

Long-term follow up of pediatric Philadelphia positive acute lymphoblastic leukemia treated with the EsPhALL2004 study: high white blood cell count at diagnosis is the strongest prognostic factor

Approximately 3-5% of children with acute lymphoblastic leukemia (ALL) present with Philadelphia chromosome (Ph⁺). In the pre-tyrosine kinase inhibitors (TKIs) era, Ph⁺ ALL was associated with a poor prognosis.¹⁻¹⁰ The outcome improved with the use of TKIs.¹¹⁻¹⁴ Results of the EsPhALL2004 study (EudraCT 2004-001647-30 and *clinicaltrials.gov* identifier 00287105) with the use of imatinib mesylate were published with a median follow-up time of 3.1 years;^{14,15} we now report on long-term outcomes and the impact of prognostic factors in this study. This update shows that event-free survival (EFS) is stable after about 4 years, and that white blood cell (WBC) count at diagnosis is the most relevant prognostic factor, in a setting where the large majority of patients receives imatinib and hematopoietic stem cell transplantation (HSCT).

Between January 2004 and December 2009, 160 Ph⁺ ALL patients were enrolled in the EsPhALL2004 study in the centers of 10 national study groups. Any subject aged 1-17 years, recruited into national frontline trials for ALL, who showed evidence of t(9;22)(q34;q11) translocation was eligible for the study. Ninety patients were defined as good risk (GR; prednisone good response at day 8, or $\leq 25\%$ bone marrow [BM] blast cells at day 15, or $< 5\%$ BM blast cells at day 21, and in complete remission (CR) at the end of induction and were randomized to receive

(N=46) or not (N=44) imatinib. The remaining 70 poor risk (PR) patients were all given imatinib. Thus, 128 patients overall were given intermittent imatinib, for a total of 10 weeks if transplanted or 18 weeks if treated with chemotherapy only. In total, 130/160 patients underwent HSCT, and some received imatinib post-HSCT according to local guidelines.

EFS was calculated as the time from diagnosis to first failure, including resistance, relapse, death or second malignant neoplasm (SMN). Disease-free survival (DFS) in GR patients was defined as the time from randomization until relapse, death or SMN, whichever occurred first. Overall survival (OS) considered death as event. Observation periods were censored at the date of last contact when no event was observed. The final follow up was on June 30, 2014. DFS analysis in GR patients was done by intention to treat; a secondary analysis was done of patients as treated. EFS, DFS and OS curves were estimated according to Kaplan-Meier (with Greenwood standard error) and compared with the log-rank test. Kaplan-Meier plots for comparison of HSCT with chemotherapy alone were adjusted to account for the waiting time to transplantation. To deal with lack of proportional hazards, univariate comparison was performed at predefined time points based on log-log transformation. The impact of age and WBC count on EFS was analyzed with a Cox model, stratified by risk group (GR and PR). The cumulative incidence of relapse (CIR) and death (CID) were estimated adjusting for competing risks of other events and compared with the Gray test. All tests were two sided. All analyses were performed with SAS software (version 9.2).

Table 1. Outcome by risk group and treatment performed in EsPhALL2004.

	Good Risk N=90		Poor Risk N=70		Overall N=160		
	Chemo N=20	HSCT CR1 N=70	Chemo N=10	HSCT CR1 N=60	Chemo N=30	HSCT CR1 N=130	Both N=160
Deaths in CCR	1	5	2	7	3	12	15 (9.4%)
Infections	1	0	2	0	1	0	1
Other ^s	0	0	0	0	2	0	2
SCT-related	–	5	–	7	–	12	12
Time from diagnosis to death in CCR							
≤ 5 months	1	0	2	0	3	0	3
6-12 months	0	2	0	2	0	4	4
>12 months	0	3	0	5	0	8	8
Relapses [deaths after relapse]	10 [6]	13 [9]	6 [4]	21 [14]	16 [10]	34 [23]	50 (31.2%) [33]
BM isolated	7	11	4	18	11	29	40
BM combined*	1	0	1	2	2	2	4
Extramedullary [†]	2	2	1	1	3	3	6
Time from diagnosis to relapse [‡]							
very early [§]	5	8	5	12	10	20	30
early	3	4	1	3	4	7	11
late	2	1	0	6	2	7	9
CCR	9	52	2	32	11	84	95 (59.4%)

Chemo: chemotherapy only; HSCT CR1: hematopoietic stem cells transplantation in first complete remission; CCR: continuous complete remission; BM: bone marrow; ^sOther causes of death were capillary leak syndrome (n=1) and cardiac failure during chemotherapy (n=1). *BM combined relapses are with central nervous system (N=2, both in Poor Risk patients) and testes (N=2, one in a Good Risk and the other one in a Poor Risk patient). [†]extramedullary relapses were in central nervous system (N=5, N=3 in Good Risk and N=2 in Poor Risk patients) or testes (N=1, in a Good Risk patient). [‡]time from diagnosis to relapse is defined as very early (if relapse occurred <18 months), early (18 to 30 months) and late (≥ 30 months). [§]three relapses were observed before the median waiting time to transplant (5 months from CR1, all counted in the chemotherapy-only group).

With a median follow-up time of 6.6 years (range, 0.3 – 10.5 years), the 5-year and 7-year EFS were 60.3% (SE 3.9) and 58.2% (SE 4.0). Corresponding OS figures were 71.5% (SE 3.6) and 68.2% (SE 3.9). The CIR at 5 and 7 years was 30.9% (SE 3.7) and 31.9% (SE 3.8, Figure 1, Panels A and B). Curves by risk group (Figure 1, Panels C-F) showed 5-year EFS and OS of 68.5% (SE 4.9) and 78.5% (SE 4.4) in GR patients, and of 49.7% (SE 6.0) and 62.9% (SE 5.8) in PR patients. The outcome in GR patients by ITT showed a 5-year DFS in the imatinib arm of 75.5% (SE 6.4) vs. 61.4% (SE 7.3) in the no imatinib arm (P -value 0.20); the difference was higher in the analysis “as treated”, corresponding figures being 77.2% (SE 5.6) vs. 54.8% (SE 8.9, P -value=0.04). As shown in Table 1, of the 130 patients transplanted in CR1, there were 12 deaths in CCR (9.2%) and 34 relapses (26.2%), including 31 medullary and 20 very early. In the small subgroup of 30 patients treated with chemotherapy only, 6 failed within the median waiting time to transplant (5.1 months; 3 died and 3 relapsed) and 13 of the remaining 24 patients relapsed later on. Patients treated with chemotherapy had a 5-year DFS of 46.8% (SE 10.6) and a 5-year OS of 68.8% (SE 8.9). All events occurred within 3 years of diagnosis. Transplanted patients had a 5-year DFS of 66.1% (SE 4.2) and a 5-year OS of 74.2% (SE 3.9). DFS was 20% higher in HSCT (P -value at 5 years=0.07) compared to chemotherapy, while survival curves were

superimposable (P -value at 5 years=0.56).

In total, 101 patients were male and 59 female; their 5-year EFS was 60.8% (SE 4.9) vs. 59.3% (SE 6.4) and 5-year OS 72.8% (SE 4.5) vs. 69.4% (SE 6.0), with non-significant differences, respectively. In the analysis by age, the outcome in the 85 patients <10 years was compared with that of 75 patients \geq 10 years: their 5-year EFS was 55.9% (SE 5.4) vs. 65.1% (SE 5.5, P -value =0.22) and 5-year OS 70.1% (SE 5.0) vs. 73.0% (SE 5.2, P -value =0.63). At diagnosis, 52 patients had a WBC count of $<20 \times 10^9/L$, 48 of $20-100 \times 10^9/L$ and 60 of $\geq 100 \times 10^9/L$. As shown in Figure 4, their 5-year EFS was 86.3% (SE 4.8), 58.3% (SE 7.1) and 39.6% (SE 6.4; P -value <0.0001) and 5-year OS was 91.9% (SE 3.9), 72.9% (SE 6.4) and 53.2% (SE 6.5; P -value <0.0001), respectively. These differences remained statistically significant in GR patients (Figure 4, panels C and D). In PR patients, the outcome was equally unfavorable for patients with WBC count of $20-100 \times 10^9/L$ and $\geq 100 \times 10^9/L$ (Figure 4, panels E and F). A Cox regression model, stratified by risk group and including age and WBC count as covariates, revealed that WBC count was the only independent risk factor: high WBC count $\geq 100 \times 10^9/L$ was associated with poor prognosis (HR 5.26, 95% CI 2.37-11.70, P -value <0.0001), as well as WBC count of 20 to $100 \times 10^9/L$ (HR 2.90, 95% CI 1.28-6.61, P -value=0.0111), as compared to WBC count $<20 \times 10^9/L$. When the analysis was repeated on the sub-

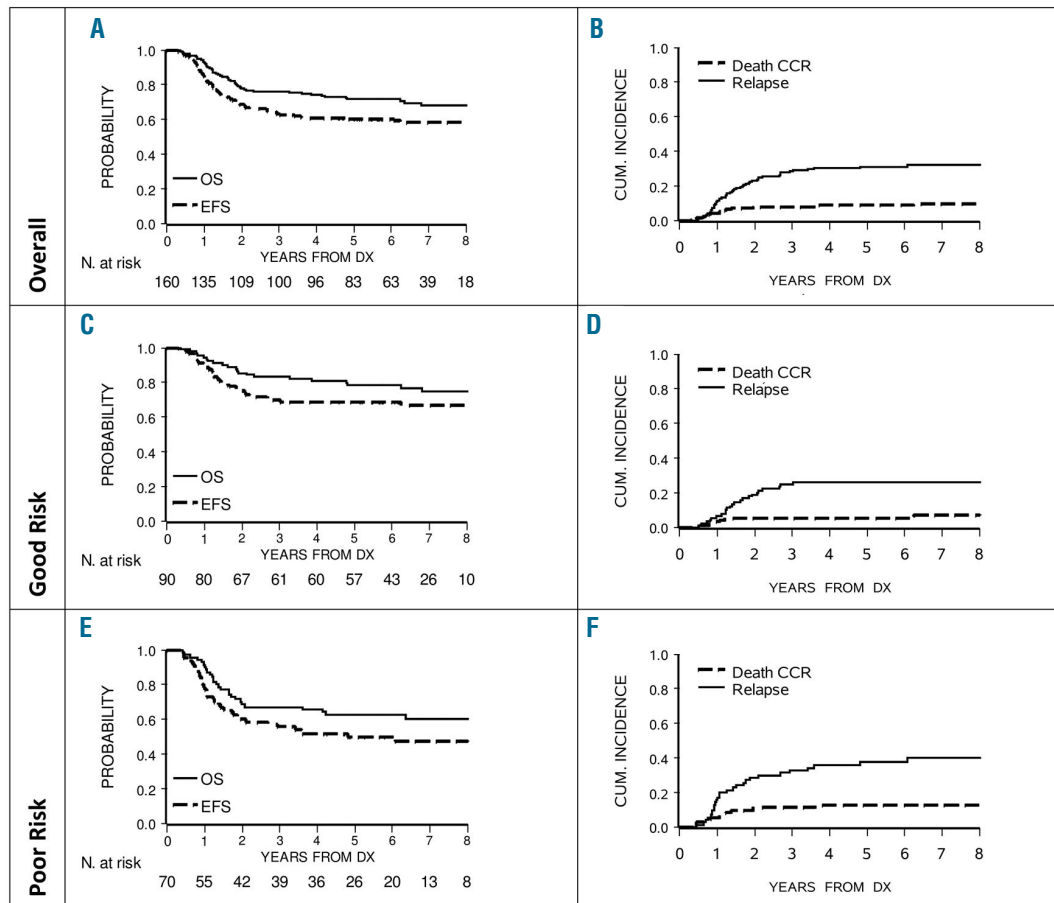


Figure 1. Event-free survival (EFS), overall survival (OS), cumulative incidence of relapse and death of 160 Ph⁺ALL patients, overall (Panels A and B) and by risk group (Good Risk in Panels C and D, Poor Risk in Panels E and F). CCR: continuous complete remission.

group of 128 patients who received imatinib, results did not change substantially; HR being of 3.22 (95% CI 1.17 – 8.82) for WBC count of 20 to $100 \times 10^9/L$ and 4.84 (95% CI 1.81 – 12.94) for WBC count $\geq 100 \times 10^9/L$. Restricting the analyses to the small subgroup of 76 patients with MRD levels available at the end of Protocol IB, WBC count $\geq 100 \times 10^9/L$ was borderline significant (HR 2.85, 95% CI 1.01- 8.10, P -value=0.0490) and MRD was not significant (HR 2.09, 95% CI 0.76- 5.72 of positive vs. negative MRD, P -value=0.15). The 4-year survival after relapse was 34.2% (SE 6.7), with very similar outcome for the 34 patients who relapsed after HSCT (23 deaths) and for the 16 patients who were previously treated with chemotherapy only (10 deaths).

The long-term follow up of the EsPhALL2004 study shows that the EFS and OS, previously reported at 4 years (EFS 61.9% and OS 72.1%), decreased slightly, with 7-year figures of 58.2% and 68.2%, respectively. After 4 years, no events occurred in the few patients who received chemotherapy only, while 3 events occurred in transplanted patients (2 relapses and 1 death in CCR due to pulmonary GvHD). Overall, these data confirm that the introduction of imatinib decreased the rate of relapses, and conveyed a marked improvement in EFS when compared to the outcome reported for Ph⁺ALL before the

advent of TKI therapy.⁴ These results are in keeping with those reported for the COG studies AALL0031¹² and AALL0622¹³: their 5-year OS of 81% (SE 6) and 86% (SE 5), respectively, support the use of increased exposure to imatinib (and other TKIs) and decreased need for HSCT in the next generation of studies. The comparison of outcome by treatment (chemotherapy vs. transplant) in EsPhALL2004 is very difficult, considering that 81% of patients was transplanted. Although survival was similar, a higher rate of relapse was observed in the few patients who did not undergo transplant, suggesting that intermittent and short exposure to imatinib is inadequate for patients treated with chemotherapy only. In transplanted patients, this treatment strategy was associated with low relapse rate in GR patients (18.6%) and a higher risk in PR patients (35.0%). Interestingly, a high proportion of relapses were very early (30/50). Of note, in this long-term evaluation, the most relevant unfavorable prognostic factor was high WBC count at diagnosis ($\geq 100 \times 10^9/L$), while neither MRD nor age at diagnosis showed a prognostic impact in the multivariable analysis. As already reported, in EsPhALL2004 only early MRD negativity, which occurred in a small proportion of patients (10%), was associated with good prognosis.¹⁵ Interestingly, with

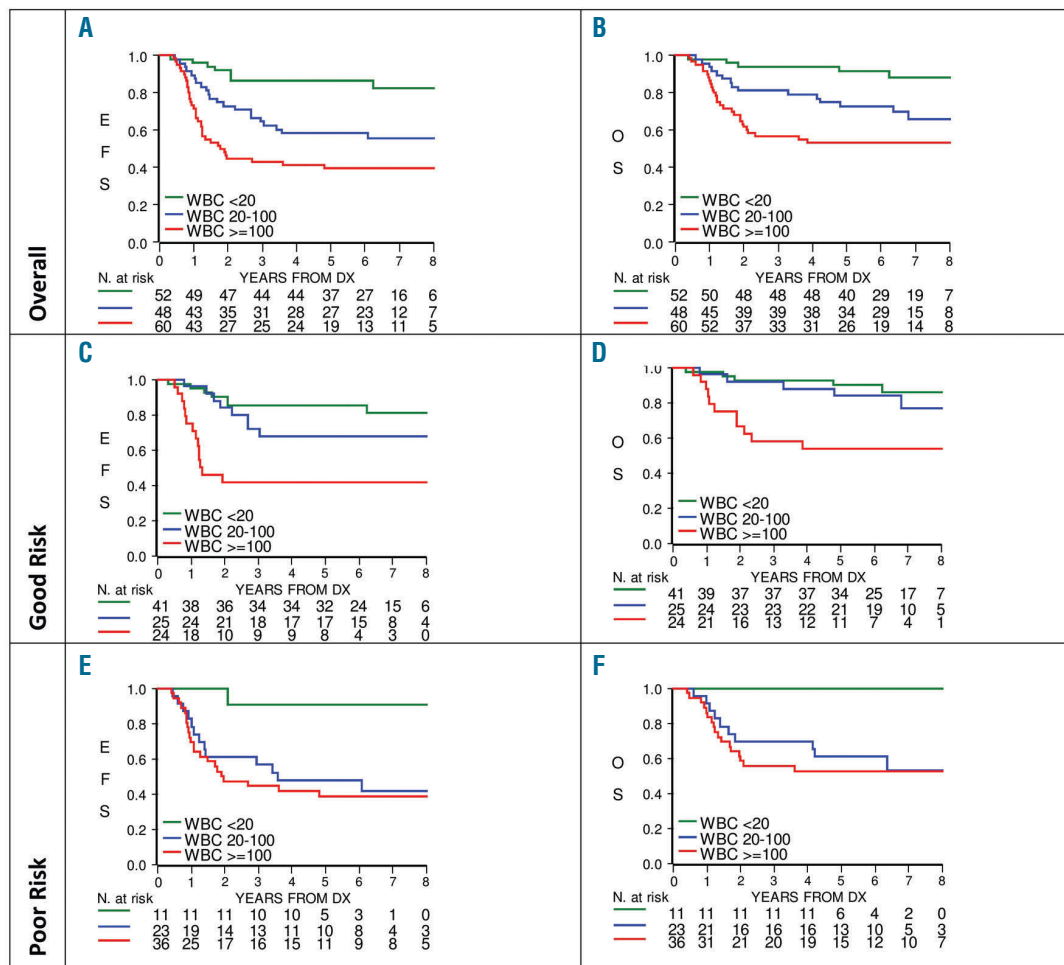


Figure 2. Event-free survival (EFS) and overall survival (OS) by white blood cell count (WBC) count at diagnosis, overall (Panels A and B), in Good Risk (Panels C and D) and Poor Risk patients (Panels E and F).

the use of imatinib and HSCT, age is no longer a relevant prognostic factor. The main findings of this update are that EFS, DFS and OS are stable after about 4 years from diagnosis. Moreover, WBC count at diagnosis emerged as the most relevant independent prognostic factor. This observation should, however, be tested in other studies with longer exposure to TKI and fewer transplanted patients.

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