

OECD QSAR Toolbox v.4.2

An example illustrating RAAF Scenario 4 and related
assessment elements

Outlook

- **Background**
- Objectives
- Specific Aims
- Read Across Assessment Framework (RAAF)
- The exercise
- Workflow

Background

- This is a step-by-step presentation designed to take the Toolbox user through the workflow of a data gap filling exercise and assessing of the outcome whether read across is scientifically acceptable or not
- The read-across prediction will be justified by fulfilling all information requirements according to the Read Across Assessment Framework (RAAF).

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Objectives

This presentation demonstrates a number of functionalities of the Toolbox:

- Define a target endpoint;
- Relevancy of profiles and data availability;
- Searching of analogues accounting for metabolism;
- A category consistency check;
- Selection of a RAAF scenario;
- Filling in the report sections related to each read across assessment element.

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Specific Aims

- To familiarize the user with the Read Across Assessment Framework (RAAF) and more specifically with Scenario 4;
- To introduce to the user the read across assessment elements;
- To introduce to the user the report basket;
- To provide sufficient information allowing a scientific assessment of the outcome;
- To explain to the Toolbox user the rationale behind each step of the exercise.

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Read Across Assessment Framework (RAAF) Overview

- RAAF has been developed by ECHA as an internal tool providing a framework for a consistent and structured assessment of grouping and read across approaches under REACH.
- The outcome of the assessment is a conclusion on whether the read across is scientifically acceptable or not.
- The RAAF defines different scenarios for different read-across approaches.
- Each scenario is associated with a particular aspects (assessment elements, AEs).
- Total six scenarios are available: two for an analogue approach and four for a category approach

Read Across Assessment Framework (RAAF)

Selection of a RAAF scenario

SCENARIO	APPROACH	READ-ACROSS HYPOTHESIS BASED ON	QUANTITATIVE VARIATIONS
1	Analogue	(Bio)transformation to common compound(s)	Property of the target substance predicted to be quantitatively equal to those of the source substance or prediction based on a worst-case approach.
2	Analogue	Different compounds have qualitatively similar properties	Properties of the target substance predicted to be quantitatively equal to those of the source substance or prediction based on a worst-case approach.
3	Category	(Bio)transformation to common compound(s)	Variations in the properties observed among source substances. Prediction based on a regular pattern or on a worst-case approach.
4	Category	Different compounds have qualitatively similar properties	Variations in the properties observed among source substances. Prediction based on a regular pattern or on a worst-case approach.
5	Category	(Bio)transformation to common compound(s)	No relevant variations in properties observed among source substances and the same strength predicted for the target substance.
6	Category	Different compounds have qualitatively similar properties	No relevant variations in properties observed among source substances and the same strength predicted for the target substance

*Read-Across Assessment Framework (RAAF) available at https://echa.europa.eu/documents/10162/13628/raaf_en.pdf

Read Across Assessment Framework (RAAF)

Selection of a RAAF scenario

1. Distinguish whether an analogue or a category approach is decided based on the number (N) of analogues:
 - a) N of analogues ≤ 3 is an Analogue approach (scenario 1-2)
 - b) N of analogues > 3 is a Category approach (scenario 3-6)
2. To identify the basis of the read across hypothesis
 - a) (Bio)transformation to (a) common compound(s) – the read across hypothesis is that different substances give rise to (the same) common compounds to which the organism is exposed to
 - b) Different compounds have the same type of (an) effect(s) – the read across hypothesis is that the organism is not exposed to common compounds but rather, as a result of similarity, that different compounds have similar (eco)toxicological and fate properties. These compounds may be the **source** and **target substances themselves** or **one or more of their (bio)transformation products**.
3. For a category approach (scenario 3-6) there is a need to take further account whether or not quantitative variations in the properties are observed among the category members:
 - a) There is a quantitative variation in the (eco) toxicity when it is more than 1 log units* (scenario 3 and 4)
 - b) A quantitative variation is not expected in the (eco) toxicity when it is less or equal to 1 log unit (scenario 5-6)

* The threshold for the number of analogues which distinguishes an analogue from a category approach is proposed by LMC

**The quantitative variation in the (eco)toxicity of 1 log unit is proposed by LMC due to empirically observations.

Read Across Assessment Framework (RAAF)

Selection of a RAAF scenario

- Each scenario consists of a pre-defined set of assessment elements (AEs) that, when taken together, covers all of the essential scientific aspects that need to be addressed in the read-across approach for a particular scenario.*
- Each AE reflects a critical scientific aspect of a read-across.
- The AEs could be:
 - **common** for all scenario within one approach - common AEs for Scenario 1 and 2 (analogue approach) and common AEs for Scenario 3, 4, 5 and 6 (category approach)
 - **specific** – addressing a specific scenario.

*Read-Across Assessment Framework (RAAF) available at https://echa.europa.eu/documents/10162/13628/raaf_en.pdf

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- Workflow

The Exercise

- In this exercise we will predict *Repeated Dose Toxicity (RDT)* of 3,5-dimethyl-aniline [CAS# 108-69-0], which will be the “target” chemical.
- The category will be defined by the aniline functionality causing a *Hemolytic anemia with a methemoglobinemia* accounting for an *in vivo* Rat metabolism;
- The read across approach will be used for the prediction. The read-across will be based on a category approach expressed as a common underlying mechanism for metabolites of source and target substances;
- Read across assessment elements will be included to the report;
- Examples for the possible content of each of AEs will be provided.

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Workflow

- **The Toolbox has six modules which are used in a sequential workflow:**
 - Input
 - Profiling
 - Data
 - Category Definition
 - Data Gap Filling
 - Report

Input Overview

- This module provides the user with several means of entering the chemical of interest or the target chemical.
- Since all subsequent functions are based on a chemical structure, the goal here is to make sure the molecular structure assigned to the target chemical is the correct one.

Input

Input the target chemical by CAS#

The screenshot shows the QSAR TOOLBOX software interface. The top toolbar contains icons for various functions, with the 'CAS#' icon highlighted by a red box and a callout '1'. The main menu bar includes 'Document', 'Single Chemical', 'Chemical List', 'Search', and 'Target Endpoint'. The 'Search' menu is active, and a search dialog box is open. The dialog box has a search field containing '108690' and a 'Search' button, with callout '2' pointing to the search field and callout '3' pointing to the 'Search' button. Below the search field, there are buttons for 'Select All', 'Unselect All', and 'Invert Selection', and a status indicator 'Selected 1 of 1'. The search results are displayed in a table with columns for 'CAS', 'SMILES', 'CS Relation', 'Substance', and 'Composition'. The first result is selected, and its chemical structure is shown on the right. Callout '4' points to the 'OK' button in the dialog box.

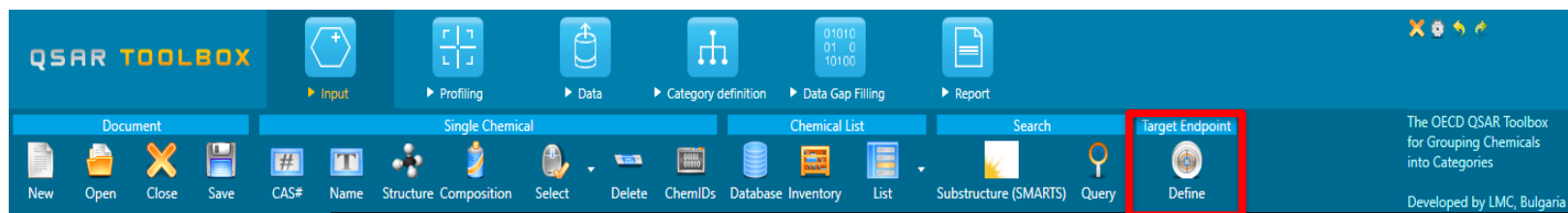
CAS	SMILES	CS Relation	Substance	Composition
108-69-0	<chem>Cc1cc(C)cc(N)c1</chem>	High	Mono constituent	3,5-dimethyl-aniline 3,5-Dimethyl-phenylamine 3,5-Dimethylaniline

1. Click **CAS#**;
2. Enter the CAS# *108-69-0* in the blank field;
3. Click **Search**;
4. When the structure appears, click **OK**.

Input

Define the target endpoint

Defining of the endpoint allows entering the endpoint of interest e.g. EC3, LOEL, LC50 etc., along with specific metadata information. Based on the metadata, different relevancy scores for profiles could be provided for the same endpoint.



Input

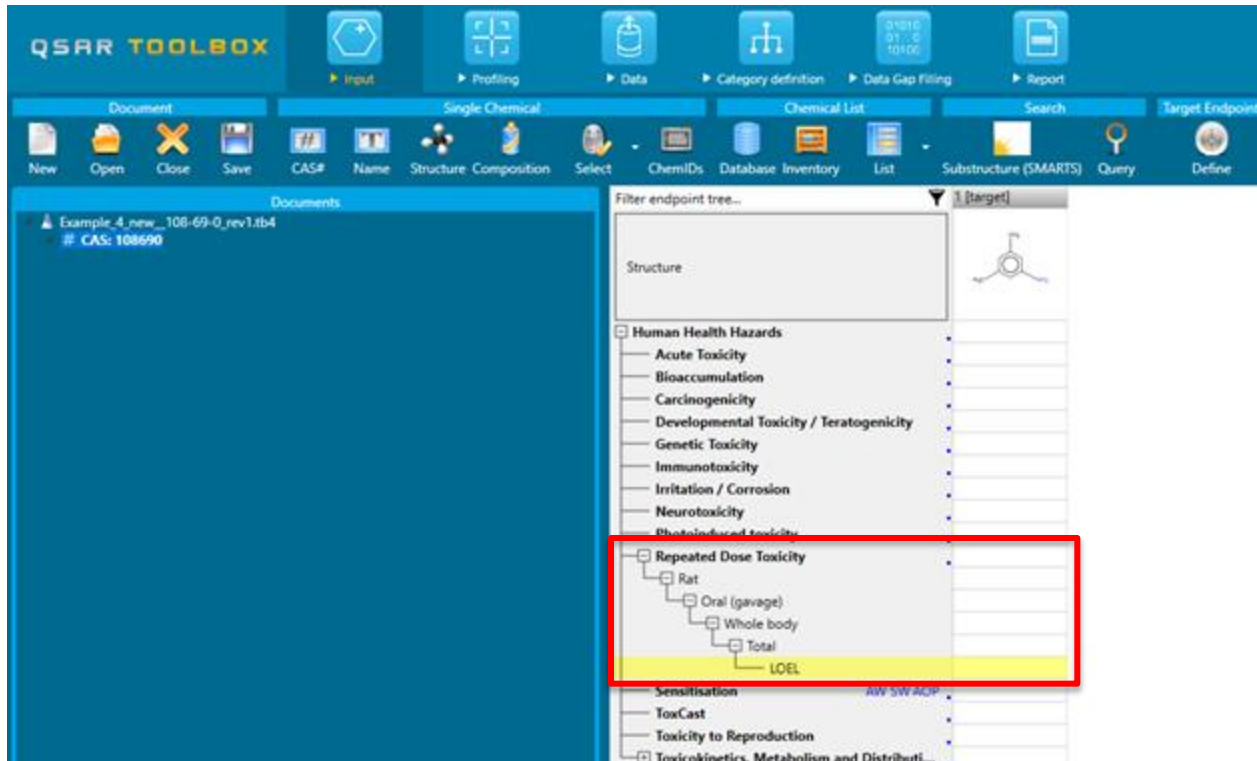
Define the target endpoint

Click on **Define** (1), select **Repeated Dose Toxicity** (2) and then click **Next** (3). Select **LOEL** as an endpoint from the drop-down menu and then consecutively the following metadata: *Effect: Total, Organism(tissue): Whole body, Test organism(species): Rat, Route of administration: Oral (gavage)* (4). Finally click on **Finish** (5).

Input

Define the target endpoint

Once the endpoint is defined along with its metadata, they appear in the endpoint tree and the corresponding row of the data matrix is yellow highlighted.



Profiling Overview

- “Profiling” refers to the electronic process of retrieving relevant information on the target compound, other than environmental fate, ecotoxicity and toxicity data, which are stored in the Toolbox database.
- Available information includes likely mechanism(s) of action, as well as observed or simulated metabolites.

Profiling

Profiling the target chemical

The screenshot displays the QSAR Toolbox interface. At the top, the 'Profiling' tab is active. Below it, the 'Apply' button is highlighted with a red box and a callout '2'. The 'Profiling methods' list on the left has 'Repeated dose (HESS)' checked, highlighted with a red box and callout '1'. The 'Metabolism/Transformations' list also has 'in vivo Rat metabolism simulator' checked, highlighted with a red box and callout '1'. The 'Filter endpoint tree...' on the right shows a tree structure with 'Repeated Dose Toxicity' selected. A chemical structure of a substituted benzene ring is shown in the top right.

1. Tick the checkboxes of the suitable profile - *Repeated dose (HESS)* and of *in vivo Rat metabolism simulator*;
2. Click on **Apply**;

Profiling

Profiling the target chemical

The screenshot displays the QSAR Toolbox interface with the following components:

- Top Bar:** Navigation icons for Input, Profiling, Data, Category definition, Data Gap Filling, and Report.
- Left Panel:**
 - Documents:** Profiling methods and Metabolism/Transformations sections with various simulation options.
 - Profiling methods:** Includes 'Suitable' (Repeated dose (HESS)) and 'Plausible' (Aquatic toxicity classification by ECOSAR, etc.).
 - Metabolism/Transformations:** Includes 'Plausible' (Dissociation, Hydrolysis, in vivo Rat metabolism) and 'Unclassified' (Autoxidation, Microbial metabolism) simulators.
- Center Panel:** 'Filter endpoint tree...' window showing a tree structure. A callout box labeled '1' points to the 'Repeated dose (HESS)' endpoint under the 'Toxicological' category.
- Right Panel:** Displays the chemical structure of Aniline (Nc1ccccc1) and a list of alerts:
 - 1 x Anilines (Hemolytic anemia with methemoglobinemia) Rank A
 - 1 x Anilines (Hepatotoxicity) Rank C
 - 1 x p-Aminophenols (Renal toxicity) Rank B
 - 2 x o-/ p-Aminophenols (Hemolytic anemia with methemoglobinemia) Rank B
 - 4 x Not categorized

Anilines (Hemolytic anemia with methemoglobinemia) Rank A alert is identified in the target chemical as a parent as well as after a metabolic activation (1)

Profiling

Profiling the target chemical

The screenshot displays the QSAR Toolbox interface for profiling a target chemical. The 'Filter endpoint tree...' panel shows a tree structure with 'Repeated Dose Toxicity' highlighted. The 'Structure' panel shows the chemical structure of the target. The 'Profile' panel shows the results of the profiling, with two red boxes highlighting specific alerts. A callout box labeled '1' points to the 'Repeated dose (HESS)' entry in the 'Profile' panel, which lists alerts for Anilines (Hemolytic anemia with methemoglobinemia) Rank A and Anilines (Hepatotoxicity) Rank C. Another callout box labeled '2' points to the 'Repeated dose (HESS)' entry in the 'Metabolism/Transformations' panel, which lists alerts for Anilines (Hemolytic anemia with methemoglobinemia) Rank A, Anilines (Hepatotoxicity) Rank C, p-Aminophenols (Renal toxicity) Rank B, o-/p-Aminophenols (Hemolytic anemia with methemoglobinemia) Rank B, and Not categorized.

Two same alerts: *Anilines (Hemolytic anemia with methemoglobinemia) Rank A* and *Anilines (Hepatotoxicity) Rank C* are identified in the target chemical as a parent as well as after a metabolic activation. *Rank A* label is assigned for the alerts that have a documented mechanism.

Data Overview

- “Data” refers to the electronic process of retrieving the environmental fate, eco-toxicity and toxicity data that are stored in the Toolbox.
- Data gathering can be executed in a global fashion (i.e., collecting all data for all endpoints) or on a more narrowly defined basis (e.g., collecting data for a single or a limited number of endpoints).

Data

Collecting experimental data

The screenshot shows the QSAR Toolbox interface. The 'Filter endpoint tree' on the right lists various hazard categories, with 'Repeated Dose Toxicity' expanded to show 'Oral (gavage)', 'Whole body', and 'Total'. A callout labeled '1' points to the 'Total' sub-entry. The 'Databases' panel on the left lists various data sources, with 'Repeated Dose Toxicity HESS' circled in red and a callout labeled '2' pointing to it. The 'Gather' button in the top toolbar is highlighted with a callout labeled '3'.

1. Select the cell corresponding to the target endpoint
2. Check **Repeated dose (HESS) database;**
3. Select **Gather**.

Data

Collecting experimental data

- Toxicity information on the target chemical is electronically collected from the selected dataset(s).
- It should be kept in mind that the search for data and analogues is performed only among the chemicals which are listed in the selected database(s), which in this example is *Repeated dose (HESS)*.
- Two experimental data related to the defined target endpoint are found.

Based on the observed data (60 mg/kg bw/d) the target chemical is classified as *Category 2* regarding GHS classification ¹

Table 3.9.2: Guidance values to assist in Category 2 classification

Route of exposure	Units	Guidance value range (dose/concentration)
Oral (rat)	mg/kg bw/d	10 - 100
Dermal (rat or rabbit)	mg/kg bw/d	20 - 200
Inhalation (rat) gas	ppm/6h/d	50 - 250
Inhalation (rat) vapour	mg/litre/6h/d	0.2 - 1.0
Inhalation (rat) dust/mist/fume	mg/litre/6h/d	0.02 - 0.2

See on the next slide

¹ Globally Harmonized System of Classification and Labeling of Chemicals (GHS):

http://www.unece.org/fileadmin/DAM/trans/danger/publi/ghs/ghs_rev04/English/ST-SG-AC10-30-Rev4e.pdf

Data

Collecting experimental data

The screenshot displays the QSAR Toolbox interface. On the left, the 'Databases' panel shows 'Repeated Dose Toxicity HESS' selected. The 'Filter endpoint tree...' window on the right shows a list of endpoints. The 'LOEL' endpoint is highlighted in blue and circled in red, with a callout bubble containing the number '1'. The 'LOEL' entry shows a value of '1/2 M: 60 mg/kg bdwt/d'. The 'Total' entry above it shows '1/2 M: 60 mg/kg bdwt/d'. The 'NOEL' entry below it shows '1/2 M: 10 mg/kg bdwt/d'. The 'Tremor/Convulsion' entry shows '1/2 M: 360 mg/kg bdwt/d'. The 'Vocalization' entry shows '1/2 M: 360 mg/kg bdwt/d'. The 'Sensitisation' entry shows 'AW SW AOP'. The 'ToxCast' entry is empty. The 'Toxicity to Reproduction' entry is empty. The 'Toxicokinetics, Metabolism and Distributi...' entry is empty. The 'Profile' entry is empty.

1. The extracted data for *LOEL Repeated dose toxicity* is displayed on the data matrix; Both experimental data for target chemical are the same (*60 mg/kg bdwt/d*).

Category definition

Overview

- This module provides the user with several means of grouping chemicals into a toxicologically meaningful category that includes the target molecule.
- This is the critical step in the workflow.
- Several options are available in the Toolbox to assist the user in refining the category definition.
- As the RDT is a systemic endpoint the metabolism could take place. The primary category in the current example will be defined accounting for an *in vivo rat* metabolism.

Category Definition

Searching for analogues accounting for an *in vivo* Rat metabolism

Endpoint	Dose	Route
Emaciation	1/2 M: 360 mg/kg	bdwt/d
Food Consumption†	1/2 M: 360 mg/kg	bdwt/d
Food Consumption‡	1/4 M: 360 mg/kg	bdwt/d
Hyperthermia	1/2 M: 360 mg/kg	bdwt/d
Hypothermia	1/2 M: 360 mg/kg	bdwt/d
Lacrimation	1/2 M: 360 mg/kg	bdwt/d
Locomotor Activity†	1/2 M: 360 mg/kg	bdwt/d
Locomotor Activity‡	1/2 M: 360 mg/kg	bdwt/d
Miosis	1/2 M: 360 mg/kg	bdwt/d
Muscle Tone‡	1/2 M: 360 mg/kg	bdwt/d
Mydriasis	1/2 M: 360 mg/kg	bdwt/d
Other Findings	1/2 M: 360 mg/kg	bdwt/d
Piloerection	1/2 M: 360 mg/kg	bdwt/d
Ptosis/Palpebral Closure	1/2 M: 360 mg/kg	bdwt/d
Salivation	1/4 M: 360 mg/kg	bdwt/d
Straub Tail	1/2 M: 360 mg/kg	bdwt/d
Total		
LOEL	M: 60 mg/kg	bdwt/d
NOEL	M: 60 mg/kg	bdwt/d
Tremor/Convulsion	1/2 M: 10 mg/kg	bdwt/d
Vocalization	1/2 M: 360 mg/kg	bdwt/d

1. Go to the **Category definition** module; 2. Click **Define with metabolism**; 3. Select ***in vivo* Rat metabolism simulator**; 4. Click **OK**.

Category Definition

Searching for analogues accounting for an *in vivo* Rat metabolism

1. Select a **profile** option for the package "parent & metabolites";
2. Select "Repeated dose (HESS)" profile;
3. Click the **Edit** button. Remove all categories except *Anilines (Hemolytic anemia with a methemoglobinemia) Rank A** category by double click or using "Down" button;

*the categories with Rank A are supported with training sets chemicals having reliable experimental data

Category Definition

Searching for analogues accounting for an *in vivo* Rat metabolism

Grouping options (in vivo Rat metabolism simulator)

All queries At least one

Chemical	Query	Criteria
Parent <chem>Cc1ccc(N)cc1</chem>	none	No criteria.
Metabolite 1 <chem>Cc1ccc(O)cc1</chem>	none	No criteria.
All chemicals		
Parent & Metabolites	profile	Profiler: Repeated dose (HESS) Options: Edit

Alert performance

Scales

Calculate

Target

Anilines (Hemolytic anemia with methemoglobinemia) Rank A

Down Up Reset Options

Profiles (N/A)

- 2-Acetylaminofluorene (Hepatotoxicity) Alert
- 2-Amino-4,5-diphenyl thiazole (Renal toxicity) Alert
- 2-Bromoethylamine (Renal Toxicity) Alert
- 3-Methylcholantrene (Hepatotoxicity) Alert
- 4,4'-Diethylaminoethoxyhexestrol (Hepatotoxicity) Alert
- 4,4'-Methylenedianilines/benzidines (Hepatobiliary toxicity) Rank B
- 4-Aminopyrazolopyrimidine (Hepatotoxicity) Alert
- 5-Azacytidine (Renal Toxicity) Alert
- Acetamide (Renal Toxicity) Alert

Combine profiles

Invert result

AND OR Strict

1

2

OK

OK Cancel

1. Click **OK** to confirm the defined search criteria.
2. Click **OK** in Map similarity options window to execute the search.

In this way we will search for analogues that have this alert as a parent or as a metabolite.

Filter data matrix

The screenshot shows the QSAR Toolbox software interface. The 'Filter endpoint tree...' dialog box is open, displaying a list of endpoints with checkboxes. The 'Total' checkbox is checked, and the 'LOEL' checkbox is also checked. The 'OK' button is highlighted. A red box highlights the dialog box, and three callout boxes (1, 2, 3) point to the 'Advanced filter' icon, the dialog box, and the 'OK' button respectively.

All information on data matrix which is not needed at the current moment could be removed using a filter. Click the **Advanced filter** icon (1). A window with the endpoint tree organization appears. Select only the nodes which you want to see in the data matrix and confirm by clicking OK (3).

Data Gap Filling

Apply Read across

The screenshot displays the QSAR Toolbox interface during a 'Data Gap Filling' workflow. The 'Read across' button in the top toolbar is highlighted with a red circle and a callout '2'. Below, a data matrix table shows chemical structures and their associated data. One cell in the table, containing the text 'M: 60 mg/kg bd...', is circled in red and labeled with a callout '1'. A dialog box titled 'Possible data inconsistency' is open, showing options for 'Route of administration' (Oral (Gavage) and Oral (gavage)) and 'Gap filling scale/unit' (log(1/mol/kg bdwt/d) and mg/kg bdwt/d). The 'OK' button in the dialog is circled in red and labeled with a callout '3'. The 'Filter endpoint tree' on the left shows a selected endpoint: 'Repeated Dose Toxicity -> Rat -> Oral (gavage) -> Whole body -> Total -> LOEL'.

After using of the *Advanced filter* only selected information appears on the data matrix.
 1. Click the cell corresponding to the target chemical in the row with the defined endpoint;
 2. Click **Read-across**; 3. **Possible data inconsistency** window appears, select **OK**.

Data Gap Filling Subcategorizations

Go to **Select/filter data** > **Subcategorize** and consecutively subcategorize by: 1) *US-EPA New Chemical Categories*; 2) *Chemical elements*; 3) *OECD HPV Chemical Categories*.

Data Gap Filling Subcategorizations

The observed *Total LOEL* data among the category members is more than 1 log unit.

Observed data variation of Total LOEL repeated dose data across members of the category

1. Click **Accept prediction** (25.6 mg/kg bdw/d); 2. Click **Yes** to confirm the prediction.

Filter endpoint tree...	1 [target]	21	22	23	27	29
Structure						
LOEL	6/13	M: 60 mg/kg bd...	M: 12 mg/kg bd...	M: 10 mg/kg bd...	M: 50 mg/kg bd...	M: 160 mg/kg b...
Profile						
Predefined						
OECD HPV Chemical Categories						
US-EPA New Chemical Categories						
Empiric						
Chemical elements						
		Dimethylaniline	Dimethylaniline	Dimethylaniline	Dimethylaniline	Dimethylaniline
		Anilines (Acute t...	Anilines (Acute t...	Anilines (Acute t...	Anilines (Acute t...	Anilines (Acute t...
		Group 14 - Carb...	Group 14 - Carb...	Group 14 - Carb...	Group 14 - Carb...	Group 14 - Carb...

Data Gap Filling

RA prediction for CAS 108-69-0

Results

Observed data:

Total LOEL - 60 mg/kg bdw/day

Prediction:

Total LOEL – 25.6 mg/kg bdw/day

Based on the predicted data (for Total LOEL) the target chemical is classified as *Category 2* regarding GHS classification ¹. The RA results is in accordance with the observed data.

Table 3.9.2: Guidance values to assist in Category 2 classification

Route of exposure	Units	Guidance value range (dose/concentration)
Oral (rat)	mg/kg bw/d	10 - 100
Dermal (rat or rabbit)	mg/kg bw/d	20 - 200
Inhalation (rat) gas	ppm/6h/d	50 - 250
Inhalation (rat) vapour	mg/litre/6h/d	0.2 - 1.0
Inhalation (rat) dust/mist/fume	mg/litre/6h/d	0.02 - 0.2

¹ Globally Harmonized System of Classification and Labeling of Chemicals (GHS):

http://www.unece.org/fileadmin/DAM/trans/danger/publi/ghs/ghs_rev04/English/ST-SG-AC10-30-Rev4e.pdf

Report Overview

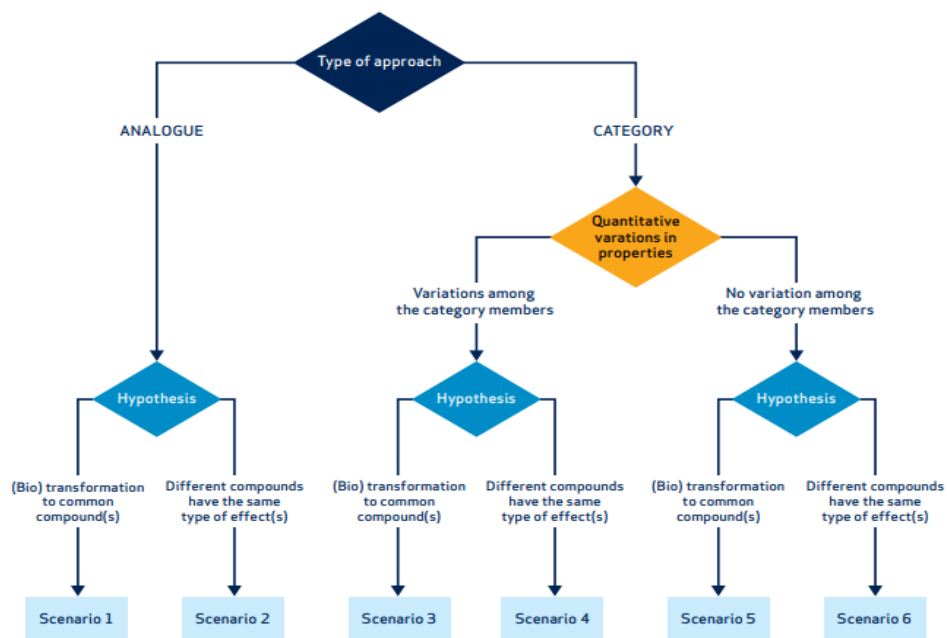
- Report module allows generating a report for any of the predictions performed within the Toolbox.
- The report module contains a predefined report template which the users can customize.
- Additionally a specific RAAF scenario could be chosen. Selection of one of the scenarios will append automatically the related assessment elements related to the corresponding report sections.

Report

Selection of a RAAF scenario

To select the applicable RAAF scenario for assessment, the following aspect should be identified*:

- the type of approach applied - an analogue approach or a category approach;
- the read-across hypothesis;
- For category approach - whether quantitative variations in the properties are observed among the category members must be considered.



*Read-Across Assessment Framework (RAAF) available at https://echa.europa.eu/documents/10162/13628/raaf_en.pdf

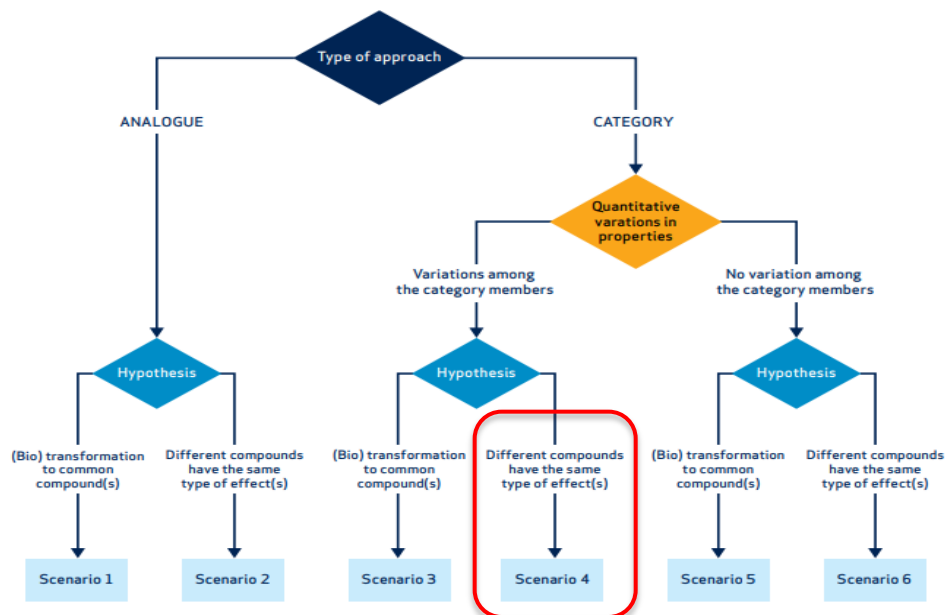
Report

Selection of a RAAF scenario

For the current example:

- the type of approach applied - a **category approach is used** (a threshold of >3 analogues is proposed by LMC for the category approach);
- the read-across hypothesis - **different compounds with a common underlying mechanism for metabolites of source and target substances**;
- For a category approach - **The observed quantitative variation of *Total LOEL* among the category members is more than 1 log unit***.

Scenario 4 was selected for the current example based on the RAAF selection criteria .



*The range of quantitative variation in the (eco)toxicity among the category members of 1 log unit is proposed by LMC based on empirically observations.

Reporting

Report Generation according to RAAF-Scenario 4

1. Go to the **Report module** and click on the cell with the prediction (**R:25.6 mg/kg bdwt/d**); 2. Click the **Prediction** button; 3. Check the box at the top to add a **RAAF scenario**; 4. Select **Scenario 4** from the drop-down menu.

Reporting

Report Generation according to RAAF-Scenario 4

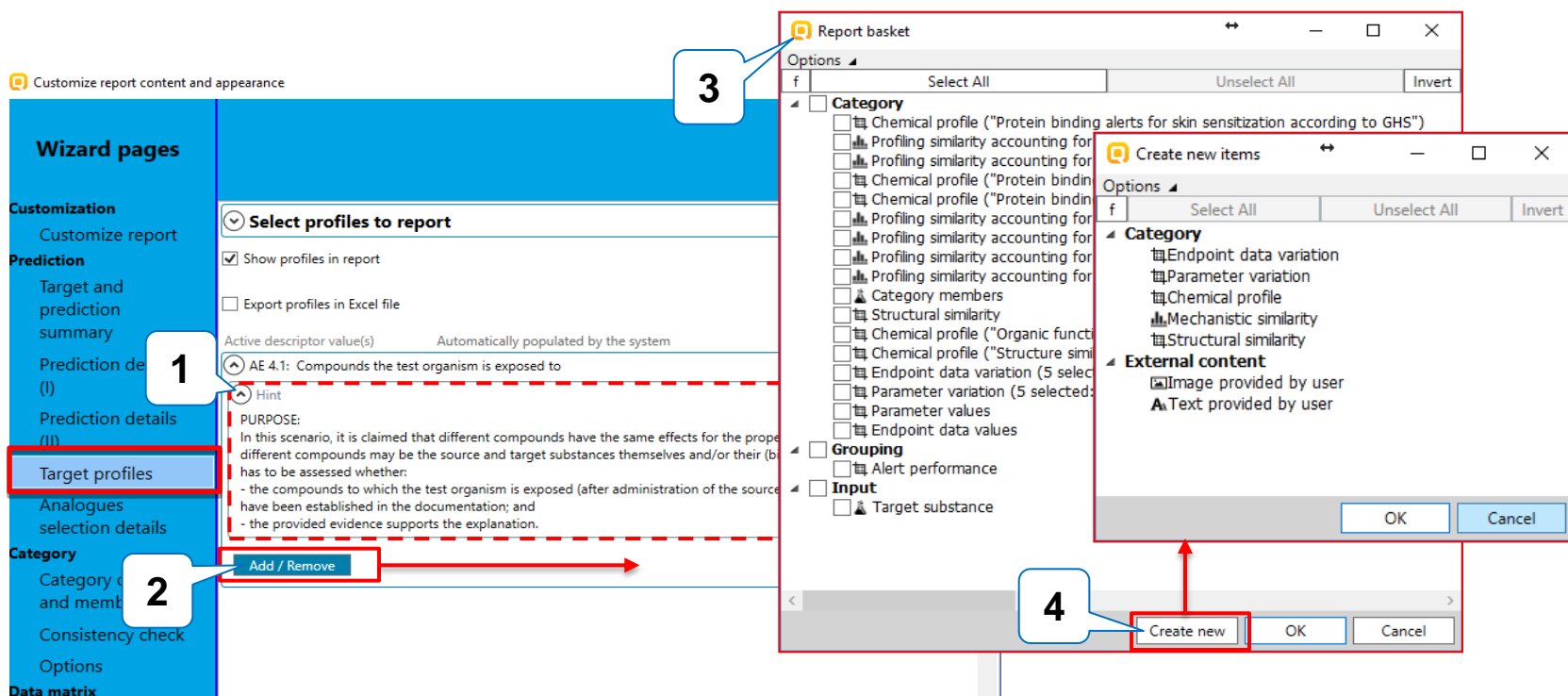
All AEs of **Scenario 4** are distributed as follows: one AE is associated with the *Target profiles* (1), one AE is associated with the *Category definition and members* (2), and nine AEs are associated with the *Consistency check* (3).

Once the RAAF scenario is selected the assessment elements (AEs) related to it will be appended to the corresponding sections of the report automatically. AEs appear in the following report sections: **Target profiles**, **Category definition and members** and **Consistency check**.

Each of the AEs will be considered in the next slides.

Reporting

Report Generation according to RAAF-Scenario 4



A hint for each of the assessment elements is available (1). Information can be included by the **Add/Remove** button (2) located below the corresponding AE. The *Add/Remove* button invokes so called "**Report basket**" (3). The latter contains different items triggered by the actions of the user during the workflow (e.g. Alert performance calculation, applying of category elements, etc.).

Additionally, new items (including items with external content) can be created (4).

Items with an external content (picture and text) will be added for **AE 4.1. Compounds the test organism is exposed to**

Reporting

Report Generation according to RAAF-Scenario 4

- Target profiles section

Customize report content and appearance

Wizard pages

- Customization
 - Customize report
- Prediction
 - Target and prediction summary
 - Prediction details (I)
 - Prediction details (II)
 - Target profiles**
 - Analogues selection details
- Category
 - Category definition and members
 - Consistency check
- Data matrix
 - Options

Select profiles to report

Show profiles in report

Export profiles in Excel file

Active descriptor value(s) Automatically populated by the system

AE 4.1: Compounds the test organism is exposed to

Hint

PURPOSE:
 In this scenario, it is claimed that different compounds have the same effects for the property under consideration. Such different compounds may be the source and target substances themselves and/or their (bio)transformation products. It has to be assessed whether:

- the compounds to which the test organism is exposed (after administration of the source and the target substances) have been established in the documentation; and
- the provided evidence supports the explanation.

Add / Remove

Items with an external content (picture and text) will be added for AE 4.1. Compounds the test organism is exposed to

Back Next Cancel Create report

Possible content of AE 4.1: Compounds the test organism is exposed to

Target A	Source B	Source C	Source D	Source E	Source F
108-69-0	95-78-3	95-68-1	95-64-7	87-62-7	87-59-2
<chem>Cc1cc(C)cc(N)c1</chem>	<chem>Cc1ccc(C)c(N)c1</chem>	<chem>Cc1ccc(N)c(C)c1</chem>	<chem>Cc1ccc(N)cc1C</chem>	<chem>Cc1ccc(C)c1N</chem>	<chem>Cc1ccc(N)c1C</chem>

- There are target substance A and five source substances (B, C, D, E and F)
- The source substances (analogues) B, C, D, E and F have the same aniline functionality as the target substance
- A primary group is defined based on aniline functionality identified either in the parent itself or as metabolites according to RDT profiler accounting for the rat in vivo metabolism.

See how to add the textual content of AE 4.1 along with an illustrative picture of the target and source substances in the next slides.

Reporting

Report Generation according to RAAF-Scenario 4

- Target profiles section

The screenshot displays the 'Customize report content and appearance' window. On the left, the 'Wizard pages' sidebar is visible, with 'Target profiles' selected. The main area shows 'Select profiles to' with a list of categories and sub-categories. A red box highlights 'AE 4.1: Compounds the' (1). Below it, the 'Add / Remove' button is highlighted (2). A 'Report basket' dialog is open, showing a list of categories. The 'Text provided by user' option under 'External content' is selected (4). A 'Create new items' dialog is also open, showing the same list of categories. A text entry dialog (5) is shown with the following text: 'Enter your text here: Target substance A and five source substances (B, C, D, E and F) Source substances (analogues) B, C, D, E and F have the same aniline functionality as the target substance A Primary group is defined based on aniline functionality identified either in the parent itself or as metabolites according to RDT profiler accounting for rat in vivo metabolism'. The 'OK' button is highlighted (6).

In order to add text information to the report: expand the AE (1), click **Add/Remove** button (2), click **Create new** (3) in Report basket window, click **Text provided by user** (4), write in or paste the text in the empty field (5), click **OK** (6).

Reporting

Report Generation according to RAAF-Scenario 4

- Target profiles section

The screenshot shows the 'Customize report content and appearance' wizard. The 'Report basket' dialog is open, showing the 'External content' section with a checked item. A 'Text provided by user' dialog is also open, showing the text to be added. Numbered callouts (1-5) indicate the steps: 1. Ticking the check box, 2. Clicking OK, 3. The item appearing in the report basket, 4. Clicking Edit, and 5. Clicking Preview. A red box highlights the text content in both dialog boxes.

The entered text is listed in the **Report basket** under *External content* section and the check box is ticked (1). Click **OK** (2). The new item is added under the corresponding AE (3). There are two options for each of the report items - **edit** (4) (if you want to change the content) and **preview** (5) (if you want to see the information provided by this item).

Reporting

Report Generation according to RAAF-Scenario 4

- Target profiles section

The screenshot displays the 'Customize report content and appearance' window. On the left, the 'Wizard pages' sidebar is visible, with 'Target profiles' selected under the 'Prediction' section. The main area shows the 'Select profiles to report' section, where the 'External content' checkbox is checked, and the 'Image provided by user' option is selected. A red circle highlights the 'Add / Remove' button (1). Below it, the 'Create new items' dialog box is open, with the 'Image provided by user' option selected (3) and the 'OK' button highlighted (4). A second 'Create new items' dialog box is also visible, with the 'Image provided by user' option selected (3) and the 'OK' button highlighted (4). A third dialog box, 'Select your image here:', is open, showing a grid of image thumbnails (Target A, Source B, Source C, Source D, Source E, Source F) with the 'Image provided by user' option selected (3) and the 'OK' button highlighted (6). The 'Image width, %' field is set to 75.

In order to also add an image: click **Add/Remove** button again (1), *create a new item* (2) and select **Image provided by user** (3) and click **OK** (4). A new window appears where you can add your custom picture by Copy/Paste or browsing (5) to the directory in your PC where the desired picture is saved. Finally confirm by **OK** (6).

Reporting

Report Generation according to RAAF-Scenario 4

- Category definition and members section

#	CAS	Name	SMILES	Structure
1	108-69-0	3,5-xylylidine	Cc1cc(C)cc(N)c1	
2	95-78-3	2,5-Xylylidene	Cc1ccc(C)c(N)c1	
3	95-68-1	2,4-xylylidine	Cc1ccc(N)c(C)c1	
4	95-64-7	3,4-xylylidine	Cc1ccc(N)cc1C	
5	87-62-7	2,6-xylylidine	Cc1ccc(C)c1N	

One AE (1) related to the characterization of the category members (target and source substances) is included in the *Category definition and members* section. The **AE C.1** is automatically filled in by the system (2) by using the available items in the *Report basket*. If impurities/additives of the used analogues are available, they will be also included. The current analogues have no additives/impurities. Example on how the AE C.1. will look in the generated report is shown in right (3).

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Customize report content and appearance

Wizard pages

Customization
Customize report

Prediction
Target and prediction summary
Prediction details (I)
Prediction details (II)
Target profiles
Analogues selection details

Category
Category definition and members
Consistency check
Options

Data matrix
Options

AE 4.4: Exposure to other compounds than to those linked to the prediction

Hint

PURPOSE:
Other compounds than those linked in the hypothesis to the prediction may be formed via other (bio)transformation pathways or may be intermediates/metabolites of the identified pathway. In addition, the impurity profiles associated with the source and target substances may have an impact on the prediction. The other compounds may have been identified by the hypothesis, but not linked to the prediction. Another possibility is that the occurrence of such compounds has been identified by the assessing expert. It has to be assessed whether:

- other compounds that those linked to the prediction may be formed (e.g. via another (bio)transformation pathway or as intermediates) or are present as impurities (see AE A.1); and
- indications are available that such compounds could influence the prediction of the property under consideration.

Add / Remove

AE 4.5: Occurrence of other effects than covered by the hypothesis and justification

AE C.3: Link of structural similarity and differences with the proposed regular pattern

AE C.6: Bias that influences the prediction

Physicochemical similarity based on calculated parameters

Selected 2D/3D parameters for category members

Physicochemical similarity based on experimental data

Selected physicochemical properties for category members

Comments on physicochemical similarity

Back Next Cancel Create report

Possible content of AE 4.4: Exposure to other compounds than to those linked to the prediction

- Target substance A and source substances B, C, D, E and F all have common reactivity pattern. They all formed anilines responsible for the toxicity effects (refer to Appendix Metabolites/Profiling):
- The metabolism in vivo shows formation of other reactive groups (reference to Appendix Metabolites/Profiling) such as:
 - ✓ p-Aminophenols (Rank B)
 - ✓ o/p-Aminophenols (Rank B)
 - ✓ Acetaminophen (Alert)
- Being less reliable (Rank B, C or Alert only) these categories gave enough prove that they do not have significant impact over the influence of the toxic effect
- Our assumption is that Anilines is the functionality responsible for the toxic effect.

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Possible content of AE 4.5: Occurrence of other effects than covered by the hypothesis and justification

- The target substance A and the source substances B,C, D, E and F have a common reactivity pattern based on Anilines functionality
- Additional alerts for repeated dose toxicity have been identified in the parents and their metabolites. The additional mechanisms are with Rank B and Rank C.
- Rank A is assigned only to the used *Anilines (Hemolytic anemia with methemoglobinemia)* category. The categories with Rank A are supported with training sets chemicals having reliable experimental data.
- It is assumed that the additional mechanism will not affect the prediction for the property under consideration.

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Wizard pages

- Customization
 - Customize report
- Prediction
 - Target and prediction summary
 - Prediction details (I)
 - Prediction details (II)
 - Target profiles
 - Analogue selection details
- Category
 - Category definition and members
 - Consistency check
 - Options
- Data matrix

Report basket

Options: Select All, Unselect All, Invert, About, Options

- Category
 - Chemical profile ("Repeated dose (HESS)")
 - Profiling similarity accounting for metabolism ("in vivo Rat metabolism simulator" and "Repeated dose (HESS)")
 - Endpoint data variation (1 selected: Human Health Hazards#Repeated Dose)
 - Category members
 - Structural similarity
 - Endpoint data values
- External content
 - Image provided by user (image from clipboard No.1)
- Input
 - Category purity/impurity
 - Target substance

Buttons: Create new, OK, Cancel

Table summarizing number of metabolites including parent with specific alerts

Alert	P1	P2	P3	P4	P5	P6
Repeated dose (HESS) 108-69-0						
Anilines (Hemolytic anemia with methemoglobinemia) Rank A	2	3	3	3	2	3
Anilines (Hepatotoxicity) Rank C	2	3	3	3	2	3
Bromfenac (Hepatotoxicity) Alert	0	0	0	0	1	0
Mefenamic Acid (Hepatotoxicity) Alert	0	0	0	3	1	6
Menadione (Hepatotoxicity) Alert	0	0	0	0	1	1
Not categorized	4	6	6	5	3	2
o- / p-Aminophenols (Hemolytic anemia with methemoglobinemia) Rank B	2	2	1	2	1	2
p-Aminophenols (Renal toxicity) Rank B	1	1	0	0	1	1
Toluene (Renal toxicity) Alert	0	1	1	0	1	1

Additionally to the entered text, the profiling similarity could be also included. To do this click **Add/Remove** button and check the box of **Profiling similarity** (2). This item stored in the report basket, is triggered by the used simulators and the profiling scheme for the primary grouping. Right click and preview the item (3). Tables with generated metabolites for each parent along with the profiling result will be provided. A table summarizing all profiling results for each of the packages "parent and metabolites" is provided (4) at the end.

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Customize report content and appearance

Wizard pages

- Customization
 - Customize report
- Prediction**
 - Target and prediction summary
 - Prediction details (I)
 - Prediction details (II)
 - Target profiles
 - Analogue selection details
- Category**
 - Category definition and members
 - Consistency check**
 - Options
- Data matrix**
 - Options

AE 4.4: Exposure to other compounds than to those linked to the prediction

AE 4.5: Occurrence of other effects than covered by the hypothesis and justification

AE C.3: Link of structural similarity and differences with the proposed regular pattern

Hint

PURPOSE:
It has to be assessed whether:

- the documentation provides an explanation why the category members should behave in a predictable manner (e.g. based on no absorption due to molecular-weight considerations, or lacking reactivity towards biological material, regular pattern in increasing strength of effect due to kinetic differences);
- it is likely that all category members follow the proposed explanation and where the boundaries of the category are in this respect; and
- the provided evidence supports the explanation.

Add / Remove

AE C.6: Bias that influences the prediction

Physicochemical similarity based on calculated parameters

Selected 2D/3D parameters for category members

Physicochemical similarity based on experimental data

Selected physicochemical properties for category members

Comments on physicochemical similarity

Structural similarity

Back Next Cancel Create report

Possible content of AE C.3: Link of structural similarity and differences with the proposed regular pattern

- The category is structurally defined as a target (A) and the five source substances (B,C, D, E and F) all have a common reactivity pattern;
- They all form *anilines* either as a parent or after a metabolic activation (an in vivo rat metabolism) that are responsible for the toxic effect.

Reporting

Report Generation according to RAAF-Scenario 4

- Consistency check section

Customize report content and appearance

Wizard pages

Customization

Customize report

Prediction

Target and prediction summary

Prediction details (I)

Prediction details (II)

Target profiles

Analogues selection details

Category

Category definition and members

Consistency check

Options

Data matrix

Options

AE 4.4: Exposure to other compounds than to those linked to the prediction

AE 4.5: Occurrence of other effects than covered by the hypothesis and justification

AE C.3: Link of structural similarity and differences with the proposed regular pattern

AE C.6: Bias that influences the prediction

Hint

PURPOSE:
It has to be assessed whether:

- it is clear from the documentation how the source substance(s) have been chosen, for example, what methods/tools have been used to map the field of potential source substance(s), which other substances have been considered and why they have been discarded;
- there are additional, structurally-similar substances which are currently not used in the analogue approach and which arguably could be used;
- there is readily-available information from these additional substances;
- this information is biologically significantly different for relevant properties in comparison with the existing analogue (s); and
- these differences decrease the confidence in the prediction (possibility of underestimation of hazard).

Add / Remove

Physicochemical similarity based on calculated parameters

Selected 2D/3D parameters for category members

Physicochemical similarity based on experimental data

Selected physicochemical properties for category members

Comments on physicochemical

Back Next Cancel Create report

Possible content of AE C.6: Bias that influences the prediction

- The used source chemicals have been found based on a common underlying mechanism for repeated dose toxicity accounting for *in vivo* Rat metabolism;
- The most reliable category was selected (with Rank A);
- The primary group was refined by applying of the following subcategorizations: 1) US-EPA New Chemical Categories, 2) Chemical elements; 3) OECD HPV Chemical Categories.

A chemical expert can provide additional literature search of similar analogues with similar effects

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Customize report content and appearance

Wizard pages

- Customization
 - Customize report
- Prediction
 - Target and prediction summary
 - Prediction details (I)
 - Prediction details (II)
 - Target profiles
 - Analogues selection details
- Category
 - Category definition and members
 - Consistency check**
 - Options
- Data matrix
 - Options

Justification for selected structure similarity profilers

Comments on structural similarity

AE C.2: Structural similarity and structural differences within the category

Hint

PURPOSE:
The aim of this AE is to verify that all category members indeed meet the criteria for structural similarities and allowed structural differences used for the category description. It has to be assessed whether:

- the structural similarities identified apply to all category members; and
- there are structural differences which are allowed within the category.

Add / Remove

Mechanistic similarity

Justification for selected mechanistic similarity profiles/metabolisms

Comments on mechanistic similarity

AE 4.2: Common underlying mechanism, qualitative aspects

Additional endpoints

Tree position:
Human Health Hazards#Repeated Dose Toxicity

Data filters:

Back Next Cancel Create report

Possible content of AE C.2: Structural similarity and structural differences within the category

- The structural similarity between the Target substance A and the five source substances (B, C, D, E and F) according to Str. similarity profiler is in the range of [33-78%]
- Target A and substances B, F have the same reactivity pattern with respect to the OFG profiler
- The source substances C and D have the same reactivity pattern as source E, with one additional group: precursor quinoid compound

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Structural similarity
Options
Mode: Hologram, CombineAllFeatures
Measure:
-Dice
Molecular features:
-AtomCenteredFragments
Atom characteristics:
-AtomType
-CountHAttached
-Hybridization

	Chemical 1	Chemical 2	Chemical 3	Chemical 4	Chemical 5	Chemical 6
Chemical 1	100%	55.6 %	55.6 %	55.6 %	33.3 %	33.3 %
Chemical 2	55.6 %	100%	100 %	77.8 %	66.7 %	77.8 %
Chemical 3	55.6 %	100 %	100%	77.8 %	66.7 %	77.8 %
Chemical 4	55.6 %	77.8 %	77.8 %	100%	77.8 %	66.7 %
Chemical 5	33.3 %	66.7 %	66.7 %	77.8 %	100%	77.8 %
Chemical 6	33.3 %	77.8 %	77.8 %	66.7 %	77.8 %	100%

Two additional items have to be added in order to support the textual information: A *structural similarity* item and an item with the results of the OFG profiler.

Click **Add/Remove** button (1) and check the *Structural similarity* item (2) which is stored in the *Report basket*. Right click and preview the item (3). A table providing structural similarity between each of the category members is shown (4).

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

The screenshot displays the 'Customize report content and appearance' wizard. The left sidebar shows the 'Consistency check' section highlighted. The main window shows a 'Report basket' with a 'Create new items' dialog box open. The dialog box has a 'Create new' button (1), a list of categories including 'Chemical profile' (2), and an 'OK' button (3). A 'Profiler' dropdown menu (4) is also visible, showing a list of profilers including 'Organic functional groups'.

In order to create an item with the OFG profiling results click the **Create new** button (1), select Chemical profile (2) and click OK (3). Select Organic functional groups profiler from the drop-down menu (4) and confirm by OK.*

***In the current example the category elements are not applied. If the consistency of the category is checked then this item will be automatically generated by the system.**

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

Customize report content and appearance

Wizard pages

Customization
Customize report

Prediction
Target and prediction summary
Prediction details (I)
Prediction details (II)
Target profiles
Analogues selection details

Category
Category definition and members

Consistency check

Data matrix
Options

Additional endpoints
 Tree position: Human Health Hazards#Repeated Dose Toxicity
Data filters:
 Category values for selected additional endpoints

Comments on mechanistic similarity
AE 4.2: Common underlying mechanism, qualitative aspects
Hint
PURPOSE: The hypothesis/justification has to explain how the compounds the test organism is exposed to lead to the same type of effects/absence of effects. It has to be assessed whether:
- the documentation has established a common underlying mechanism;
- this mechanism links the structures of these compounds under consideration with the possibility to predict qualitatively similar type of effects for the target substance for the property under consideration; and
- the provided evidence supports the explanation.

Buttons: Back, Next, Cancel, Create report

Possible content of AE 4.2. A common underlying mechanism, qualitative aspects

- The Target substance A and the source substances B, C, D, E and F all react via a common underlying mechanism according to the RDT profiler
- They all have anilines functionality either as a parent or after a metabolic activation
- The similarity with respect to the metabolic pattern could be seen in **AE 4.5.** above.

Additionally, metabolic maps (for each of the analogues), produced by external software or found in the literature, could be included to this AE in order to support the mechanistic similarity of the category.

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section

The screenshot displays the 'Customize report content and appearance' wizard. On the left, the 'Wizard pages' sidebar has 'Consistency check' highlighted. The main window shows the 'Report basket' with a tree view where 'Endpoint data variation (1 selected: Human Health Hazards#Repeated Dose Toxicity)' is checked. Below this, the 'AE 4.3: Common underlying mechanism, quantitative aspects' section is expanded, showing a 'Hint' and 'PURPOSE' text. The 'PURPOSE' text reads: 'Under this scenario, quantitative differences for the same type of effects are expected caused by the underlying mechanism. It has to be assessed whether: - the documentation established that the strengths of the same type of effects vary in predictable manner; - the prediction is derived from the relation between an observed property and the independent variable which determines the order within the category (prediction model); - the prediction model is consistent with the common mechanism; and - the provided evidence supports the explanation.'

Possible content of AE 4.3. A common underlying mechanism, quantitative aspects

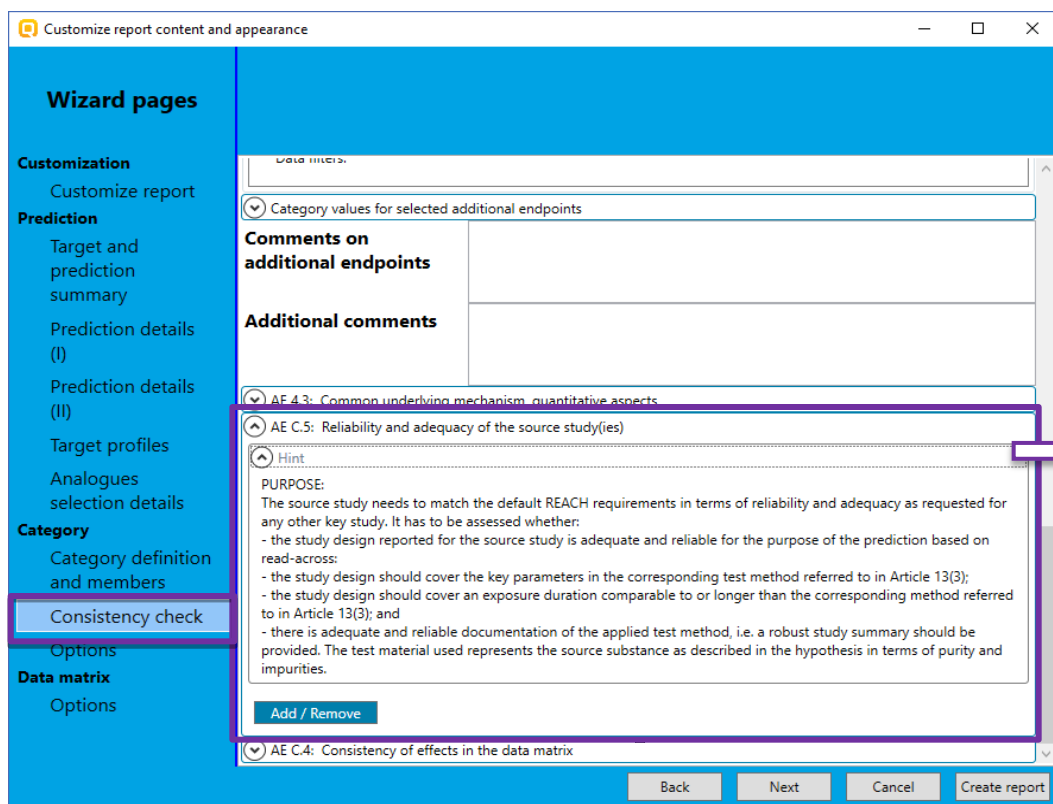
- The target substance A and the five source substances have a common reactivity pattern.
- They all formed aniline functionality either as parents or as metabolites responsible for the toxicity effects
- Similar toxic effects observed in source substances support the prediction for the target
- Toxic effects of all source substances and target are supported by the identified additional RDT data (references could be included).
- The range of variation of the LOEL experimental data for all category members is shown below:

After the last bullet include the Endpoint data variation item stored in the report basket (1).

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section



Possible content of AE C.5: Reliability and adequacy of the source study(ies)

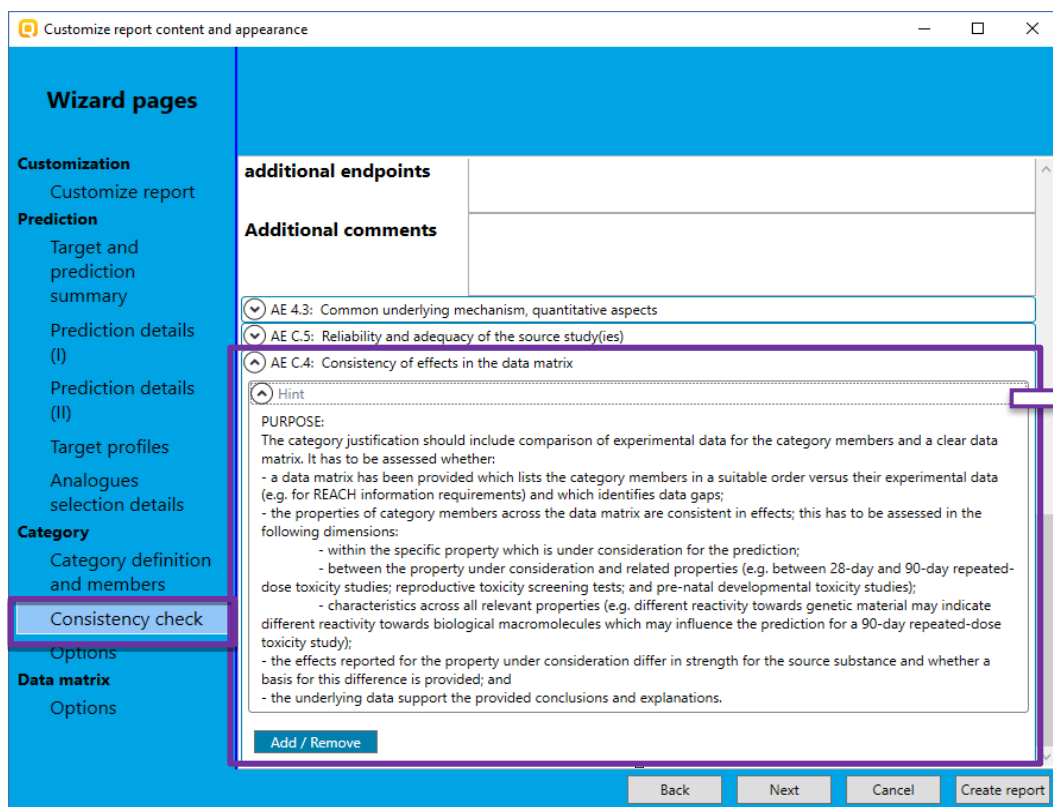
- The target substance has been tested according to test guideline 407
- All of the five source substances with one exception (substance E) has been tested based on test guideline 407: Repeated does 28-day oral toxicity study in Rodents
- For the source substances E and B the study was based on report NTP Long term and OECD: Combined Repeated Dose Toxicity Study with the Reproduction/Developmental Toxicity Screening Test

A snapshot from *Filter points by test conditions* could be provided.

Reporting

Report Generation according to RAAF-Scenario 4

- A consistency check section



Possible content of AE C.4: Consistency of effects in the data matrix

- The target substance A and the five source substances (B, C, D, E and F) show indication for a repeated dose effect especially for reduce red blood cell. The Total LOEL read-across prediction in this case is around 30 mg/kg bdw/day which classify the target chemical in the range of Category 2 according to GHS classification
- The latter is supported by the experimental data found for all of used source substances for the investigated endpoint and other similar properties

Here should be provided the data matrix snapshot or reference to the *Data matrix report*.

Report

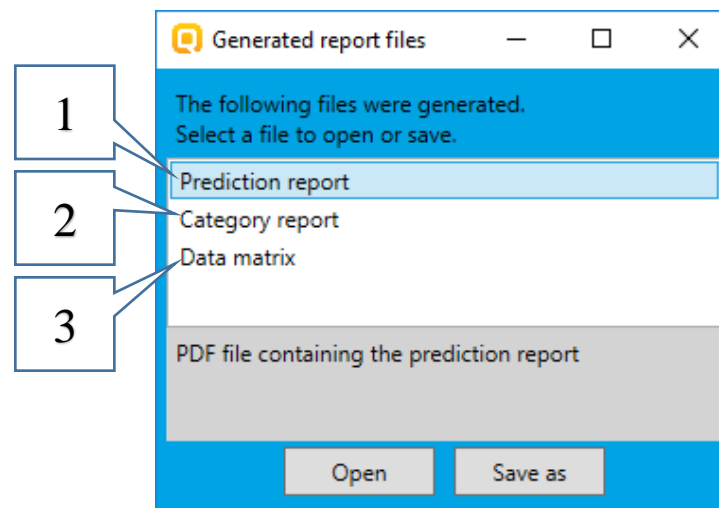
Report Generation

After the click on the Create report button, The *Generated report files* window appears. It contains three type of files:

- 1) Prediction report** - a PDF file containing the prediction information related to the target.
- 2) Category report** - a PDF file containing information for the consistency of the final category (target plus used analogues)
- 3) Data matrix** - a MS Excel file containing chemicals used for prediction along with their data for selected parameters, profiles and endpoint tree positions.

RAAF AEs are included in the first two files.

All generated files should be provided when submit a prediction.



Report Generated report files

Prediction report

QSAR Toolbox prediction for single chemical

(In accordance with RAAF scenario 4)

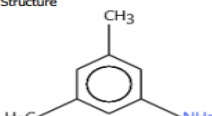
The selected RAAF scenario is specified in the first page

Category report

QSAR Toolbox report for category

(In accordance with RAAF scenario 4)

Date: 26 Mar 2018
Author(s):
Contact details:

Target information		
Structural information	Numerical identifiers	Chemical names
SMILES: Cc1cc(C)cc(N)c1	CAS#: 108-69-0 Other: EC Number:3777173	3,5-dimethyl-aniline 3,5-Dimethyl-phenyla mine 3,5-Dimethylaniline
Structure 		

1. Category definition

1.1. Category definition

Not provided by the user

manually editable field

Ranges for selected physicochemical properties and calculated parameters

Not provided by user

1.2. Covered (target) endpoint(s)

- Human Health Hazards/Repeated Dose Toxicity: LOEL, Total, Whole body, Oral (gavage), Rat

manually editable field

1.3. Category hypothesis

Not provided by the user

manually editable field

1.4. Profiles/Metabolisms

manually editable field

Profile/subcategorization:

"in vivo Rat metabolism simulator" Combined parent and products requirements: Anilines c anemia with methemoglobinemia) Rank A (Repeated dose (HESS)) (primary grouping)

New Chemical Categories (subcategorization)

Elements (subcategorization)

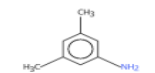
PV Chemical Categories (subcategorization)

Members

manually editable field

Profile characterization





Members

Name	SMILES	Structure
3,5-xylidine	Cc1cc(C)cc(N)c1	

QSAR TOOLBOX

TPRF v4.2

Data matrix report

	Neighbour #2	Neighbour #3	Neighbour #4	Neighbour #5
Structure				
CAS number	108-69-0	95-78-3	95-68-1	95-64-7
Chemical name	3,5-xylidine	2,5-Xylidine	2,4-xylidine	3,4-xylidine
Other identifier				
SMILES	Cc1cc(C)cc(N)c1	Cc1cc(C)cc(N)c1	Cc1cc(N)cc(C)c1	Cc1cc(C)cc(N)C
Parameters	unit			
Profiles				
Profiles used for grouping/subcategorization	Parent and 7 metabolites; Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Has the following additional Anilines (Acute toxicity); Group 14 - Carbon C; Group 15 - Nitrogen N	Parent and 10 metabolites; Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Has the following additional Anilines (Acute toxicity); Group 14 - Carbon C; Group 15 - Nitrogen N	Parent and 9 metabolites; Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Has the following additional Anilines (Acute toxicity); Group 14 - Carbon C; Group 15 - Nitrogen N	Parent and 11 metabolites; Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Has the following additional Anilines (Acute toxicity); Group 14 - Carbon C; Group 15 - Nitrogen N
Repeated dose (HESS)	Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Anilines (Hepatotoxicity) Rank C	Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Anilines (Hepatotoxicity) Rank C; Toluene (Renal toxicity) Alert	Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Anilines (Hepatotoxicity) Rank C; Toluene (Renal toxicity) Alert	Anilines (Hemolytic anemia with methemoglobinemia) Rank A; Anilines (Hepatotoxicity) Rank C
Measured and predicted data				
Environment	endpoint	value	unit	species, duration, test type, type of method, assay, strain, test guideline
Repeated Dose Toxicity	LOEL	60	mg/kg bw/d/d	Rat 28 d
Repeated Dose Toxicity	LOEL	12	mg/kg bw/d/d	Rat 28 d

Congratulation

- You have now been introduced to the RAAF scenario;
- You have now been introduced to the *Report basket*.
- You have now been introduced to the AEs related to Scenario 4.
- Note proficiency comes with practice.