

<b>Scientific Title</b>	Recommendations for risk adapted treatment of acute promyelocytic leukemia in the AML-SAL-Study group - an observational study -
<b>Short Title</b>	AIDA2009
<b>Id KN/ELN</b>	LN_SAL_2010_370
<b>Trialgroup</b>	SAL
<b>Type of Trial</b>	multicentric
<b>Disease</b>	Acute myeloid leukemia( AML) FAB M3 (APL)
<b>Stage of Disease</b>	de novo/non-treated
<b>Aim</b>	<ul style="list-style-type: none"><li>- registration of number and basic characteristics of patients, remission rates, toxicity and survival</li><li>- long term remission, tolerance</li></ul>
<b>Outcomes</b>	<ul style="list-style-type: none"><li>- CR-Rate, Remissionsdauer, Überleben CR rates, duration of remission, survival (Primary Outcome)</li><li>- Toxizität toxicity</li></ul>
<b>Inclusion Criteria</b>	<ul style="list-style-type: none"><li>- Diagnosis of acute promyelocytic leukemia, based on morphological result and confirmatory cytogenetic/molecular finding of t(15;17) and/or PML/RAR</li><li>- no contraindication for chemotherapy</li><li>- informed consent</li></ul>
<b>Exclusion Criteria</b>	<ul style="list-style-type: none"><li>- contraindications for intensive chemotherapy, e.g. cardiac co-morbidities</li></ul>
<b>Age</b>	>= 18 years
<b>Status</b>	Closed
<b>start of Recruitment</b>	19.10.2009
<b>Recruiting countries</b>	Germany
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<b>Remark</b>	ATRA and chemotherapy followed by Maintenance therapy with chemotherapy and ATRA; <a href="http://www.sal-aml.org">www.sal-aml.org</a>