

Customer: .....

To: Chelab srl

**REQUEST**

We ask for the execution of an in use test for the following samples on 20 volunteers.  
The aim is to assess the product compatibility with human skin - intended as absence of skin irritation when applying the product for the first time - and not to define the intrinsic irritant potential of the tested product.

**SAMPLE IDENTIFICATION**

Sample 1	batch nr.
Sample 2	batch nr.
Sample 3	batch nr.
Sample 4	batch nr.
Sample 5	batch nr.
Sample 6	batch nr.
Sample 7	batch nr.
Sample 8	batch nr.
Sample 9	batch nr.
Sample 10	batch nr.
Sample 11	batch nr.
Sample 12	batch nr.
Sample 13	batch nr.
Sample 14	batch nr.
Sample 15	batch nr.
Sample 16	batch nr.
Sample 17	batch nr.
Sample 18	batch nr.
Sample 19	batch nr.
Sample 20	batch nr.
Sample 21	batch nr.
Sample 22	batch nr.
Sample 23	batch nr.
Sample 24	batch nr.
Sample 25	batch nr.
Sample 26	batch nr.
Sample 27	batch nr.
Sample 28	batch nr.
Sample 29	batch nr.
Sample 30	batch nr.

The products are delivered:

- in the selling package complete of the ingredient INCI declaration, intended use and application
- in packages different from the selling one; in this case please enclose:
  - o intended use and application (specify if it is a “leave on” or “rinse off” product and if the quantity of surfactants is superior or inferior to 10%)
  - o the ingredient list according to the INCI list in order of decreasing weight

Chelab keeps the remaining product for 3 months after the report emission. The analytical results refer only to the tested sample.

### DECLARATION

**The undersigned – authorized by the company..... declares that:  
the product and its components:**

- **do not contain any substance which is forbidden by the EC legislation in force as far as the use of cosmetic and personal hygiene products is concerned.**
- **have been analysed by a qualified expert to guarantee that the existing information on the product and its components justify the exposure on man**
- **any significant risks for the volunteers are to be expected in the offered study conditions and at the expected exposure level**
- **the aim of the study is only to confirm those conclusions.**

We agree that the testing procedure shall be immediately interrupted if it is necessary to preserve users from possible adverse risks; in this case we will be informed of the interruption.

Date

.....

Customer

.....  
(stamp & signature)