

Draft Guidance on Adapalene

This draft guidance, when finalized, will represent the current thinking of the Food and Drug Administration (FDA, or the Agency) on this topic. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations. To discuss an alternative approach, contact the Office of Generic Drugs.

Active ingredient: Adapalene

Dosage Form; Route: Lotion; topical

Recommended studies: One study

Type of study: Bioequivalence (BE) Study with clinical endpoint

Design: Randomized, double blind, parallel, placebo controlled, in vivo

Strength: 0.1%

Subjects: Males and nonpregnant females with acne vulgaris

Additional comments: Specific recommendations are provided below

Analytes to measure (in appropriate biological fluid): Not Applicable (N/A)

Bioequivalence based on (90% CI): Clinical endpoint

Waiver request of in vivo testing: N/A

Dissolution test method and sampling times: N/A

Additional comments regarding the BE study with clinical endpoint:

1. The Office of Generic Drugs (OGD) recommends conducting a BE study with a clinical endpoint in the treatment of acne vulgaris comparing adapalene lotion, 0.1% test product versus the reference listed drug (RLD) and placebo (vehicle) control, each administered as one application once a day for 12 weeks.
2. The recommended two primary endpoints of the study are: 1) mean percent change from baseline to Week 12 (Day 84) in the inflammatory (papules and pustules) lesion count and 2) mean percent change from baseline to Week 12 in the non-inflammatory (open and closed comedones) lesion count.
3. The protocol should clearly define papules, pustules, open comedones, closed comedones, nodules and cysts. When counting facial acne lesions, it is important that all lesions be counted, including those present on the nose. Counts of nodules and cysts should be reported separately and not included in the inflammatory or non-inflammatory lesion counts.

4. A placebo (vehicle) control arm is recommended to demonstrate that the test product and RLD are active and as a parameter that the study is sufficiently sensitive to detect differences between products.
5. Subjects should be instructed to cleanse the face with a mild or soapless, non-medicated cleanser, pat dry and then apply a thin layer of the product to the entire face, avoiding contact with the eyes, lips, angles of the nose, and mucous membranes and washing hands before and after applications. Subjects should be instructed to not apply the product to cuts, abrasions, eczematous skin, or sunburned skin, not apply the product more than once daily, not use more than the recommended amount and not use “waxing” as a depilatory method on skin treated with the product.
6. Subjects should not apply moisturizers, new brands of make-up, creams, lotions, powders or any topical product other than the assigned treatment to the treatment area. Subjects should be instructed to minimize exposure to sunlight, including sunlamps, while using the product. Use of sunscreen products and protective clothing over treated areas is recommended when sun exposure cannot be avoided.
7. Inclusion Criteria (the sponsor may add additional criteria)
 - a. Male or nonpregnant female aged ≥ 12 and ≤ 40 years with a clinical diagnosis of acne vulgaris.
 - b. On the face, ≥ 25 non-inflammatory lesions (i.e., open and closed comedones) AND ≥ 20 inflammatory lesions (i.e., papules and pustules) AND ≤ 2 nodulocystic lesions (i.e., nodules and cysts).
 - c. Investigator’s Global Assessment (IGA) of acne severity Grade 2, 3, or 4 (per Table 1).

Table 1. Sample IGA Scale for Acne Vulgaris¹

Grade	Description
0	Clear skin with no inflammatory or noninflammatory lesions
1	Almost clear; rare noninflammatory lesions with no more than one small inflammatory lesion
2	Mild severity; greater than Grade 1; some noninflammatory lesions with no more than a few inflammatory lesions (papules/pustules only, no nodular lesions)
3	Moderate severity; greater than Grade 2; up to many noninflammatory lesions and may have some inflammatory lesions, but no more than one small nodular lesion
4*	Severe; greater than Grade 3; up to many noninflammatory lesions and may have some inflammatory lesions, but no more than a few nodular lesions

* The Case Report Forms for acne studies can allow for reporting by investigators of lesion worsening beyond Grade 4 with treatment. It is recommended that enrollment of acne vulgaris subjects not include subjects with nodulocystic acne. Subjects who worsen beyond Grade 4 are to be described in the safety evaluation.

- d. Willing to refrain from use of all other topical acne medications or antibiotics during the 12-week treatment period.
 - e. If female of childbearing potential, willing to use an acceptable form of birth control during the study.
8. Exclusion Criteria (the sponsor may add additional criteria)
- a. Pregnant, breast feeding or planning a pregnancy.
 - b. Presence of any skin condition that would interfere with the diagnosis or assessment of acne vulgaris (e.g., on the face: rosacea, dermatitis, psoriasis, squamous cell carcinoma, eczema, acneform eruptions caused by medications, steroid acne, steroid folliculitis, or bacterial folliculitis).
 - c. Excessive facial hair (e.g. beards, sideburns, moustaches, etc.) that would interfere with diagnosis or assessment of acne vulgaris.
 - d. History of hypersensitivity or allergy to adapalene, retinoids and/or any of the study medication ingredients.
 - e. Use within 6 months prior to baseline or during the study of oral retinoids (e.g. Accutane®) or therapeutic vitamin A supplements of greater than 10,000 units/day (multivitamins are allowed).
 - f. Use for less than 3 months prior to baseline of estrogens or oral contraceptives; use of such therapy must remain constant throughout the study.
 - g. Use on the face within 1 month prior to baseline or during the study of 1) cryodestruction or chemodestruction, 2) dermabrasion, 3) photodynamic therapy, 4) acne surgery, 5) intralesional steroids, or 6) x-ray therapy.

¹ U.S. Department of Health and Human Services, Food and Drug Administration, Center for Drug Evaluation and Research. Draft Guidance for Industry: Acne Vulgaris: Developing Drugs for Treatment. Clinical/Medical. September 2005. Accessed at <http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM071292.pdf>

- h. Use within 1 month prior to baseline or during the study of 1) systemic steroids, 2) systemic antibiotics, 3) systemic treatment for acne vulgaris (other than oral retinoids, which require a 6-month washout), or 4) systemic anti-inflammatory agents.
 - i. Use within 2 weeks prior to baseline or during the study of 1) topical steroids, 2) topical retinoids, 3) topical acne treatments including over-the-counter preparations, 4) topical anti-inflammatory agents, or 5) topical antibiotics.
9. The protocol should include a list of the prescription and over-the-counter drug products, procedures, and activities that are prohibited during the study, such as:
 - a. Any other topical products applied to face.
 - b. Medicated soaps used on face.
 - c. Spironolactone.
 - d. Oral retinoids, therapeutic vitamin A supplements of greater than 10,000 units/day (multivitamins are allowed) or other systemic treatment for acne vulgaris.
 - e. Systemic (e.g., oral or injectable) antibiotics.
 - f. Systemic steroids, systemic anti-inflammatory agents or immunosuppressive drugs.
 - g. Antipruritics, including antihistamines, within 24 hours of study visits.
 - h. Use on the face of 1) cryodestruction or chemodestruction, 2) dermabrasion, 3) photodynamic therapy, 4) acne surgery, 5) intralesional steroids, or 6) x-ray therapy.
 - i. Use of hormonal contraceptives should not be initiated or changed during the study.
 - j. Use of tanning booths, sunbathing, or excessive exposure to the sun.
10. To establish bioequivalence, the 90% confidence interval of the test/reference ratio of the mean percent change from baseline to Week 12 in the inflammatory (papules and pustules) lesion counts and in the non-inflammatory (comedones) lesion counts should be contained within [0.80, 1.25], using the per protocol (PP) population.
11. The dichotomized IGA severity scale should be treated as a secondary endpoint for supportive evidence. This secondary endpoint should be evaluated as the proportion of subjects with a clinical response of “success” at Week 12. Success should be defined as an IGA score that is at least 2 grades less than the baseline assessment. Failure should be defined as an IGA score that is the same, higher or one grade lower than the baseline assessment.
12. Application site reactions such as erythema, dryness, burning/stinging, erosion, edema, pain and itching are to be recorded at each visit to allow a comparison between treatment groups. A descriptive analysis comparing the application site reactions for each treatment group is recommended. It is important to ensure that the test product is not worse than the reference product regarding the expected and unexpected application site reactions.
13. These recommendations are specific to this product and may not be appropriate for bioequivalence studies of any other product, including any other dosage form or strength of adapalene.
14. Please refer to the product-specific guidance on Adapalene; Benzoyl peroxide topical gel 0.3%; 2.5% (“[Draft Guidance on Adapalene; Benzoyl Peroxide](#)”) for a recommended

approach to statistical analysis and study design for bioequivalence studies with clinical endpoints.

15. Study data should be submitted in a standardized format. Please refer to the study data standards published at www.fda.gov²

² Study Data Standards for Submission to CDER and CBER available at:
<https://www.fda.gov/Drugs/DevelopmentApprovalProcess/FormsSubmissionRequirements/ElectronicSubmissions/ucm248635.htm>