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Analysis of Practice Patterns Among Experts and Community Healthcare Providers for the Treatment of Acute Myeloid Leukemia

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Disclosures

Ravandi: MacroGenics: Research Funding; AbbVie: Consultancy, Honoraria, Research Funding; Jazz Pharmaceuticals: Consultancy, Honoraria, Research Funding; Xencor: Consultancy, Honoraria, Research Funding; Orsenix: Consultancy, Honoraria, Research Funding; Astellas: Consultancy, Honoraria, Research Funding; Amgen: Consultancy, Honoraria, Research Funding; Celgene: Consultancy, Honoraria; BMS: Consultancy, Honoraria, Research Funding; AstraZeneca: Consultancy, Honoraria.

Smith: Agios: Consultancy, Membership on an entity's Board of Directors or advisory committees; Celgene: Consultancy, Membership on an entity's Board of Directors or advisory committees; Pfizer: Consultancy, Membership on an entity's Board of Directors or advisory committees; Novartis: Consultancy, Membership on an entity's Board of Directors or advisory committees; Jazz: Consultancy, Membership on an entity's Board of Directors or advisory committees.

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Off Label Disclosure: In this report of results from a treatment decision support tool for AML, some of the drugs selected by the experts are recommended in off-label applications (eg, IDH2 inhibitor enasidenib in frontline AML therapy).



Background

- Acute myeloid leukemia (AML) treatment options have expanded rapidly, with 9 new agents FDA approved since 2017
- This has provided many new strategies for both patients and healthcare providers (HCPs), but it has also introduced new challenges in treatment selection
- To address this, CCO developed an online AML decision support tool designed to provide HCPs with expert guidance for optimal individualized patient treatment selection



AML Tool Development

- 5 experts identified a simplified set of key patient/disease characteristics on which they based treatment recommendations for patients with AML
 - **Experts:** Jeffrey E. Lancet, MD; Farhad Ravandi, MD; B. Douglas Smith, MD; Roland P. Walter, MD, PhD; Eunice S. Wang, MD
 - **Patient/disease characteristics:** disease setting, age and fitness, secondary AML, previous HMAs, cytogenetic/ molecular risk factors, biomarkers, others
- The expert panel provided treatment recommendations in February 2019 for 330 distinct case scenarios in ND (n = 150) and R/R (n = 180) AML



Using the AML Tool

- HCPs are prompted to select defined patient/disease characteristics from drop-down menus and then are asked to provide their Tx choice
- They then receive expert recommendations for that case scenario

Your Patient Case

What is your patient's disease setting? Newly diagnosed

What is your patient's age and fitness? Fit (and ≤ 75 years)

Does your patient have secondary AML? Yes, from prior MDS or therapy related

Has your patient received prior HMA therapy for hematologic disease? No

What is your patient's cytogenetic/molecular risk category? Intermediate

Is your patient positive for any of the following biomarkers? None of the above

Which treatment are you considering for your patient with newly diagnosed AML? Uncertain

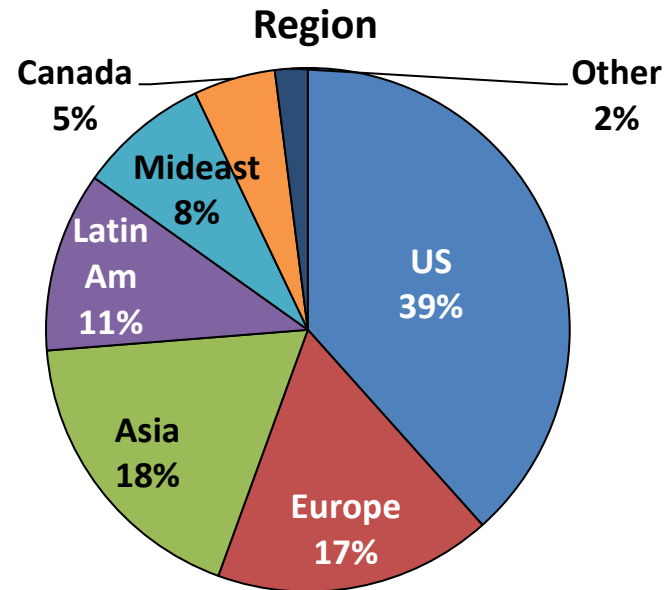
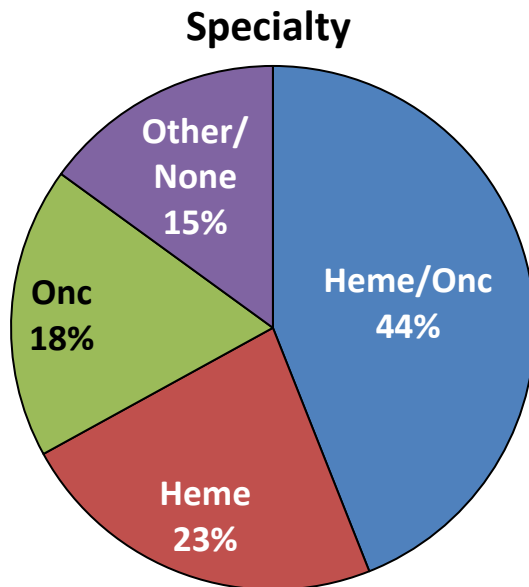
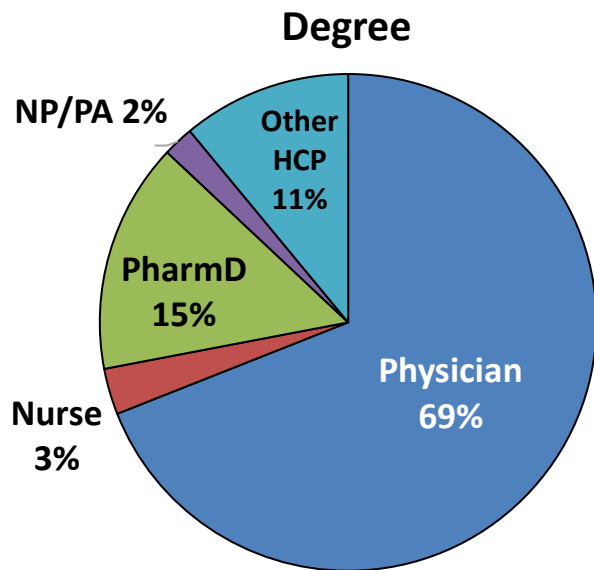
Recommendations

	Recommendations	Additional Treatment
Expert 1	CPX-351	AlloSCT in CR1
Expert 2	Venetoclax plus HMA	None
Expert 3	CPX-351	None
Expert 4	CPX-351	AlloSCT in CR1
Expert 5	CPX-351	AlloSCT



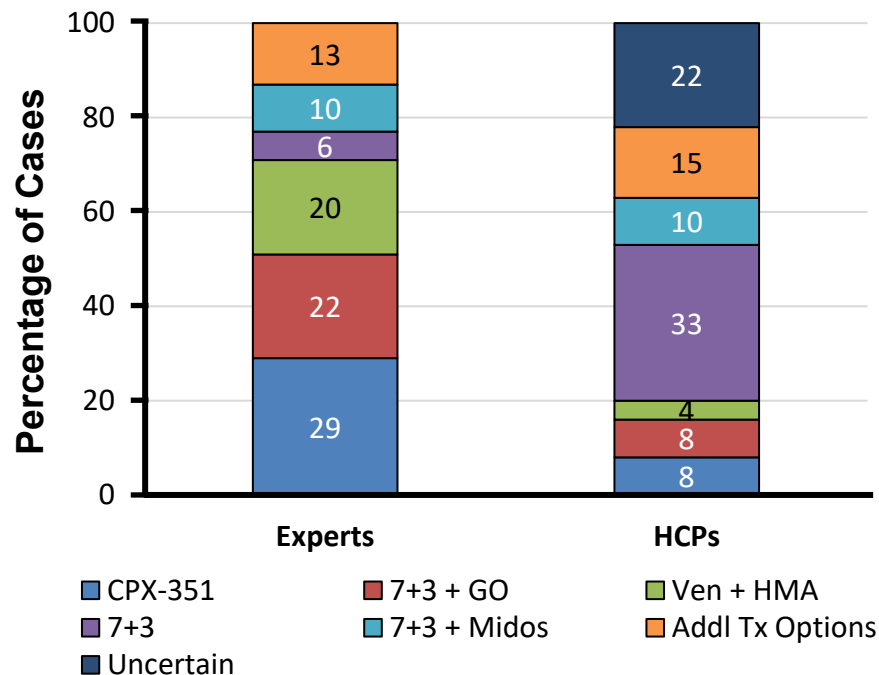
Demographics of AML Tool Participants

- A total of 417 HCPs entered 934 patient scenarios from June 2019 - July 2020

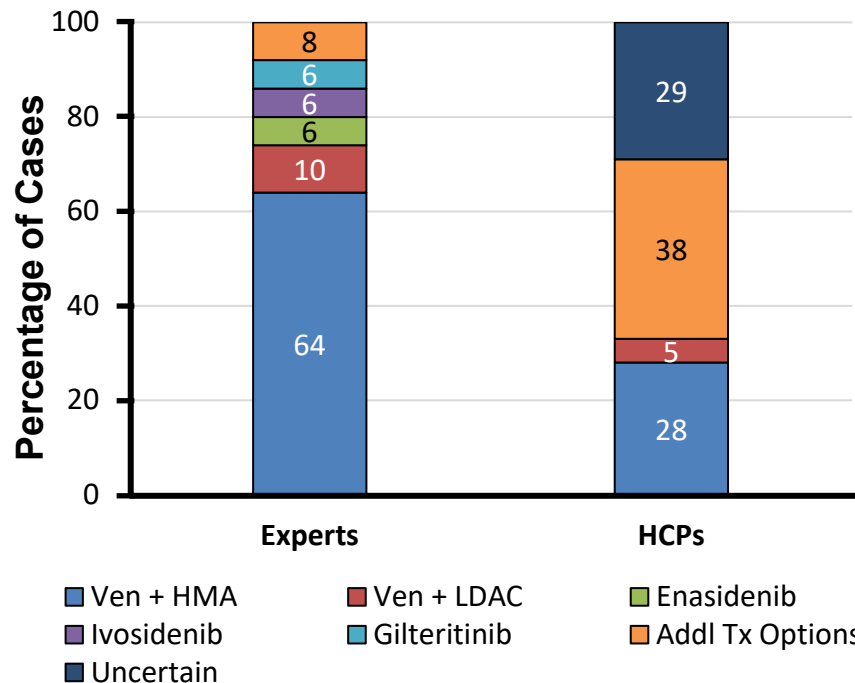


Treatment Recommendations for Cases of ND AML

≤ 75 yrs of age, fit (n = 535)

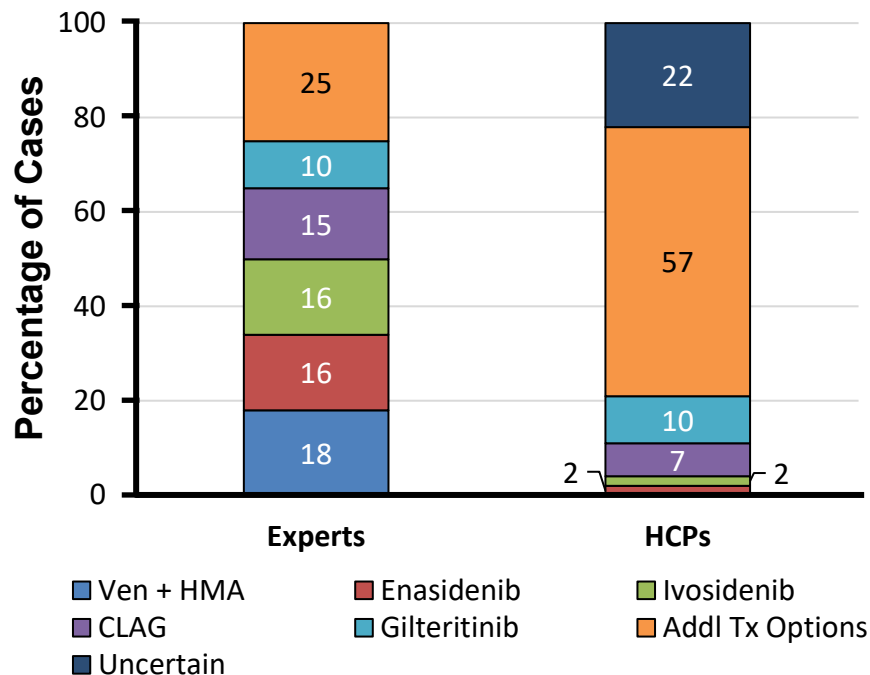


> 75 yrs of age, less fit (n = 213)

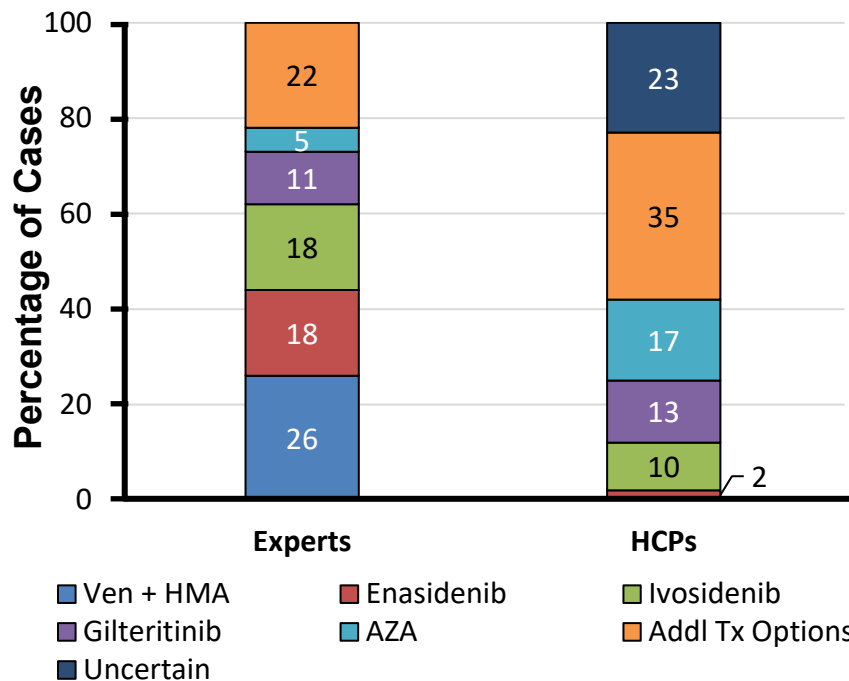


Treatment Recommendations for Cases of First Relapse AML

≤ 75 yrs of age, fit (n = 134)

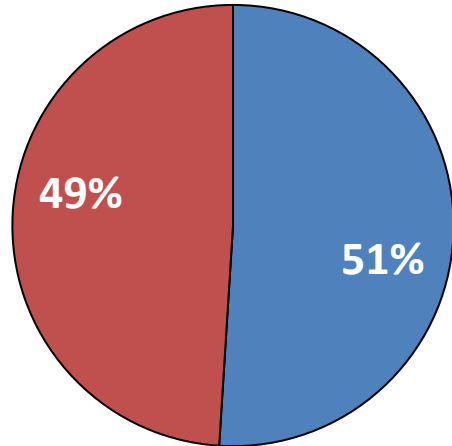


> 75 yrs of age, less fit (n = 52)



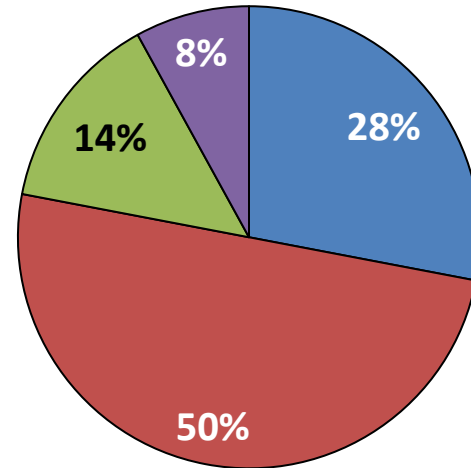
Case Disposition and Recommendation Impact

**Submitted Patient Case
(n = 174)**



■ Actual patient ■ Hypothetical patient

**Expert Rec Impact on Tx Choice
(n = 193)**



■ Changed Tx choice ■ Confirmed Tx choice
■ Barriers to Tx change ■ Undecided



Conclusions

- Data analysis showed expert consensus regarding Tx strategies for AML, including:
 - Venetoclax plus HMAs for older, less fit patients with ND AML
 - Targeted therapies for patients with AML and *FLT3* or *IDH1/2* mutations
- HCP practice patterns differed considerably from the experts for most cases
 - Fit, younger ND AML: CPX-351, 7+3 plus gemtuzumab ozogamicin, or venetoclax + HMA selected by experts in 71% of cases vs 20% for HCPs
 - Older, less fit ND AML: venetoclax plus HMA or LDAC selected by experts in 74% of cases vs 33% for HCPs
 - In first relapse AML cases with *FLT3* or *IDH1/2* mutations, experts chose targeted therapies in 87% of cases vs 41% for HCPs
- Differences between HCPs and experts suggest continued educational need to increase HCP awareness of best practices in AML

