



# ID Card

## Hexachloroplatinic acid

Version 22 September 2016

**Notes:**

- This ID card is used to support the substance sameness discussions in SIEFs and to describe the substance to the best of the SIEF members' knowledge.
- It also aims at grouping communications relevant to the request of available data or information, the approval of the proposed Lead Registrant and the registration strategy with the SIEF.
- It is the responsibility of each individual registrant to identify their substance and to report company-specific identity in their Registration Dossier (section 1 of IUCLID).

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### 1. Identification of the substance

**Table 1. Identification of the substance**

	Original (in EC inventory)
Name	Hexachloroplatinic acid
EC number	241-010-7
CAS number	16941-12-1
Description	Not available
Composition type	Mono-constituent substance

### 2. Synonyms and other identifiers of the substance

**Table 2. Synonyms and other identifiers of the substance**

IUPAC name	Dihydrogen hexachloroplatinate(2-)
CAS name	
Abbreviations	
Other commercial, brand or international names	Chloroplatinic acid Chloroplatinic(IV) acid Platinic chloride Speier's catalyst
Other identity codes	PubChem ID: 61859



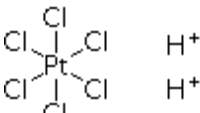
### 3. Substances (with core identifiers) also falling under this substance (with justification)

Table 3. Substances also falling under this substance

Name	EC number	CAS number	Justification
Hexachloroplatinic acid hydrate		26023-84-7 18497-13-7	According to Annex V(6) of the REACH Regulation, hydrates of a substance are exempted from Registration provided that the anhydrous form has been registered by the manufacturer or importer using this exemption.

### 4. Information related to molecular and structural formula of the substance

Table 4. Information related to molecular and structural formula of the substance

Molecular formula	Cl <sub>6</sub> Pt.2H (anhydrous basis)
Structural formula	
Smiles notation	[H+].[H+].Cl[Pt-2](Cl)(Cl)(Cl)(Cl)Cl
Optical activity	Not applicable
Typical ratio of (stereo) isomers	
Molecular Weight / Molecular Weight range	409,81 g/mol (anhydrous basis)

### 5. Typical composition of the substance

Hexachloroplatinic acid can be placed on the market in anhydrous and hydrated form. All forms of Hexachloroplatinic acid will be addressed in the same Registration Dossier but are reported individually in IUCLID section 1.2.

#### 5.1. Hexachloroplatinic acid (anhydrous)

Table 5. Typical composition

	Name	Symbol / Formula	Min & Max concentrations (%) <sup>§</sup>	Typical concentration (%) <sup>§§</sup>
Main constituent(s)*	Hexachloroplatinic acid	H <sub>2</sub> Cl <sub>6</sub> Pt	99 - 100	> 99
Impurity(ies) <sup>#</sup>	Several minor (especially metallic) impurities which do not affect the classification of the substance because of their non-hazardous nature or because they do not exceed the classification cut-off limits in the substance	e.g. Ag, Au, Cu, Ir, Pb, Pd, Ru	0 - 1	< 1

\* ≥ 80 % (w/w) for mono-constituent substances; ≥ 10 % (w/w) and < 80 % (w/w) for multi-constituent substances.



# An impurity is an unintended constituent present in a substance, as produced. It may originate from the starting materials or be the result of secondary or incomplete reactions during the production process. While impurities are present in the final substance, they were not intentionally added.

§ Concentration ranges define the substance sameness criteria agreed by all EPMF Members in preparation of the communication with other SIEF members.

§§ Typical concentration refers to the representative sample used for testing.

The composition given above is typical and should therefore represent the majority of Hexachloroplatinic acid (anhydrous) as manufactured and/or imported in the EEA market.

## 5.2. Hexachloroplatinic acid hydrate

**Table 6. Typical composition**

	Name	Symbol / Formula	Min & Max concentrations (%) <sup>§</sup>	Typical concentration (%) <sup>§§</sup>
<b>Main constituent(s)*</b>	Hexachloroplatinic acid hydrate	H <sub>2</sub> Cl <sub>6</sub> Pt.n H <sub>2</sub> O	99 - 100	> 99
<b>Impurity(ies)#</b>	Several minor (especially metallic) impurities which do not affect the classification of the substance because of their non-hazardous nature or because they do not exceed the classification cut-off limits in the substance	e.g. Ag, Au, Cu, Ir, Pb, Pd, Ru	0 – 1	< 1

\* ≥ 80 % (w/w) for mono-constituent substances; ≥ 10 % (w/w) and < 80 % (w/w) for multi-constituent substances.

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§§ Typical concentration refers to the representative sample used for testing.

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## 6. Information on appearance, physical state and properties of the substance

**Table 7. Appearance / physical state / properties of the solid substance**

<b>Physical state</b>	Solid
<b>Physical form*</b>	Amorphous
<b>Appearance</b>	Brownish-red/orange solid
<b>Particle size**</b>	Fine powder / Coarse powder / Massive object
<b>Does the solid hydrolyse?#</b>	No
<b>Is the solid hygroscopic?§</b>	Yes

\* Crystalline form: solid material whose constituent atoms, molecules, or ions are arranged in an ordered pattern extending in all three spatial dimensions. Amorphous form: solid material whose constituent atoms, molecules, or ions are randomly arranged.

\*\* Nanoform: particles in the size range 1 - 100 nm (for full definition of a nanomaterial, see <http://ec.europa.eu/environment/chemicals/nanotech/index.htm#definition>). Fine powder: particles in the size range 100 – 2.500 nm. Coarse powder: particles in the size range 2.500 nm – 1 mm. Massive object: particles in the size range > 1 mm.

# Hydrolysis: decomposition (cleavage of chemical bonds) by the addition of water.

§ Hygroscopic substance: readily attracts moisture from its surroundings in open air, through either absorption or adsorption. Cf. also water/moisture content in Table 5.



## 7. Analytical data

Annex VI of REACH requires the registrant to describe the analytical methods and/or to provide the bibliographical references for the methods used for identification of the substance and, where appropriate, for the identification of impurities and additives. This information should be sufficient to allow the methods to be reproduced.

**Table 8. Analytical methods for identification of the substance**

Parameter / Method	Recommended for substance identification and sameness check	Applicable	Not applicable or not recommended
<b>Elemental analysis</b>			
ICP (ICP-MS or ICP-OES)	X		
Atomic absorption spectroscopy (AAS)			
Glow discharge mass spectrometry (GDMS)			
<b>Molecular analysis</b>			
Infrared (IR) spectroscopy	X		
Raman spectroscopy			
<b>Mineralogical analysis</b>			
X-Ray Fluorescence (XRF)		X	
X-Ray Diffraction (XRD)	X		
<b>Morphology and particle sizing</b>			
Electron microscopy (SEM, TEM, REM)* #			
Laser diffraction* #	X		
Particle size by other means (e.g. sieve analysis)#			
Surface area by N-BET* #	X		
<b>Other</b>			

\* Analytical techniques particularly (but not exclusively) relevant for nanomaterials.

# The choice of the technique for particle size depends on the size of the material as manufactured/imported/placed on the market/used.

## 8. Lead Registrant

Johnson Matthey (United Kingdom) volunteers to be the Lead Registrant for Hexachloroplatinic acid. The EPMF will provide support to the Lead Registrant as laid down in the EPMF Agreement.



## 9. REACH Strategy

The table below presents the overall Registration Strategy for Hexachloroplatinic acid based on the information available to the EPMF by the date given above on the document.

The Registration Dossier will be prepared for the highest substance status (information requirements associated to a substance or Article 10 Registration being higher than an intermediate handled under strictly controlled conditions or Article 17 or 18 one) and associated tonnage band.

The recap below therefore reflects the scope of work of the EPMF for Hexachloroplatinic acid and sets the minimum and maximum set of information that will be gathered and/or produced when preparing the Registration Dossier for Hexachloroplatinic acid as described in this ID Card.

If higher information requirements are necessary, these can be included in the Registration dossier (if EPMF is made aware of these additional requirements in-time) as an update to the already submitted dossier.

**Table 9. REACH strategy for the substance (basis for REACH Registration preparation)**

Item	Description
<b>REACH category</b>	Mono-constituent substance
<b>Highest status</b>	Substance
<b>Highest tonnage band</b>	10-100 t/a
<b>Information requirements</b>	Available / Existing + Annex VII + Annex VIII
<b>Existing classification*</b>	Met. Corr. 1 (H290) Acute tox. 2 (H300) (oral) Skin corr. 1B (H314) Skin sens. 1B (H317) Eye dam. 1 (H318) Resp. Sens. 1A (H334) STOT RE1 (H372) Aquatic acute 1 (H400) Aquatic chronic 1 (H410) Acute M-factor 10 Chronic M-factor 10
<b>Registration deadline</b>	2018

\* For the pure form, as in the REACH registration dossier

## 10. Scope of the Registration Dossier

The uses included in this Registration Dossier are listed on the [EPMF website](#).

## 11. Analytical reference information

Below the results of Raman analysis of a reference sample (solution).

- Instrument: Renishaw inVia
- Measurement head: Confocal microscope
- Objective: x50 L
- Laser: 785 nm
- Grating: 1200 lines/mm
- Laser power: 1 %
- Resolution: 1 cm<sup>-1</sup>
- Sample scans: 1
- Scan time: 10 s



- Scan range: 50 - 3200  $\text{cm}^{-1}$

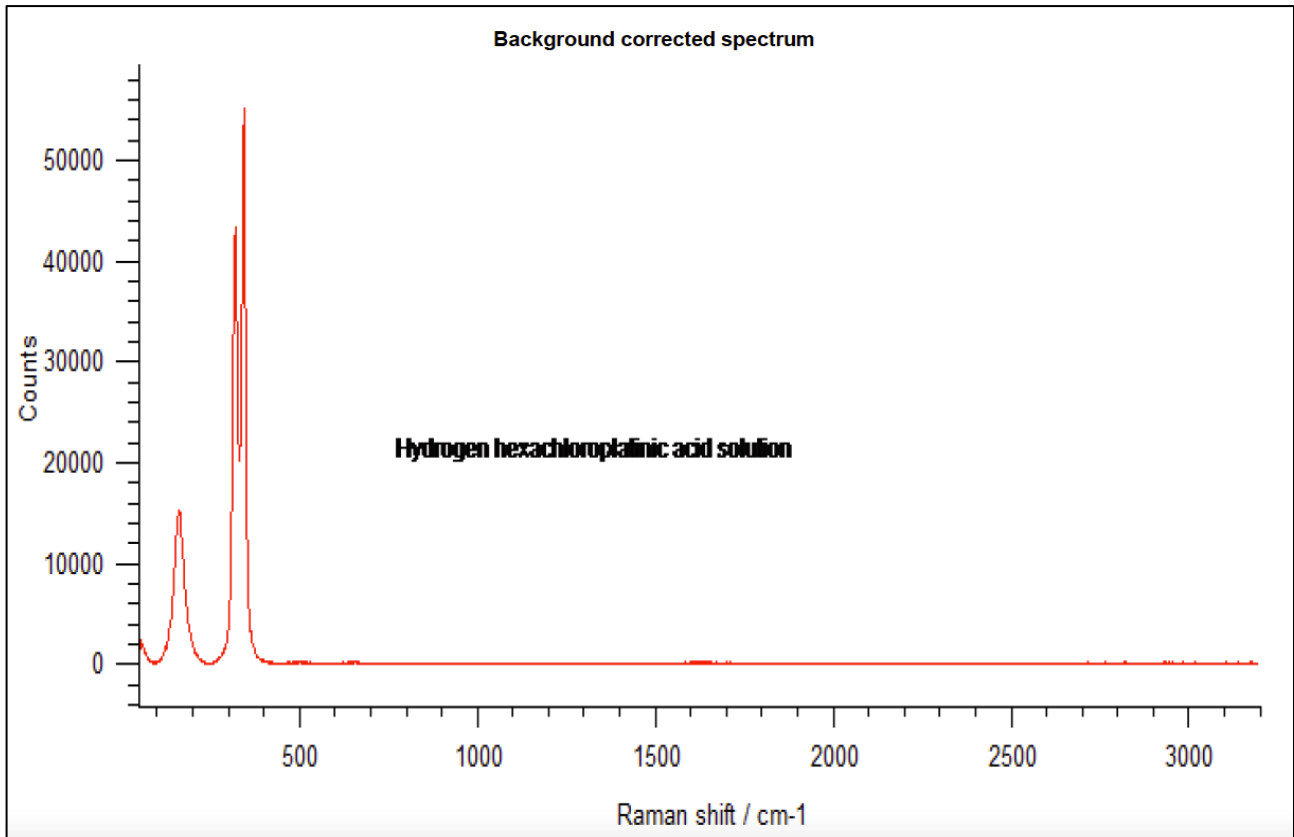


Figure 1. Raman spectrum of Hexachloroplatinic acid solution