

Food Ingredients Stakeholder Forum Tuesday, August 4, 2009 USP Headquarters

Recommendations and Actions

Quality Standards for Medicines, Supplements, and Food Ingredients throughout the World

2. Strategic Directions

- USP needs to better understand the food industry
 Continue stakeholder dialogues
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- USP could provide industry with what CFR is lacking
- USP could review all GRAS Notified substances and assure they are included in FCC
- FCC could further expand to include products allowed outside of US
- Consider harmonizing with FAO and others
- Update FDA recognition of FCC



3. Food Additive Legislative Review

 The Dietary Supplement Verification Program should be made more well known to regulators, industry, and consumers



4. Reference Standards

- USP should consider providing a RS for known adulterants
- Because less complex methods are preferred by some companies, RS that are complex may be challenging
- Complex testing, however, can deter "bad actors."
- Can more than one analysis be listed for each RS; a more complex analysis and a less complex analysis
- Some customers don't know when RSs expire, and this is important especially when calculations are tied to a RS.



5. Stakeholder Roundtable

- See other slides, to be integrated into this presentation
- Consider survey stakeholder survey to help provide improvement feedback for future Stakeholder Forums



6. Adulteration





• USP encourages industry to include as much data as possible in monograph submissions.







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