
Tackling Environmental Science Issues

The Industry continues to push before the Food and Drug Administration (FDA) predictable policies and timelines in household regulations.

In February 2020, during a CLP dialogue with Sen. Koko Pimentel the FDA committed to fast-track the process with the caveat that registrants must submit complete documents. After that dialogue, speed in the processing was achieved. To further achieve a common undertaking of GHS compliance, CLP sponsored a webinar on GHS Bridging Principles.

One notable gain of CLP's advocacy with FDA is the decrease in quantity of Amount of Reference Standards from 1 gram to 250 milligrams, a request that FDA granted.