



*POST-ASH* Issue 4, 2016

**Ponatinib versus Allogeneic  
Stem Cell Transplant for Patients  
with CML or Ph+ ALL with the  
T315I Mutation**

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## CME INFORMATION

### OVERVIEW OF ACTIVITY

Each year, thousands of clinicians, basic scientists and other industry professionals sojourn to major international oncology conferences, like the American Society of Hematology (ASH) annual meeting, to hone their skills, network with colleagues and learn about recent advances altering state-of-the-art management in hematologic oncology. These events have become global stages where exciting science, cutting-edge concepts and practice-changing data emerge on a truly grand scale. This massive outpouring of information has enormous benefits for the hematologic oncology community, but the truth is it also creates a major challenge for practicing oncologists and hematologists.

Although original data are consistently being presented and published, the flood of information unveiled during a major academic conference is unmatched and leaves in its wake an enormous volume of new knowledge that practicing oncologists must try to sift through, evaluate and consider applying. Unfortunately and quite commonly, time constraints and an inability to access these data sets leave many oncologists struggling to ensure that they're aware of crucial practice-altering findings. This creates an almost insurmountable obstacle for clinicians in community practice because they are not only confronted almost overnight with thousands of new presentations and data sets to consider but they are also severely restricted in their ability to review and interrogate the raw findings.

To bridge the gap between research and patient care, this CME activity will deliver a serial review of the most important emerging data sets on novel agents and therapeutic options for the treatment of newly diagnosed and relapsed/refractory acute and chronic leukemias, myelodysplastic syndromes and myeloproliferative neoplasms from the latest ASH meeting, including expert perspectives on how these new evidence-based concepts may be applied to routine clinical care. This activity will assist medical oncologists, hematologists, hematology-oncology fellows and other healthcare professionals in the formulation of optimal clinical management strategies and the timely application of new research findings to best-practice patient care.

### LEARNING OBJECTIVES

- Appraise emerging clinical research findings on the efficacy and safety of novel antibody-based therapies for acute lymphoblastic leukemia.
- Assess the activity of the multitargeted kinase inhibitors midostaurin and sorafenib with chemotherapy for FLT3-mutated acute myeloid leukemia.
- Compare the risks and benefits associated with discontinuing imatinib therapy for patients with chronic myeloid leukemia (CML) who have achieved a deep molecular response.
- Recall recent data on the activity and tolerability of second- and third-generation tyrosine kinase inhibitors for the treatment of CML.
- Evaluate the efficacy and safety of ruxolitinib alone or in combination with other therapies for patients with myelodysplastic syndromes and myeloproliferative neoplasms.

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**FACULTY** — The following faculty (and their spouses/partners) reported relevant conflicts of interest, which have been resolved through a conflict of interest resolution process:

Richard M Stone, MD  
Director, Adult Leukemia Program  
Dana-Farber Cancer Institute  
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This activity is supported by educational grants from Celgene Corporation, CTI BioPharma Corp/Baxalta Inc, Jazz Pharmaceuticals Inc, Novartis Pharmaceuticals Corporation, Seattle Genetics and Takeda Oncology.

**Hardware/Software Requirements:**

A high-speed Internet connection

A monitor set to 1280 x 1024 pixels or more

Internet Explorer 7 or later, Firefox 3.0 or later, Chrome, Safari 3.0 or later

Adobe Flash Player 10.2 plug-in or later

Adobe Acrobat Reader

(Optional) Sound card and speakers for audio

Last review date: May 2016

Expiration date: May 2017



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The May 17th historic FDA approval (accelerated) of the anti-PD-1 antibody nivolumab for the treatment of classical Hodgkin lymphoma that has relapsed or progressed after autologous stem cell transplantation and post-transplant brentuximab vedotin is a vivid reminder of just how far we have come in moving new options for patients into practice. Nivo's endorsement was predicated on an evaluation of efficacy in just 95 patients, but because the drug as monotherapy produced useful responses in approximately two thirds of individuals with limited treatment options, the FDA's decision is great news to oncologists, and it could be that pembrolizumab, which also has strong but similarly limited supporting data, could soon follow.



**Richard M Stone, MD**

However, while it is comforting to see new therapies that are clearly effective become rapidly available, in reality many of the "steps forward" in clinical research continue to come as a result of large randomized trials evaluating standard systemic regimens alone or in combination with novel agents, many of which demonstrate marginal but clearly statistically significant advantages. What makes these kinds of findings and subsequent regulatory approvals so challenging is that they often come in diseases with dismal outcomes and few options for treatment, and usually the drug in question has limited single-agent activity and, most significantly, the benefit provided is far from a home run.

These factors create an environment of constant debate about the relevance of a therapy and how it fits into the risk/benefit/value equation. Recent examples in the hem/onc world include in multiple myeloma the use of panobinostat as an add-on to bortezomib/dexamethasone and the incorporation of elotuzumab with lenalidomide/dexamethasone. The same issue arises in chronic lymphocytic leukemia with the addition of obinutuzumab (as opposed to rituximab) to chlorambucil and in follicular lymphoma to bendamustine.

This is precisely the scenario that is emerging in the world of acute myeloid leukemia (AML) related to the combination of multikinase tyrosine kinase inhibitors (TKIs) with

induction chemotherapy and maintenance for the approximately 35% of patients with AML who have FLT3 genomic alterations — long known to be associated with a particularly adverse prognosis, which has led to the current common practice of following induction treatment with allogeneic stem cell transplantation.

The FLT3 AML debate began during a 2014 ASH plenary presentation of a large German study that demonstrated a progression-free survival advantage with the addition of the multikinase TKI sorafenib to induction chemotherapy and its continuation for a year as maintenance. A key and very provocative finding (and to this point pretty much unexplained) was that the benefit was observed in patients with and without FLT3 abnormalities.

At the ASH 2015 meeting another plenary talk focused on this strategy — specifically the presentation by Dana-Farber's Dr Richard Stone of the international Phase III RATIFY trial led by the CALGB that blindly randomly assigned 717 patients with 1 of the 3 FLT3 alterations, including tyrosine kinase domain mutations and both low and high internal tandem duplication allele burdens, to another multikinase TKI, midostaurin, which, like other similar compounds, has limited single-agent activity but can be safely combined with intensive chemotherapy.

The study added midostaurin to daunorubicin/cytarabine induction and high-dose ara-C consolidation and then administered it as maintenance therapy in patients age 18 to 60 and met its primary endpoint of improving overall survival, although the 23% relative reduction in risk of death translates to a somewhat modest improvement in 4-year survival from 44.2% to 51%. Of great interest is that the benefit was observed regardless of whether allotransplant occurred or which of the 3 major predetermined FLT3 subtypes were present, which caused some in the audience to wonder if, as with the sorafenib trial, the benefit might accrue to a larger segment of patients.










I met with Dr Stone for his take on these data, what they mean to patient care and whether this is the beginning of the end of an era in AML lasting more than a decade in which the only regulatory action has been in the wrong direction (withdrawal of approval of gemtuzumab ozogamicin).

To make a long story short, the key message from our conversation is that although some caveats exist, Dr Stone believes that these findings are meaningful and the drug should be incorporated into practice. This sentiment appears to be echoed by the FDA, which recently granted midostaurin a breakthrough therapy designation, but we shall see whether it will become another in the recent unprecedented flurry of novel agents entering practice.





Either way, a host of other new small molecules with anti-FLT3 activity such as gilteritinib (formerly ASP2215) are marching on in the research development process with the hope of greater specificity resulting in increased efficacy and more favorable tolerability.

Of course, midostaurin was definitely not the only relevant news coming out of ASH. For that reason and to keep you up to date on what happened, we have created the following 26 slide sets across a number of diseases (AML, myelodysplastic syndromes, acute lymphocytic leukemia, myeloproliferative neoplasms and chronic myeloid leukemia), which review highlights from the data and provide Dr Stone's insights into their relevance and meaning.









## Acute myeloid leukemia (AML)/chronic myelomonocytic leukemia (CMML)/myelodysplastic syndromes (MDS)

-  (PLENARY) Up-front use of the multikinase inhibitor midostaurin with a "7 plus 3" chemotherapy regimen in FLT3-mutated AML
-  Sorafenib with chemotherapy improves the overall survival of older adults with FLT3-ITD-mutated AML
-  Antileukemic activity and tolerability of the multikinase inhibitor gilteritinib in FLT3-mutated AML
-  Up-front treatment with venetoclax and decitabine or azacitidine for older patients with AML
-  Pracinostat with azacitidine in elderly patients with AML; oral azacitidine monotherapy in AML
-  Allogeneic stem cell transplantation after high- versus reduced-intensity conditioning in MDS and AML
-  Safety and efficacy of AG-221, a potent inhibitor of mutant IDH2 in advanced hematologic cancers
-  Azacitidine with lenalidomide or vorinostat versus azacitidine monotherapy in MDS and CMML
-  Eltrombopag for the treatment of thrombocytopenia of low and intermediate-1 IPSS risk MDS; luspatercept in low or intermediate-1 risk MDS






## Acute lymphoblastic leukemia (ALL)

-  (PLENARY) Rituximab and chemotherapy in adults with CD20-positive, Philadelphia chromosome-negative, B-cell precursor ALL
-  Dose-intensified pegylated asparaginase pediatric regimen in adults with untreated ALL
-  Blinatumomab in adult patients with relapsed/refractory Philadelphia chromosome-positive ALL
-  Front-line inotuzumab ozogamicin combination with low-intensity chemotherapy for older patients with ALL

## Chronic myeloid leukemia (CML)

-  PPT Long-term follow-up of the French 1 Stop Imatinib study in CML
-  PPT Personalized daily doses of imatinib by therapeutic drug monitoring in CML
-  PPT Clinical significance of early imatinib-induced ABCB1 overexpression in CML
-  PPT Dose-optimized nilotinib in newly diagnosed CML
-  PPT Impact of age on efficacy and toxicity of nilotinib in patients with CML
-  PPT Dasatinib and peginterferon alpha-2b as up-front treatment for CML
-  PPT Ponatinib versus allogeneic stem cell transplant in patients with CML/acute lymphoblastic leukemia and the T315I mutation
-  PPT Next-generation sequencing versus conventional sequencing to detect BCR-ABL mutations in CML

## Myeloproliferative neoplasms (MPN)

-  PPT Long-term efficacy and safety of ruxolitinib (RUX) in myelofibrosis (MF)
-  PPT Pacritinib in MF
-  PPT The antifibrotic agent PRM-151 in MF
-  PPT 5-azacytidine with RUX in MDS/MPN; sonidegib with RUX in patients with MF; RUX with pomalidomide in MF; RUX and buparlisib in MF
-  PPT Interferon alpha-2 with a JAK1/2 inhibitor in Philadelphia chromosome-negative MPN

Neil Love, MD

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# **Ponatinib versus Allogeneic Stem Cell Transplant for Patients with CML or Ph+ ALL with the T315I Mutation**

**Presentation discussed in this issue**

Nicolini FE et al. **The impact of ponatinib versus allogeneic stem cell transplant (SCT) on outcomes in patients with chronic myeloid leukemia (CML) or Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ ALL) with the T315I mutation.** *Proc ASH 2015*; **Abstract 480**.

**Slides from a presentation at ASH 2015 and transcribed comments from a recent interview with Richard M Stone, MD (2/16/16)**

## **The Impact of Ponatinib versus Allogeneic Stem Cell Transplant (SCT) on Outcomes in Patients with Chronic Myeloid Leukemia (CML) or Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia (Ph+ ALL) with the T315I Mutation**

**Nicolini FE et al.**

*Proc ASH 2015*; Abstract 480.

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## Ponatinib versus Allogeneic Stem Cell Transplant (Allo SCT) for Chronic Myeloid Leukemia (CML) and Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia (Ph+ ALL) with the T315I Mutation

- Indirect comparison of ponatinib, a multitargeted tyrosine kinase inhibitor (using data from the Phase II PACE trial) to allo SCT (using data from the EBMT database)
- N = 184 patients with CML and Ph+ ALL with the T315I mutation: N = 128 receiving ponatinib and N = 56 receiving allo SCT
- **Primary endpoint:** Overall survival (OS) for patients with the T315I mutation:

Leukemia type	Median OS ponatinib (mo)	Median OS allo SCT (mo)	HR	p-value
CP CML (N = 64, 26)	NR	103.3	0.37	0.013
AP CML (N = 18, 8)	NR	55.6	0.90	0.889
BP CML (N = 24, 17)	7.0	10.5	2.29	0.026
Ph+ ALL (N = 22, 5)	6.7	32.4	2.77	0.119

HR = hazard ratio; CP = chronic phase; NR = not reached; AP = accelerated phase; BP = blast phase

Nicolini FE et al. *Proc ASH* 2015;Abstract 480.

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## Ponatinib versus Allo SCT: Conclusions

- OS by leukemia type:
  - CP CML: Significantly longer with ponatinib; promising strategy
  - AP CML: Similar
  - BP CML and Ph+ ALL: Longer with allo SCT
- Results warrant consideration of ponatinib as an alternative to allo SCT for patients with T315I mutation-positive CP CML.
- Further study is needed.

Nicolini FE et al. *Proc ASH* 2015;Abstract 480.

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## **Investigator Commentary: Ponatinib versus Allo SCT for Patients with CML or Ph+ ALL Harboring the T3151 Mutation**

Because of the high rate of efficacy in terms of deep and durable remissions with tyrosine kinase inhibitors in CML, the use of allo SCT, formerly the standard approach for this condition, has declined sharply. However, some patients require SCT for CML, including those who present in blast crisis (and possibly accelerated phase) and those who do not respond to the available tyrosine kinase inhibitors.

Of particular difficulty is a secondary mutation of the BCR-ABL fusion gene. This mutation, the T3151 mutation, is inhibited by ponatinib, a "third-generation" tyrosine kinase inhibitor that is associated with worrisome vasculopathy. It is currently unknown whether patients who have a T3151 mutation should undergo allo SCT or should receive ponatinib. The retrospective study presented by Dr Nicolini and colleagues suggests that in comparison to allo SCT ponatinib therapy produces longer survival for patients with CML in chronic phase but is inferior for those in blast crisis with a T315I mutation.

***Interview with Richard M Stone, MD, February 16, 2016***

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