# Certificate of Compliance

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**Code:** 20395PF
**INCI Name:** Butylene Glycol & Water & Spiraea Ulmaria Extract
**INCI Status:** Conforms
**CAS #:** 107-88-0 & 7732-18-5 & 84775-57-5
**EINECS #:** 203-529-7 & 231-791-2 & 283-866-3
**China NMPA #:** 168810-02809-1313

**Below is a list of processing aids used, but not declared on the ingredient label:**

| INCI Name | CAS# | EINECS# | Percentage (%) | Function |
|---|---|---|---|---|
| Lactobacillus Ferment | 1686112-36-6 (or) 68333-16-4 | N/A (or) N/A | 2.00% | Natural Antimicrobial |

The following information on regulatory clearances is believed to be accurate and is given in good faith as a guide to a global use of our ingredients in cosmetic applications. No representation or warranty as to its competences or accuracy is made. Information is offered for use in general cosmetic applications and may vary in particular applications. Users are responsible for determining the suitability of these products for their own particular use. All regulatory decisions should be made on the advice of your regulatory group or legal counsel.

| Country / Regulatory Body | Status of Product |
|---|---|
| EU (CosIng) | Compliant |
| USA (TSCA) | Compliant |
| Australia (AICS) | Compliant |
| Japan (METI) | Compliant |
| Canada (DSL) | Compliant |
| China (IECIC) | Compliant |
| Brazil (ANVISA) | Compliant |
| Korea (KECI) | Compliant |
| Philippines (PICCS) | Contact Us |
| Mexico (COFEPRIS) | Compliant |




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# AC Sebum Control Enzyme PF
## Code: 20395PF

Attention must be paid to the use of AC Sebum Control Enzyme PF in the equivalent of OTC formulations (eg. quasi-drugs in Japan, or therapeutic goods in Australia). Some countries maintain restricted inventories of raw materials that can be used in those applications so more detailed guidance may be required.

AC Sebum Control Enzyme PF and its components and impurities are in compliance with the rules governing cosmetic products in the European Union (Directive 76/768/ECC & Regulation No. 1223/2009). The recommended use levels for AC Sebum Control Enzyme PF is 2.00 – 5.00%.

The Nagoya Protocol provides a scheme for the fair and equitable sharing of benefits derived from Genetic Resources. Information regarding the Nagoya Protocol and Access and Benefit Sharing (ABS) is available at https://www.cbd.int/abs/. The agreement focusses on wild taxa and excludes most commercially cultivated crops. For the signatories to the agreement, responsibility for Benefit Sharing falls on the entity exporting or extracting the resource from the signatory country. Active Concepts audits its suppliers to conform compliance with the Nagoya Protocol where applicable.

AC Sebum Control Enzyme PF is considered a non-hazardous material. All significant toxicological routes of absorption have been considered as well as the systemic effects and margin of safety (MoS) based on a no observed adverse effects level (NOAEL). Due to the restriction placed on animal testing of cosmetic raw materials, and Active Concepts, LLC's internal non-animal testing policy, this product was not tested for NOAEL.

AC Sebum Control Enzyme PF was tested using *in vitro* dermal and ocular irritation models. This product was found to be non-irritating in both models.

To our knowledge the above material is free of CMR (*) substances, as defined according to Regulation (EC) No 1272/2008 and Cosmetic Regulation (EC) No 1223/2009 as amended. Products supported for Personal Care applications will not be classified as CMR (*), as defined by (EC) 1272/2008 on the Classification, Labelling and Packaging of Substances and Mixtures, unless supported by a positive SCCS opinion.

(*) Carcinogenic, Mutagenic, toxic for Reproduction

Active Concepts, LLC certifies that AC Sebum Control Enzyme PF is produced in compliance with the programs/regulations set by the following agencies in the State of California:

- California Proposition 65
- California Safe Cosmetics Act (SB 484)

Active Concepts, LLC certifies that AC Sebum Control Enzyme PF does not contain any materials prohibited by Halal laws.

As of February 3, 2026, AC Sebum Control Enzyme PF does not contain any substances present on the so called "Candidate List" of Substances of Very High Concern (SVHC) provided by the European Chemicals Agency (ECHA). We further certify that this material has not been manufactured using any of the species listed in the CITES Appendices as of February 3, 2026.

Active Concepts, LLC certifies that AC Sebum Control Enzyme PF does not contain Diethylene Glycol (DEG) and Ethylene Glycol (EG) up to the threshold limit indicated in the USP 38 monograph (0.10% maximum).

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# AC Sebum Control Enzyme PF
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AC Sebum Control Enzyme PF is REACH Compliant and the following ingredients are not intentionally added or produced, so we do not expect trace amounts to be present:

- 1,4-Dioxane
- Alcohol/ethanol
- Aluminum
- Ammonia
- Butylated hydroxyanisole (BHA)
- Butylated hydroxytoluene (BHT)
- Butylphenyl methylpropional (Lilial)
- Conflict minerals
- Dyes
- Formaldehyde or formaldehyde donors
- Gluten
- Lactose
- Latex
- Melamine
- Microplastics
- Nanoparticles
- Nitrosamines
- Palm oil/palm kernel oil (or derivatives
- Para-aminobenzoic acid (PABA)
- Parabens
- Paraffin
- Per- and polyfluoroalkyl substances (PFAS)
- Pesticide residues
- Phthalates
- Polyethylene glycol (PEG)
- Residual solvents
- Salicylates
- Silicone
- Sulfites
- Sulfates
- Volatile organic compounds

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# Raw Component Regulations

Please note that the below are global regulations for the raw materials used to manufacture AC Sebum Control Enzyme PF and are not for the product itself.

AC Sebum Control Enzyme PF contains 50.00% Butylene Glycol. See below for a list of regulations:

Butylene Glycol:

- USA: Maximum Authorized Concentration: Safe up to 89.00%
  *Journal Citation: JACT 4(5):223-48, 1985 confirmed 02/04 IJT 25(S2), 2006