

## Column E Explanation

This form is intended as an aid to completing the Column E explanation. It is not an official form and its use is voluntary. Names, addresses, protocols, veterinary care programs, and the like, are not required as part of an explanation. A Column E explanation must be written so as to be understood by lay persons as well as scientists.

1. Registration Number. 33-R-0090
2. Number 128 of animals used in this study.
3. Species (common name) Rabbit (NZW) of animals used in the study.
4. Explain the procedure producing pain and/or distress.  
 Acute dermal Toxicity and D.O.T. Corrosivity Route of administration is topical. 10% of body surface dosing site is occluded for 4 to 24 hours. Sometimes topical irritation is observed. Maximum duration of study is 14 days.
5. Provide scientific justification why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results. (For Federally mandated testing, see Item 6 below)  
 The test is intended to determine if the test compound is absorbed by the skin and if so will it cause a toxic reaction other than mild irritation.
6. What, if any, federal regulations require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g., APHIS, 9 CFR 113.102):

Agency EPA 40 CFR 160 and 792  
Department of Transportation

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1. Registration Number: 33 R-0090
2. Number 135 of animals used in this study.
3. Species (common name) Rabbit (NZW) of animals used in the study.
4. Explain the procedure producing pain and/or distress.  
 Eye Irritation Rabbit.  
 Test compound is instilled into the rabbit eye.  
 The eye is anesthetized (Tetracaine) prior to testing.  
 Maximum duration of study is 21 days.  
 Severe irritants or corrosive compounds are not tested.
5. Provide scientific justification why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results. (For Federally mandated testing, see Item 6 below)  
 Local anesthesia is used to eliminate discomfort or distress.
6. What, if any, federal regulations require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g., APHIS, 9 CFR 113.102):

Agency EPA CFR 40 CFR 160 and 792