



COUNCIL OF PRODUCERS & DISTRIBUTORS OF AGROTECHNOLOGY

**APPLICATION
FOR
CPDA SPRAY ADJUVANT CERTIFICATION**

The undersigned party submits this application for certification of the designated product under the terms and conditions of the "Labeling and Performance Standards for Spray Adjuvants" adopted by the Council of Producers & Distributors of Agrotechnology.

PART 1

Please provide the following information:

1. Name of product: _____
- 2(a). Primary Certification Sub-Certification (check one)
- 2(b). If sub-certification, list the name of the Certified Product.

3. I have submitted the label of product to be certified. Yes
4. I have submitted a 1-2 page summary report of applicable acute toxicity studies. Yes
(Oral Toxicity, Dermal Toxicity, and Eye Irritation studies required)
5. I have submitted an SDS. Yes
6. Complete section below and pages 2 and 3 of this application form.

Licensee:

Company: _____ Application Date: _____

Address: _____

Submitted by: _____

Signature: _____

Title: _____

Phone: _____ Fax: _____ E-Mail: _____

Sub-Certification: (List name and address of sub-certifying company as they appear on the label.)

Name: _____ Address: _____

Part 2 of Application begins on Page 2

**PART 2
APPLICATION
For
CPDA SPRAY ADJUVANT CERTIFICATION**

1. Name of product: _____ Referenced Standards (S-1)
- 2(a). List functionality claim(s) defined in ASTM E-1519 or ASTM E-609.

- 2(b). List each ASTM Standardized Method(s) used to document functionality claims. (S-12)
 ASTM Method Used: _____ ASTM Method Not Available: _____
 ASTM Method Used: _____
 ASTM Method Used: _____
3. Product type (NIS, COC, etc.): _____
4. Is product labeled for, and/or intended for use on food crops? (S-3)
 Yes No
5. Are all product components listed in 40 CFR part 180 or applicable sections of 21 CFR? (S-3)
 Yes No
6. Is product labeling and packaging compliant with applicable U.S. DOT regulations? (S-4)
 Yes No
7. Does product contain OSHA regulated hazardous materials? (S-5)
 Yes No
 (If yes, acute toxicity studies are required for inhalation and/or skin corrosion/irritation as applicable to the nature of the hazard.)
8. Provide the name and phone number of 24 hour data service. (S-6)
 Name: _____ Phone: _____
9. Identify the Product Hazard Signal Word (based on acute toxicity studies) (S-7)(S-8)
 _____ Danger _____ Warning
- | Toxicity Study Category | (If required based on Item 7 response) |
|--------------------------------|---|
| Oral Toxicity: _____ | Inhalation Toxicity: _____ |
| Eye Irritation: _____ | Skin Corrosion: _____ |
| Dermal Toxicity: _____ | Skin Irritation: _____ |
10. Applicable precautionary and safety statements included on the label? (S-7)(S-9)
 Yes
11. Does product label include recommendation for use in aquatic applications? (S-10)
 Yes No
 (If yes, aquatic toxicity data are required.)
12. The Principal Functioning Agent statement lists only formulation components that contribute to the claimed functions of the product? (S-11)
 Yes

Part 2 continues on Page 3

PART 2 Continued

13(a). If intended functionality includes reference to surfactant, spreader, or wetting agent, list % surfactant guarantee. _____% N/A (S-13)
(Must be separately listed on the label)

13(b). Each formulation component from 13(a) is water dispersible and reduces the surface tension of water to 45 dynes/cm or less at 0.5% w/w per ASTM method D-1331? Yes (S-13)

14. Describe the type(s) of oil utilized. (If two or more types are used, list them in descending order of their compositional content.) 1 _____ 2 _____ N/A 3 _____ 4 _____ (S-14)

Provide the Un sulfonated Oil Residue (UR) Value. _____% N/A (S-15)

15. All state labels bearing the Certification Mark must be identical to the approved national label with respect to all applicable Certification Standards.

(a) Is this product currently registered in any state?

Yes No

(b) Will this product bear the Certification Mark on the registered state label(s)?

Yes No

(If yes to 16(b), then the state label(s) must be included in the application materials.)*

CPDA reserves the right to request any additional data or information it deems necessary to complete the certification process.

*The Certification Mark cannot be used on any label that has not been reviewed by the Certification Committee. If a product receives certification, existing state labels must be reviewed by the Committee prior to the mark being added to the state label(s). If a product has been certified and will bear new state labels in the future, the Certification Mark cannot be used on the label until the label is reviewed by the Committee.

Return completed form to CPDA