



# Cosmetic Product Safety Report

## Kohana Ombre Gel

Responsible person:

Kohana LTD

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This document has been prepared in compliance with:

- Regulation (EC) No 1223/2009 regarding cosmetic products
- COMMISSION IMPLEMENTING DECISION of 25 November 2013 on Guidelines on Annex I to Regulation (EC) No 1223/2009 of the European Parliament and of the Council on cosmetic products.
- SCCS Notes of Guidance for the testing of cosmetic ingredients and their safety evaluation - 12th revision – SCCS/1647/22

## Qualitative and quantitative composition of the cosmetic product

### Qualitative composition

**Ingredients:** Acrylates/Carbamate Copolymer, Acrylates/VP Copolymer, Acrylates Copolymer, +/-: CI 77891, CI 77491, CI 77492, CI 77499, CI 77742

**The quantitative composition of the cosmetic product, including chemical identity of the substances (INCI, CAS, EINECS/ELINCS, where possible) and their intended function.**

Trade name	INCI	Function	Concentration [%]	CAS NO	Annex/Ref
Acrylates/Carbamate Copolymer	Acrylates/Carbamate Copolymer	Conditioning	65	24938-16-7	-
Acrylates/VP Copolymer	Acrylates/VP Copolymer	Bonding, film forming	32	26589-26-4	-
Acrylates Copolymer	Acrylates Copolymer	Film forming	1	25035-69-2	-
CI77891	CI 77891	Colorant	1,2	13463-67-7	IV/143
CI77491	CI 77491	Colorant	0,1	1345-27-3	IV/135
CI77492	CI 77492	Colorant	0,2	51274-00-1	IV/136
CI77499	CI 77499	Colorant	0,2	12227-89-3	IV/137
CI77742	CI 77742	Colorant	0,3	10101-66-3	IV/140

### Description of the name and code number of the composition and the identity of the supplier.

This formulation does not contain a synthetic fragrance and therefore a fragrance safety evaluation as per IFRA code of practice is not applicable to this product.

### Undesirable effects and serious undesirable effects

Kohana Ombre Gel is a product placed on the market. In accordance with the requirements set out in Art. 23 of Regulation EC 1223/2009, the responsible person is obliged to document and report an adverse reaction caused by a cosmetic product. Information on the adverse reaction must be kept up to date and made available to the safety assessor for possible changes to the safety report.