



Heraklion, 25/02/2017

Instructions

Dear Customer and Downstream User,

Plastika Kritis S.A., in order to facilitate your necessary actions for using the KRITILEN® masterbatches containing DCC's PY.34 and PR.104, provides the following Annexes:

Annex II: DCC's REACH Authorization for C.I. Pigment Yellow 34 and C.I. Pigment Red 104: For User of Masterbatches (Downstream Users) based on DCC pigments (PY.34 and PR.104) Checklist

Annex III: DCC's REACH Authorization Application for C.I. Pigment Yellow 34 and C.I. Pigment Red 104

Annex IV: USE 5: Verification of DCC PY.34 and PR.104 RMM and OC. Summary of Blood lead, Air Monitoring and RPE

Annex IV can be downloaded in a convenient format for easiness of use at our web site at <http://www.plastikakritis.com/en/kritilen-downloads>.

For any questions please contact your sales person or the undersigned.

Yours faithfully,

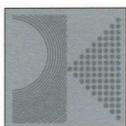
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ANNEX II:

DCC's REACH Authorization for C.I. Pigment Yellow 34 and C.I. Pigment Red 104: For User of Masterbatches (Downstream Users) based on DCC pigments (PY.34 and PR.104) Checklist

| Action | Comments | Dates and Deadlines |
|--|---|--|
| 1) Obtain latest Safety Data Sheet (SDS) for DCC PY.34 & PR.104 and PLASTIKA KRITIS relevant masterbatches | Contact DCC representative and/or visit DCC website/ Ask a Plastika Kritis salesperson | As soon possible |
| 2) Update SDS and labels with Authorization numbers | | As soon possible |
| 3) Obtain latest Extended Safety Data Sheets (eSDS) for PY.34 and PR.104 | Contact DCC: Ms Hella Schulpen (REACH Only Representative) at hschulpen@dominioncolour.nl | As soon possible |
| 4) Identify use to the supplier/distributor to ensure all relevant exposure scenarios are taken into account | | As soon possible |
| 5) Maintain a personal protective equipment program on the selection / use / maintenance / training (including respiratory protection equipment) | | As soon possible |
| 6) Notify your use to ECHA | DCC REACH IT guidance is available | Within three (3) months of first delivery after Official Publication (14 th September 2016) |
| 7) Verify the risk management measures and operational conditions are in line with those outlined in DCC's eSDS of PY.34 & PR.104 | Use DCC's PY.34 & PR.104 Verification/Notification template Information submitted to ECHA will be provided to DCC. However, DCC requires this information in order to prepare updated reports to ECHA/EC by 2017 and 2020. It is recommended to provide this information to DCC as soon as possible | Submission to ECHA via the Use Notification webform (see DCC REACH IT guidance) *Submit completed forms every March (i.e. 2016 data to be provided by March 2017) to DCC- Ms Hella Schulpen (REACH Only Representative) at hschulpen@dominioncolour.nl |
| 8) Provide ECHA with: a) Blood lead data b) Chromium in air data Surveillance record for each worker exposed to DCC PY.34 & PR.104 (if the exposure data is greater than the limits set by EC) | | |
| 9) Notify the supplier/DCC if risk management measures assigned in the eSDS are not appropriate and/or new information on hazardous properties is available | | As soon as possible |
| 10) Prepare Assessment of Alternatives by 30th June 2017: a) Define end use of PY.34 & PR.104 b) Identify the alternatives associated with <i>each</i> end use c) Explain why the alternatives are not a sufficient replacement (technical & economic considerations) | Use DCC Assessment of End Use and Alternates template Information submitted to ECHA will be provided to DCC. However, DCC requires this information to prepare its updated report to the EC by 31 December 2017. It is recommended to provide this information to DCC as soon as possible | Submit to ECHA by 30 June 2017 via the Use Notification webform (see DCC REACH IT guidance) **Submit completed forms as soon as possible to DCC – Ms Lucy Gibbons (Field Chemist) at lgibbons@dominioncolour.com |



ANNEX III:

DCC's REACH Authorisation Application for C.I. Pigment Yellow 34 and C.I. Pigment Red 104

ECHA notification of your uses covered under DCC's Authorisation

Dear Customer and Downstream User,

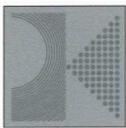
Following the official approval of DCC's Authorisation request for the continued use of PY.34 and PR.104 in the EEA¹ as communicated on 27 September 2016, as our valued customer we would like to draw your attention to the notification obligation to ECHA described under REACH Article 66.

According to Article 66, all downstream users of an authorised substance (such as PY.34 and PR.104) must notify ECHA within **three months** of the first delivery of an authorised substance following the publication in the Official Journal of the European Union (OJ). In its "News Alert"² of 16 September 2015 ECHA already made reference to this notification requirement for downstream users. This notification is free from any charge.

DCC's Authorisation was published on the Official Journal of the European Union on September 14th 2016. As such all DCC PY.34 and PR.104 received after this date must be notified to ECHA within three months of its receipt.

As downstream users may have stock of DCC PY.34/PR.104, we would advise you to act pro-actively by submitting your notification to ECHA as soon as possible.

In the following sections, we will provide you with some background information and a short practical guidance on how to submit your notification to ECHA.



What are your obligations as a Downstream User of an Authorised substance

a. *Communication down the supply chain*

DCC's REACH communication letter of 27 September 2016 summarised the assigned Authorisation numbers, one for each use. These authorisation numbers will be disseminated on DCC's SDS's and bag labels.

As a downstream user, these Authorisation numbers must be communicated further down the supply chain by incorporating them on your SDS's and (bag) labels as well.

b. *Notification to ECHA*

Besides the above supply chain communication, you also have to notify ECHA of your intended use of DCC PY.34 and PR.104.

Notification needs to be done via the REACH-IT³ portal. This notification requires specific details on:

- the identity of your company,
- the authorisation number(s) which is (are) linked to your specific use(s) and
- contact information.

Note that information about your typical annual volume and the number of staff using the substance is not mandatory. However, this information can be submitted on a voluntary basis.

How to submit your information to ECHA

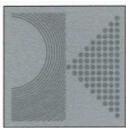
Before starting the process you must have an active REACH-IT account to be able to submit your notification. In the following steps we will guide you through the process step by step:

Step 1: Create a REACH-IT account

In case you already have an active REACH-IT account, go to step 2.

- Follow the link <https://reach-it.echa.europa.eu/reach/public/welcome.faces>
- Click 'Register a company'





- Complete the form and click on 'Save'

Create Account

User Information

* First Name:

* Last Name:

* Email:

* Username:

Phone:

Password

* Password:

The password must have at least 8 characters and contain three of the following character types: uppercase letter, lowercase letter, number and non-alphabetical. The password must not contain your username, your first name or your last name. The password cannot be the same as a previously used password. The password cannot be changed more than once a day.

* Confirm Password:

Security

* Security Question:

* Security Answer:

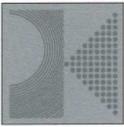
- You will receive an email at the indicated email address.



Follow the link in your email or copy it in your browser and your email address will be verified.

- When completed, you receive the message below. Click on 'login' to continue with the process





- Login with your personal defined Username and Password

ECHA
EUROPEAN CHEMICALS AGENCY

Username

Password

[LOGIN](#)

Help

- > SIC6 accounts
- > Forgot your password?
- > Forgot your username?
- > CHANGE password

Not a user yet?

- > Sign up

- Click on '*Account, Users & Parties management*' to complete your company details

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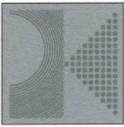
Please select one of the following Applications

In order to finalize the sign-up process (e.g. by entering your Company details and adding contacts) or to subsequently manage your Company's users and details, please select

- > **Account, Users & Parties management**

If you have successfully completed all the steps for signing up your Company and you want to log into ECHA applications, please select one of the options below

- > SIC6/IT submission tool
- > SIC6
- > IC submission tool



- For the purpose of this notification process, click 'Encode Legal Entity'

ECHA
EUROPEAN CHEMICALS AGENCY

Create Legal Entity

In order to use ECHA's submission systems you need to provide your Legal Entity's details, correct information, users, etc. Please continue by selecting either encode legal entity or import EICIS format file below.

Import EICIS format file: If you already have a legal entity definition file in EICIS exchange format (also known as the EICIS file format), and if you want to copy this same EICIS file your legal entity definition as for your EICIS EICIS, you should import your EICIS. Upon successful validation of the imported file, you will be redirected to the legal entity profile screen to review / update / import details as well as to enter additional details.

Encode Legal Entity: If you do not have an EICIS format file or do not want to update, you may encode your legal entity information.

[Import EICIS Data](#) [Encode Legal Entity](#)

- Complete the form. Billing information is required, but this is just a formality. There is no fee charged for this notification.

Add identification

Please provide the required details. Fields marked with an asterisk (*) are mandatory. Having user's ITI main status helps information. Note: you will not be able to modify "country" after pressing the "Save" button.

General details

* Legal entity name ❗ This field is mandatory and must be filled.

Country

VAT number

Remarks

General Contact information

* Phone

* Email (Example: name@domain.com)

Website

Fax

Address information

* Street

Street 2

* Postal code

* City

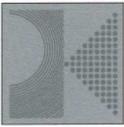
Region/Country

* Country Select...

Billing address

Same information as per Legal Entity's name and address

Not the same



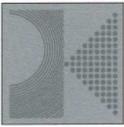
- After saving, your Legal Entity creation under REACH-IT is complete and you can close the website.

Step 2: Log in at the ECHA Notification webform

- Follow the link <https://echa.europa.eu/support/dossier-submission-tools/reach-it/downstream-user-authorized-use>
- Click 'Downstream user notification webform' at the bottom of the page

The screenshot shows the ECHA website interface. The main heading is "Submitting downstream user notification of authorised uses". Below the heading, there are several sections of text providing instructions and information. At the bottom of the page, a button labeled "Downstream user notification webform" is highlighted with a red circle. The page also features a navigation menu on the left, a search bar at the top, and social media icons on the right.

- Log in with your personal defined Username and password



Step 3: Complete the ECHA webforms

- Form 1 ('Notification type'): tick the box 'This is a new notification' ; tick 'Next'

The screenshot shows the 'Downstream User Notification of Authorised Uses' webform. The 'Notification type' section is visible, with a red circle around the 'This is a new notification' checkbox. Below this, there are several other checkboxes for 'Authorised uses' which are currently unchecked. The ECHA logo is visible in the bottom right corner.

- Form 2 ('Authorised uses'): tick those box(es) relevant to your specific use(s). Note there are separate Authorisation numbers for PY.34 and PR.104. For details on the assigned Authorisation numbers, see DCC's REACH communication letter of 27 September 2016. Information on site volume and staff is not mandatory.

The screenshot shows the 'Downstream User Notification of Authorised Uses' webform, specifically the 'Authorised uses' section. A red circle highlights this section, which contains a list of various uses with corresponding checkboxes. The ECHA logo is visible in the bottom right corner.



- Form 3 ('Sites'): check the information and complete where needed. To re-confirm, tick again the boxes identified for your specific use(s). Tick 'Next'.

The screenshot shows the 'Downstream User Notification of Authorized Uses' form on the ECHA website. The form is divided into several sections, including 'User Information' and 'Uses'. A red oval highlights the 'Uses' section, which contains a list of authorized uses with checkboxes for selection. The ECHA logo is visible in the bottom right corner.

- Form 4 ('Send'): tick the declaration box and send. You will receive a confirmation number and a copy of your submission for your records.

The screenshot shows the 'Downstream User Notification of Authorized Uses' form on the ECHA website, specifically the 'Send' section. The form contains a declaration box with a red oval around it, and a green 'Send Notification' button. The ECHA logo is visible in the bottom right corner.



What happens next

As far as known, ECHA will not share these use notification details with DCC. However you are encouraged to keep us informed as this could help DCC when reviewing the Authorisation dossier.

This DCC guideline describes the process of notifying your use to ECHA. Following the official publication in the Official Journal, each downstream user also has an obligation to provide monitoring data to ECHA (blood Pb values / air measurements of Cr) as well as a justification of use report by the end of June 2017. ECHA requests these documents to be attached to this use notification webform. To add these documents:

- login again to the ECHA webform as described under Step 2 via <https://echa.europa.eu/support/dossier-submission-tools/reach-it/downstream-user-authorized-use>
- In Form 1 ('Notification type') tick 'This is an update to an earlier notification' and include the earlier received notification number.

The screenshot shows the ECHA webform interface. The title is 'Downstream User Notification of Authorized Users'. Below the title, there are several sections. The 'Notification type' section is highlighted with a red circle, and it contains a checkbox labeled 'This is an update to an earlier notification'. Other sections include 'Product name', 'CAS number', and 'ECHA registration number'. The ECHA logo is visible in the bottom right corner.

- Attach the documents in Form 4 ('Send') by browsing your computer and send the notification

The screenshot shows the ECHA webform interface. The title is 'Downstream User Notification of Authorized Users'. Below the title, there are several sections. The 'Send' section is highlighted with a red circle, and it contains a green button labeled 'Send'. Other sections include 'Justification of use', 'Monitoring data', and 'Additional information'. The ECHA logo is visible in the bottom right corner.



Further background information on the general requirements as a downstream user under REACH can be found in the ECHA Guidance:

http://echa.europa.eu/documents/10162/13634/du_en.pdf



ANNEX IV:

USE 5: Verification of DCC PY.34 and PR.104 RMM and OC Summary of Blood lead, Air Monitoring and RPE

Dear Customer and Down Stream User,

We are pleased to inform you that DCC has successfully applied for the continued use of Pigment Yellow 34 and Pigment Red 104. In order to ensure safe handling of these pigments, the European Commission has included the assessment of biomonitoring data, air monitoring data and RPE effectiveness by the Authorisation holder.

As per Article 3(e) of the final EC decision, C(2016) 5644 of 07.09.2016, all customers and downstream users are obligated to provide blood lead information and air monitoring results for chromium to ECHA. DCC Maastricht B.V. OR will receive data from ECHA as per Article 3(f) for the purpose of preparing a review reports.

In order to complete your notification to ECHA, please refer to Table 1 which can be found in each eSDS to identify the worker functions and the activities that are covered per use under DCC's Authorisation. Table 1 indicates the specific activities that a worker is permitted to perform under the specified technical and organisational conditions. The table below summarises the uses and the assigned Authorisation numbers for DCC PY.34 and PR.104.

| DCC eSDS (Use) | Description of USE for Paint Sector | DCC PY.34 Authorisation Numbers: | DCC PR.104 Authorisation Numbers: |
|----------------|--|----------------------------------|-----------------------------------|
| Use 1 | Use: distribution and mixing of pigment powder in an industrial environment into solvent-based paints for non-consumer use | REACH/16/3/0 | REACH/16/3/6 |
| Use 2 | Use: industrial application of paints on metal surfaces (such as machines vehicles, structures, signs, road furniture, coil coating, etc.) | REACH/16/3/1 | REACH/16/3/7 |
| Use 3 | Use: professional, non-consumer application of paints on metal surfaces (such as machines, vehicles, structures, signs, road furniture, etc.) or as road marking | REACH/16/3/2 | REACH/16/3/8 |

| DCC eSDS (Use) | Description of USE for Plastic Sector | DCC PY.34 Authorisation Numbers: | DCC PR.104 Authorisation Numbers: |
|----------------|--|----------------------------------|-----------------------------------|
| Use 4 | Use: distribution and mixing pigment powder in an industrial environment into liquid or solid premix to colour plastic/plasticised articles for non-consumer use | REACH/16/3/3 | REACH/16/3/9 |
| Use 5 | Use: industrial use of solid or liquid colour premixes and pre-compounds containing pigment to colour plastic or plasticised articles for non-consumer use | REACH/16/3/4 | REACH/16/3/10 |
| Use 6 | Use: professional use of solid or liquid colour premixes and pre-compounds containing pigment in the application of hot melt road marking | REACH/16/3/5 | REACH/16/3/11 |

DCC recommends using the attached form **for each employee** involved with DCC PY.34 and/or PR.104 to verify and monitor worker exposures. This form can be submitted to ECHA and to DCC (hschulpen@dominioncolour.nl) by March, 31st of each year, summarising the results of the previous year.

If you have any questions regarding the use of DCC PY.34 and/or PR.104, please feel free to contact your DCC representative.



I. General Information:

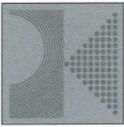
| | | | |
|---|--|------------------|--|
| Company Name: | | Contact Name: | |
| Address: | | Contact Phone #: | |
| Postal Code | | Contact e-mail: | |
| City, Country: | | Date: | |
| # Workers handling/exposed to PY.34 and/or PR.104 | | | |
| Annual volume PY.34/PR.104 used: | | | |
| Comments: | | | |

II. Company specific: Use and Legal Requirements

| REACH Authorisation Number: Tick the boxes that apply to your company | % PY.34/PR.104 in product (ranges are acceptable, but within $\pm 10\%$ of the contents) | Member State Regulation related to Health Surveillance | Health Surveillance | | Air Monitoring for Chromium |
|---|--|---|--|--|---|
| | | | Action Level ¹ | Suspension Level ² | Comments: |
| PY.34 <input type="checkbox"/> USE 1: REACH/16/3/0 <input type="checkbox"/> USE 2: REACH/16/3/1 <input type="checkbox"/> USE 3: REACH/16/3/2 <input type="checkbox"/> USE 4: REACH/16/3/3 <input type="checkbox"/> USE 5: REACH/16/3/4 <input type="checkbox"/> USE 6: REACH/16/3/5 | | <input type="checkbox"/> Directive 98/24/EC <input type="checkbox"/> Directive 2004/37/EC <input type="checkbox"/> Other. | <input type="checkbox"/> 40 μg lead/100 ml blood <input type="checkbox"/> Other. | <input type="checkbox"/> 70 μg lead/100 ml blood <input type="checkbox"/> Other. | Please provide Member State Regulation for Air Monitoring: Please provide Sampling Procedure for air monitoring: |
| PR.104 <input type="checkbox"/> USE 1: REACH/16/3/6 <input type="checkbox"/> USE 2: REACH/16/3/7 <input type="checkbox"/> USE 3: REACH/16/3/8 <input type="checkbox"/> USE 4: REACH/16/3/9 <input type="checkbox"/> USE 5: REACH/16/3/10 <input type="checkbox"/> USE 6: REACH/16/3/11 | | If other apply, please specify: | If other apply, please specify: | If other apply, please specify: | Please provide analytical method for air monitoring: |
| | | | Please provide Sampling test Method for Blood Lead: | | |

¹ Action level: triggers employer to carry out investigation, review control measures and take steps to reduce employee's blood lead concentration below the action level so far as reasonably practicable

² Suspension level: concentration at which employees are normally taken off work which exposes them to lead. Return to work/modification of work exposure is based on Doctor's recommendations



III. Worker / Employee specific

Table 1. Overview of tasks per function for Use 5 (ES 7, 9)

| Company: | | | Employee (internal) Id ³ : | | | | Approximate volume handled by employee over last 6-12 months: | | | Approximate volume handled by employee/per day: | | | | | |
|---|----------------------------------|--|--|--|------------------------------------|------------------------|---|-----------|-----------------|---|----------------------|--------------|---------------------------|-----------------------------------|--------------|
| Comments: | | | <input type="checkbox"/> Smoker smoker <input type="checkbox"/> Non-smoker | | | | | | | | | | | | |
| Description of Worker Functions and Tasks | | | Summary of RMM for USE 5 | | | | Verification of RMM | | | Results of Monitoring Programs | | | | | |
| Function | Workplace instruction card (WIC) | Tasks (description of Contributing Scenario) | Technical RMM | | | Organizational RMM | RPE | Site RMM | Max. hours/week | Type RPE used | Blood lead Testing | | Personal Air Monitoring | | |
| | | | A. Containment of source | B. Personal enclosure (PE) / segregation (SEG) | C. Local exhaust ventilation (LEV) | Maximum hours per week | Minimum Respiratory | A, B or C | hours | APF | µg lead/100 ml blood | Date of test | pigment µg/m ³ | Cr µg/m ³ ⁵ | Date of Test |
| Operator | WIC30 | <input type="checkbox"/> Charging/discharging of coloured plastic granules | | | Y | 12 | - | | | | | | | | |
| | | <input type="checkbox"/> Mixing coloured plastic granules in closed mixing vessel | Y | | | 9 | - | | | | | | | | |
| | | <input type="checkbox"/> Production of plastic articles by extrusion and injection moulding or other processes | Y | Y (SEG) | | 21 | - | | | | | | | | |
| | | Total | | | | 40 ⁶ | - | | | | | | | | |
| General worker # 1 | WIC31 | <input type="checkbox"/> Delivery, storage and handling of coloured plastic granules | | | | | | | | | | | | | |

³ Only list non-confidential details. Personal data to be maintained by the customer/downstream user.

⁴ Level of respiratory protection that can realistically be expected to be achieved in the workplace by 95% of adequately trained and supervised wearers using a properly functioning and correctly fitted respiratory protective device.

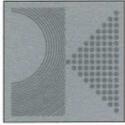
⁵ The exposure related to chromium can be calculated based on a general 15% Cr-content of PY.34/PR.104

⁶ The limit value for the two pigments is based on an exposure duration of 40 hours per working week. Therefore the maximum duration for any function is 40 even though for this function the summed duration per task can be over 40 hours



| Company: | | | Employee (internal) Id ⁵ : | | | | Approximate volume handled by employee over last 6-12 months: | | | | | | | | |
|---|----------------------------------|--|---|--|------------------------------------|------------------------|---|-----------|-----------------|--------------------------------|----------------------|--------------|---------------------------|----------------------|--------------|
| Comments: | | | <input type="checkbox"/> Smoker <input type="checkbox"/> Non-smoker | | | | Approximate volume handled by employee/per day: | | | | | | | | |
| Description of Worker Functions and Tasks | | | Summary of RMM for USE 5 | | | | Verification of RMM | | | Results of Monitoring Programs | | | | | |
| Function | Workplace instruction card (WIC) | Tasks (description of Contributing Scenario) | Technical RMM | | | Organizational RMM | RPE | Site RMM | Max. hours/week | Type RPE used | Blood lead Testing | | Personal Air Monitoring | | |
| | | | A. Containment of source | B. Personal enclosure (PE) segregation (SEG) | C. Local exhaust ventilation (LEV) | Maximum hours per week | Minimum Respiratory protective equipment (APF ⁶) | A, B or C | hours | APF | µg lead/100 ml blood | Date of test | pigment µg/m ³ | Cr µg/m ³ | Date of Test |
| | | <input type="checkbox"/> Delivery, storage and handling of packed plastic premix or pre-compound | Y | | | 10 | - | | | | | | | | |
| | | <input type="checkbox"/> Transfer of articles and single coloured granules (non-dedicated facility) or | Y | - | | 15 | 10 | | | | | | | | |
| | | <input type="checkbox"/> Transfer of articles and single coloured granules (dedicated facility) | Y | Y (SEG) | | 12 | - | | | | | | | | |
| | | <input type="checkbox"/> Charging/discharging of coloured plastic premix or pre-compound or | Y | | | 6 | 10 | | | | | | | | |
| | | <input type="checkbox"/> Transfer of mixed coloured granules and articles | Y | | | 12 | - | | | | | | | | |
| | | Total | | | | 38⁷ | | | | | | | | | |

⁷ The limit value for the two pigments is based on an exposure duration of 40 hours per working week. Therefore the maximum duration for any function is 40 even though for this function the summed duration per task can be over 40 hours, which exposes them to lead. Return to work/modification of work exposure is based on Doctor's recommendations



| Company: | | | Employee (internal) Id ⁷ : | | | | Approximate volume handled by employee over last 6-12 months: | | | | | | | | |
|---|----------------------------------|---|---|--|------------------------------------|------------------------|---|-----------|------------------|--------------------------------|----------------------|--------------|---------------------------|----------------------|--------------|
| Comments: | | | <input type="checkbox"/> Smoker <input type="checkbox"/> Non-smoker | | | | Approximate volume handled by employee per day: | | | | | | | | |
| Description of Worker Functions and Tasks | | | Summary of RMM for USE 5 | | | | Verification of RMM | | | Results of Monitoring Programs | | | | | |
| Function | Workplace instruction card (WIC) | Tasks (description of Contributing Scenario) | Technical RMM | | | Organizational RMM | RPE | Site RMM | Max. hours /week | Type RPE used | Blood lead Testing | | Personal Air Monitoring | | |
| | | | A. Containment of source | B. Personal enclosure (PE) / segregation (SEG) | C. Local exhaust ventilation (LEV) | Maximum hours per week | Minimum Respiratory protective | A, B or C | hours | APF | µg lead/100 ml blood | Date of test | pigment µg/m ³ | Cr µg/m ³ | Date of Test |
| QC | WIC32 | <input type="checkbox"/> Transfer of mixed coloured granules and articles | | | | 12 | 10 | | | | | | | | |
| | | <input type="checkbox"/> Charging/discharging of coloured plastic premix or pre-compound (non-dedicated facility) or | Y | - | - | 6 | 10 | | | | | | | | |
| | | <input type="checkbox"/> Charging/discharging of coloured plastic premix or pre-compound (dedicated filling line) | - | Y (SEG) | Y | 6 | - | | | | | | | | |
| | | <input type="checkbox"/> Roll application and heat curing of coloured plastic paste | | | | 18 | - | | | | | | | | |
| | | Total | | | | 36 | | | | | | | | | |
| General worker # 2 | WIC33 | <input type="checkbox"/> Handling and manipulation of pigment plastic articles and plastic coated textiles | | | | 40 | - | | | | | | | | |
| Maintenance worker in industrial setting | WIC34 | <input type="checkbox"/> Handling and manipulation of pigment plastic articles, plastic coated textiles and coloured road marking | | | | 12 | 10 | | | | | | | | |
| | | <input type="checkbox"/> High energy manipulation of pigment plastic articles, plastic coated textiles and coloured road marking using abrasive techniques like mechanical cutting, grinding, drilling or sanding | | | Y | 12 | 100 | | | | | | | | |
| | | Total | | | | 24 | | | | | | | | | |

⁷ Only list non-confidential details. Personal data to be maintained by the customer/downstream user.

⁸ Level of respiratory protection that can realistically be expected to be achieved in the workplace by 95% of adequately trained and supervised wearers using a properly functioning and correctly fitted respiratory protective device.

⁹ The exposure related to chromium can be calculated based on a general 15% Cr-content of PY.34/PR.104

