abbvie

EFPIA Disclosure Code
2020 Self-Certification Scheme

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

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, AbbVie hereby confirms that its disclosures of transfers of value (ToV) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

AbbVie certifies that:

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

AbbVie
1 North Waukegan Road
North Chicago, IL 60064

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

AbbVie certifies that:

- Data collection complies with the requirements of the EFPIA Disclosure Code;
- Actions are taken to encourage individual disclosure for HCP's and HCO's transfers of value (each as defined in the EFPIA Disclosure Code).

Aggregate disclosures are limited to Research and Development ToV and such ToV 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 Disclosure 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 section.

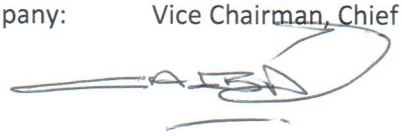
Ensuring compliance with Data Privacy Obligations

AbbVie certifies that its disclosure complies with the applicable privacy and data protection law.

Date: June 8, 2020

Name of signatory: Carlos Alban

Position in the Company: Vice Chairman, Chief Commercial Officer

Signature: 

almirall
feel the science

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Almirall hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Almirall certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Almirall certifies that:

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

almirall
feel the science

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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.

Peter Guenter
Chief Executive Officer

25th June 2020

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 Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Amgen hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Amgen certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Amgen certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: 6/4/2020

Name of signatory: Murdo Gordon

Position in the Company: Executive Vice President Global Commercial Ops

Signature: DocuSigned by:
Murdo Gordon
Signer Name: Murdo Gordon
Signing Reason: I approve this document
Signing Time: 6/4/2020 | 11:25:30 AM PDT
461AC89E462847C9A2D3F0A5002098E9

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

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 and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Astellas Pharma Europe Ltd certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Astellas Pharma Europe Ltd certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: 23rd June 2020

Name of signatory: Mr. Dirk Kosche

Position in the Company: President of Established Markets- Commercial

Signature:

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, AstraZeneca hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

AstraZeneca certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

AstraZeneca certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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

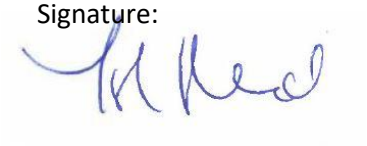
AstraZeneca certifies that its disclosure complies with the Data Privacy obligations.

Date: 24/06/2020

Name of signatory: Iskra Reic

Position in the Company: Executive Vice President Europe

Signature:



EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Bayer AG hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Bayer AG certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Bayer AG certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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,

DocuSigned by:
6D6E3CEAE949498...

Stefan Oelrich
Member of the Board of Management
President Pharmaceuticals Division

Berlin, 6/17/2020 | 3:22:14 PM CEST

DocuSigned by:
173EFABB48DE48F...

Dr. Stefan Gehring
Law, Patents and Compliance
Business Partner Pharmaceuticals

Boehringer Ingelheim

Hubertus von Baumbach

EFPIA
Leopold Plaza Building
Rue du Trone
1050 Bruxelles
BELGIUM

29 June 2020

EFPIA Disclosure Code
2020 Self-Certification Scheme

BMD Chairman

Phone +49 6132 77-2899
Fax +49 6132 77-2899

C.H. Boehringer Sohn
AG & Co. KG
Binger Strasse 173
55216 Ingelheim am Rhein
Germany

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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Boehringer Ingelheim hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Limited Partnership

Registered Office
Ingelheim am Rhein
Commercial Register Mainz
HR A 21732

General Partner
Boehringer AG

Board of Managing Directors
Hubertus von Baumbach
(Chair)
Carinne Brouillon
Dr Michel Pairet
Jean Scheftsik de Szolnok
Michael Schmelmer

Chair of the Supervisory Board
Christian Boehringer

Registered Office
Ingelheim am Rhein
Commercial Register Mainz
HR B 23354

Disclosure quality

Boehringer Ingelheim certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Boehringer Ingelheim certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Name of signatory: Hubertus von Baumbach
Position in the Company: Chairman of the Board of Managing Directors

Date: June 29, 2020

Signature: 

Bial

PORTELA & Cª, S. A.

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

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 and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Bial-Portela & Cª, SA certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Bial-Portela & Cª, SA certifies that:

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

À AV. DA SIDERURGIA NACIONAL • 4745-457 CORONADO (S. Romão e S. Mamede) • PORTUGAL • Tel.: +351 22 986 6100 • Fax: +351 22 986 6190 • 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 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 Disclosure 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

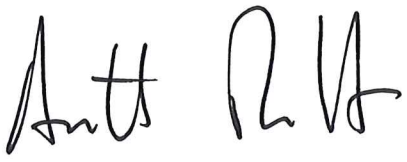
Bial-Portela & Cª, SA certifies that its disclosure complies with the Data Privacy obligations.

Date: 2020.06.30

Name of signatory: António Portela

Position in the Company: Chief Executive Officer

Signature:



EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Biogen hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Biogen certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Biogen certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: 30th June 2020

Name of signatory: Johanna Friedl-Naderer

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

Signature: DocuSigned by: Johanna Friedl-Naderer E770471D5B234DF...

Bristol Myers Squibb™

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Bristol-Myers Squibb Company, LLC 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Bristol-Myers Squibb Company, LLC hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Bristol-Myers Squibb Company, LLC certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Bristol-Myers Squibb Company, LLC certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 Company, LLC certifies that its disclosure complies with the Data Privacy obligations.

Date:

Name of signatory: Christopher Boerner

Position in the Company: EVP, Chief Commercialization Officer

Signature:

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Celgene Corporation 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Celgene Corporation hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Celgene Corporation certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Celgene Corporation certifies that:

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

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

Celgene Corporation certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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

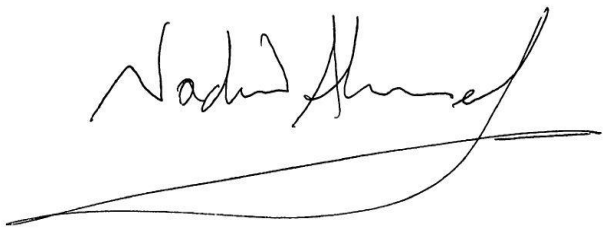
Celgene Corporation certifies that its disclosure complies with the Data Privacy obligations.

Date: 6/10/2020

Name of signatory: Nadim Ahmed

Position in the Company: EVP & President of Hematology

Signature:



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

People and ideas for innovation in healthcare

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

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 and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Chiesi Farmaceutici S.p.A. certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Chiesi Farmaceutici S.p.A. certifies that:

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

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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 22nd, 2020

Name of signatory: Alberto Chiesi

Position in the Company: President

Signature: A. Chiesi

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.daichi-sankyo.eu

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Daiichi Sankyo 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Daiichi Sankyo hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Daiichi Sankyo certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Daiichi Sankyo certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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

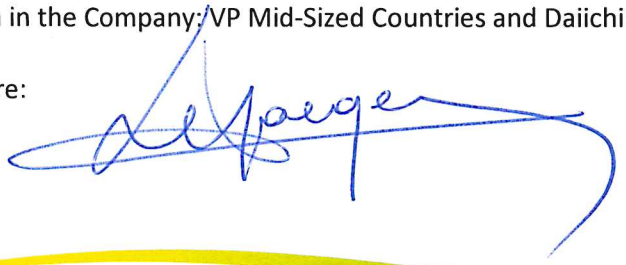
Daiichi Sankyo certifies that its disclosure complies with the Data Privacy obligations.

Date: July 2, 2020

Name of signatory: Curd Lejaegere

Position in the Company: VP Mid-Sized Countries and Daiichi Sankyo Representative to EFPIA

Signature:



EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, (Eisai Europe Limited) hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

(Eisai Europe Limited) certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

(Eisai Europe Limited) certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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

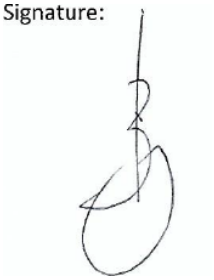
(Eisai Europe Limited) certifies that its disclosure complies with the Data Privacy obligations.

Date: 16.06.2020

Name of signatory: Nick Burgin

Position in the Company: President & COO EMEA & President General Value & Access

Signature:



ESTEVE
Torre Esteve
Pg. de la Zona Franca, 109, 4ª planta
08038 Barcelona (Spain)
T. +34 93 446 60 00
www.esteve.com

EFPIA Disclosure Code
2020 Self-Certification Scheme

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

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Esteve Pharmaceuticals, S.A. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Esteve Pharmaceuticals, S.A. certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Esteve Pharmaceuticals, S.A. certifies that:

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

ESTEVE
Torre Esteve
Pg. de la Zona Franca, 109, 4ª planta
08038 Barcelona (Spain)
T. +34 93 446 60 00
www.esteve.com

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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

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

19/06/2020

Staffan Schüberg
Chief Executive Officer

Signature:

DocuSigned by:
7638F633C7714C8...

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Grünenthal 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Grünenthal hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Grünenthal certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Grünenthal certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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.

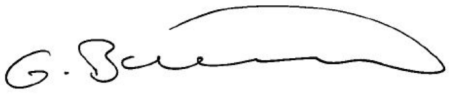
Ensuring compliance with Data Privacy Obligations

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

Date: 23.06.2020

Name of signatory: Gabriel Baertschi

Position in the Company: Chief Executive Officer



Signature (Gabriel Baertschi)

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom GlaxoSmithKline 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, GlaxoSmithKline hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

GlaxoSmithKline certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

GlaxoSmithKline certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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

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

Date: 01 June 2020

Name of signatory: Luke Miels

Position in the Company: President Global Pharmaceuticals - GlaxoSmithKline

Signature: 

IPSEN
Innovation for patient care

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Ipsen hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Ipsen certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Ipsen certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: June 29, 2020

Name of signatory: Aymeric Le Chatelier

Position in the Company: Chief Executive Officer

Signature:

Janssen Pharmaceutica NV
Turnhoutseweg 30
B-2340 Beerse, Belgium

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Janssen Pharmaceutica hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Janssen Pharmaceutica certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Janssen Pharmaceutica certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: 2 June 2020

Name of signatory: Kris Sterkens

Position in the Company: Company Group Chairman Janssen EMEA

Signature:

Dermatology
beyond the skin

LEO Pharma A/S
Industriparken 55
2750 Ballerup
Denmark

Main +45 4494 5888
Fax +45 7226 3321

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

EFPIA Disclosure Code
2020 Self-Certification Scheme

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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, LEO Pharma hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

LEO Pharma certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

LEO Pharma certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: June 22, 2020

Name of signatory: **Catherine Mazzacco**

Position in the Company: **President & 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
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Lilly hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Lilly certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Lilly certifies that:

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

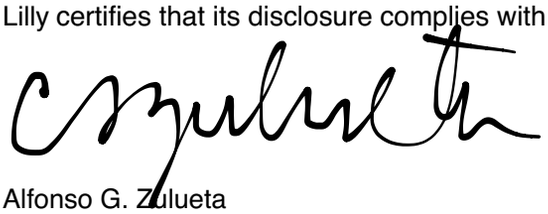
Aggregate disclosures are limited to Research and Development ToVs and such ToVs that cannot be disclosed on an individual basis for legal reasons

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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

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



Alfonso G. Zulueta

Senior Vice-President and President of Lilly International

May 2020

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Lundbeck hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Lundbeck certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Lundbeck certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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 8th 2020

Deborah Dunsire
CEO, H. Lundbeck A/S

A. MENARINI
INDUSTRIE FARMACEUTICHE RIUNITE

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

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 and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

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

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

A. MENARINI
INDUSTRIE FARMACEUTICHE RIUNITE

Methodology used for the collection and organisation of ToVs is in line with the EFPIA Disclosure 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 Disclosure Code;
- Actions are taken to ensure individual disclosure for HCPs and HCOs' transfers of values (each as defined in the EFPIA Disclosure Code).

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: 29 June 2020

Name of signatory: Eric Cornut

Position in the Company: Chairman of the Board of Directors

Signature:

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Merck hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Merck certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Merck certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: May 26, 2020

Name of signatory: Stefan Oschmann

Position in the Company: Chairman of the Executive Board & CEO

Signature:

Frank K. Clyburn

Chief Commercial Officer
Merck & Co., Inc.

Merck & Co., Inc.
2000 Galloping Hill Road
Kenilworth, NJ 07033 M
U.S.A
Phone: 1.908-740-4000
merck.com

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Merck & Co., Inc. 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Merck & Co., Inc. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Merck & Co., Inc. certifies that:

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

Frank K. Clyburn

Chief Commercial Officer
Merck & Co., Inc.

Merck & Co., Inc.
2000 Galloping Hill Road
Kenilworth, NJ 07033 M
U.S.A
Phone: 1.908-740-4000
merck.com

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

Merck & Co., Inc. certifies that:

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

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

Merck & Co., Inc. certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 & Co., Inc. certifies that its disclosure complies with the Data Privacy obligations.

Date: June 22, 2020

Name of signatory: Frank K. Clyburn

Position in the Company: Chief Commercial Officer, Merck & Co., Inc.

Signature: 

Novartis Pharma AG
Communication
Novartis Campus
Postfach
4002 Basel
Switzerland
T: +41 61 324 1321

EFPIA Disclosure Code - 2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Novartis Pharma AG hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Novartis Pharma AG certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Novartis Pharma AG certifies that:

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

NOVARTIS

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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

The collection, processing and disclosure of transfers of value have been made in accordance with the Data Privacy laws applicable in the respective countries.

Date: 8. 6 - 2020

Name of signatory:

Susanne Schaffert

Position in the Company:

President Novartis Oncology

Signature:

EFPIA Disclosure Code
2020 Self-Certification Scheme

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

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Novo Nordisk hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Novo Nordisk certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Novo Nordisk certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Date: 04. June 2020

Name of signatory: Maziar Mike Doustdar

Position in the Company: Executive Vice President for International Operations

Signature:

Otsuka
Otsuka-people creating new products for better health worldwide

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Otsuka Pharmaceutical Europe Ltd., its affiliate companies and subsidiaries ("Otsuka") 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Otsuka hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Otsuka certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Otsuka certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Date: 10th June 2020

Name of signatory: Mel Walker

Position in the Company: RVP, Innovation, Business Development & Patient Access

Signature: 

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Pfizer Inc. 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Pfizer Inc. hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Pfizer Inc. certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Pfizer Inc. certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 Inc. certifies that its disclosure complies with the Data Privacy obligations.

Date: 6/9/2020

Name of signatory: Angela Hwang

Position in the Company: Group President, Pfizer Biopharmaceuticals Group

Signature: 

Pierre Fabre
Médicament

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom PIERRE FABRE MEDICAMENT 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, PIERRE FABRE MEDICAMENT hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

PIERRE FABRE MEDICAMENT certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

PIERRE FABRE MEDICAMENT certifies that:

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

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

PIERRE FABRE MEDICAMENT certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 MEDICAMENT certifies that its disclosure complies with the Data Privacy obligations.

Date: 17 JUIN 2020

PIERRE FABRE MEDICAMENT

Jean-Luc LOWINSKI
President

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs aims to build 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Roche hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Roche certifies that:

- its disclosures are made in each country where it operates;
- its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Roche certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: May 27, 2020

Name of signatory: Bill Anderson

Position: CEO Roche Pharmaceuticals

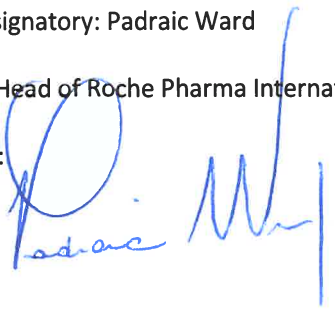
Signature:

Date: June 9, 2020

Name of signatory: Padraic Ward

Position: Head of Roche Pharma International

Signature:



EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organizations (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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Sanofi hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Sanofi certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 organization of ToVs is in line with the EFPIA Disclosure Code's requirements and applicable codes

Sanofi certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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

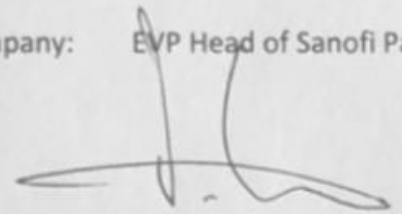
Sanofi certifies that its disclosure complies with the Data Privacy obligations.

Date: 3.6.2020

Name of signatory: David Loew

Position in the Company: EVP Head of Sanofi Pasteur

Signature:



EFPIA Disclosure Code
2020 Self-Certification Scheme

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

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, SERVIER hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

SERVIER certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

SERVIER certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Date: June 19th, 2020

Name of signatory: M. Olivier Laureau

Position in the Company: President

Signature:

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Shire, now part of 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Shire, now part of Takeda Pharmaceuticals International AG hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Shire, now part of Takeda Pharmaceuticals International AG certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Shire, now part of Takeda Pharmaceuticals International AG certifies that:

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

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

Shire, now part of 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 Disclosure 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

Shire, now part of Takeda Pharmaceuticals International AG certifies that its disclosure complies with the Data Privacy obligations.

Date: 20 May 2020

Name of signatory: Giles Platford

Position in the Company: President Europe and Canada

Signature:

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

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 and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Takeda Pharmaceuticals International AG certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Takeda Pharmaceuticals International AG certifies that:

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

Aggregate disclosures are limited to Research and Development ToVs and such 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 Disclosure 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: 20 May 2020

Name of signatory: Giles Platford

Position in the Company: President Europe and Canada

Signature:

teva

EFPIA Disclosure Code
2020 Self-Certification Letter
Teva Pharmaceuticals Europe BV

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Teva Pharmaceutical Europe BV 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Teva Pharmaceutical Europe BV hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure scope:

Teva Pharmaceutical Europe BV certifies that its disclosures of ToVs:

- have been completed in each EFPIA country where Teva Pharmaceuticals Europe BV operates,
- include direct and indirect ToVs, as defined in the codes and associated guidance issued by EFPIA as well as associated national codes/clarifications, and
- are further described in the respective country's Methodological Note.

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

Teva Pharmaceutical Europe BV certifies that:

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

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

Teva Pharmaceutical Europe BV certifies that aggregate disclosure is limited to the following topics:

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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;
- Decisions of local code authorities in response to the COVID19 pandemic (ABPI in the UK).

Ensuring compliance with data privacy obligations

Teva Pharmaceutical Europe BV also certifies that its disclosure complies with relevant data privacy obligations.

Date: June 26, 2020

Name of signatory: Richard Daniell

Position in the Company: Executive Vice President
Teva Pharmaceuticals Europe BV

Signature: 

Inspired by patients.
Driven by science.

EFPIA Disclosure Code
2020 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, UCB hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

UCB certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

UCB certifies that:

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

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

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

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

Ensuring compliance with Data Privacy Obligations

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

Date: 22/06/2020

Name of signatory: Jean-Christophe Tellier

Position in the Company: Head of UCB, CEO

Signature:

EFPIA Disclosure Code
2020 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Vifor Pharma Group 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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. Despite the exceptional circumstances related to COVID-19, Member Companies made their best efforts to disclose the relevant information for 2019.

Except in countries where disclosure is prescribed by laws, Vifor Pharma Group hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2019 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Vifor Pharma Group certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Vifor Pharma Group certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 Pharma Group certifies that its disclosure complies with the Data Privacy obligations.

Date: 24.06.2020

Name of signatory: Andreas Walde

Position in the Company: General Secretary

Signature: 

VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA

VIFOR PHARMA

EFPIA Disclosure Code
2019 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Vifor Pharma Group 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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 Pharma Group hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2018 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Vifor Pharma Group certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Vifor Pharma Group certifies that:

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 Pharma Group certifies that its disclosure complies with the Data Privacy obligations.

Date: 20 June 2019

Name of signatory: Dr. Oliver P. Kronenberg

Position in the Company: Group General Counsel

Signature:

Vifor Pharma Ltd.

Name of signatory: Dr. Andreas Walde

Position in the Company: General Secretary

Signature:

VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA

VIFOR PHARMA

EFPIA Disclosure Code
2018 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Vifor Pharma Group 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.

Throughout the medicines life cycle pharmaceutical companies work with scientists and HCPs. These collaborations are essential in addressing patient needs. Industry and HCPs collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

EFPIA and its member associations have adopted codes and guidelines to ensure that the interactions of their member companies with HCPs and HCOs meet the high standards of integrity and transparency. Building greater transparency to the relationships between pharmaceutical companies and HCPs/HCOs 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 Pharma Group hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2017 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Vifor Pharma Group certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Vifor Pharma Group certifies that:

- Data collection complies with the requirements of the EFPIA Disclosure Code;

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

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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 Pharma Group certifies that its disclosure complies with the Data Privacy obligations.

Date: 30 June 2018

Name of signatory: Dr. Oliver P. Kronenberg

Position in the Company: Group General Counsel

Signature:

Name of signatory: : Dr. Andreas Walde

Position in the Company: General Secretary

Signature:

VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA

VIFOR PHARMA

EFPIA Disclosure Code
2016 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Vifor Pharma Group 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.

Throughout the medicines life cycle pharma companies work with scientists and healthcare professionals. These collaborations are essential in addressing patient needs. Industry and healthcare professionals collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

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

Vifor Pharma Group hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2016 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Vifor Pharma Group certifies that:

- Its disclosures are made in each country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes 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 Disclosure Code's requirements and applicable codes

Vifor Pharma Group certifies that:

- Data collection complies with the requirements of the EFPIA Disclosure Code;
- Actions were taken to seek consent from HCPs and HCOs (each as defined in the EFPIA Disclosure Code), where applicable.

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure 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.

Seeking the consent of Recipients

Vifor Pharma Group certifies that it has used all reasonable steps for the purpose of obtaining consent to individual disclosure where such consent is required by applicable law.

Date: 5 July 2017

Name of signatory: Stefan Schulze

Position in the Company: President of the Executive Committee and Chief Operational Officer Vifor Pharma

Signature:

Name of signatory: Oliver P. Kronenberg

Position in the Company: General Counsel Vifor Pharma

Signature:

Vifor Pharma

EFPIA Disclosure Code
2015 Self-Certification Scheme

Healthcare professionals (HCPs) and healthcare organisations (HCOs) with whom Vifor 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.

Throughout the medicines life cycle pharma companies work with scientists and healthcare professionals. These collaborations are essential in addressing patient needs. Industry and healthcare professionals collaborate in a range of activities from clinical research to sharing best clinical practice and exchanging information on how new medicines fit into the patient pathway.

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

Vifor Pharma hereby confirms that its disclosures of transfers of value (ToVs) to HCPs and HCOs made in 2015 have been reported in application of the EFPIA Disclosure Code following key principles:

Disclosure quality

Vifor Pharma certifies that:

- Its disclosures are made for each EFPIA country where it operates;
- Its disclosures include direct and indirect ToVs, as defined in the codes and associated guidance issued by EFPIA;
- Its Methodological Notes describe 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 Disclosure Code's requirements and applicable codes

Vifor Pharma certifies that:

- Data collection complies with the requirements of the EFPIA Disclosure Code;
- Actions were taken to seek consent from HCPs and HCOs (each as defined in the EFPIA Disclosure Code), where applicable.

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

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

- Research and Development Transfers of Value (as defined in the EFPIA Disclosure Code);
- Transfers of Value to Recipients that have opposed to the publication on grounds of the protection of their private data;

Vifor Pharma

- 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.

Seeking the consent of Recipients

Vifor Pharma certifies that it has used all reasonable steps for the purpose of obtaining consent to individual disclosure where such consent is required by applicable law.

Date: 5 July 2016

Name of signatory: Oliver P. Kronenberg

Position in the Company: General Counsel Vifor Pharma

Signature:

Name of signatory: Beatrix Benz

Position in the Company: Head of Global Communications & Public Affairs

Signature: