EFPIA Code Disclosure
2023 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom AbbVie works to 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 2022 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;

AbbVie Europe
10 rue d'Arcueil
94150 Rungis
France
Tél : +33 (0)1 45 60 15 50

Succursale Française AbbVie Biopharmaceuticals Sarl G
RC de Zoug. CH-170.4.011.236-4
Siège social : Neuhofstrasse 23,6341 Baar
RCS Créteil 752 319 343

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

AbbVie Europe
10 rue d'Arcueil
94150 Rungis
France
Tél : +33 (0)1 45 60 15 50

Succursale Française AbbVie Biopharmaceuticals Sarl G
RC de Zoug. CH-170.4.011.236-4
Siège social : Neuhofstrasse 23,6341 Baar
RCS Créteil 752 319 343

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: _____

Name of signatory: Jerome Bouyer

Position in the Company: Senior Vice President, Europe – AbbVie

Signature: jerome bouyer _____

Electronically signed by: jerome bouyer
Reason: Management Approval
Date: Jun 16, 2023 11:45 GMT+2

AbbVie Europe
10 rue d'Arcueil
94150 Rungis
France
Tél : +33 (0)1 45 60 15 50

Succursale Française AbbVie Biopharmaceuticals Sarl G
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Siège social : Neuhofstrasse 23,6341 Baar
RCS Créteil 752 319 343

EFPIA Code Disclosure 2023 Self-Certification Scheme

Final Audit Report 2023-06-16

Created:	2023-06-16
By:	Audra Marmu (audra.marmu@abbvie.com)
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"EFPIA Code Disclosure 2023 Self-Certification Scheme" History

Document created by Audra Marmu (audra.marmu@abbvie.com)
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almirall
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EFPIA Code Disclosure
2023 Self-Certification Scheme

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

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

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

Except in countries where disclosure is prescribed by laws, Almirall 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

Almirall 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 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).

Almirall, S.A.
Rda. General Mitre, 151
08022 Barcelona, Spain
P +34 93 291 30 00
F +34 93 291 31 80
almirall.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

Almirall 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

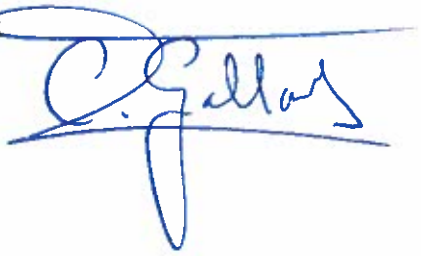
Almirall certifies that its disclosure complies with the Data Privacy obligations.

Date: 7/6/2023

Name of signatory: Carlos Gallardo

Position in the Company: CEO

Signature:



Almirall, S.A.
Rda. General Mitre, 151
08022 Barcelona, Spain

P +34 93 291 30 00
F +34 93 291 31 80

almirall.com

EFPIA Code Disclosure
2023 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 2022 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: 3/28/2023

Name of signatory: Murdo Gordon

Position in the Company: Executive Vice President Global Commercial Operations

Signature: DocuSigned by:

Signer Name: Murdo Gordon
Signing Reason: I approve this document
Signing Time: 3/28/2023 | 3:41:37 PM GMT
461AC89E462847C9A2D3F0A5002098E9

EFPIA Code Disclosure
2023 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 2022 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: 22 juin 2023

Name of signatory: Damien Bailly

Position in the Company: President - Established Markets

Signature: DocuSigned by: Damien Bailly 4F5270BA99FC459...

EFPIA Code Disclosure
2023 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 2022 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: Jun 30, 2023

Name of signatory: Stefan Woxström

Position in the Company: SVP Europe & Canada

Signature: DocuSigned by:
073ED573BFBC4CC...

EFPIA Code Disclosure
2023 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 2022 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).

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, June 2, 2023

Stefan Oelrich
Member of the Board of Management
President Pharmaceuticals Division

Berlin, June 1, 2023

ppa. Königer

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

EFPIA Code Disclosure
2023 Self-Certification Scheme

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

Bial-Portela & Cª, 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 & Cª, 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).

À AV. DA SIDERURGIA NACIONAL • 4745-457 CORONADO (S. Romão e S. Mamede) • PORTUGAL • Tel.: +351 22 986 6100 • www.bial.com • info@bial.com
Sociedade Anónima • Cons. Reg. Com. Trofa Matrícula nº 500 220 913 • NIPC 500 220 913 • Capital Social €50.000.000

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 & Cª, 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 & Cª, SA certifies that its disclosure complies with the Data Privacy obligations.

Date: 2023.04.26

Name of signatory: António Portela

Position in the Company: Chief Executive Officer

Signature:

À AV. DA SIDERURGIA NACIONAL • 4745-457 CORONADO (S. Romão e S. Mamede) • PORTUGAL • Tel.: +351 22 986 6100 • www.bial.com • info@bial.com
Sociedade Anónima • Cons. Reg. Com. Trofa Matrícula nº 500 220 913 • NIPC 500 220 913 • Capital Social €50.000.000

EFPIA Code Disclosure
2023 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: 15th June 2023

Name of signatory: Wolfram Schmidt

Position in the Company: President Europe, Canada & Partner Markets

Signature: DocuSigned by: Wolfram Schmidt DC9F2170B9DC435...

Boehringer Ingelheim

EFPIA Code Disclosure
2023 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, I hereby confirm 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

I certify that:

- The disclosures are made in each country covered by the EFPIA Code where Boehringer Ingelheim operates;
- The 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

I certify 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.

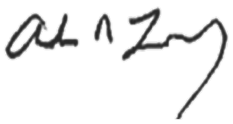
Ensuring compliance with Data Privacy Obligations

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

Date: June 12, 2023

Name of signatory: Adam Lenkowsky

Position in the Company: EVP Chief Commercialisation Officer

Signature: 

EFPIA Code Disclosure
2023 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 2022 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

I certify 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.

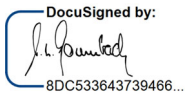
Ensuring compliance with Data Privacy Obligations

I certify that Boehringer Ingelheim's disclosure complies with the Data Privacy obligations.

Date: 01-Jun-2023

Name: Hubertus v. Baumbach

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

Signature: 

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

EFPIA Code Disclosure
2023 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 2022 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 12th, 2023

Name of signatory: Alberto Chiesi

Position in the Company: President

Signature: A. Chiesi

CSL Vifor

EFPIA Code Disclosure
2023 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 2022 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: 18-Apr-23

Name of signatory: Hervé Gisserot

Position in the Company: General Manager CSL Vifor

Signature: DocuSigned by: Hervé Gisserot FCA3E999854C41E...

Passion for Innovation.
Compassion for Patients.™

Daiichi-Sankyo

Daiichi Sankyo Europe GmbH
Zielstattstrasse 48
81379 Munich · Germany
Phone +49 89 78080
Fax +49 89 7808267
service@daiichi-sankyo.eu
www.daiichi-sankyo.eu

EFPIA Code Disclosure
2023 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 2022 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Daiichi Sankyo Europe GmbH certifies that:

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

Passion for Innovation.
Compassion for Patients.™

Daiichi-Sankyo

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

Daiichi Sankyo Europe GmbH certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Date:	June 15, 2023	Juni 15, 2023
Name of signatory:	Curd Lejaegere	ppa. Martin Fürle
Position in the Company:	VP Mid-Size Countries Daiichi Sankyo Europe Efpia Representative	GC & CCO Europe
Signature:	DocuSigned by: Curd Lejaegere 15BA0ABE3AC1406...	DocuSigned by: Martin Fürle AF72B0726EF9440...

EFPIA Code Disclosure
2023 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Eisai Europe Limited 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 Limited 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

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

Date: 5th July 2023

Name of signatory: Nick Burgin

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

Signature: DocuSigned by: Nick Burgin DF29518349B8416...

GILEAD
Creating Possible

EFPIA Code Disclosure
2023 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 2022 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: 16 May 2023

Name of signatory: Rudolf Ertl

Position in the Company: Senior Vice President, ACE Region

Signature:

IPSEN
Innovation for patient care

EFPIA Code Disclosure
2023 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 2022 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).

.../...

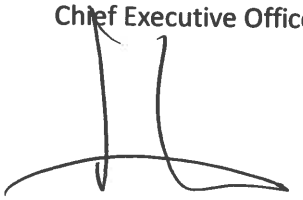
Aggregate disclosures are limited to Research and Development ToVs and such HCPs/HCOs' ToVs that cannot be disclosed on an individual basis for legal reasons

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:	08 June 2023
Name of signatory:	David Loew
Position in the Company:	Chief Executive Officer
Signature:	

Janssen Pharmaceutica NV
Turnhoutseweg 30
B-2340 Beerse, Belgium

EFPIA Code Disclosure
2023 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Janssen Pharmaceutica 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 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

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

Date: Beerse, 3 May 2023

Name of signatory: Kris Sterkens

Position in the Company: Company Group Chairman Janssen EMEA

Signature:

EFPIA Code Disclosure
2023 Self-Certification Scheme

Dermatology
beyond the skin

LEO Pharma A/S
Industriparken 55
2750 Ballerup
Denmark

Main +45 4494 5888

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

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 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, LEO Pharma 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

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: 14.06.2023

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
2023 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 2022 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).

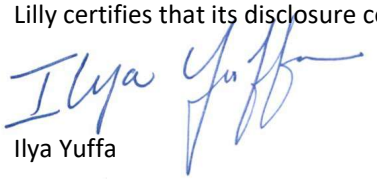
Aggregate disclosures are limited to Research and Development ToVs and such ToVs that cannot be disclosed on an individual basis for legal reasons

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 2023

EFPIA Code Disclosure
2023 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 2022 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: June 20TH 2023

Name of signatory: Deborah Dunsire

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

Signature:

A. MENARINI
INDUSTRIE FARMACEUTICHE RIUNITE

EFPIA Code Disclosure
2023 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom A. Menarini Industrie Farmaceutiche Riunite S.r.l. 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, A. Menarini Industrie Farmaceutiche Riunite S.r.l. 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

A. Menarini Industrie Farmaceutiche Riunite S.r.l. 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

A. Menarini Industrie Farmaceutiche Riunite S.r.l. certifies that:

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

A. MENARINI
INDUSTRIE FARMACEUTICHE RIUNITE

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

A. Menarini Industrie Farmaceutiche Riunite S.r.l. 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

A. Menarini Industrie Farmaceutiche Riunite S.r.l. certifies that its disclosure complies with the Data Privacy obligations.

Date: 6 June 2023

Name of signatory: Luca Lastrucci

Position in the Company: General Manager Medico Marketing and Sales, Pharmaceuticals

Signature:

EFPIA Code Disclosure
2023 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 2022 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

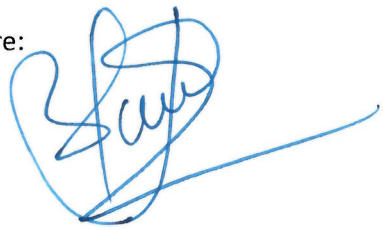
Merck certifies that its disclosure complies with the Data Privacy obligations.

Date: 18.04.2023

Name of signatory: Belén Garijo

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

Signature:



Joseph Romanelli
President, Human Health International

MSD International Business GmbH
Tribschenstrasse 60
6005 Lucerne, Switzerland

www.msd.com

EFPIA Code Disclosure
2023 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 2022 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.

Joseph Romanelli
President, Human Health International

MSD International Business GmbH
Tribschenstrasse 60
6005 Lucerne, Switzerland

www.msd.com

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: May 26, 2023

Name of signatory: Joseph Romanelli

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

Signature:

NOVARTIS | Reimagining Medicine

EFPIA Code Disclosure: 2023 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 2022 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

Date: 21. June 2023

Name of signatory:
Marie-France Tschudin

Position in the Company:
President, IM Int & Chief Commercial Officer

Signature:

EFPIA Code Disclosure
2023 Self-Certification Scheme

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

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

Except in countries where disclosure is prescribed by laws, Novo Nordisk A/S hereby confirms that its disclosures of transfers of value (ToVs) to HCPs, HCOs and POs made in 2022 have been reported in application of the EFPIA Code following key principles:

Disclosure quality

Novo Nordisk A/S certifies that:

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

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

Novo Nordisk A/S certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Date: 26 MAY 2023

Name of signatory: **Lars Fruergaard Jørgensen**

Position in the Company: **President & Chief Executive Officer**

Signature: 

EFPIA Code Disclosure
2023 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 2022 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: 12 June 2023

Name of signatory: Angela Hwang

Position in the Company: Chief Commercial Officer, President, Global Biopharmaceuticals Business.

Signature:

Pierre Fabre
Médicament

EFPIA Code Disclosure
2023 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 2022 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

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

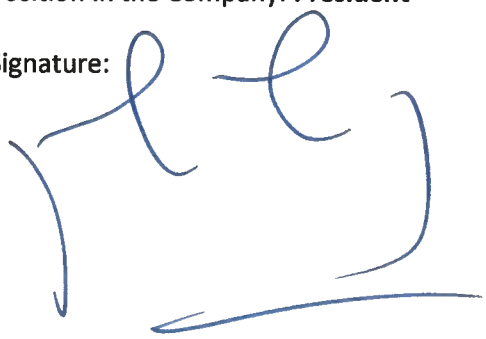
Pierre Fabre Médicament certifies that its disclosure complies with the Data Privacy obligations.

Date: 16.05.2023

Name of signatory: **Jean Luc LOWINSKI**

Position in the Company: **Président**

Signature:



EFPIA Code Disclosure
2023 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom F. Hoffmann –La Roche (hereinafter "Roche") 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, Roche 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

Roche 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

Roche 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

Roche 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

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

Date: 09-May-2023	Date: 09-May-2023
Name of signatory: Teresa Graham	Name of signatory: Padraic Ward
Position in the Company: CEO Roche Pharma	Position in the Company: Head of Pharma International
Signature: Teresa Graham	Signature: Padraic Ward

sanofi

EFPIA Code Disclosure
2023 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 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, Sanofi 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

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).

sanofi

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.

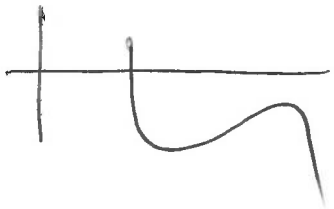
Ensuring compliance with Data Privacy Obligations

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

Date: May 4, 2023

Name of signatory: Olivier Charmeil

Position in the Company: EVP, General Medicines

Signature: 

EFPIA Code Disclosure
2023 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 2022 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: June 14th 2023

Name of signatory: Ricardo Marek

Position in the Company: President Europe and Canada

Signature:

EFPIA Code Disclosure
2023 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 2022 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).

...2/

Teva Pharmaceuticals Europe B.V.
Piet Heinkade 107, P.O. Box 16416, 1001 RM AMSTERDAM – The Netherlands
Phone: +31(0)20 2193 000 – Fax: +31 (0)20 2193 299 – 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: June 8th, 2023

Name of signatory: Richard Daniell

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

Signature: 

Inspired by patients.
Driven by science.

EFPIA Code Disclosure
2023 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 2022 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).