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Food and Drug Administration
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Re: Notice of Withdrawal / Suitability Petition and Amendment (FDA Docket No. 2006P-0369; Regulations.gov ID No. FDA-2006-P-0514)
ANDA Suitability Petition for Alclometasone Dipropionate Lotion, 0.05% and
Subsequent Amendment

Dear Sir or Madam:

We are writing to formally withdraw the above-referenced Suitability Petition for Alclometasone Dipropionate Lotion, 0.05% (FDA Docket No. 2006P-0369; Regulations.gov ID No. FDA-2006-P-0514). A Suitability Petition was filed on September 1, 2006 requesting that the Commissioner of the Food and Drug Administration (FDA) make a determination that a new dosage form of a topical alclometasone dipropionate drug product, Alclometasone Dipropionate Lotion, 0.05%, is suitable for filing under an abbreviated new drug application (ANDA). An amendment requesting a waiver from the pediatric assessment requirements of the Pediatric Equity Act was subsequently filed on November 30, 2006.

The requested determinations are no longer needed by our client. For this reason, we respectfully request that the submitted Suitability Petition and subsequent Amendment be withdrawn.

Please do not hesitate to contact us with any questions.

Sincerely,



Rebecca L. Dandeker

FDA-2006-P-0514

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