

# SCIP IT user group

3 December 2019





### Validation rules for SCIP

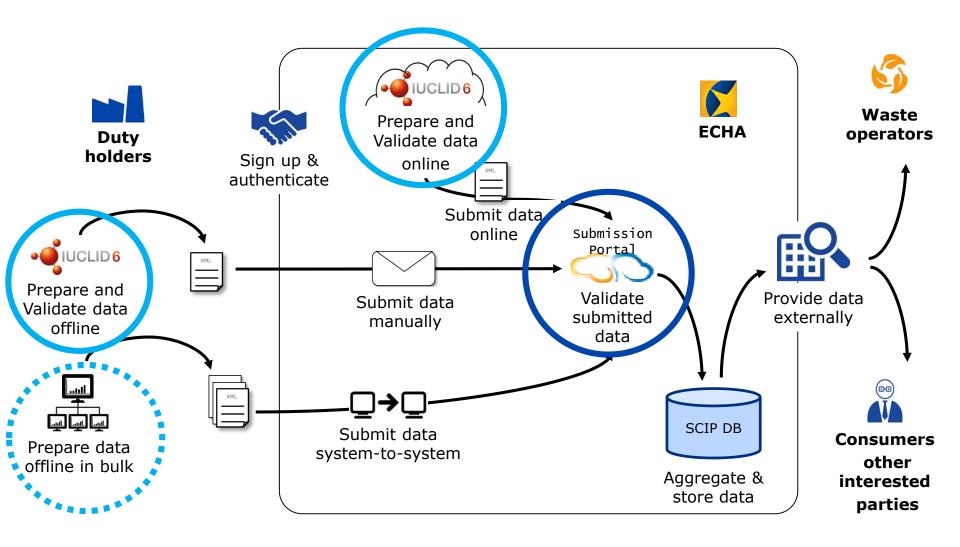
SCIP IT User group

3/12/2019

Saara Sumiala Data quality team



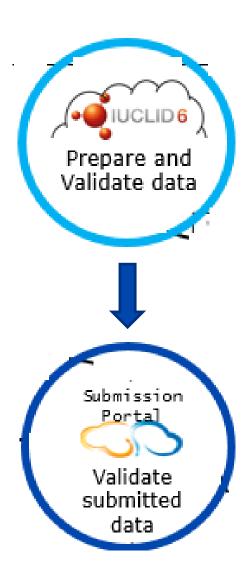
# **SCIP** General Overview





## **Validation steps**

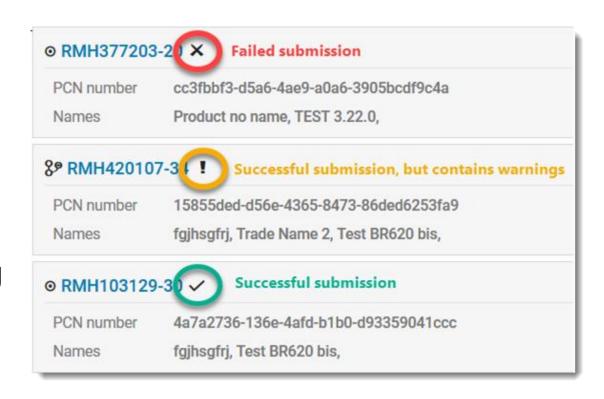
- Validation rules in IUCLID
  - Rules that can be checked at dossier level ('Validation assistant rules')
- Validation rules in ECHA
   Submission Portal
  - All the rules which were checked at IUCLID (Validation assistant rules) + Contextual rules ('Portal rules')





# Failures (BR) and Warnings (QLT)

- Business rule
   BR failure
- Quality ruleQLT warning



Failed – dossier not moved for next step (it is discarded)

Successful submission – dossier moved for next step



#### **Validation outcome**

- All rules are automatically checked
   →Validation results in seconds
- If notification fails validation rulesnew version of the dossier can be submitted immediately



#### **Estimate of the number of rules:**

#### **IUCLID Validation rules**

- **BR** 17
- **QLT** 12

#### ECHA Submission portal rules

- BR 1
- **QLT** 1

#### ~ 20 rules in total



Example 1:

Minimum information is provided:

'Article category' must be indicated



Example 2:

### Information is complete:

If in characteristics 'Height' was provided then the adjacent 'Unit' must be provided as well.



Example 3:

#### Information is consistent:

Same substance can not be indicated to be both **no longer present** and **present** in the same 'Article' record



Example 4:

#### Provided format is correct:

If 'ECHA SCIP ID' was indicated as 'Identifier type' then the provided value must be in UUID format



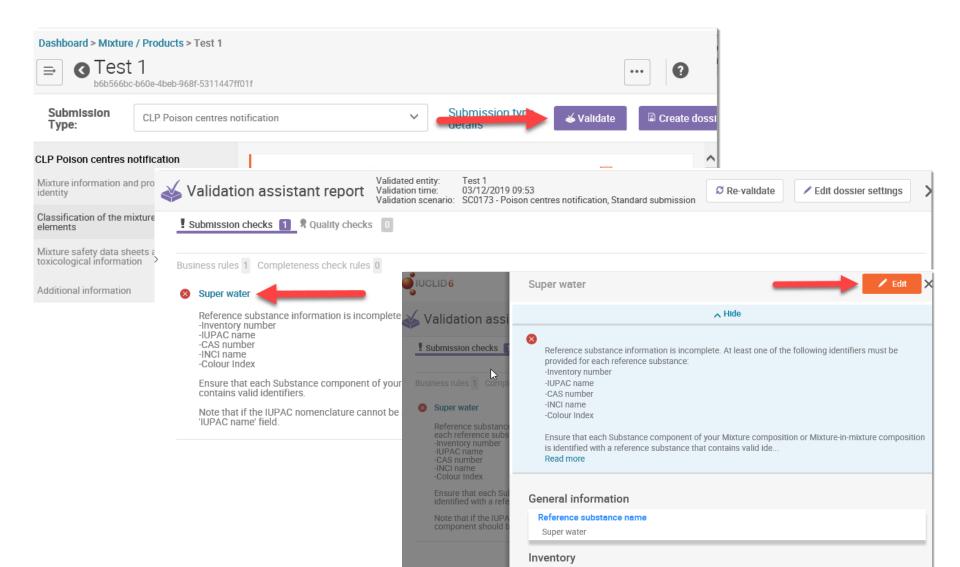
Example 5:

Attachment sizes are not too big:

Attachments must be less than 10MB per 'Article' record

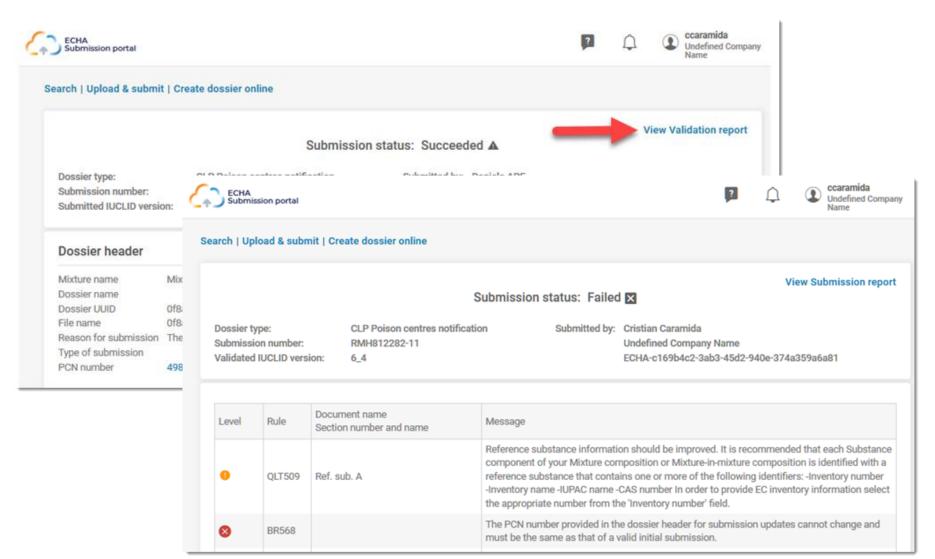


#### **Validations in IUCLID**





#### Validations in 'ECHA Submission Portal'





# Help that we provide

Document listing the rules

Safe use instructions	
BR708	Either 'Safe use instructions' must be reported or 'No need to provide safe use information' must be ticked. Both cannot be selected.
QLT706	If <b>Disassembling instructions</b> are provided then the <b>language</b> must be indicated.
BR709	If <b>Disassembling instructions</b> are provided then the attachment must be provided in <b>PDF format</b> .

CONCERN ELEMENT	
BR713	'Candidate list version' must be indicated.
BR703	All the 'Concern element' blocks provided should be complete: 'Reference substance' must be indicated.
BR715	For all the 'Reference substances' at least one relevant identifier must be provided.  • EC number  • CAS number  • IUPAC name  • ECHA Substance ID
QLT701	All the 'Reference substances' included must be mentioned in 'Candidate list'.



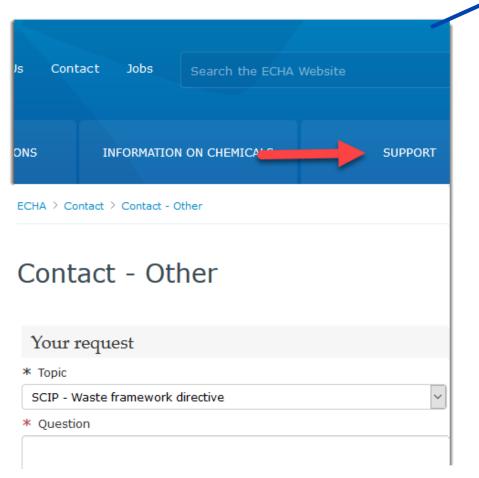
### Help that we provide (system-to-system)

# Document listing the rules + field paths

CONCERN ELEMENT		
BR713	'Candidate list version' must be indicated.	
	ARTICLE.ConcernElements.CandidateListVersion	
BR703	All the 'Concern element' blocks provided should be complete: 'Reference substance' must be indicated.	
	ARTICLE.ConcernElements.ConcernElement.CandidateListSubstanceLink	
BR715	For all the 'Reference substances' at least one relevant identifier must be provided.  • EC number  • CAS number  • IUPAC name/Chemical name  • ECHA Substance ID	
	EC number: REFERENCE_SUBSTANCE.Inventory.InventoryEntry	
	CAS number: REFERENCE_SUBSTANCE.ReferenceSubstanceInfo.CASInfo.CASNumber	
	IUPAC name and Chemicals name:  REFERENCE_SUBSTANCE.ReferenceSubstanceInfo.IupacName	
	Other: ECHA Submission ID:	

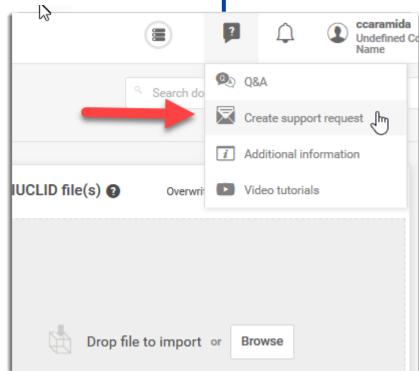


# Help that we provide



ECHA HelpDesk

**IUCLID Cloud support request** 





#### **Timeline**

- IUCLID Validation rules in beginning of 2020
  - Possible changes provided when IUCLID is updated