

**DISCLOSURE OF TRANSFERS OF VALUE FROM PHARMACEUTICAL COMPANIES TO HEALTHCARE PROFESSIONALS AND HEALTHCARE ORGANISATIONS**

Publication Date: 30.06.2017

*Article 2 - Section 2.03*

	Full Name  <i>(Art. 1.01)</i>	HCPs: City of Principal Practice HCOs: city where registered  <i>(Art. 3)</i>	Country of Principal Practice  <i>(Schedule 1)</i>	Principal Practice Address  <i>(Art. 3)</i>	Unique country local identifier <b>OPTIONAL</b>  <i>(Art. 3)</i>	Donations and Grants to HCOs <i>(Art. 3.01.1.a)</i>	Contribution to costs of Events <i>(Art. 3.01.1.b &amp; 3.01.2.a)</i>			Fee for service and consultancy <i>(Art. 3.01.1.c &amp; 3.01.2.c)</i>		Transfers of Value for Research & Development as defined <i>(Art. 3.04)</i>	
							Sponsorship agreements with HCOs / third parties appointed by HCOs to manage an Event	Registration Fees	Travel & Accommodation	Fees	Related expenses agreed in the fee for service or consultancy contract		
<b>HCPs</b>	<i>INDIVIDUAL NAMED DISCLOSURE - one line per HCP (i.e. all transfers of value during a year for an individual HCP will be summed up; itemization should be available for the individual Recipient or public authorities' consultation only, as appropriate)</i>												
	<i>OTHER, NOT INCLUDED ABOVE - where information cannot be disclosed on an individual basis for legal reasons</i>												
	<b>Aggregate amount attributable to transfers of value to such Recipients - Art. 3.2</b>						N/A	N/A	0	0	4.150,00 €	413,00 €	N/A
	<b>Number of Recipients (named list, where appropriate) - Art. 3.2</b>						N/A	N/A	0	0	3	2	N/A
<b>% of total transfers of value to individual HCPs - Art. 3.2</b>						N/A	N/A	0%	0%	100%	100%	N/A	
<b>HCOs</b>	<i>INDIVIDUAL NAMED DISCLOSURE - one line per HCO (i.e. all transfers of value during a year for an individual HCO will be summed up; itemization should be available for the individual Recipient or public authorities' consultation only, as appropriate)</i>												
	<i>OTHER, NOT INCLUDED ABOVE - where information cannot be disclosed on an individual basis for legal reasons</i>												
	<b>Aggregate amount attributable to transfers of value to such Recipients - Art. 3.2</b>						0	0	0	0	0	0	N/A
	<b>Number of Recipients (named list, where appropriate) - Art. 3.2</b>						0	0	0	0	0	0	N/A
<b>% of total transfers of value to individual HCOs - Art. 3.2</b>						%	%	%	%	%	%	N/A	
<b>R&amp;D</b>	<b>AGGREGATE DISCLOSURE</b>											4.115.916,00 €	

Daiichi Sankyo Europe GmbH  
Methodology for Disclosure of Transfer of Value

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## **I. GENERAL**

### **1. Cross Border Payments**

All disclosures are made in the country in which the HCP practices or in which the HCO is located.

Daiichi Sankyo Europe GmbH is the European Headquarter of Daiichi Sankyo in Europe. As such we publish only transfer of value to HCOs and HCPs based in European countries other than those where Daiichi Sankyo has established affiliate organizations.

### **2. Currency**

Where payments were made in a currency other than Euro, the exchange rate will vary according to the date on which the conversion calculation was made. For general purposes, the conversion date should be regarded as the average monthly exchange rate when the event took place.

### **3. Added value tax**

All values provided in our report are net values, i.e. do not include value added tax.

### **4. Co-marketing projects**

Where Daiichi Sankyo jointly markets a product with another pharmaceutical company, Daiichi Sankyo will only declare those payments made directly from Daiichi Sankyo. Transfers of value made by its co-marketing partners will be disclosed separately by those organizations.

### **5. Reporting date**

Daiichi Sankyo will disclose the details of the payment on the date the transfer of value to HCPs/HCOs is actually made. This may mean that some projects taking place at the end of 2015 will be disclosed as part of the next annual reporting period because the payment may not occur until January when the invoice has been received and settled.

## 6. **Intermediaries**

### 6.1 Intermediaries acting on behalf of Daiichi Sankyo

All intermediary (third parties) that represent or act on behalf of Daiichi Sankyo, are subject to written contract and are obliged to provide Daiichi Sankyo with any contribution made to HCP or HCO. If this information cannot be provided due to the nature of the contribution (e.g. market research), it is in intermediaries responsibility to disclose the costs of contributions.

### 6.2 Intermediaries acting on behalf of HCO/HCP

Where the intermediary is a professional conference organizer (PCO), Daiichi Sankyo declares the Transfers of Value in the appropriate category in the name of the sponsored HCO/HCP.

### 6.3 Private companies and associated charities

The payment received by the contracting entity – which may be a HCP, a legal entity owned by a HCP (which is then a HCO) or a HCO – will be disclosed as a Transfer of Value made to that entity.

## **II. DATA PRIVACY**

### 1. **Informed consent**

Data Privacy law requires that Daiichi Sankyo obtain permission from individual HCPs prior to disclosing personal data such as individual transfers of value. Daiichi Sankyo has made every effort to secure and retain a record of the necessary permissions.

Where permission has not been obtained or where the individual HCP has refused permission, Daiichi Sankyo has declared the total spend as an aggregate figure within the relevant disclosure category.

### 2. **Partial consent**

Where only partial permissions has been granted to disclose transfer of value by an HCP, the entire transfer of value to this particular HCP is disclosed as aggregate

### **III. REPORTING CATEGORIES**

According to EFPIA Guidance and country local disclosure code

Travel and accommodation expenses related to advisory boards and other consulting meetings are published under “Fee for service and consultancy”

### **IV. RESEARCH AND DEVELOPMENT**

#### **1. Definition**

Transfers of Value to HCPs or HCOs related to the planning or conduct of (i) non-clinical studies (as defined in OECD Principles on Good Laboratory Practice); (ii) clinical trials (as defined in Directive 2001/20/EC); or (iii) non-interventional studies that are prospective in nature and that involve the collection of patient data from or on behalf of individual, or groups of, HCPs specifically for the study.

#### **2. Composition of R&D transfer of value**

The aggregate R&D transfer of value includes:

- Contribution to costs of Investigator Meetings and Committees
- Investigator fees for patient visits paid directly to clinical trial site staff or to CROs as an intermediary. Delayed or preliminary payments by CROs to clinical trial site staff are not considered

The aggregate R&D transfer of value does not include fees paid to Clinical Research Organizations (CROs)