

# DR·AUGUST·WOLFF



## **Clinical Trial Synopsis**

The presented information is the property of Dr. August Wolff GmbH & Co.KG Arzneimittel. Commercial use or reproduction of all or part of this document is prohibited without prior written permission.

The content and information given in this clinical trial synopsis does not replace medical advice provided by a patient's physician or other qualified healthcare providers. Patients should always consult their physician before making conclusions regarding their treatment. Healthcare professionals are advised to refer to the approved labelling and prescribing information. Results from a single study may not reflect the overall results of a drug and need to be interpreted in the context of all relevant scientific data.

<b>Name of Sponsor/ Company:</b> Dr. August Wolff GmbH & Co. KG Arzneimittel	Individual Study Table Referring to Part of the Dossier  Volume:  Page:	<i>(For National Authority Use only)</i>
<b>Name of Finished Product:</b> Alpicort F		
<b>Name of Active Ingredient:</b> Estradiolbenzoate / Prednisolone / Salicylic acid		
<b>Title of study:</b>  A Phase III, single-center, randomized, double-blind, parallel group, placebo-controlled study of Alpicort F in the treatment of chemotherapy induced alopecia in women.		
<b>Identifiers:</b>  Sponsor study code: AFCT-14/2009  EudraCT number: 2009-017415-16  NCT number: -		
<b>Study Status:</b> This study was cancelled with no patients enrolled.		