PPDC EPWG Meeting #5 Notes

Date: 3/11/2021

- May 13 initial report date- Need to finalize comments for this meeting
- Challenges with List N (Kristin Willis): supplemental product issues: resources and supplemental distributer labels (state monitored) not matching federal labels
- Questions/comments for Kristen-
- Is there a way for the registrants to own list N additions?
- Is there anything to learn from state monitoring departments for labels? (Software, etc.)
- Add a narrative to list N explaining that supplemental products exist?
- Seth suggests: Is it worth the agency's resources to make these types of improvements? If the registrants want supplemental products on list N they should get it registered.
- Elaine counters: The issue is the confusion of the end user.

Continuation of Charge Question #2

- What if any EPA guidance should have increased flexibilities to respond to supply chain challenges in times emergency etc?
- Expanding beyond list N
- Not just active ingredient, all inert ingredients as well were impacted
- Shortage of bottles/caps etc- could just be a manufacturing issue (expand beyond "emergency")
- Tajah- if we broaden too much, anything can be determined as an emergency focus on the pandemic and apply to other situations when time comes quantifiable emergency.
- Having a metric is an unbiased way to determine qualifications mock-up.
- Create a template for expedited reviews
- Who declares the pandemic: CDC (side note: EPA should post online)?
- Language in EVP triggering the EVP pandemic guidance
- Aftermath of EVP: Maybe another charge question? Build in pandemic type guidance to normal guidance
- Virus availability (time frame-late spring)
- Komal will create sub-categories so group can expand thoughts and have group incorporate thoughts on what categories should be