

CLP Annex VIII- Get ready to notify

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ABOUT RB- WHERE WE ARE

We are truly global

60

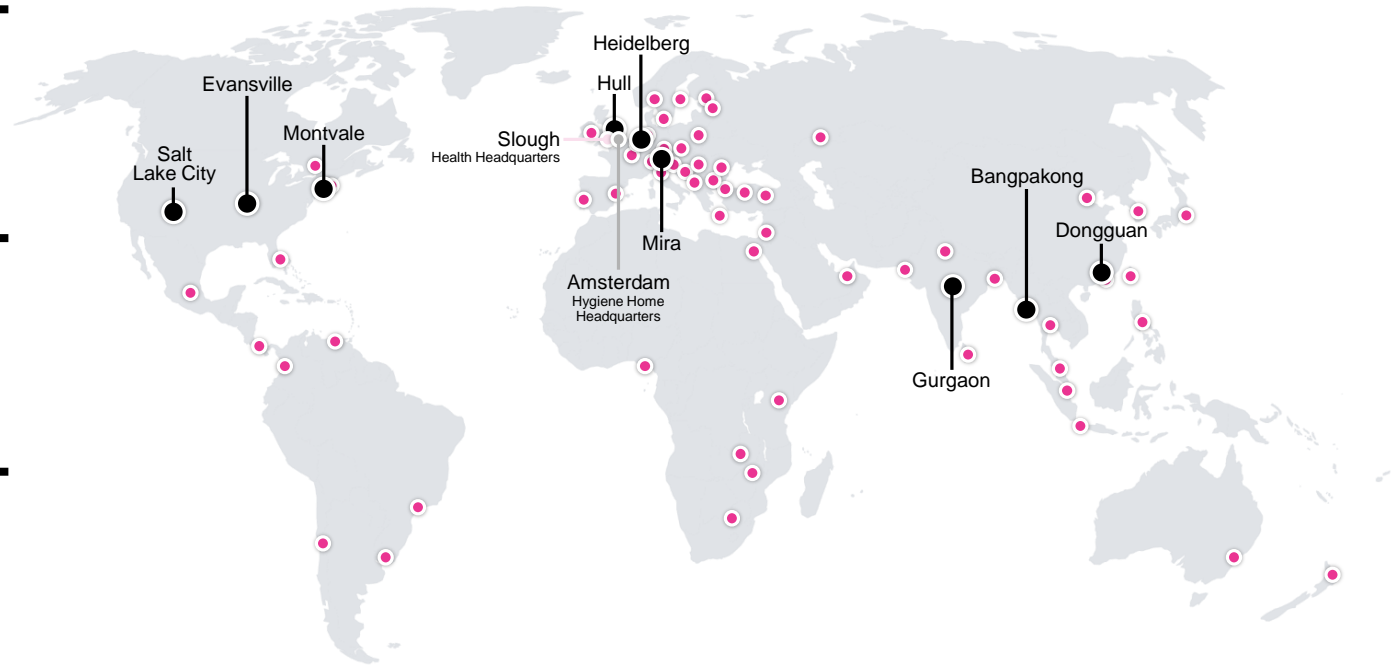
countries across six continents

190+

countries selling our products

09

Centres of Excellence

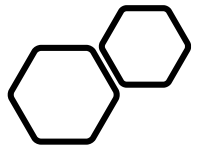


ABOUT RB- GOOD HOUSE

Long standing, trusted brands

1819

| | | | | | | | | | | | |
|-----------------------------|----------------------------|----------------------------|--------------------------------|------------------------------|----------------------------|-------------------------------|-------------------------------|-----------------------------|------------------------------------|-------------------------------|------------------------------|
| Mortein launched 1880 | Scholl launched 1904 | Harpic launched 1923 | Dettol launched 1932 | Air Wick launched 1943 | Finish launched 1953 | Strepsils launched 1958 | Clearasil launched 1959 | Nurofen launched 1983 | Mucinex launched 2002 | Enfinitas launched 2016 | Botanica launched 2020 |
| Lysol launched 1889 | Veet launched 1922 | Durex launched 1929 | Nutramigen launched 1942 | Woolite launched 1951 | Calgon launched 1956 | Enfamil launched 1959 | Gaviscon launched 1965 | Vanish launched 1983 | Cillit Bang launched 2004 | Neuriva launched 2019 | |



Agenda



RECAP ON ANNEX VIII



HISTORY AND LATEST
DEVELOPMENTS



NOTIFIER
PERSPECTIVE



CONCLUSIONS

Recap



Why do we notify

Addresses **accidental exposure** to hazardous chemicals at home and workplace

Crucial for medical staff to have **access to information** about chemicals



Key principles

Industry responsible to provide information about their products:

- to enable unambiguous identification (Name, UFI, etc)
- to enable appropriate response (Composition, Hazard information, etc)

Poison Centres responsible for processing the data and providing adequate response to public in case of emergency

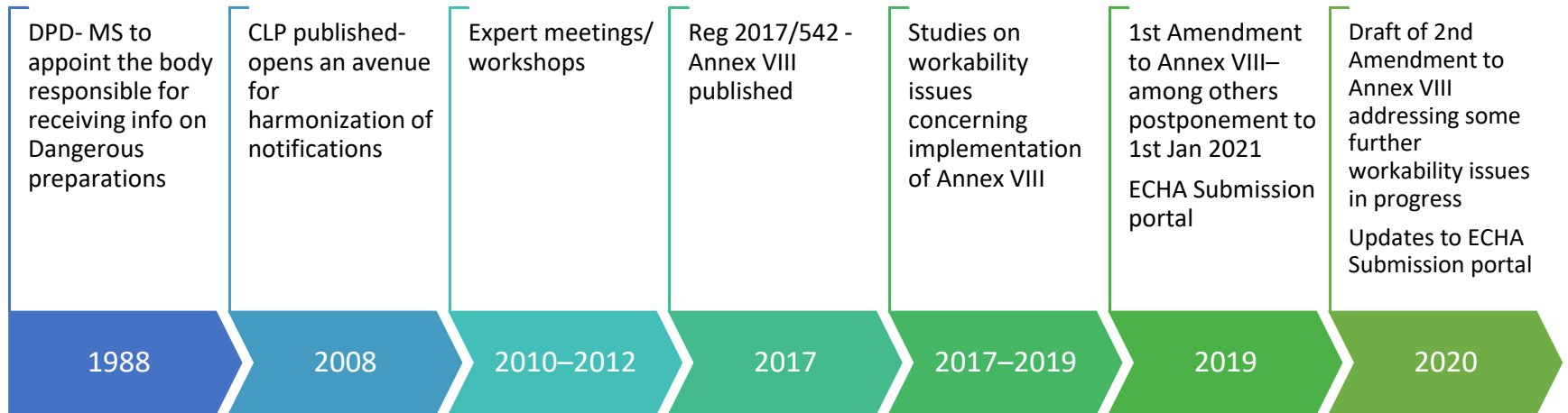


Deadlines

1st Jan 2021- Consumer and Professional mixtures

1st Jan 2024- Industrial mixtures

We have come a long way... but updates to ECHA submission portal still expected. Industry will need to adapt quickly



Selected workability issues for producers of detergents



Insufficient time

Insufficient time prior to deadline to assimilate notification tools

Postponement achieved to 1st Jan 2021 (1st amendment), however until 2nd amendment published, industry cannot fully complete notification infrastructure



Regular product variations

Technically equivalent raw materials from different sources used interchangeably.

Those RMs can vary in their discrete compositions

Very low disclosure thresholds introduced by Annex VIII would result in multiple & frequent changes of UFI for the same product

ICG concept (drafted in 2nd amendment) expected to help



UFI location

If required on all packaging layers would cause unnecessary complexity & not consumer relevant

Also flexibility required to print either on label or packaging itself

Positive clarity introduced in 1st amendment to Annex VIII



GPI

Generic Product identifier only for non hazardous fragrances. Most fragrances are hazardous within detergent industry

Increase number of components disclosed (Fragrance can contain up to 200 components), which will lead to more frequent PCN updates

Questions to ask



DO I UNDERSTAND ALL REQUIREMENTS?



DO I KNOW WHEN DO I NEED TO START COMPLYING?



ARE ALL MY PRODUCTS IMPACTED?



WHAT IS MY ROLE IN SUPPLY CHAIN AND RESULTING OBLIGATIONS?



DO I KNOW ALL NECESSARY INFORMATION ABOUT MY PRODUCTS?



DO I HAVE RIGHT SYSTEM/PROCESS TO HANDLE NOTIFICATIONS?



DO I HAVE FULL ACCESS TO MY MIXTURE CHEMISTRY?

Understanding requirements- hints

1st Jan 2021 or 1st Jan 2024?

- detergent Industry suppliers also need to comply by 1st Jan 2021, if applicable

Does all products need to be notified?

- Products excluded (Non classified, classified for environmental or supplemental hazard endpoints only)
- Voluntary submissions are possible

When do I need to amend notification?

- not all changes to notified information drive need for immediate updates (e.g. concentration within notified range, packaging, notifier address)
- UFI changes only when composition changes. Changes within permitted ranges are not considered a change.
- MiM UFI change only drives product UFI change if it was due to composition changes

Communicating within supply chain even more important

Verify

Verify if you have gaps in knowledge of your Raw materials full composition
Some suppliers claim trade secret
Check if supplier UFI is linked with Notification before you use it. Voluntary disclosure of UFI (without notification) on label is possible

Remember

Remember that use of MiM UFI is conditional:

- Disclosure of full composition of MiM takes precedence
- MiM must have been notified for your use type
- MiM must have been notified in all markets that you intend to sell

Engage

Sign NDAs to obtain compositional information
Obtain MiM UFI ahead of implementation deadlines, if needed
Discuss substance hazard classification within supply chain. Not all substance classifications are harmonized. This may limit your ability to use multiple sources interchangeably

Case 1

RM A is a non classified mixture. Supplier disclosed hazardous component downstream on voluntary basis

| Composition of RM | CLP classification | Conc. |
|-------------------|--------------------|-----------|
| Substance a, b, c | Not classified | Total 98% |
| Substance d | Skin Irritant | 2% |



Consumer product incorporate RM A. It is a hazardous product

| Composition of RM | CLP classification | Conc. |
|-------------------|--------------------|-------|
| Substance x | Not classified | 48% |
| Substance y | Eye Irritant | 50% |
| RM A | Not classified | 10% |

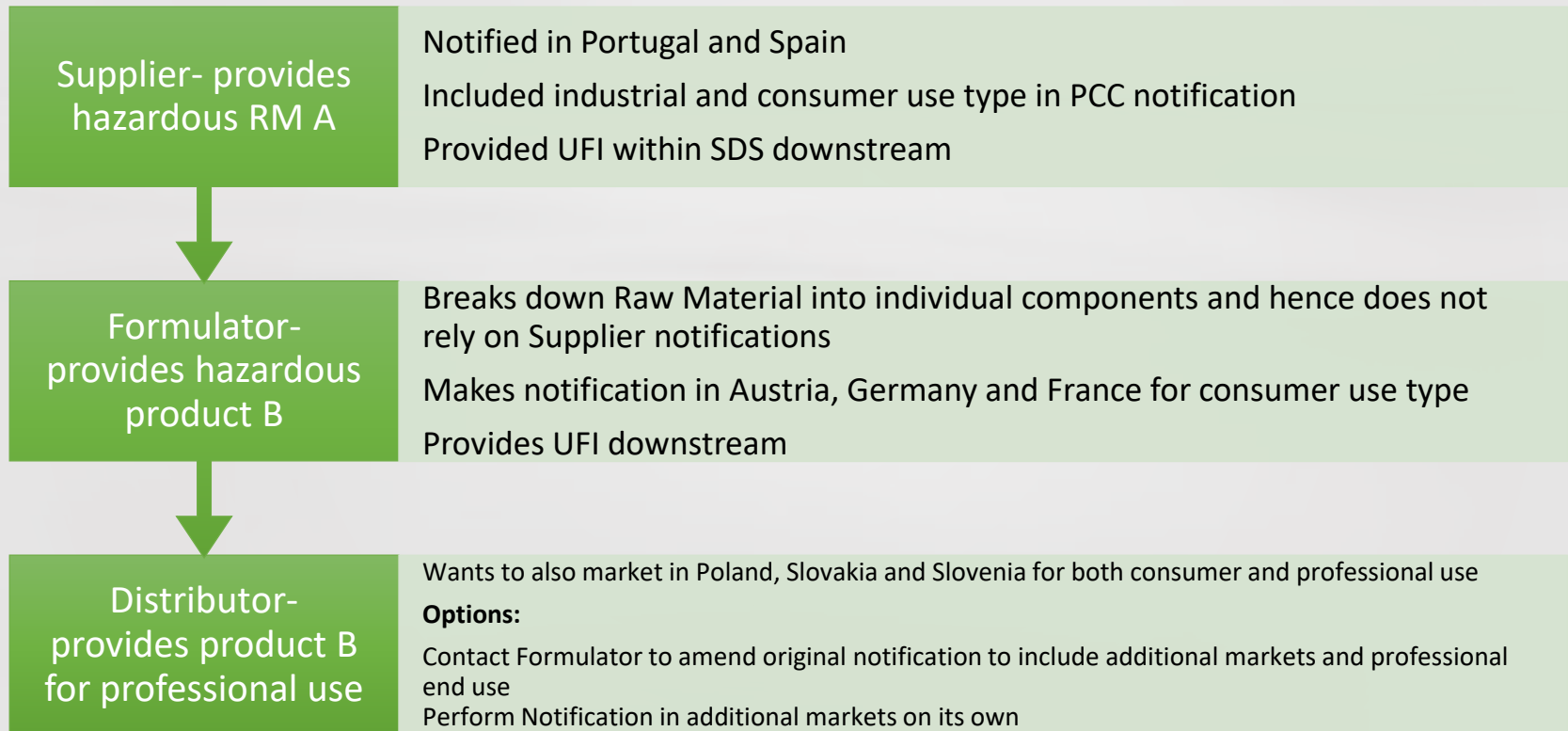
- **Supplier of RM A**

- has no obligation to notify PCC as RM A is not classified
- does not have to provide SDS for non classified RM
- claims Trade secret on the composition

- **DU Options:**

- According to Annex VIII rules formulator of consumer product need to include RM A breakdown in notification to PCC
- **Option 1:** Ask your supplier for voluntary PCC submission in all member states that you market your product. Use Supplier MiM UFI
- **Option 2:** Ask for SDS. For mixture that contains >1% of hazardous component Supplier should provide SDS on request. Disclose available information from SDS

Case 2



Make strategic decisions



UFI is considered supplemental label information and Annex VIII provides relative freedom as to its location and placement.

printed online on production line means investment in printing capabilities/ Quality check included as part of artwork means additional cost when UFI needs changing



It is industry responsibility to ensure notification data and UFI matches with their internal records

company integrated systems means significant investment
ECHA notification portal may mean loss of efficiency



Design a process fit for your organisation

Who will be responsible for generation of UFI and Notification forms
How the information will flow within your internal systems and departments

Start planning implementation early

- Map out your impacted products, those existing ones and future ones
- Make use of transitional period, where it makes sense
- Try to combine PCC notifications with other usual business activities (e.g. new product developments, formula reworks, rebranding activities)
- Good planning will minimise economic impact to your business, but also environmental waste



Conclusions



PCC NOTIFICATIONS
HARMONISATION WILL IMPROVE
EMERGENCY HEALTH RESPONSE



CENTRAL USE OF ECHA
NOTIFICATION PORTAL COULD
DRIVE SOME EFFICIENCY.
HOWEVER NUMBER OF
NOTIFICATION IS EXPECTED TO
SIGNIFICANTLY INCREASE



THE KEY FOR INDUSTRY IS TO
UNDERSTAND THE IMPACT,
ENGAGE WITH SUPPLIERS, ADAPT
SYSTEMS AND PROCESSES AND
PLAN IMPLEMENTATION IN
ADVANCE



Thank You