



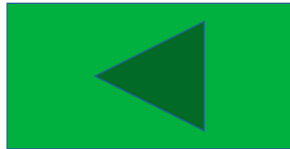
# Insights Into Philadelphia Chromosome-Positive (Ph<sup>+</sup>) Acute Lymphoblastic Leukemia (ALL) and Chronic Myeloid Leukemia (CML)

March 20, 2023

# How to Navigate This Report










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## STUDY OBJECTIVES

- > Gain perspectives on community treatment practices in Ph+ ALL and CML
- > Gain insight into the influence of recent data on community treatment practices

# Report Snapshot: Session Overview



A moderated roundtable discussion was held virtually with oncologists in the United States on **March 20, 2023**

Disease state and data presentations were led and moderated by **Dr Elias Jabbour** from MD Anderson Cancer Center, in conjunction with content developed by the Aptitude Health clinical team

Insights were obtained on **therapies for Ph+ ALL and CML** in the community setting

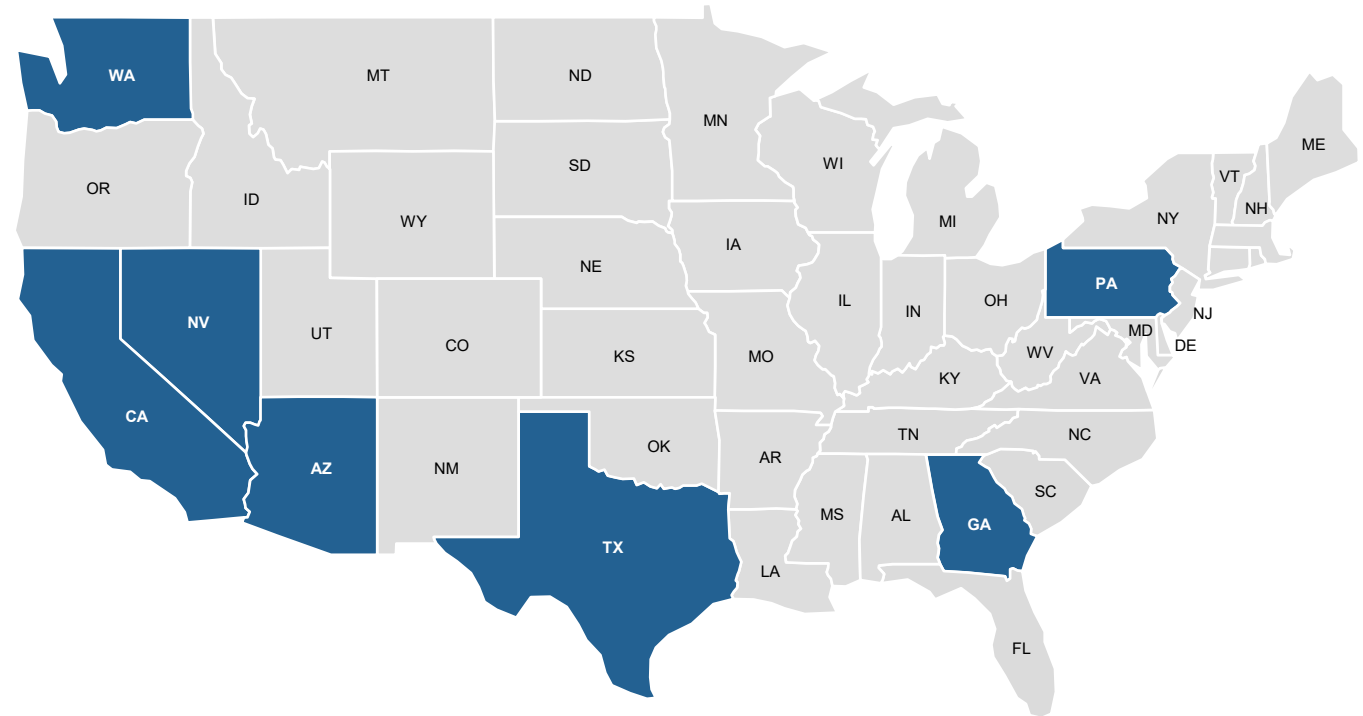
Data collection was accomplished through use of audience response system (ARS) questioning and in-depth moderated discussion

# Report Snapshot: Attendee Overview



- > The group of physicians comprised 9 oncologists from across the United States
  - Attendees of the roundtable represented community oncologists from Arizona, California, Georgia, Nevada, Pennsylvania, Texas, and Washington

INSTITUTION	CITY	STATE
Cancer Transplant Institute	Scottsdale	AZ
City of Hope	Duarte	CA
The Oncology Institute of Hope and Innovation	South Pasadena	CA
Blood and Marrow Transplant of Georgia	Atlanta	GA
Comprehensive Cancer Centers of Nevada	Las Vegas	NV
Allegheny Health Network	Wexford	PA
The Center for Cancer and Blood Disorders	Benbrook	TX
The Center for Cancer and Blood Disorders	Fort Worth	TX
University of Washington/Fred Hutchinson Cancer Center	Seattle	WA



# Report Snapshot: Agenda



Time (ET)	Topic
6.00 PM – 6.10 PM (10 min)	<b>Introduction</b> <ul style="list-style-type: none"><li>• Program overview</li></ul>
6.10 PM – 6.25 PM (15 min)	<b>ARS Questions</b>
6.25 PM – 7.00 PM (35 min)	<b>Management Options in Ph+ ALL and CML</b> <ul style="list-style-type: none"><li>• Overview of current data in Ph+ ALL<ul style="list-style-type: none"><li>– Induction regimens<ul style="list-style-type: none"><li>▪ Choice of TKI</li><li>▪ PhALLCON</li></ul></li><li>– Postremission therapy</li><li>– MRD and R/R disease</li></ul></li><li>• Overview of current data in CML<ul style="list-style-type: none"><li>– Primary treatment<ul style="list-style-type: none"><li>▪ Choice of TKI</li><li>▪ Setting treatment goals</li></ul></li><li>– Response milestones and next steps</li></ul></li><li>• Moderated discussion</li></ul>
7.00 PM – 7.50 PM (50 min)	<b>Discussion</b>
7.50 PM – 8.00 PM (10 min)	<b>Key Takeaways and Meeting Evaluation</b>



## Discussion

Management Options in CML



## CML – INSIGHTS AND DATA

*"I use [ponatinib] after, if they have the T315I mutation. . . . I have used ponatinib after they failed like, I generally start off*

1. Treatment success in frontline BCR-ABL

The overall success rate is very high. This is not necessarily because this is a curable disease, it is not really curable. . . . I would not use a treatment option after that using TKI or TKI, and I would not start the disease for less than 1 year. . . . I believe in that TKI is important if there is significant toxicity with the treatment, and usually going from something else.

2. Data needed to switch from TKI to frontline

That is all a lot of things have been done, nothing is better than BCR-ABL and TKI. . . . I would not use a treatment option after that using TKI or TKI, and I would not start the disease for less than 1 year. . . . I believe in that TKI is important if there is significant toxicity with the treatment, and usually going from something else. . . . I would not use a treatment option after that using TKI or TKI, and I would not start the disease for less than 1 year. . . . I believe in that TKI is important if there is significant toxicity with the treatment, and usually going from something else.

# Discussion: Management Options in CML (2/2)



## CML – INSIGHTS AND DATA

*“In the community, I am not able to get the mutational analysis, like the way I would want. Sometimes I get an arbitrary*

<p>1. Treatment success in frontline BCR-ABL</p>	<p>The overall survival data was not clear. This is not necessarily because there is a survival benefit, it is not clear overall survival. I would like to see a breakdown analysis with using TKI or TKI, and I would like to know the duration time with TKI. I believe as there is a reported 100% of significant toxicity with the treatment, and overall drug that something is available.</p>
<p>2. Data needed to confirm front BCR-ABL</p>	<p>That is all a lot of things have been done, nothing is better than BCR-ABL and TKI. It is really hard with how BCR-ABL patients do in TKI. I would like to see a breakdown of the data with TKI or TKI or something like that. I want something that can show me and see how they are TKI. The toxicity was not very severe. I think a reported rate of 100% or better would be something that I would be looking at. Overall survival data, that is clear, but in this disease with TKI a report is coming by, so you do have to use some surrogate of efficacy. So I do think that a lot of people have the survival rate of data, so there is something going to start showing the use of TKI. TKI is not TKI.</p>



## Advisor Key Takeaways

# Advisor Key Takeaways



ADVISOR		ADVISOR	
	<ul style="list-style-type: none"> <li>&gt; Would love to use ponatinib up front, but need more data</li> </ul>		<ul style="list-style-type: none"> <li>&gt; Use of MRD testing in ALL at 3 months</li> </ul>

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## ARS Results

Management Options in Ph+ ALL

# All Physicians Treated at Least 6 Newly Diagnosed ALL Patients in the Past Year

Approximately how many newly diagnosed ALL patients have you treated in the past 12 months? (n = 7\*)

FOR EXAMPLE PURPOSES ONLY

# All Physicians Treated at Least 2 Newly Diagnosed Ph+ ALL Patients in the Past Year

In the past 12 months, approximately how many of your newly diagnosed ALL patients were Ph+? (n = 7\*)

FOR EXAMPLE PURPOSES ONLY

# Achieving MRD Negativity and Getting to Transplant Are Physicians' Primary Treatment Goals in Frontline Ph+ ALL

In frontline Ph+ ALL, what are your treatment goals?  
(Select all that apply.) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.





# When Choosing a TKI for Frontline Ph+ ALL, Physicians Are Most Influenced by Efficacy Data

Which of the following factors most influences your choice of TKI in frontline Ph+ ALL?  
(n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.



# Physicians Have Most Commonly Used TKI With High-Intensity Chemotherapy in the Past Year



Which of the following induction regimens have you used in Ph+ ALL in the past 12 months? (Select all that apply.) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.



# 88% of Physicians Believe It Is “Crucially Important” to Achieve MRD Negativity in Ph+ ALL

How important is it to achieve MRD negativity? (n = 8\*)

100%

FOR EXAMPLE PURPOSES ONLY

# The Majority of Physicians Check for MRD at the End of Induction

When do you prefer to perform an initial assessment for MRD? (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.



# The Majority of Physicians Do Not Face Challenges When Ordering MRD Testing

What challenges do you face in ordering MRD testing? (Select all that apply.) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

# Two-Thirds of Physicians Were Familiar With PhALLCON



Which of the following Ph+ ALL studies are you familiar with? (Select all that apply.) (N = 9)

FOR EXAMPLE PURPOSES ONLY



# Physicians Are Aware of Some PhALLCON Data, Such as the MRD-Negativity Rate and MRD-Negative CR Rate

Which of the following are true of PhALLCON? (Select all that apply.) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.





# ARS Results

Management Options in CML



# Physicians See Varying Numbers of CML Patients

How many CML patients have you personally managed in the past 12 months? (N = 9)

FOR EXAMPLE PURPOSES ONLY

# Physicians Have Varying Levels of Experience in Later Lines of Treatment

How many CP-CML patients have you treated through  $\geq 3$  lines of therapy? (N = 9)

FOR EXAMPLE PURPOSES ONLY

# Physicians Primarily Consider Mutation Status, Prior AEs, and Comorbidities When Selecting Treatment After First-Generation TKI Failure



FOR EXAMPLE PURPOSES ONLY



# Dasatinib Is the Most Commonly Recommended Treatment for Patients Who Are Resistant or Intolerant to First-Generation TKIs

What would you recommend for patients with CP-CML who are resistant or intolerant to first-generation TKIs? (N = 9)

FOR EXAMPLE PURPOSES ONLY

# Physicians More Strongly Consider Resistance and Response to Prior Agents When Selecting Treatment After Second-Generation TKI Failure Compared With First-Generation Failure

Which of the following patient characteristics do you consider most important when selecting treatment following second-generation TKI failure for your CML patients? (Select your top 3.) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.



# Ponatinib Is the Most Commonly Recommended Treatment for Patients Who Are Resistant or Intolerant to Second-Generation TKIs

What would you recommend for patients with CP-CML who are resistant or intolerant to second-generation TKIs? (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

\*One physician did not respond.



# Mutational Status, Safety Profile, and Efficacy Profile Are the Strongest Drivers of Therapy Choice in R/R CP-CML

What factors drive your choice of therapy for R/R CP-CML? (*Select all that apply.*) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

# All Physicians Would Use Ponatinib for a Patient With a *T315I* Mutation

Which agent(s) do you use for patients with a *T315I* mutation? (Select all that apply.) (N = 9)

FOR EXAMPLE PURPOSES ONLY



# All Physicians Are Comfortable With TKI Dose-Reduction Strategies, With 67% Identifying as “Very Comfortable”

How comfortable are you utilizing TKI dose-reduction strategies to manage toxicity?  
(N = 9)

FOR EXAMPLE PURPOSES ONLY



# Insurance/Reimbursement Denials and AE Management Are the Most Common Challenges Physicians Face in CML

What are the major challenges you encounter in the management of your CML patients?  
(Select all that apply.) (n = 8\*)

FOR EXAMPLE PURPOSES ONLY

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