

	<b>Study Number:</b> D6130C00003	<i>Confidential Document</i>
		<b>As applicable:</b> <b>Trial Master File Zone:</b> Trial Management <b>Artifact:</b> 1. 1. 10

**Title/Reference:** A Phase I/II, Open-Label, Multicentre 2-Part Study to Assess the Safety, Tolerability, Pharmacokinetics, and Efficacy of AZD2811 as Monotherapy or in Combination in Treatment-Naïve or Relapsed/Refractory Acute Myeloid Leukaemia Patients Not Eligible for Intensive Induction Therapy

**Topic:** Results not posted on the EudraCT Registry

**Summary:** The above mentioned study has been submitted to the authorities, however the study was stopped without any patients being recruited, therefore results will not be posted in the EudraCT registry.

Anneke Ravensbergen, Group Director Study Operations

Full Name, Position

Date

DocuSigned by Anneke Ravensbergen



I am the author of this document  
17 January 2022 | 08:47 AST

F646F5C31FEA49C4B953A354FC4B32D3

17 January 2022 | 08:47 AST

**Certificate Of Completion**

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Anneke Ravensbergen anneke.ravensbergen@acerta-pharma.com Group Associate Director Study Operations Security Level: Email, Account Authentication (Required)	  Signature Adoption: Pre-selected Style Signature ID: F646F5C3-1FEA-49C4-B953-A354FC4B32D3 Using IP Address: 147.161.172.170  With Signing Authentication via DocuSign password With Signing Reasons (on each tab): I am the author of this document	Sent: 1/13/2022 2:31:15 PM Viewed: 1/13/2022 4:01:56 PM Signed: 1/17/2022 12:47:22 PM

**Electronic Record and Signature Disclosure:**

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ID: 619990e1-7e85-45b6-b1f8-5995d12c9362

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**Editor Delivery Events**

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**Certified Delivery Events**

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Payment Events	Status	Timestamps
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