENARINI

INDUSTRIE FARMACEUTICHE RIUNITE

EFPIA Code Disclosure 2024 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 2023 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 S.R.L. - SEDE: VIA SETTE SANTI, 3 - 50131 FIRENZE - TEL. +39 055 56801 - FAX +39 055 582771 - WWW.MENARINI.IT
C.P. 4083 - 50135 FIRENZE - CAP. SOC. € 150.000.000,00 I.V. - C. F., P. I.V.A. E REG. IMPRESE 06395270481 C.C.P. 15447501 - R.E.A. 7874 - R.E.C. ISCRIZ. N.87678 DEL 18.9.92

Aziende del Gruppo Menarini

Italia: MALESCI - Firenze, F.I.R.M.A. - Firenze, CODIFI - Firenze, A. MENARINI FARMACEUTICA INTERNAZIONALE - Firenze, A. MENARINI INDUSTRIE FARMACEUTICHE RIUNITE - Firenze, A. MENARINI MANUFACTURING LOGISTICS AND SERVICES - Firenze, L'Aquila e Pisa, MENARINI RICERCHE - Firenze e Pomezia, MENARINI BIOTECH - Pomezia, GUIDOTTI - Pisa, LUSOFARMACO - Milano, LUSOCHIMICA - Pisa e Lomagna (Lecco), RESEARCH TOXICOLOGY CENTRE - Pomezia.

London: ALBANIA - Tirana, ARGENTINA - Buenos Aires, ARMENIA - Yerevan, AUSTRALIA e NUOVA ZELANDA - Sydney, AUSTRIA - Vienna, AZERBAIGIAN - Baku, BELGIO - Bruxelles, BIELORUSSIA - Minsk, BOSNIA-ERZEGOVINA - Sarajevo, BULGARIA - Sofia, CINA - Pechino e Shanghai, COREA DEL SUD - Seul e Yongin, COSTARICA - San Jose', CROAZIA - Zagabria, DANIMARCA - Copenhagen, EL SALVADOR - San Salvador, ESTONIA - Tallinn, FILIPPINE - Manila, FINLANDIA - Helsinki, FRANCIA - Parigi, GEORGIA - Tbilisi, GERMANIA - Berlino e Dresda, GRECIA - Atene, GUATEMALA - Città del Guatemala, HONDURAS - Tegucigalpa, HONG KONG - Hong Kong, INDIA - Ahmedabad, Mumbai e Nuova Delhi, INDONESIA - Bekasi e Jakarta, IRLANDA - Dublino e Shannon, KAZAKISTAN - Almaty, KIRGHIZISTAN - Bishkek, LETTONIA - Riga, LITUANIA - Vilnius, LUSSEMBURGO - Lussemburgo, MALAYSIA - Kuala Lumpur, MESSICO - Città del Messico, MOLDAVIA - Chisinau, MONTENEGRO - Podgorica, NICARAGUA - Managua, OLANDA - Amsterdam, PANAMA - Panama, POLONIA - Varsavia, PORTOGALLO - Lisbona, REGNO UNITO - Londra, REPUBBLICA Ceca - Praga, ROMANIA - Bucarest, RUSSIA - Mosca, SERBIA - Belgrado, SINGAPORE - Singapore, SLOVACCHIA - Bratislava, SLOVENIA - Lubiana, SPAGNA - Barcellona, SUD AFRICA - Bryanston, SVIZZERA - Zurigo, TAILANDIA - Bangkok, TAIWAN - Taipei, TURCHIA - Istanbul, TURKMENISTAN - Ashgabat, UCRAINA - Kiev, UNGHERIA - Budapest, UZBEKISTAN - Tashkent, VIETNAM - Hanoi e Ho Chi Minh.

Diagnostica: AUSTRIA - Vienna, BELGIO - Zaventem, FRANCIA - Parigi, GERMANIA - Berlino, GRECIA - Atene, ITALIA - Firenze, OLANDA - Valkenswaard, PORTOGALLO - Lisbona, REGNO UNITO - Londra, SPAGNA - Barcellona, SVEZIA - Malmö, SVIZZERA - Zurigo.



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: 3rd June 2024

Name of signatory: Elcin Barker Ergun

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

Signature: _____

A. MENARINI INDUSTRIE FARMACEUTICHE RIUNITE S.R.L. - SEDE: VIA SETTE SANTI 3 - 50131 FIRENZE - TEL. +39 055 56801 - FAX +39 055 582771 - WWW.MENARINI.IT
C.P. 4083 - 50135 FIRENZE - CAP. SOC. € 150.000.000,00 I.V. - C. F., P. I.V.A. E REG. IMPRESE 06395270481 C.C.P. 15447501 - R.E.A. 7874 - R.E.C. ISCRIZ. N.87678 DEL 18.9.92

Aziende del Gruppo Menarini

Italia: MALESCI - Firenze, F.I.R.M.A. - Firenze, CODIFI - Firenze, A. MENARINI FARMACEUTICA INTERNAZIONALE - Firenze, A. MENARINI INDUSTRIE FARMACEUTICHE RIUNITE - Firenze, A. MENARINI MANUFACTURING LOGISTICS AND SERVICES - Firenze, L'Aquila e Pisa, MENARINI RICERCHE - Firenze e Pomezia, MENARINI BIOTECH - Pomezia, GUIDOTTI - Pisa, LUSOFARMACO - Milano, LUSOCHIMICA - Pisa e Lomagna (Lecco), RESEARCH TOXICOLOGY CENTRE - Pomezia.

Mondo: ALBANIA - Tirana, ARGENTINA - Buenos Aires, ARMENIA - Yerevan, AUSTRALIA e NUOVA ZELANDA - Sydney, AUSTRIA - Vienna, AZERBAIGIAN - Baku, BELGIO - Bruxelles, BIELORUSSIA - Minsk, BOSNIA-ERZEGOVINA - Sarajevo, BULGARIA - Sofia, CINA - Pechino e Shanghai, COREA DEL SUD - Seul e Yonjin, COSTARICA - San Jose', CROAZIA - Zagabria, DANIMARCA - Copenhagen, EL SALVADOR - San Salvador, ESTONIA - Tallinn, FILIPPINE - Manila, FINLANDIA - Helsinki, FRANCIA - Parigi, GEORGIA - Tbilisi, GERMANIA - Berlino e Dresda, GRECIA - Atene, GUATEMALA - Città del Guatemala, HONDURAS - Tegucigalpa, HONG KONG - Hong Kong, INDIA - Ahmedabad, Mumbai e Nuova Delhi, INDONESIA - Bekasi e Jakarta, IRLANDA - Dublino e Shannon, KAZAKISTAN - Almaty, KIRGHIZISTAN - Bishkek, LETTONIA - Riga, LITUANIA - Vilnius, LUSSEMBURGO - Lussemburgo, MALESIA - Kuala Lumpur, MESSICO - Città del Messico, MOLDAVIA - Chisinau, MONTENEGRO - Podgorica, NICARAGUA - Managua, OLANDA - Amsterdam, PANAMA - Panama, POLONIA - Varsavia, PORTOGALLO - Lisbona, REGNO UNITO - Londra, REPUBBLICA Ceca - Praga, ROMANIA - Bucarest, RUSSIA - Mosca, SERBIA - Belgrado, SINGAPORE - Singapore, SLOVACCHIA - Bratislava, SLOVENIA - Lubiana, SPAGNA - Barcellona, SUD AFRICA - Bryanston, SVIZZERA - Zurigo, TAILANDIA - Bangkok, TAIWAN - Taipei, TURCHIA - Istanbul, TURKMENISTAN - Ashgabat, UCRAINA - Kiev, UNGHERIA - Budapest, UZBEKISTAN - Tashkent, VIETNAM - Hanoi e Ho Chi Minh.

Diagnostica: AUSTRIA - Vienna, BELGIO - Zaventem, FRANCIA - Parigi, GERMANIA - Berlino, GRECIA - Atene, ITALIA - Firenze, OLANDA - Valkenswaard, PORTOGALLO - Lisbona, REGNO UNITO - Londra, SPAGNA - Barcellona, SVEZIA - Malmö, SVIZZERA - Zurigo.

EFPIA Code Disclosure 2024 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 2023 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.

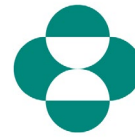
Date: 04.06.2024

Name of signatory: Belén Garijo

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

Signature:





EFPIA Code Disclosure 2024 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 2023 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 24, 2024

Name of signatory: Joseph Romanelli

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

Signature: 

**EFPIA Code Disclosure
2024 Self-Certification Scheme**

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom 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, Novartis Pharma AG 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:

1) 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.

2) 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).

3) 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.

4) Ensuring compliance with Data Privacy Obligations

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

After the spin-off between Novartis and Sandoz in November 2023, it has been decided that Novartis will publish the full Sandoz Transfer of Value (ToV) for the year 2023. From 2025 onwards, only the Novartis ToV will be published on the Novartis page.

Date: June 14, 2024

Name of signatory: Patrick Horber

Position in the Company: Novartis President, International

Signature:





EFPIA Code Disclosure 2024 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Novo Nordisk 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 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

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

Date: 17 June 2024

Name of signatory: Lars Fruergaard Jørgensen

Position in the Company: President & Chief Executive Officer

Signature:





EFPIA Code Disclosure 2024 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 2023 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: June 24th 2024

Name of signatory: Alexandre de Germay

Position in the Company: Chief International Commercial Office

Signature:

A handwritten signature in grey ink, appearing to read 'A. de Germay', is positioned below the 'Signature:' label.

EFPIA Code Disclosure 2024 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Pierre Fabre Médicament 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 Médicament 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

Pierre Fabre Médicament 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 Médicament 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).

Pierre Fabre Médicament

Siège social : Les Cauquillous – 81500 Lavaré Cédex
Société Anonyme au capital de 242 080 969 €
Siren : 326 118 502 RCS Castres

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

Date: 28/05/2024

Name of signatory: **Jean Luc LOWINSKI**

Position in the Company: **Président**

Signature:

DocuSigned by:

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EFPIA Code Disclosure 2024 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 building 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 2023 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.

Name of signatory: Teresa Graham
Position in the Company: CEO Roche
Pharmaceuticals
Signature

Teresa Graham

Date
30-May-2024 | 17:00 CEST

Name of signatory: Padraic Ward
Position in the Company: Head of Pharma
International
Signature

Padraic Ward

Date
30-May-2024 | 18:34 CEST



EFPIA Code Disclosure 2024 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 2023 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

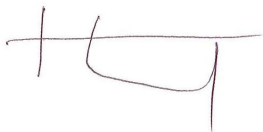
Sanofi certifies that its disclosure complies with the Data Privacy obligations.

Date: June 26, 2024

Name of signatory: OLIVIER CHARMEIL

Position in the Company: EVP, GENERAL MEDICINES

Signature:

A handwritten signature in blue ink, consisting of a stylized 'O' and 'C' followed by a vertical line.

EFPIA Code Disclosure 2024 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom STALLERGENES GREER 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 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

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

Date: June 20th, 2024

Name of signatory: Michele ANTONELLI

Position in the Company: Chief Executive Officer

Signature:





EFPIA Code Disclosure 2024 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 2023 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

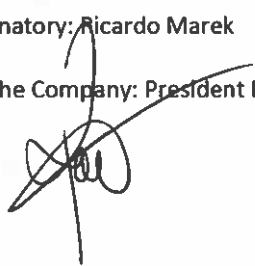
Takeda Pharmaceuticals International AG certifies that its disclosure complies with the Data Privacy obligations.

Date: 13.06.2024

Name of signatory: Ricardo Marek

Position in the Company: President Europe and Canada

Signature:

A handwritten signature in black ink, appearing to be 'Ricardo Marek', written over a horizontal line.



EFPIA Code Disclosure 2024 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 2023 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).

Teva Pharmaceuticals Europe B.V.

Piet Heinkade 107, P.O. Box 16416, 1001 RM AMSTERDAM - The Netherlands
Phone: +31(0)20 2193 000 - info@tevaeu.com - www.tevapharm.com

Chamber of Commerce Amsterdam 30110625, VAT NO. NL 003 973 190 B01,
Bank: Citibank: NL89CITI0266077889, BIC: CITINL2X



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

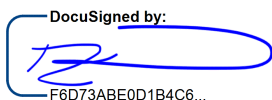
Teva Pharmaceuticals Europe B.V. certifies that its disclosure complies with the Data Privacy obligations.

Date: 20-Jun-2024 | 11:41 BST

Name of signatory: Richard Daniell

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

Signature:

DocuSigned by:

F6D73ABE0D1B4C6...

Teva Pharmaceuticals Europe B.V.

Piet Heinkade 107, P.O. Box 16416, 1001 RM AMSTERDAM - The Netherlands
Phone: +31(0)20 2193 000 - info@tevaeu.com - www.tevapharm.com

Chamber of Commerce Amsterdam 30110625, VAT NO. NL 003 973 190 B01,
Bank: Citibank: NL89CITI0266077889, BIC: CITINL2X



EFPIA Code Disclosure 2024 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 2023 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: 30-May-2024

Name of signatory:
Jean-Christophe Tellier

Position in the Company:
Head of UCB, CEO

Signature: 1B98A754E76844C...


Certificate Of Completion

Envelope Id: 693CCEFC1EC74FC1A73DED95256414C0		Status: Completed
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Contract Title:		
Source Envelope:		
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Time Zone: (UTC-08:00) Pacific Time (US & Canada)		DeniseMichelle.JONES@ucb.com
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Jean-Christophe.Tellier@ucb.com		Viewed: 5/30/2024 7:24:39 AM
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Security Level: Email, Account Authentication (None)		
Signature Adoption: Pre-selected Style		
Using IP Address: 46.178.29.34		
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Agent Delivery Events	Status	Timestamp
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Carbon Copy Events	Status	Timestamp
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Witness Events	Signature	Timestamp
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Notary Events	Signature	Timestamp
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Envelope Summary Events	Status	Timestamps
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Signing Complete	Security Checked	5/30/2024 8:30:10 AM
Completed	Security Checked	5/30/2024 8:30:10 AM

Payment Events	Status	Timestamps
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Electronic Record and Signature Disclosure

ELECTRONIC RECORD AND SIGNATURE DISCLOSURE

From time to time, UCB S.A., and its affiliates and subsidiary companies (we, us or Company) may be required by law to provide to you certain written notices or disclosures. Described below are the terms and conditions for providing to you such notices and disclosures electronically through your DocuSign, Inc. (DocuSign) Express user account. Please read the information below carefully and thoroughly, and if you can access this information electronically to your satisfaction and agree to these terms and conditions, please confirm your agreement by clicking the 'I agree' button at the bottom of this document.

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To advise UCB S.A., and its affiliates and subsidiary companies of your new e-mail address

To let us know of a change in your e-mail address where we should send notices and disclosures electronically to you, you must send an email message to us at steve.davis@ucb.com and in the body of such request you must state: your previous e-mail address, your new e-mail address. We do not require any other information from you to change your email address..

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To inform us that you no longer want to receive future notices and disclosures in electronic format you may:

- i. decline to sign a document from within your DocuSign account, and on the subsequent page, select the check-box indicating you wish to withdraw your consent, or you may;
- ii. send us an e-mail to steve.davis@ucb.com and in the body of such request you must state your e-mail, full name, IS Postal Address, telephone number, and account number. We do not need any other information from you to withdraw consent.. The consequences of your withdrawing consent for online documents will be that transactions may take a longer time to process..

Required hardware and software

Operating Systems:	Windows2000? or WindowsXP?
Browsers (for SENDERS):	Internet Explorer 6.0? or above
Browsers (for SIGNERS):	Internet Explorer 6.0?, Mozilla FireFox 1.0, NetScape 7.2 (or above)
Email:	Access to a valid email account
Screen Resolution:	800 x 600 minimum
Enabled Security Settings:	<ul style="list-style-type: none">• Allow per session cookies

	<ul style="list-style-type: none"> • Users accessing the internet behind a Proxy Server must enable HTTP 1.1 settings via proxy connection
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** These minimum requirements are subject to change. If these requirements change, we will provide you with an email message at the email address we have on file for you at that time providing you with the revised hardware and software requirements, at which time you will have the right to withdraw your consent.

Acknowledging your access and consent to receive materials electronically

To confirm to us that you can access this information electronically, which will be similar to other electronic notices and disclosures that we will provide to you, please verify that you were able to read this electronic disclosure and that you also were able to print on paper or electronically save this page for your future reference and access or that you were able to e-mail this disclosure and consent to an address where you will be able to print on paper or save it for your future reference and access. Further, if you consent to receiving notices and disclosures exclusively in electronic format on the terms and conditions described above, please let us know by clicking the 'I agree' button below.

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- I can print on paper the disclosure or save or send the disclosure to a place where I can print it, for future reference and access; and
- Until or unless I notify UCB S.A., and its affiliates and subsidiary companies as described above, I consent to receive from exclusively through electronic means all notices, disclosures, authorizations, acknowledgements, and other documents that are required to be provided or made available to me by UCB S.A., and its affiliates and subsidiary companies during the course of my relationship with you.