## Precision Medicine Initiative (PMI) Committee Meeting

November 1, 2023



## Agenda –

- Role Call
- Project Status Updates
- Project Discussion Items
- Next Steps

## **Stakeholder Representation**



## **Project Status Updates**



## Project Updates – PMI Deliverables

Deliverable	Target Release Date	Target Release Vehicle
<b>ComboMATCH changes</b> (Disease fields, histology, behavior field code)	PROD Released: July 17, 2023	Screening Protocol EC Template v2.0
Designated Labs for Combo	PROD Released: Sept 6, 2023	Screening Protocol EC Template v3.0 (UAT Release w/EA – 07/27/2023)
Re-Screening ComboMATCH	PROD Released: Sept 27, 2023	MSRP Re-Screening EC Template v1.0 (UAT Release w/EA – 07/27/2023)
Cohort Migration	PROD OPEN Released: Sept 28, 2023	Treatment Protocol Cohort Migration EC Template v1.0
	PROD EC Temp v1.0 Released: Oct 27, 2023	
MyeloMATCH Tx Protocol Crossover (for S01, CTG01 and EA02)	n/a	Part of the existing Treatment Protocol workflow; no PRODUCTION OPEN Release required; the Westat team is prepared to support this workflow as Groups begin using it.
Re-Screening MM	n/a	No PRODUCTION OPEN Release is required for MM Re-Screening activities; the Westat team is prepared to support this workflow as Groups begin using it.
	PROD RS EC Temp v2.0: Nov tbd, 2023	MSRP Re-Screening EC Template v2.0 (UAT Release w/SWOG – 10/62023; 2 <sup>nd</sup> UAT Release w/SWOG – 10/24/2023; PROD RS EC Template v2.0 date TBD)
MyeloMATCH Stratification	PROD OPEN Release: Nov 22, 2023	Part of existing Treatment Protocol Workflow.
BETA PMI Screening Protocol ALS v2.0 New Forms: STMF, CLIA, PATH GROUP	BETA Release: Dec 22, 2023	Target date for BETA Release for Groups to initiate UAT activities. (accompanied by BETA Screening Protocol Central Study ALS v2.0)
PROD PMI Screening Protocol ALS v2.0 New Forms: STMF, CLIA, PATH GROUP	PROD Release: Feb 26, 2024	Target date for PRODUCTION Release for Groups to initiate implementation activities. (accompanied by PROD Screening Protocol Central Study ALS v2.0)

## Project Updates - EC Template Release Schedule

Template	Version	UAT Release Date	PROD Release Date	Details
PMI Screening Protocol EC Template	V1.0	Released: 12/16/2022	Released: 02/15/2023	
PMI Screening Protocol EC Template	V2.0	Released: 05/01/2023	Released: 07/17/2023	Includes CM changes (Disease fields, histology, behavior field code)
PMI Screening Protocol EC Template	V3.0	Released: 07/27/2023	Released: 09/06/2023	Includes updates for Designated Labs (DLAP fields)
PMI Screening Protocol EC Template <u>Fact Sheet</u>	v3.1	n/a	Target: 10/6/2023	Administrative version/release of the Fact Sheet to capture MyeloMATCH Field Help/Instructional Text for the 'Prior therapy name' field; to be used in conjunction with the PMI Screening Protocol EC Template v3.0.
PMI Treatment Protocol EC Template	V1.0	Released: 12/16/2022	Released: 02/15/2023	
PMI MSRP Re-Screening EC Template	V1.0	Released: 07/27/2023	Released: 09/27/2023	Supports ComboMATCH MSRP.
PMI MSRP Re-Screening EC Template	V2.0	Released: 10/06/2023 Re-released: 10/24/2023	Target: 11/xx/2023	Supports MyeloMATCH MSRP; PROD release date to be determined.
PMI Treatment Protocol Cohort Migration EC Template	V1.0	Released: 09/18/2023	Released: 10/27/2023	Supports cohort migration activities for ComboMATCH; PROD release date to be determined.
PMI Treatment Protocol Cohort Migration EC Template <u>Fact Sheet</u>	v1.1	Re-Released: 10/4/2023 Released: 09/18/2023	n/a	Administrative version/re-release of the UAT Fact Sheet to capture updated Field Help/Instructional Text for the 'Cohort migration assignment' field; to be used in conjunction with the UAT PMI Treatment Protocol Cohort Migration EC Template v1.0.

## ComboMATCH Priority 1 List

4	Protocol	Protocol Title	Group	Activated	Current Status	
	EAY191	Molecular Analysis for Combination Therapy Choice (ComboMATCH)	ECOG- ACRIN	Yes	10/4: Completed and no other updates 11/1: No Updates/Issues	
	2 EAY191- N4	Molecular Analysis for Combination Therapy Choice (ComboMATCH) EAY191-N4: A Randomized Trial of Selumetinib and Olaparib or Selumetinib Alone in Patients with Recurrent or Persistent RAS Pathway Mutant Ovarian and Endometrial Cancers A ComboMATCH Treatment Trial	NRG	Yes	<ul> <li>10/4: Still waiting on the edit check with CTSU, Randonode was updated; Westat is doing internal testing</li> <li>11/1: Received edit check, waiting to be able to test the edit check; waiting on patients. Westat will provide the patients to test by EOD 11/1</li> </ul>	
	B EAY191- E4	A ComboMATCH Treatment Trial ComboMATCH Treatment Trial E4: Nilotinib and Paclitaxel in Patients with Prior Taxane-Treated Solid Tumors	ECOG- ACRIN	Yes	10/4: Working on incorporating ineligibility requirements 11/1: EA Leadership has questions around the ineligibility requirements that NCI will need to review	
2	EAY191- N2	A ComboMATCH Treatment Trial EAY191-N2: Phase 2 Trial of Fulvestrant and Binimetinib in Patients with Hormone Receptor-Positive Metastatic Breast Cancer with a Frameshift or Nonsense Mutation or Genomic Deletion in NF1	NRG	Yes	<ul> <li>10/4: Submitted OPEN request forms and new forms for STEP 1 and 2 are in OPEN.</li> <li>CSTU added a widget on Treatment model Step 2. Need patients but need to update Rave.</li> <li>11/1: Tested Cohort Migration; in the process of CTSU ineligibility edit check</li> </ul>	
!	5 EAY191- S3	Phase 2 Study of Paclitaxel + Ipatasertib in Taxane- Refractory Participants with AKT-Altered Advanced Non-Breast Solid Tumors	SWOG	Yes	<ul> <li>10/4: Westat will send over additional documentation for ineligibility implementation and will remove SSN.</li> <li>11/1: Reading through inelibility information that NCI provided; will work on impelmentaton</li> </ul>	

## ComboMATCH Priority 2 List

#	Protoco I #	Protocol Title	Group	Activated	Current Status
6	EAY191- A6	A ComboMATCH Treatment Trial: FOLFOX in Combination with Binimetinib as 2nd Line Therapy for Patients with Advanced Biliary Tract Cancers with MAPK Pathway Alterations	Alliance	Yes	10/18: Scoping out the efforts needed to implement ineligibility workflow 11/1: Working through the email for the workflow
7	EAY191- A3	Palbociclib and Binimetinib in RAS-Mutant Cancers	Alliance	No	<ul> <li>10/18: Fit for purpose testing re-scheduled for 10-26-2023, due to delay in matchbox algorithm setup, if testing passes will activate 11-15-2023.</li> <li>11/1: Working through the email for the workflow; completed FFP testing and confirmed activation on 11/15.</li> </ul>

## ComboMATCH Priority 3 List

#	Protocol #	Protocol Title	Group	Current Status
8	EAY191-A2	Olaparib Plus Low-Dose Alpelisib for Breast Cancer	Alliance	<i>9/20:</i> Dropped
9	EAY191-C1	Phase 2 Subprotocol of the Combination of a MEK Inhibitor and a Pan-RAF Inhibitor in Patients with Relapsed/Refractory Tumors Harboring Activating MAPK Pathway Mutations	COG	10/18:Waiting on finalized protocol. Have already reviewed the Ineligibility capture and have a plan to address in the upcoming study build 11/1: Waiting on protocol; CTEP still in discussions around PK discussions
10	EAY191-E5	ComboMATCH Treatment Trial E5: A Randomized Phase II Study of AMG 510 (Sotorasib) with or Without Panitumumab in Advanced Solid Tumors	ECOG- ACRIN	<b>10/4:</b> Working on OPEN setup 11/1: Continuing to work on Cohort Migration; waiting for CTEP approval. Hope to complete work by 11/30.
11	EAY191-N5	Molecular Analysis for Combination Therapy Choice (ComboMATCH) EAY191-N5: A Randomized Trial of Neratinib, A Pan-ERBB Inhibitor, Alone or in Combination with Palbociclib, a CDK4/6 Inhibitor, in Patients with HER2+ Gynecologic Cancers and Other Solid Tumors A ComboMATCH Treatment Trial	NRG	<ul> <li>10/4: Protocol was submitted on 9/15 to PIO; starting study build but hoping for an extension; date is Oct 23<sup>rd</sup> – No cohort migration, just a crossover David will follow up with NRG and the PIs; cannot verify copy number loss; my require a protocol update</li> <li>David will send Rich the language to send to the sites to update</li> <li>11/1: Protocol received an extension; OEWG is 2/20/24; prioritizing build and working on forms;</li> </ul>

## MyeloMATCH Priority 1 List

#	Protocol #	Protocol Title	Group	Current Status
1	MYELOMATCH	Master Screening and Reassessment Protocol (MSRP) for the NCI MyeloMATCH Clinical Trials	SWOG	<ul> <li>10/4: Started work on this again, target to finish is end of the month; SWOG can perform internal testing in OPEN UAT, that is okay to do that.</li> <li>11/1: Need additional details on integration testing; Westat will reach out to SWOG to work on integration testing; SWOG will need to create enrollments in UAT. Initial testing is 2-3 enrollments. SWOG will be ready today. Need clarification on how TAP works so would like to test that functionality in UAT. Westat is hoping to get through testing in a week.</li> </ul>
2	MM1YA-S01	A Randomized Phase II Study Comparing Cytarabine + Daunorubicin (7 + 3) to (Daunorubicin and Cytarabine) Liposome, Cytarabine + Daunorubicin + Venetoclax, and Azacitidine + Venetoclax in Patients Aged 59 or Younger with High-Risk (Adverse) Acute Myeloid Leukemia; A MYELOMATCH Clinical Trial	SWOG	10/4: Started work, but focusing on Screening as a priority 11/1: SWOG is pretty far ahead in this build; still focusing on Screening.
3	MM1YA-CTG01	A Measurable Residual Disease (MRD) Driven, Phase II Study of Venetoclax Plus Chemotherapy for Newly Diagnosed Younger Patients with Intermediate Risk Acute Myeloid Leukemia: A Tier 1 MYELOMATCH Clinical Trial	CCTG	11/1: Received the ineligibility workflow with scripts, starting to process that. Waiting on screening patients to come through.
6	MM1OA-EA02 (Concept)	A Randomized Phase II Study of HMA-Based Therapies for the Treatment of Older Adults with Newly Diagnosed FLT3-Mutated Acute Myeloid Leukemia: A myeloMATCH Treatment Trial	ECOG- ACRIN	11/1: Have been focusing on CM at this time; working on RAVE build and OPEN checklist. Testing treatment trials in possibly two weeks

## MyeloMATCH Priority 2 List

#	Protocol #	Protocol Title	Group	Current Status
4	MM2YA-EA01	Eradicating Measurable Residual Disease in Patients with Acute Myeloid Leukemia (AML) Prior to Stem Cell Transplantation (ERASE): A MyeloMATCH Treatment Trial	ECOG- ACRIN	11/1:
5	MM10A-S02	A Randomized Phase II Study of Targeted Agents with the Oral DNMT1 Inhibitor Cedazuridine-Decitabine (ASTX727) for the Treatment of Newly Diagnosed TP53-Mutated Acute Myeloid Leukemia in Older Patients: A myeloMATCH Treatment Trial	SWOG	11/1:
7	MM1MDS- EA03 (Concept	ERADICATE MDS with TP53 Study: Efficacy of targeted Agents Combinations with Oral DNMT1 Inhibitor C-DEC (ASTX727) for Treatment of Higher Risk Myelodysplastic Syndromes with TP53 Mutations: A Non-Comparative, Parallel, Multi-Arm Phase 2 Study	ECOG- ACRIN	11/1:

## **PMI Project Discussion Items**



## Next Steps





## Next meeting will be on 11/15/2023 at 1:00pm EST

## Agenda

- Role Call
- Project Status Update
- Group Status Update
- Review FAQs

## Communication



#### Contact the PMI Mailbox for any PMI related questions & comments

#### pmistandards@nih.gov

The project team will respond within 48 hours with a response or a follow up

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#### PMI Wiki

#### https://wiki.nci.nih.gov/display/CDISC/Precision+M edicine+Initiative

All presentations, recordings, minutes, project documents and releases will be posted on this wiki



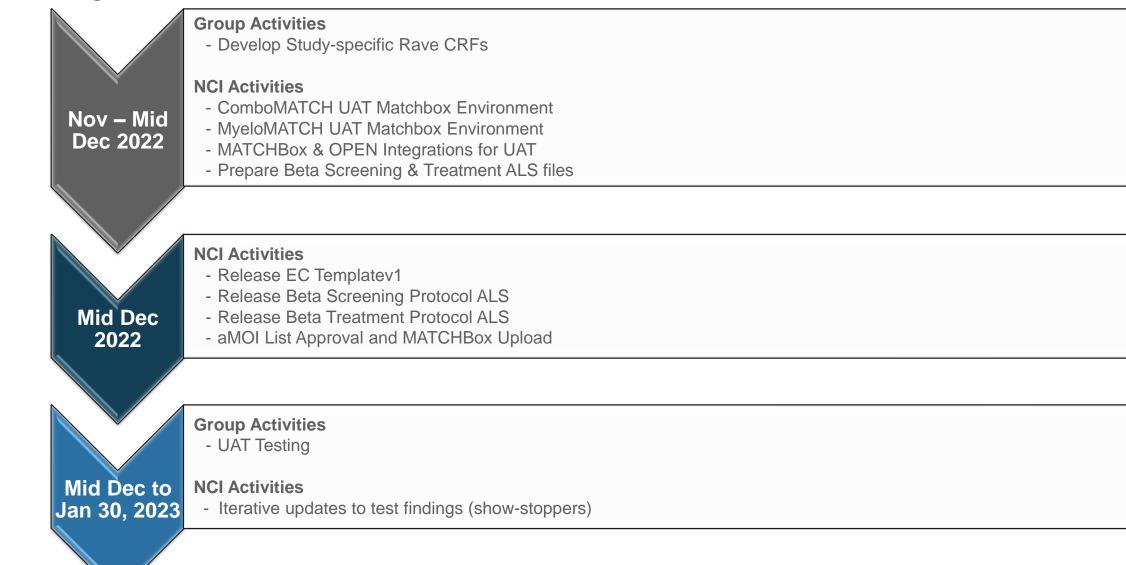
www.cancer.gov/espanol

www.cancer.gov

## Appendix



#### **Target Timeline**



## Target Timeline

Jan 31, 2023	NCI Activities - Release EC Template v1 - Release Prod Screening Protocol ALS - Release Prod Treatment Protocol ALS - Release Prod Central Study ALS - Release NCI OPEN Integrations for Prod - Release NCI Genexus Installation (MyeloMATCH)
Jan 31 to Feb 14, 2023	NCI + Group Activities - Support Group Study Builds
Feb 14,	NCI + Group Activities
2023	- Launch BOTH Initiatives
Spring	NCI Activities
2023	- Release Additional ALS(s)

## ComboMATCH Priority 1 List

Study #	Document Number	Document Title	Group	Current Status	Next Steps	Primary Agent Name (P) Other Agent Name (O)
1	EAY191	Molecular Analysis for Combination Therapy Choice (ComboMATCH)	ECOG-ACRIN	8/15/2022 Approval on Hold	Revision 3 back from CIRB with approval. On hold for DTL, funding sheet and IND filing.	
2	EAY191-N4	Molecular Analysis for Combination Therapy Choice (ComboMATCH) EAY191-N4: A Randomized Trial of Selumetinib and Olaparib or Selumetinib Alone in Patients with Recurrent or Persistent RAS Pathway Mutant Ovarian and Endometrial Cancers A ComboMATCH Treatment Trial	NRG	8/15/2022 Approval on Hold	Revision 4 back from CIRB with approval. On hold for DTL, funding sheet and IND filing.	<ul> <li>(P) Olaparib (AZD2281)</li> <li>(747856)</li> <li>(O) Selumetinib</li> <li>(AZD6244 hydrogen sulfate) (748727)</li> </ul>
3	EAY191-E4	A ComboMATCH Treatment Trial ComboMATCH Treatment Trial E4: Nilotinib and Paclitaxel in Patients with Prior Taxane-Treated Solid Tumors	ECOG-ACRIN	8/26/2022 Approval on Hold	Revision 3 back from CIRB with approval. On hold for DTL, funding sheet and IND filing.	(P) Nilotinib (747599) (O) Paclitaxel (Taxol) (673089)
4	EAY191-N2	A ComboMATCH Treatment Trial EAY191-N2: Phase 2 Trial of Fulvestrant and Binimetinib in Patients with Hormone Receptor-Positive Metastatic Breast Cancer with a Frameshift or Nonsense Mutation or Genomic Deletion in NF1	NRG	8/26/2022 Approval on Hold	Revision 3 back from CIRB with approval. On hold for DTL, funding sheet and IND filing.	(P) Binimetinib (788187) (O) Fulvestrant (Faslodex) (719276)
5	EAY191-S3	Phase 2 Study of Paclitaxel + Ipatasertib in Taxane- Refractory Participants with AKT-Altered Advanced Non-Breast Solid Tumors	SWOG	8/29/2022 Approval on Hold	Revision 4 back from CIRB with approval. On hold for DTL, funding sheet and IND filing.	(P) Ipatasertib (781451) (O) Paclitaxel (Taxol) (673089)

## ComboMATCH Priority 2 List

Stu #	dy Document Number	Document Title	Group	Current Status	Next Steps	Primary Agent Name (P) Other Agent Name (O)
6	EAY191-A6	A ComboMATCH Treatment Trial: FOLFOX in Combination with Binimetinib as 2nd Line Therapy for Patients with Advanced Biliary Tract Cancers with MAPK Pathway Alterations	Alliance	Approval on Hold 9/14/2022	10/24. CTEP currently reviewing stips.	<ul> <li>(P) Binimetinib (788187)</li> <li>(O) OXALIplatin</li> <li>(Eloxatin) (266046)</li> <li>(O) 5-Fluorouracil (5-FU)</li> <li>(19893)</li> <li>(O)Leucovorin calcium</li> <li>(3590)</li> </ul>
7	EAY191-A3	Palbociclib and Binimetinib in RAS-Mutant Cancers	Alliance	Approval on Hold 8/26/2022	Rev 2 sent to CIRB . Reviewed on 10/6, study team responding to stips	(P) Palbociclib (PD- 0332991) (772256) (O) Binimetinib (788187)

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8	EAY191-A2	Olaparib Plus Low-Dose Alpelisib for Breast Cancer	Alliance	In Review 4/4/2022	Rev 3 recv;d 10/19, currently with LR. Looks like it'll be another disapproval. Not yet sent to CIRB for initial review.	(P) Olaparib (AZD2281) (747856) (O) Alpelisib (BYL719) (801658)
9	EAY191-C1	Phase 2 Subprotocol of the Combination of a MEK Inhibitor and a Pan-RAF Inhibitor in Patients with Relapsed/Refractory Tumors Harboring Activating MAPK Pathway Mutations	COG	In Review 9/29/2022	Protocol reviewed at PRC 10/20; CR is being put together	<ul> <li>(P) Selumetinib</li> <li>(AZD6244 hydrogen sulfate) (748727)</li> <li>(O) DAY101 (TAK-580, MLN-2480) (800798)</li> </ul>
10	EAY191-E5	ComboMATCH Treatment Trial E5: A Randomized Phase II Study of AMG 510 (Sotorasib) with or Without Panitumumab in Advanced Solid Tumors	ECOG-ACRIN	In Review 2/22/2022	Waiting for revision 2 since 9/28; reminder was sent to EA 10/24	(P) AMG 510 (Sotorasib) (825510) (O) Panitumumab (742319)
11	EAY191-N5	Molecular Analysis for Combination Therapy Choice (ComboMATCH) EAY191-N5: A Randomized Trial of Neratinib, A Pan-ERBB Inhibitor, Alone or in Combination with Palbociclib, a CDK4/6 Inhibitor, in Patients with HER2+ Gynecologic Cancers and Other Solid Tumors A ComboMATCH Treatment Trial	NRG	In Review 11/22/2021	Rev 3 disapproval sent 10/15; waiting for Rev 4	(P) Neratinib (783782) (O) Palbociclib (PD- 0332991) (772256)

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2	MM1YA-S01	A Randomized Phase II Study Comparing Cytarabine + Daunorubicin (7 + 3) to (Daunorubicin and Cytarabine) Liposome, Cytarabine + Daunorubicin + Venetoclax, and Azacitidine + Venetoclax in Patients Aged 59 or Younger with High-Risk (Adverse) Acute Myeloid Leukemia; A MYELOMATCH Clinical Trial	SWOG	9/2/2022: Approval on Hold	CIRB Reviewed on 9/15, version with stips under CTEP review now.
3	MM1YA-CTG01	A Measurable Residual Disease (MRD) Driven, Phase II Study of Venetoclax Plus Chemotherapy for Newly Diagnosed Younger Patients with Intermediate Risk Acute Myeloid Leukemia: A Tier 1 MYELOMATCH Clinical Trial	CCTG	9/2/2022: Approval on Hold	CIRB Review Oct 6, responding to stips
6	MM1OA-EA02 (Concept)	A Randomized Phase II Study of HMA-Based Therapies for the Treatment of Older Adults with Newly Diagnosed FLT3-Mutated Acute Myeloid Leukemia: A myeloMATCH Treatment Trial	ECOG-ACRIN	8/29/2022: Approved, no protocol yet	Waiting for protocol

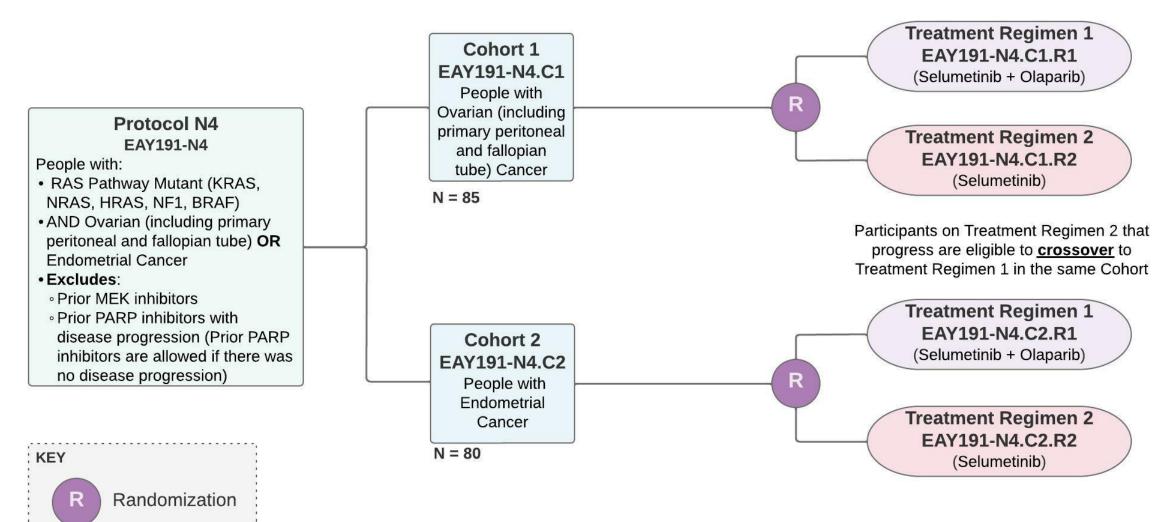
## MyeloMATCH Priority 2 List

Study #	Document Number	Document Title	Group	Current Status	Next Steps
4	MM2YA-EA01	Eradicating Measurable Residual Disease in Patients with Acute Myeloid Leukemia (AML) Prior to StEm Cell Transplantation (ERASE): A MyeloMATCH Treatment Trial	ECOG-ACRIN	1/21/2022: In Review	Reviewed by CIRB, responding to stips
5	MM10A-S02	A Randomized Phase II Study of Targeted Agents with the Oral DNMT1 Inhibitor Cedazuridine-Decitabine (ASTX727) for the Treatment of Newly Diagnosed TP53-Mutated Acute Myeloid Leukemia in Older Patients: A myeloMATCH Treatment Trial	SWOG	10/17/2022: In Review	Scheduled for PRC 11/4
7	MM1MDS- EA03 (Concept	ERADICATE MDS with TP53 Study: Efficacy of taRgeted Agents Combinations with Oral DNMT1 Inhibitor C-DEC (ASTX727) for TrEatment of Higher Risk myElodysplastic Syndromes with TP53 Mutations: A Non-Comparative, Parallel, Multi-Arm Phase 2 Study	ECOG-ACRIN	8/12/2022: Approval on hold, waiting for drug commitment	W+A6:G8aiting for Astex commitment, Genentech has provided commitment on 10/20

## **Review Schemas**



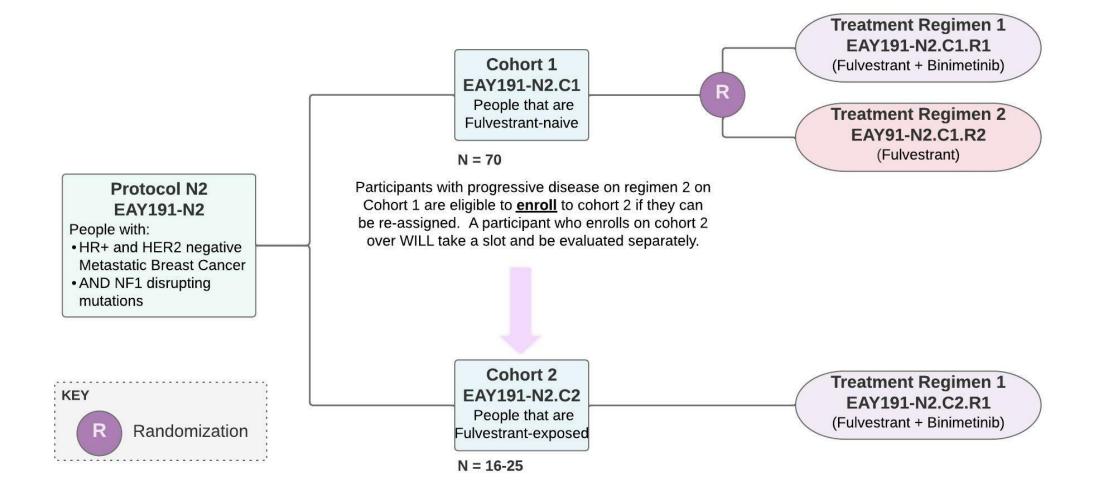
#### EAY191-N4



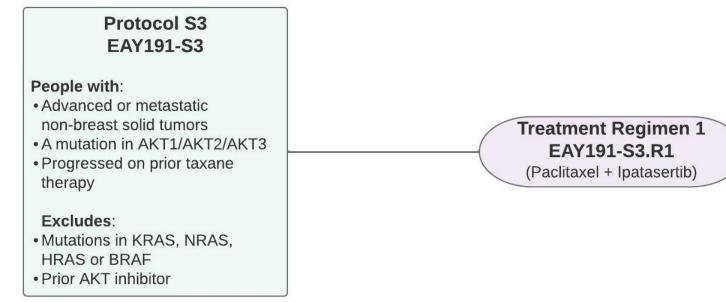
#### EAY191-E4



#### EAY191-N2- Draft



#### EAY191-S3



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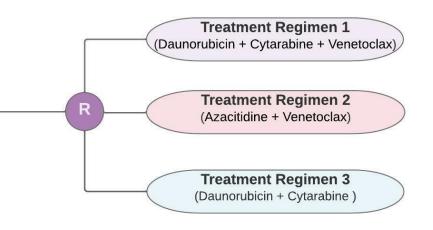
#### MMIYA-CTG01 Draft

#### Protocol MM1YA-CTG01

People who have Acute Myeloid Leukemia in the intermediate risk ELN category

#### **Excludes People with:**

- · isolated myeloid sarcoma
- prior therapy for AML (except for hydroxyurea and leukapheresis to control blood counts)
- the following categories of AML:
- Favorable cytogenetics: (t(8;21)q22;q22.1); RUNX1-RUNX1T1, inversion 16(p13.1;q22), t(16;16)(p13.1;q22);CBFB-MYH11
- CEBPA biallelic mutations
- NPM1 mutation
- AML with PML-RAR
- AML with any adverse cytogenetics, TP53 mutation, RUNX1 mutation, ASXL1, 11q23/KMT2 rearrangements
- AML with FLT3-ITD or FLT3-TKD mutations
- Therapy related AML, or AML following a diagnosis of myelodysplasia or myeloproliferative neoplasm



#### N = 153

