Precision Medicine Initiative (PMI) Committee Meeting

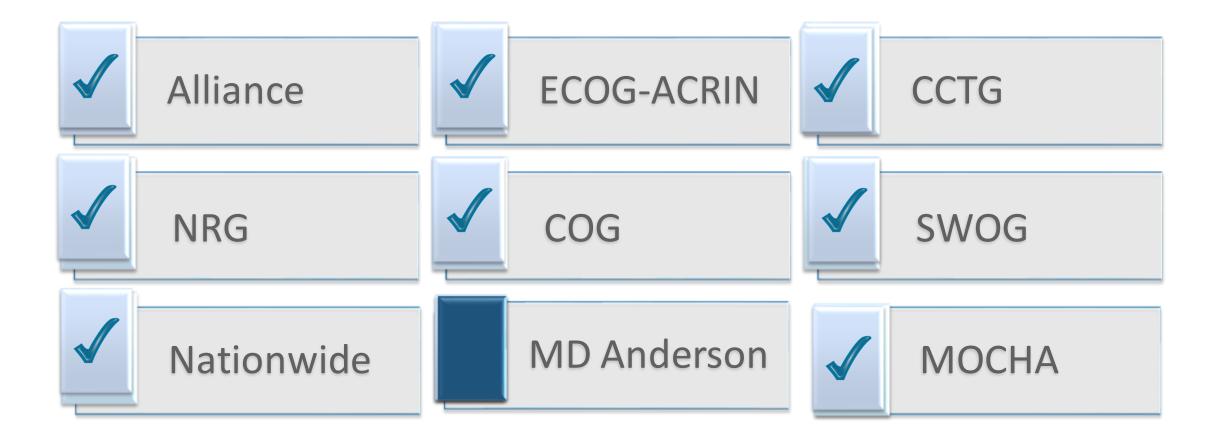
April 3, 2024



Agenda

- Roll Call
- Project Updates
- PMI Committee Call Process Updates
- Reassignment Workflow- Demo
- Open Discussion

Stakeholder Representation



Project Updates



Project Updates – PMI Deliverables

	Deliverable	Target Release Date	Target Release Vehicle
	ComboMATCH changes (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	PROD RS EC Temp v2.0 Released: Nov 30, 2023	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. MSRP Re-Screening EC Template v2.0 (UAT Release w/SWOG – 10/6/2023; 2 nd UAT Release w/SWOG – 10/24/2023; 3rd UAT Release to SWOG – 11/07/2023; PROD RS EC Template v2.0 target date 11/30/2023)
	MyeloMATCH Stratification	PROD OPEN Release: February 12, 2023	Part of existing Treatment Protocol Workflow.
	BETA PMI Screening Protocol ALS v2.0 New Forms: STMF, CLIA, PATH GROUP	Released: BETA Release: January 12, 2024	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: April 22, 2024	Target date for PRODUCTION Release for Groups to initiate implementation activities. (accompanied by PROD Screening Protocol Central Study ALS v2.0)
NIH	NATIONAL CANCER INSTITUTE		3

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 Fact Sheet	v3.1	n/a	Released: 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 Treatment Protocol EC Template Fact Sheet	V1.1	n/a	Released: 11/16/2023	Administrative version/release of the Fact Sheet to provide a stand-alone TX Protocol-specific document allows Groups to easily access the Fact Sheet information; to be used in conjunction with the PMI Treatment Protocol EC Template v1.0.
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	Re-released: 11/07/2023 Re-released: 10/24/2023 Released: 10/06/2023	Released: 11/30/2023	Supports MyeloMATCH MSRP; PROD release date confirmed.
PMI Treatment Protocol Cohort Migration EC Template	V1.0	Released: 09/18/2023	Released: 10/27/2023	Supports cohort migration activities for ComboMATCH.
PMI Treatment Protocol Cohort Migration EC Template Fact Sheet	v1.1	Re-Release: 10/4/2023	Released : 10/27/2023	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

#	Protocol #	Protocol Title	Group	Activated	Current Status
1	. EAY191	Molecular Analysis for Combination Therapy Choice (ComboMATCH)	ECOG- ACRIN	Yes	2/21: No Updates 3/20: No updates 4/3: No updates
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	2/21: No Updates 3/20: No updates 4/3: No updates
(1)	EAY191-E4	A ComboMATCH Treatment Trial ComboMATCH Treatment Trial E4: Nilotinib and Paclitaxel in Patients with Prior Taxane-Treated Solid Tumors	ECOG- ACRIN	Yes	2/21: No Updates 3/20: No updates 4/3: Temp suspended
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	2/7: Got comments from Amendment 1, need to update EC template & need updated protocol 2/21: Protocol Submitted – working on updates 3/20: No updates 4/3: Amendment approvedreleasing April 15 th .
	EAY191-S3	Phase 2 Study of Paclitaxel + Ipatasertib in Taxane- Refractory Participants with AKT-Altered Advanced Non-Breast Solid Tumors	SWOG	Yes	2/7: Removed the SSN and full name and released 2/21: No updates 3/20: No Updates 4/3: No updates 7

ComboMATCH Priority 2 List

#	; F	Protoco #	Protocol Title	Group	Activated	Current Status
e	5	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		Yes	2/21: No Updates 3/20: No Updates 4/3: Going through getting IDs into OPEN to work on ECs
7	7	EAY191- A3	Palbociclib and Binimetinib in RAS-Mutant Cancers	Alliance	Yes	2/21: No Updates 3/20: Suspend Cohort 3. 4/3: No updates, : Suspend Cohort 3.

ComboMATCH Priority 3 List

#	Protocol #	Protocol Title	Group	Current Status
8	EAY191-A2	Olaparib Plus Low-Dose Alpelisib for Breast Cancer	Alliance	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	DROPPED
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	1/10: No Updates 2/7: Active 2/21: No Updates 3/20: No updates 4/3: No updates
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	1/10: No Updates 2/7: Released to production on 2/12 2/21: No updates 3/20: No updates 4/3: No updates

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	2/21: Completed, working through step to prep for activation – Westat is reviewing the bug found in OPENWestat will provide an update EOD 3/20: Pending FFP; having trouble getting a volunteer for testing. If we do not identify someone soon, we will need to use an internal person. 4/3: Going through FFP; almost completed
2	MM1YA-SO1	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	 2/7: Ready for MATCHbox integration testing on Jan 29. 2/21: - Pending patient registration – until Westat bug is updated 3/20: Halfway through integration testing, have a few bugs they are working through. 4/3: Going through FFP; almost completed; sent email template messages out for feedback
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	 2/21: Waiting on new testing patients to complete testing – have a request in 3/20: Working on integration testing, received extra test patients. Put in a request with CTSU for a fix. Identified sites for FFP testing. 4/3: Integration testing and FFP testing; issues with Withdrawal consent and treatment but working with Westat
6	MM10A-EA02	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	 2/7: Cohort/Stratum Assignment – Redoing the checklist – won't be ready for FFP by this week but working to meet that date. 2/21: Working with randostats team to get files in place to begin testing 3/20: Working on integration testing, hoping to complete in the next few days. Reached out to FFP testing, waiting on the sites to get back to us. 4/3: FFP testing is completed; finalizing Rave build, working on validation Action Item: Leila will reach out for feedback

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	2/21: No Updates 3/20: No Updates 4/3: No updates
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	2/21: No Updates 3/20: No Updates 4/3: Begin this soon as FFP is almost done; build has started; will take about a month until integration testing
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	2/21: Withdrawn

PMI Committee Call Process Updates



PMI Committee Meeting Process Updates

- Meeting Process updates:
 - Meeting minutes will be documented from each meeting.
 - Minutes will include the action items and owners
 - Action Items will be reviewed until completed, as needed.
- PMI Committee Wiki Page Updates:
 - New Section added for Meeting Minutes
 - Currently updated with meeting minutes from the last 2 PMI calls

Meeting Minutes

- 03_20_2024 PMI Committee Meeting Minutes
- 02_21_2024 PMI Committee Meeting Minutes

Action Items from 3.20.24 Meeting

Action Item	Owner	Status/Updates
Identify a date to present the Stakeholder Feedback findings to the PMI Committee	Chioma Ani	Planned date to present findings to PMI Committee- 5/08/2024
Reassignment workflow with 2 Scenarios: Integrations team to identify a date to present the upcoming changes to the PMI Committee	Integrations Team	Item to be reviewed today 4/3/2024
Add the information for How to Suspend a Protocol for interim analysis to the user guide	Leila Abraham	Due to MM Activation Priorities, this item will be updated to the RSS User guide by 5/15/2024

Reassignment Workflow: Demo

Reassignment Workflow- DEMO

- Implemented workflow will include 2 scenarios:
 - Participant is ineligible
 - Protocol not open at site
 - If either of the 2 are selected, assignment should be regenerated excluding the initially selected assignment
- Feature enhancement is currently available in OPEN UAT
- Additional enhancements to this feature can be requested moving forward to add additional scenarios with the current base design in place
 - Planned Addition: Add a separate "TAP Assignment" button for MyeloMATCH
 - Requested enhancements will be reviewed by the Project team and PMI Leadership before addition

Next Steps



Open Discussion

- Item 1: The SWOG protocol coordinator asked if there is an allowable window for the performance status that is entered at initial registration to the MSRP. Since this field is a MATCHBox requirement, do you have a specific time that you would expect that to be done? For example, does it need to be evaluated within a certain number of days prior to the registration?
- Rich: Critical date comes in for MSRP as part of their sub study assignment. The Performance status needs to be entered immediate to be able to assigned to the treatment. Needs to be entered in the same time the labs are sent in. No date requirement.

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

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

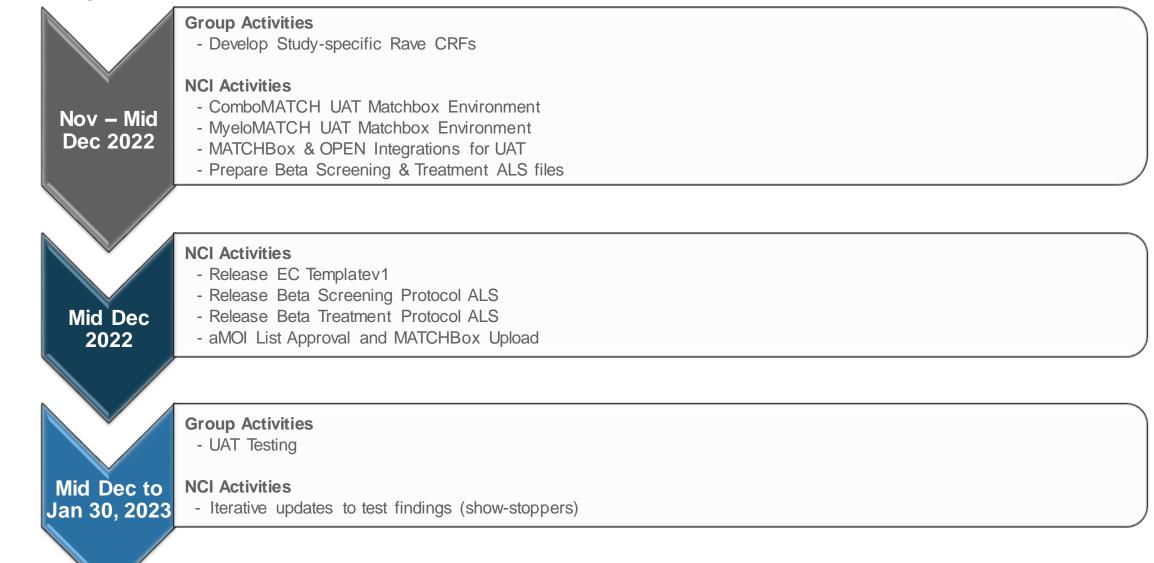
Appendix

CTEP CDISC Policy Governance Review

Compliance for Use of ALS Versions

- Standard forms built in the LPO's Global Library within 60 days of production ALS release to reflect additions/updates included in the ALS
 - Example: Release ALSv7.X on November 1, LPOs would be expected to incorporate the ALSv7.X into their Global Library by January 1
 - -ALS versions for new study builds
 - LPOs will be required to use the most current version of the ALS available at the time of each study build

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, 2023	NCI + Group Activities - Launch BOTH Initiatives
Spring 2023	NCI Activities - 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.	(P) Olaparib (AZD2281) (747856) (O) Selumetinib (AZD6244 hydrogen sulfate) (748727)
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

Study #	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.	 (P) Binimetinib (788187) (O) OXALI platin (Eloxatin) (266046) (O) 5-Fluorouracil (5-FU) (19893) (O)Leucovorin calcium (3590)
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)

ComboMATCH Priority 3 List

Study #	Document Number	Document Title	Group	Current Status	Next Steps	Primary Agent Name (P) Other Agent Name (O)
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	(P) Selumetinib (AZD6244 hydrogen sulfate) (748727) (O) DAY101 (TAK-580, MLN-2480) (800798)
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		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)

MyeloMATCH Priority 1 List

Study #	Document Number	Document Title	Group	Current Status	Next Steps
1	MYELOMATCH	Master Screening and Reassessment Protocol (MSRP) for the NCI MyeloMATCH Clinical Trials	SWOG	11/12/2021: Approval on Hold	CIRB Reviewed on 9/15, version with stips under CTEP review now.
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

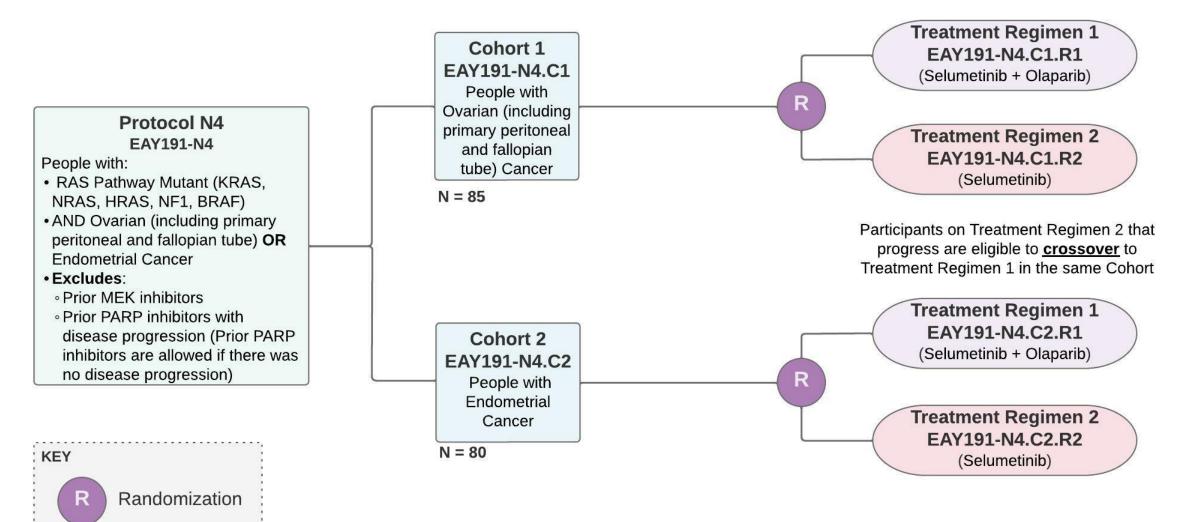
MyeloMATCH Priority 2 List

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	MM1OA-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	, , , , , ,	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



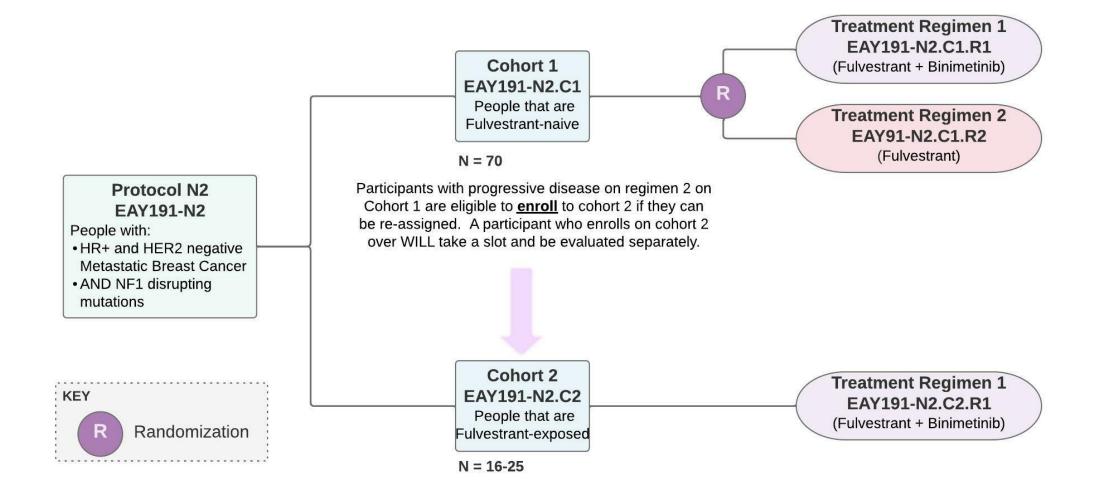
EAY191-N4



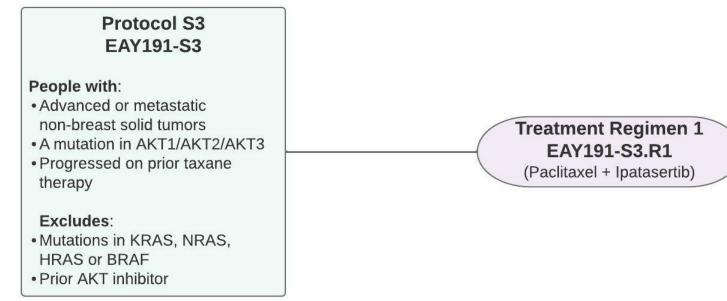
EAY191-E4



EAY191-N2- Draft



EAY191-S3



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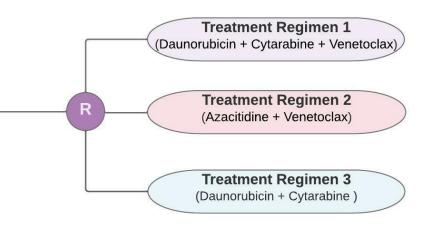
MMIYA-CTG01 Draft

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

