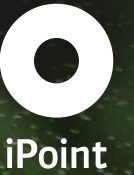
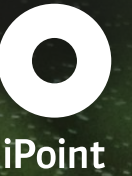


iPoint SCIP Connector



Thank you for joining!
We will be starting shortly.

iPoint SCIP Connector



Speakers



Dr. Angelika Steinbrecher

Senior Expert Compliance, Sustainability, and Innovation
iPoint Group



Sushma Kittali-Weidner

Director of Operations
iPoint Group

Agenda



1. Latest Updates on ECHA SCIP Database
2. Challenges of manual efforts for SCIP Submission
3. Demo of SCIP Database Connector Enterprise and Professional
4. Q&As

Housekeeping



1-hour session



Dedicated time for questions



Session is recorded



Audience feedback



Audience on mute



Slides and recording will be available after the webinar

About iPoint



iPoint



We are iPoint



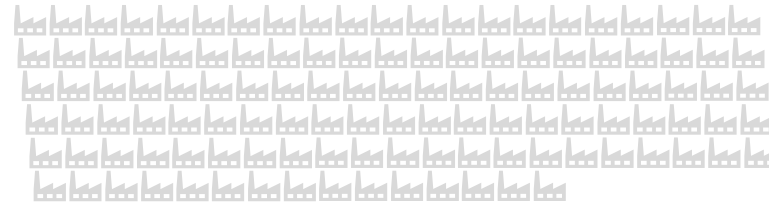
2001 
foundation iPoint



Presence in
**North America, Europe, Asia and
Australia**



170 experts
and growing



250 global
enterprises among customers

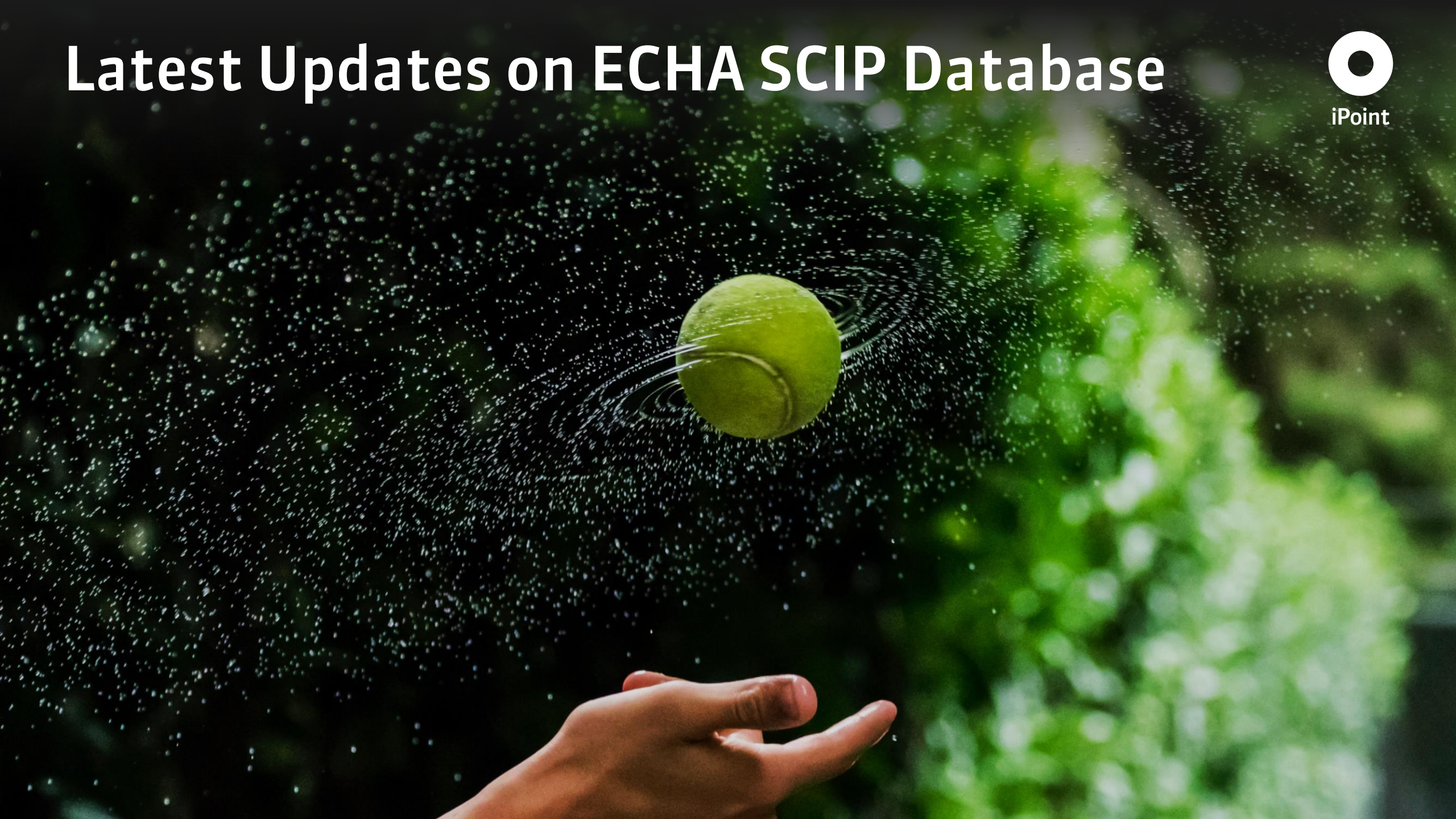
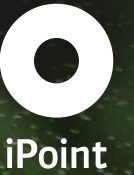


We develop software

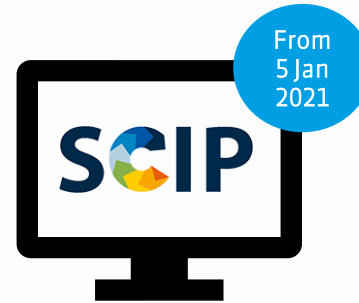
- Compliance
- Sustainability
- Risk



Latest Updates on ECHA SCIP Database

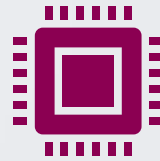


SCIP Database – in a nutshell



Who reports?

EU based companies
supplying articles
to the EU market



What is reported?

Articles with
SVHC >0.1%

(and complex objects
containing these)



Why?

- Enable proper treatment of hazardous waste
- Push for substitution
- Strengthen circular economy
- Increase authorities' information base

Latest Updates – SCIP DB go live Oct 28th



- SCIP database ready to use (since October 28th, 2020)
- In line with latest release of IUCLID version 6.5
- Update of picklists: production in EU statement, primary article identifier types
- Changes to Candidate List Package:
 - 8 substances added
 - 23 substances modified (description, name, CAS-number)

Latest Updates – Validation Rules

- SSN-SCIP number should be valid
- Non-EU companies not allowed to submit SCIP notifications
- ‚SCIP notifications‘ accepted by submission portal
- Wording modified: „no need to provide [....]“ changed to:

‘The identification of the Candidate List substance is sufficient to allow safe use of the article throughout the whole life cycle including service life, disassembly and waste/recycling stage’

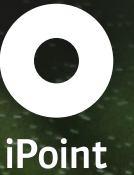
Source: [ECHA Validation Rules](#)

Latest Updates – Access and Trials



- Get your ECHA IT access and S2S key now – for each legal entity
- Contact ECHA in case of high volume of S2S key requests providing list of member companies that need S2S key
- Test environment still available to try out SCIP submissions – also via S2S

Challenges of manual efforts for SCIP Submission

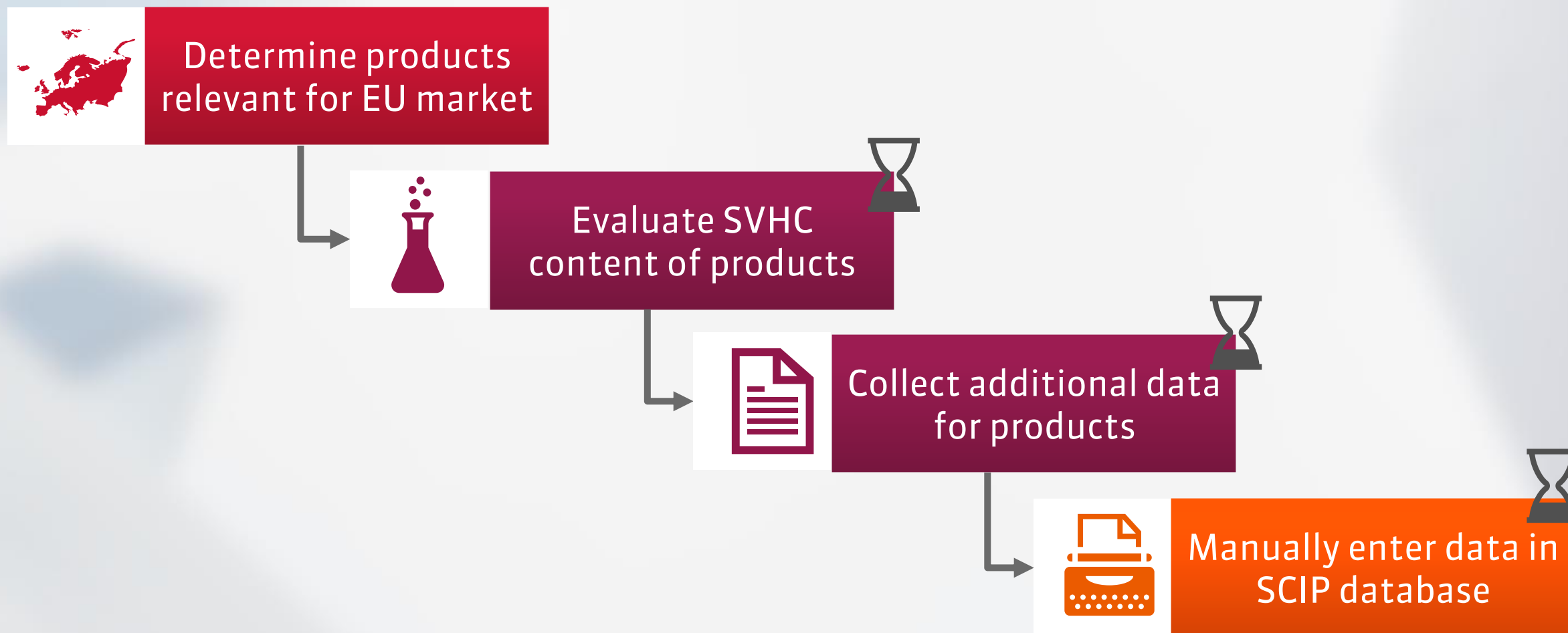


SCIP Reporting

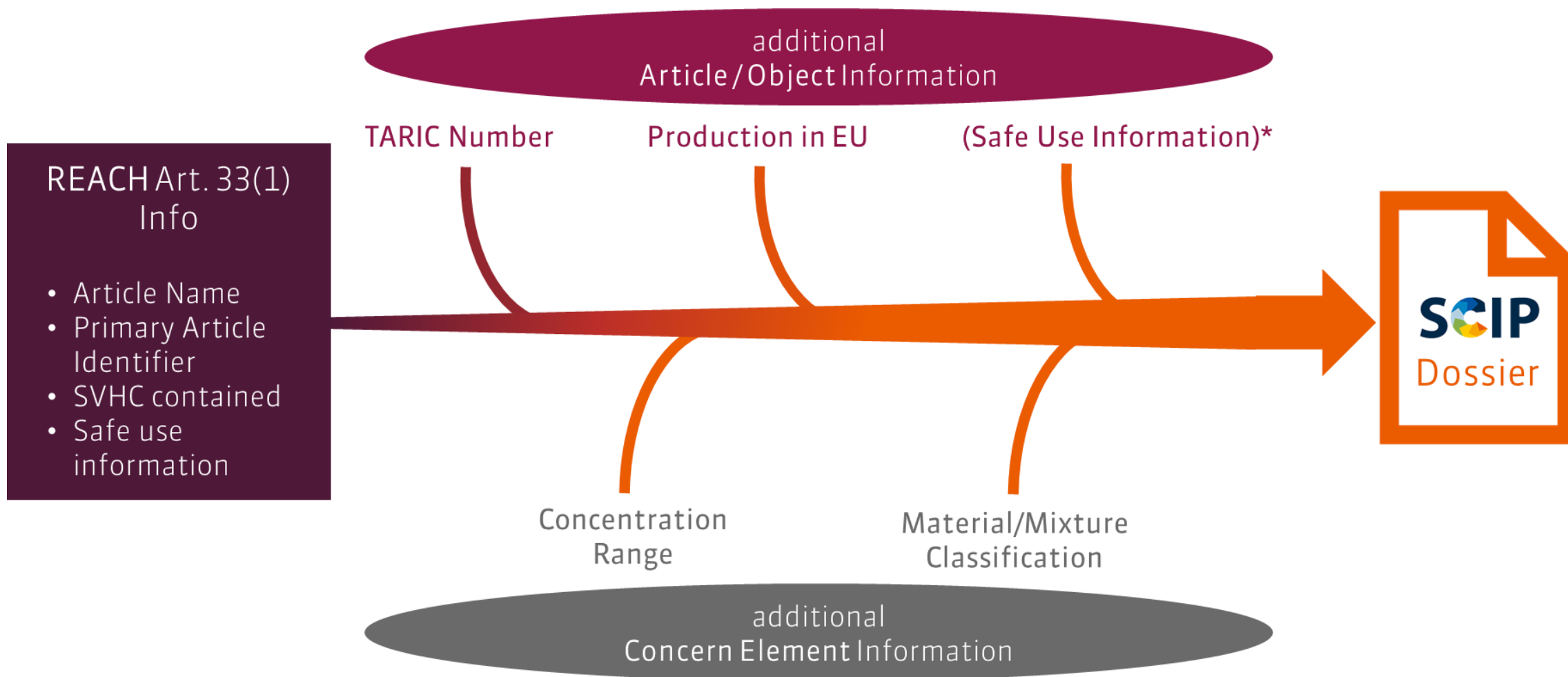
What are the challenges?

- Identification of SCIP relevant BOM structure to be reported
- Management of Article/Material Categories and SUI
- Collecting data from supply chain
- Aggregation of market relevant products and their composition
- Calculation of SVHC content in Products
- Management of Legal entities and their reporting obligations
- High volume of data with varying ways of identification

Manual SCIP Workflow

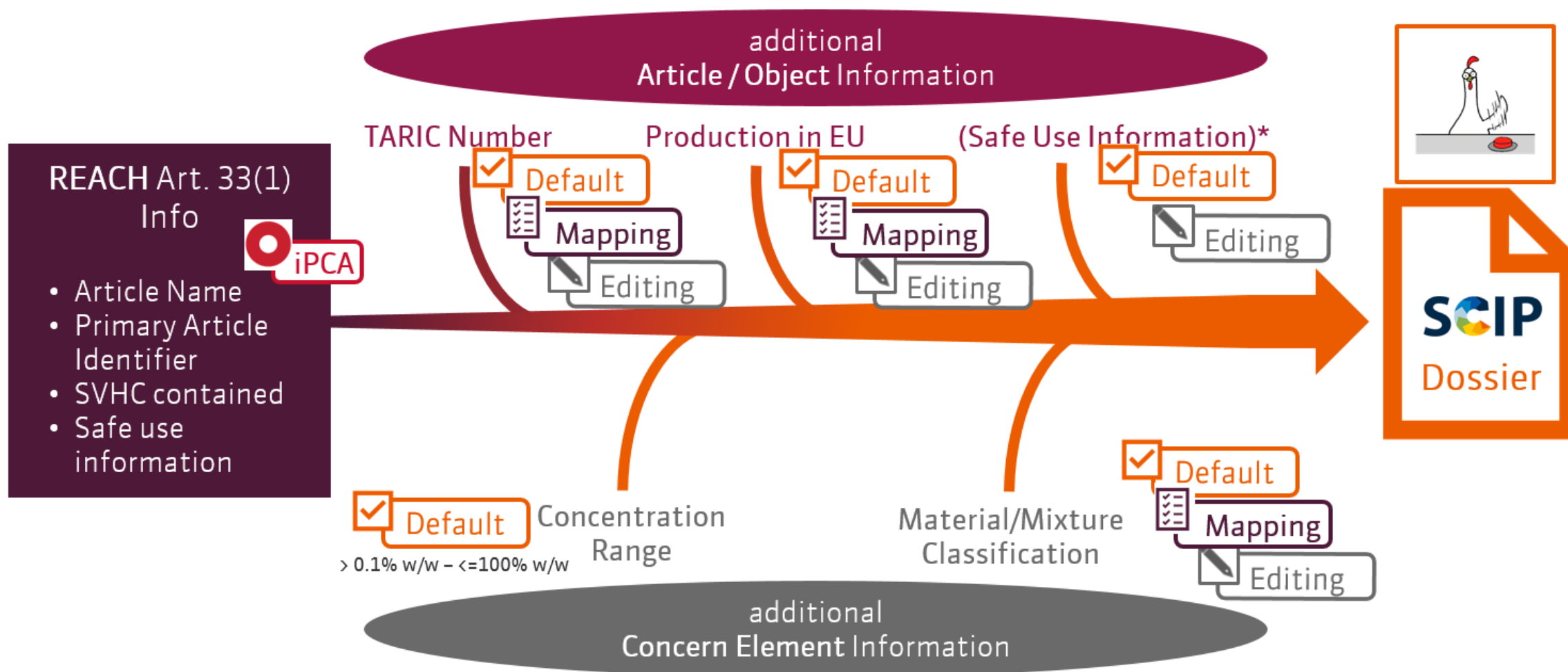


Data Sources for SCIP Submission



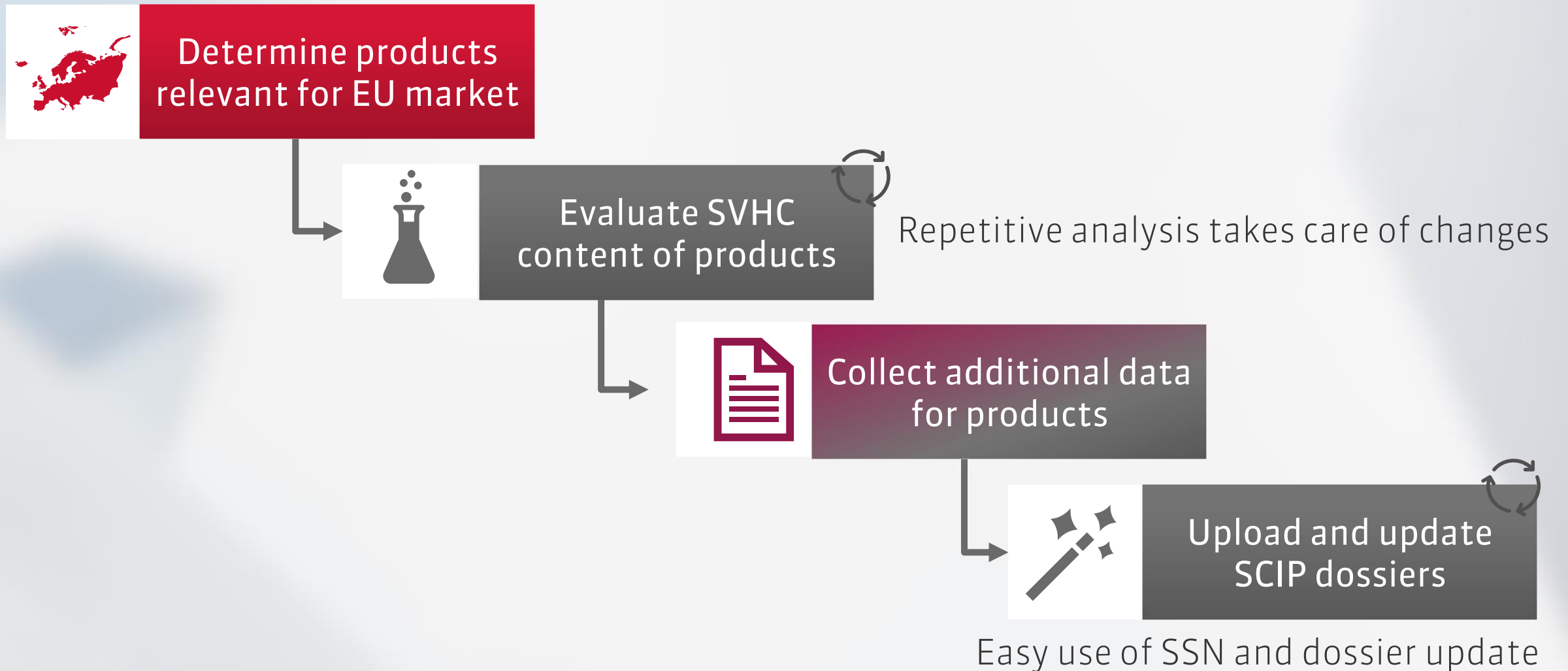
* Inclusion of additional Safe Use Information might be required in some cases

Data Sources for SCIP Submission



* Inclusion of additional safe use information might be required in some cases

SCIP Workflow with Automation



Large OEM

1 week -> 10 minutes
for 2,500+ Parts

**Determine products
relevant for EU market**



10 weeks -> 10 minutes
20% parts contain SVHC

**Evaluate SVHC
content of products**



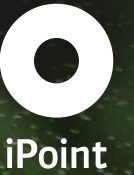
2 weeks -> 2 hours
Outreach to 500 Suppliers

**Collect additional data
for products**

**Upload and update
SCIP dossiers**

4 hours -> 10 minutes
Create 1 Dossier

SCIP Database Connector Enterprise and Professional



iPoint SCIP Solution

Features



Web Interface



Dashboard
overview



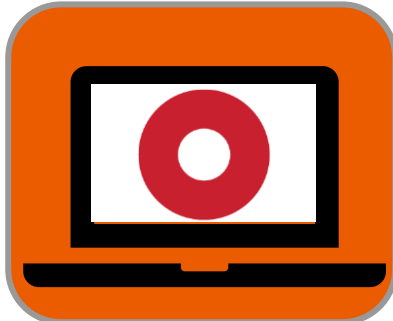
Multiple data
sources



Import of
additional data



ECHA IT Access
Mgmt. for all
L.E.



Global default
setting



Individual and
bulk handling



User
management



Bulk upload



Submission
History



Simplified
Notification

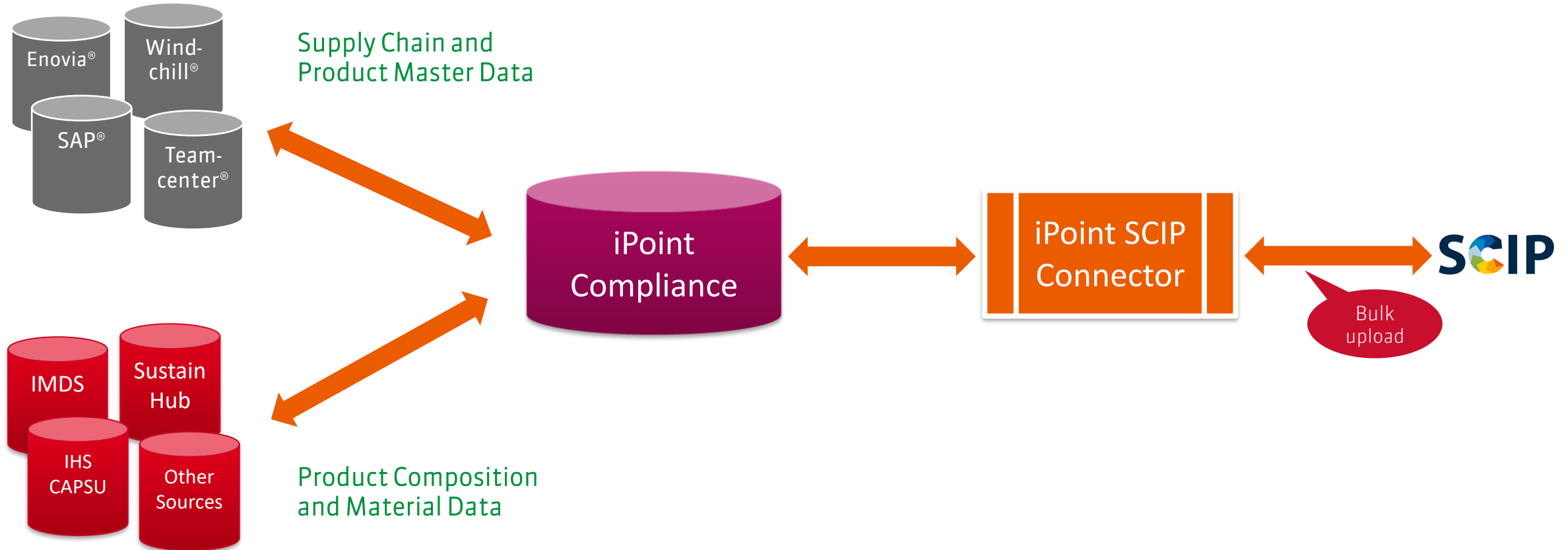
Solution Options



- SCIP Connector with iPoint Compliance Enterprise
- SCIP Connector Professional
- SCIP Turn-key data management

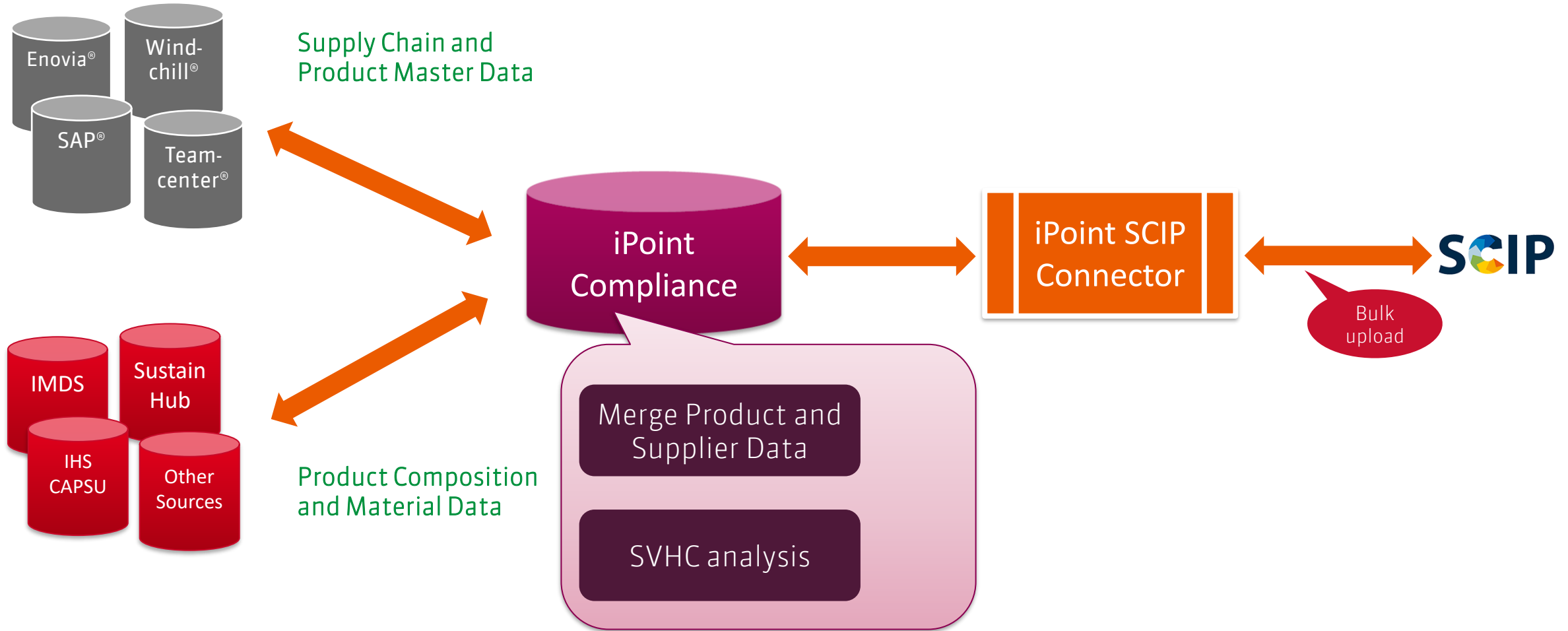
Rough workflow for SCIP dossier submission

Using iPoint's Enterprise solution



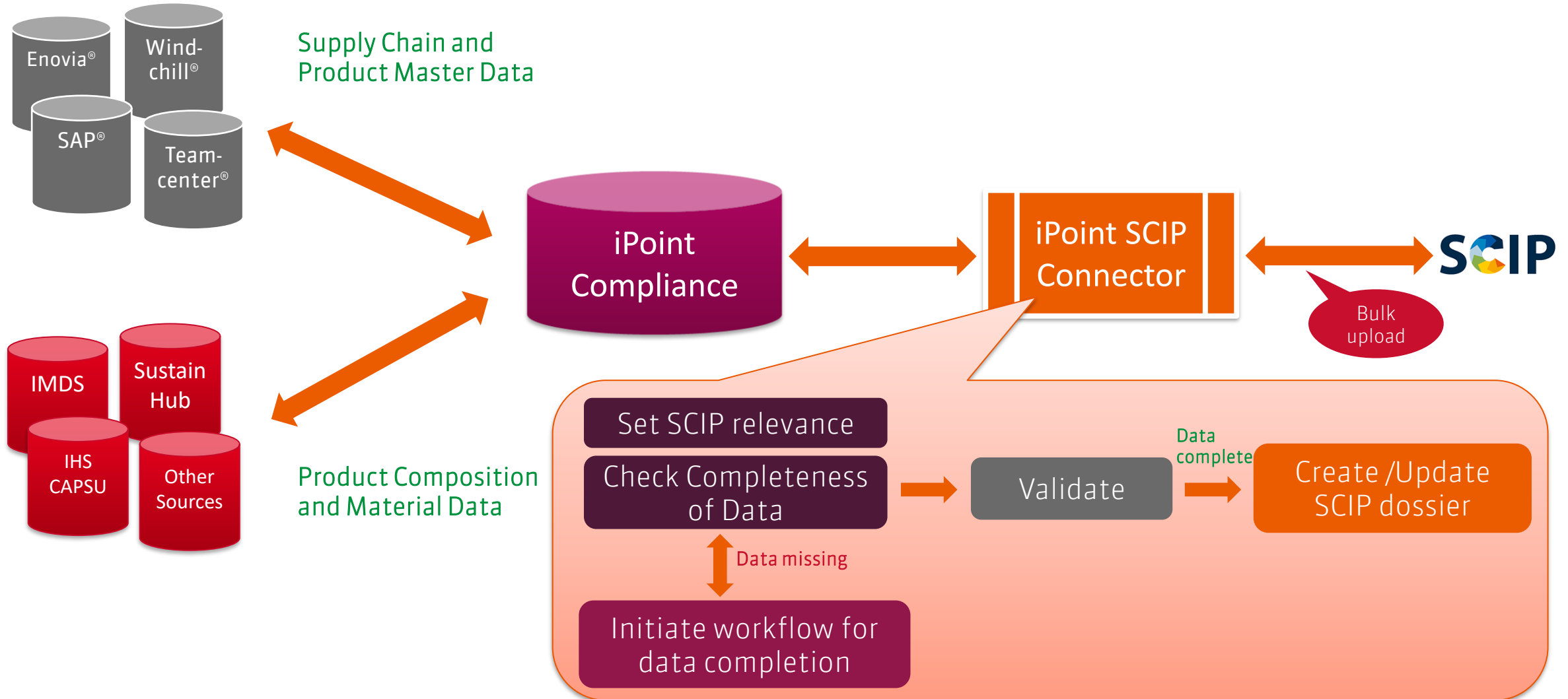
Rough workflow for SCIP dossier submission

Using iPoint's Enterprise solution



Rough workflow for SCIP dossier submission

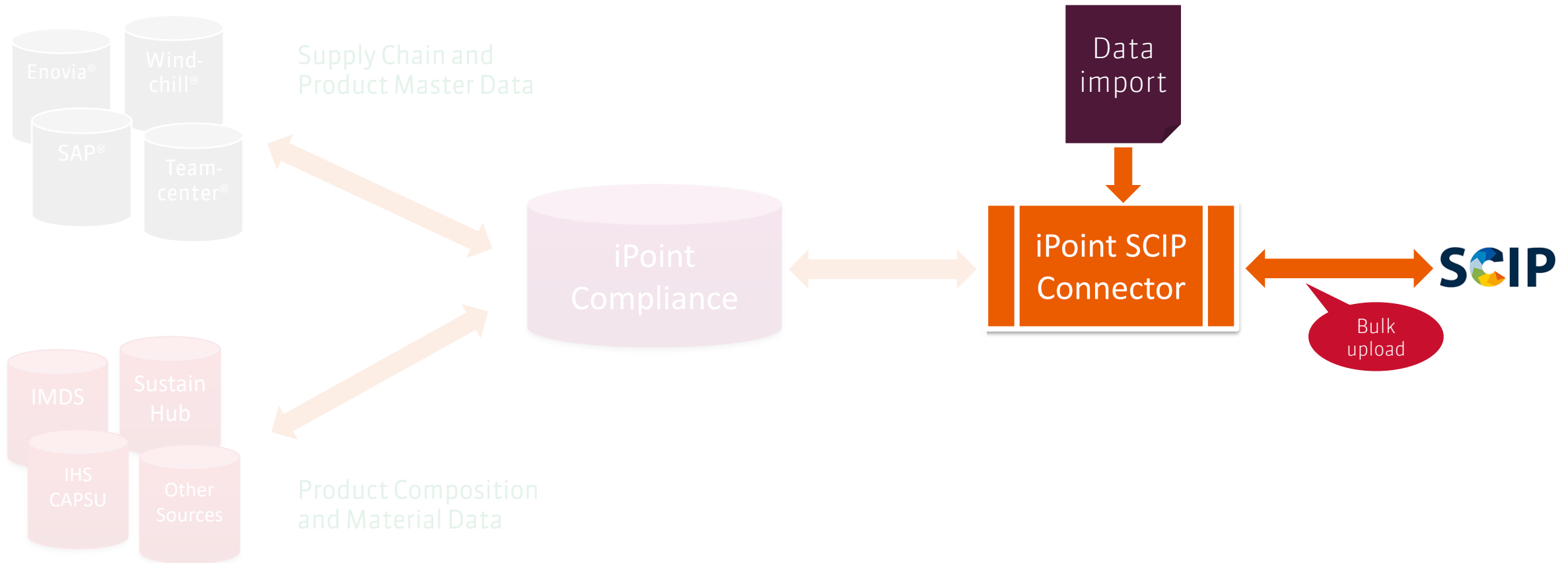
Using iPoint's Enterprise solution



Rough workflow for SCIP dossier submission



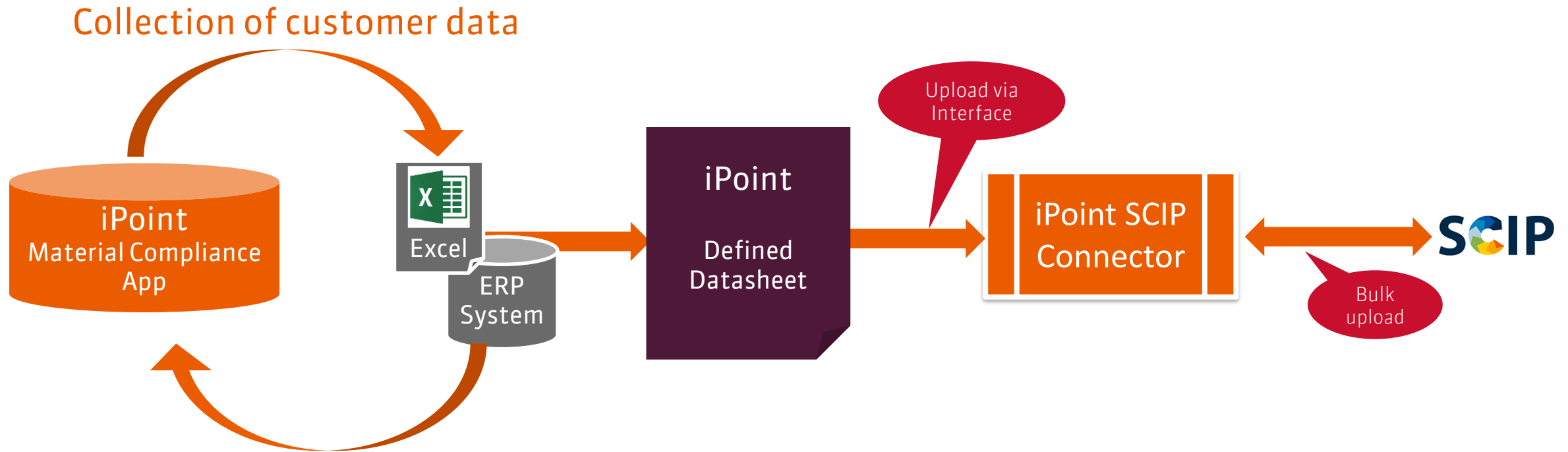
Using iPoint's Professional solution via data import



Rough workflow for SCIP dossier submission



Using iPoint's Professional solution via data import

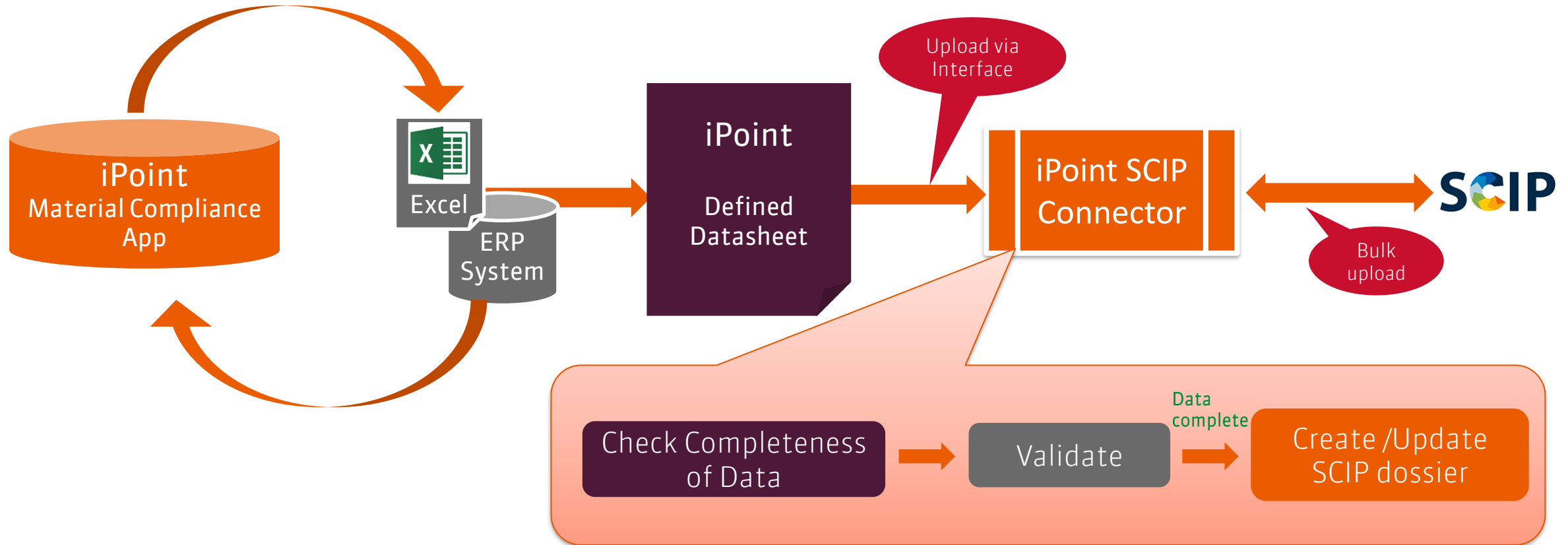


Rough workflow for SCIP dossier submission



Using iPoint's Professional solution via data import

Collection of customer data



SCIP Turn-key data management



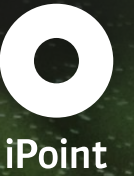
How does it work?

- **Software**
 - iPoint team will use our software to manage the service
- **SME Support**
 - Engage with iPoint SME for best practice advise
- **Data Management Team**
 - Engage iPoint Services team support for complete turn-key solution
 - Supplier training
 - Data collection
 - SCIP dossier submissions
 - Program Management for Business Units

Benefits

- **Reduces Time/Cost**
 - Eliminates manual processes
 - Makes Auditing Easier/Safer
- **Expert Advise**
 - Process and technical expertise
 - Best in class software
 - A positive impact on your company's compliance process
- **Provides Scalability**
 - Internal staff can focus on core responsibilities
 - Easily incorporate other BU/acquisitions
 - Provides a path for internal staff to take over the program later if desired

SCIP Database Connector Demo



An iterative approach to SCIP

Initial Submission – using available data

- Product Families
- Flat BOM
- Default values for:
 - Article Category (TARIC)
 - Production in Europe
 - Safe Use Information

Data collection
campaigns



1st submission refinement – case-by-case basis

- Key individual products (e.g. sales volume)
- Update due to changes in SVHC composition of product (including SVHC removal?)

2nd submission refinement

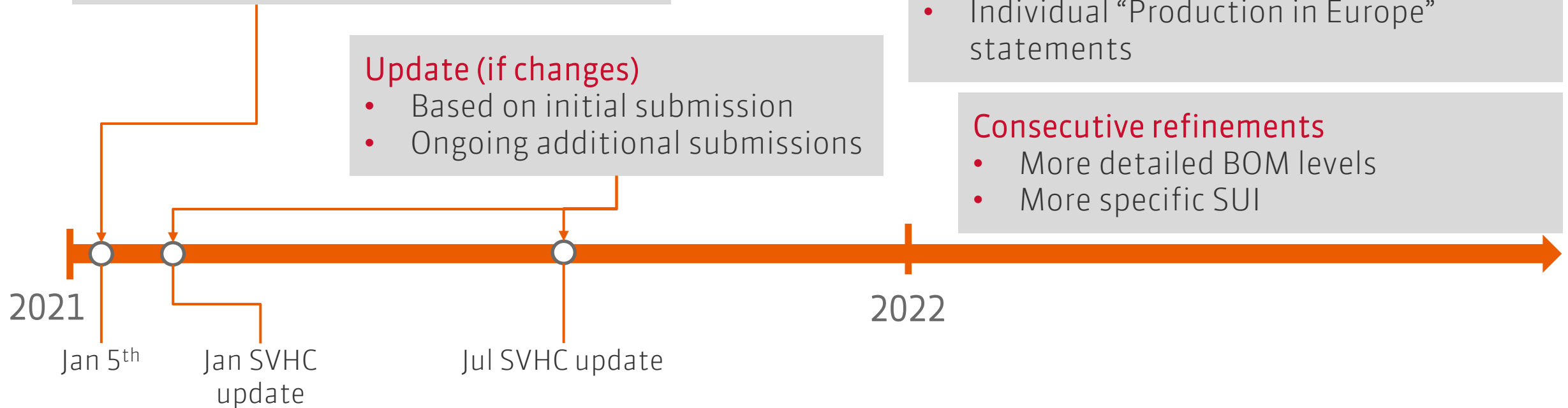
- Further individual products
- Individual “Production in Europe” statements

Consecutive refinements

- More detailed BOM levels
- More specific SUI

Update (if changes)

- Based on initial submission
- Ongoing additional submissions



Upcoming features



- Anonymize Article Descriptions
- Submissions based on grouping
- Re-use of internal/external reference submissions
- Workflow and Event Notification

Q&A



iPoint





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Senior Expert Compliance, Sustainability, and Innovation
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Sushma Kittali-Weidner

Director of Operations
iPoint Group

Thank you very much!



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