

Due to increasing restrictions and the regulation of substances in the EU and on the global market, the iPCA system offers options for detecting critical and restricted substances in the MDS database and identifying which components are affected. This means that these important results can be shared 'internally' within your company and the next steps discussed. To fulfil all the legal requirements and continue to ensure the conformity of your products, it is necessary to know how to analyse your database using the where-used list and the MDS analysis in iPCA, and to identify affected substances in your MDS database at an early stage. The necessary measures can then be initiated.

>> OBJECTIVE

You are familiar with the possibilities that the MDs analysis in iPCA offers you at the level of classification, materials, basic substances and recycle information, and you can export these to your company for presentation and evaluation. You understand the difference between the interactive analysis in iPCA in the where-used list function and the planning and setting up of an analysis job. You can confidently establish both procedures and the associated processes in your company. You know how to export and prepare the obtained results for reporting. You can independently configure and operate the influential inspector CSI and its wizard to plan and perform rule-based inspections in your account, and you know how to interpret and further process the results.

Your benefit: Practical exercises will help you apply what you've learned right away.

>> TOPICS COVERED

Introduction and contents

MDS analysis

- General and call-up
- MDS types (classification, materials, basic substances, recycle)
- Grouping, display options
- Exporting results

Where-used list

- General and call-up
- Types: MDS, substances, applications, name/number
- Difference between general and details
- Interactive analysis and analysis job
- Wizard: Details step 1 to 3
- Result evaluation and export of results

CSI analysis

- General and call-up
- Key dates: significance and management
- Creating test runs
 - Details
 - Wizard: step 1 to 3
- Retrieving results
- Difference between general and detailed results
- Evaluation and export of results

Summary, answering questions

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Teaching Method

Lecture and demonstrations
with exercises



Duration

180 Minutes
(3 Hours)



Max. Number of Participants

14

>> TARGET GROUP

Compliance management, EHS management, quality management, SCIP officers, product and project management

>> PERSONAL PREREQUISITES

Participation at our course No. 5120 "iPCA in Practice" or equivalent knowledge gained through your professional experience.

>> CERTIFICATE OF PARTICIPATION

As a participant in our authorized iPCA training courses, you will receive a personal certificate with which you can prove your iPCA qualification.

>> TRAINING DOCUMENTATION

You will receive a copy of the presentation used in class as a PDF.

>> PUBLIC TRAINING

Current prices and dates can be found on our website www.imds-professional.com

>> EXCLUSIVE TRAINING

You can also book this training exclusively. In this case, the training should be carried out on your iPCA system.

Benefits: You determine the location, date and number of participants and can set the content priorities. If you wish, we can take over the entire organization.