

# Solid Tumor Protocol List

July 2021

Ewing treatment protocols			
Study	Clinical trial name	Phase/type	Age
<b>Aflac ST1502 CHOANOME II</b>	AflacST1502: A Phase II Study of Sirolimus in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr
<b>AflacST1903</b>	A Maintenance Protocol of Sirolimus in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC# 788607, IND# 141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT0322003">https://clinicaltrials.gov/ct2/show/NCT0322003</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr

<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr
<b>COG ADVL1721</b>	A Non-randomized, Open-label, Multi-center, Phase I/II Study of PI3K Inhibitor Copanlisib in Pediatric Patients with Relapsed/ Refractory Solid Tumors or Lymphoma <a href="https://clinicaltrials.gov/ct2/show/NCT03458728">https://clinicaltrials.gov/ct2/show/NCT03458728</a>	I/II	≥6 mo to ≤21 yr
<b>Ignyta RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr
<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>COG ADVL1414</b>	ADVL1414, A Phase 1 Study of Selinexor (KPT-330, IND# 125052), A Selective XPO1 Inhibitor, in Recurrent and Refractory Pediatric Solid Tumors, including CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02323880">https://clinicaltrials.gov/ct2/show/NCT02323880</a>	I	≥12 mo to ≤21 yr
<b>COG ADVL1514</b>	ADVL1514, A Phase 1 Study of ABI-009 (Nab-Rapamycin) (IND# 133537) in Pediatric Patients with Recurrent or Refractory Solid Tumors, Including CNS Tumors as a Single Agent and in Combination with Temozolomide and Irinotecan <a href="https://clinicaltrials.gov/ct2/show/NCT02975882">https://clinicaltrials.gov/ct2/show/NCT02975882</a>	I	≥12 mo to ≤21 yr
<b>COG ADVL1921</b>	Phase 1 Study to Evaluate the Safety and Pharmacokinetics of Palbociclib (Ibrance®) in Combination with Irinotecan and Temozolomide in Pediatric Patients with Recurrent or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03709680">https://clinicaltrials.gov/ct2/show/NCT03709680</a>	I	≥2 to <21 yr
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr
<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03507491">https://clinicaltrials.gov/ct2/show/NCT03507491</a>	I	≥6 mo to ≤30 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr

<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr
<b>IMPACT</b>	Informational Meetings for Planning And Coordinating Treatment (IMPACT)	III	≥1 to <18 yr
<b>Ewing biology, supportive treatment and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj
<b>NCICOVID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG ALTE03N1</b>	Key Adverse Events after Childhood Cancer (cardiac, stroke, secondary malign)/closed to AVN patients as of 11-26-08 <a href="https://clinicaltrials.gov/ct2/show/NCT00082745">https://clinicaltrials.gov/ct2/show/NCT00082745</a>	Non-therapeutic	≤21 yr at dx
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age
<b>DFHCC ctDNA</b>	Evaluation of ctDNA as a Prognostic Biomarker for Patients with Newly Diagnosed Localized Ewing Sarcoma or Osteosarcoma	Non-therapeutic	≥12 mo
<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr
<b>Neuroblastoma treatment protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>COG ANBL1232<sup>REQ</sup></b>	Utilizing Response- and Biology-based Risk Factors to Guide Therapy in Patients with Non-high-Risk Neuroblastoma <a href="https://clinicaltrials.gov/ct2/show/NCT02176967">https://clinicaltrials.gov/ct2/show/NCT02176967</a>	III	<12 mo at diag with INRG Stage L1 <18 mo at diag with INRG Stage L2 or Stage Ms nbl
<b>COG ANBL1531</b>	ANBL1531: A Phase 3 Study of <sup>131</sup> I-Metaiodobenzylguanidine ( <sup>131</sup> I-MIBG) or Crizotinib Added to Intensive Therapy for Children with Newly Diagnosed High-risk Neuroblastoma (NBL) (IND# 134379) <a href="https://clinicaltrials.gov/ct2/show/NCT03126916">https://clinicaltrials.gov/ct2/show/NCT03126916</a>	III	≥365 days to ≤30 yr
<b>Aflac ST1502 CHOANOMEII</b>	AflacST1502: A Phase II Study of Sirolimus in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr

<b>AflacST1903</b>	A Maintenance Protocol of Sirolimus in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC# 788607, IND# 141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG ANBL1821</b>	ANBL1821: A Phase 2 Randomized Study of Irinotecan/ Temozolomide/Dinutuximab with or without Eflornithine (DFMO) (IND# 141913) in Children with Relapsed, Refractory or Progressive Neuroblastoma <a href="https://clinicaltrials.gov/ct2/show/NCT03794349">https://clinicaltrials.gov/ct2/show/NCT03794349</a>	II	≥1 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT03220035">https://clinicaltrials.gov/ct2/show/NCT03220035</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr
<b>COG ADVL1721</b>	A Non-randomized, Open-label, Multi-center, Phase I/II Study of PI3K Inhibitor Copanlisib in Pediatric Patients with Relapsed/Refractory Solid Tumors or Lymphoma <a href="https://clinicaltrials.gov/ct2/show/NCT03458728">https://clinicaltrials.gov/ct2/show/NCT03458728</a>	I/II	≥6 mo to ≤21 yr

<b>Ignyta RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr
<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>NANT 2013-01</b>	NANT 2013-01: A Phase I Dose Escalation Study of Autologous Expanded Natural Killer (NK) Cells for Immunotherapy of Relapsed Refractory Neuroblastoma with Dinutuximab +/- Lenalidomide <a href="https://clinicaltrials.gov/ct2/show/NCT02573896">https://clinicaltrials.gov/ct2/show/NCT02573896</a>	I	≤30 yr
<b>NANT 2015-02<sup>REQ</sup></b>	NANT 2015-02: Phase 1 Study of Lorlatinib (PF-06463922), an Oral Small Molecule Inhibitor of ALK/ROS1, for Patients with ALK-driven Relapsed or Refractory Neuroblastoma <a href="https://clinicaltrials.gov/ct2/show/NCT03107988">https://clinicaltrials.gov/ct2/show/NCT03107988</a>	I	≥12 mo
<b>NANT 2017-01<sup>REQ</sup></b>	NANT 2017-01: A Phase I Study of <sup>131</sup> I-MIBG with Dinutuximab for Relapsed/Refractory Neuroblastoma (IND# 137554) <a href="https://clinicaltrials.gov/ct2/show/NCT03332667">https://clinicaltrials.gov/ct2/show/NCT03332667</a>	I	≥1 to <30 yr
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr
<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT038507491">https://clinicaltrials.gov/ct2/show/NCT038507491</a>	I	≥6 mo to ≤30 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr
<b>COG ADVL1921</b>	Phase 1 Study to Evaluate the Safety and Pharmacokinetics of Palbociclib (Ibrance®) in Combination with Irinotecan and Temozolomide in Pediatric Patients with Recurrent or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03709680">https://clinicaltrials.gov/ct2/show/NCT03709680</a>	I	≥2 to <21 yr
<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr

<b>COG ANBL19P1</b>	ANBL19P1, A Pilot Study of Dinutuximab, Sargramostim (GM-CSF), and Isotretinoin in Combination with Irinotecan and Temozolomide in the Post-Consolidation Setting for High-Risk Neuroblastoma	Pilot	<31 yr
<b>MIBG Access</b>	An Open Label, Expanded Access Protocol Using <sup>131</sup> I-Metaiodobenzylguanidine ( <sup>131</sup> I-MIBG) Therapy in Patients with Refractory Neuroblastoma, Pheochromocytoma, or Paraganglioma <a href="https://clinicaltrials.gov/ct2/show/NCT03015844">https://clinicaltrials.gov/ct2/show/NCT03015844</a>	Access to MIBG therapy	≥12 mo
<b>Neuroblastoma biology, supportive treatment and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj
<b>NCICOVID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG APEC14B1</b>	APEC14B1: The Project:Everychild Protocol: A Registry, Eligibility Screening, Biology and Outcome Study <a href="https://clinicaltrials.gov/ct2/show/NCT02402244">https://clinicaltrials.gov/ct2/show/NCT02402244</a>	Non-therapeutic	≤25 yr
<b>COG APEC1621SC</b>	APEC1621SC: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) Screening Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT03155620">https://clinicaltrials.gov/ct2/show/NCT03155620</a>	NA	≥12 mo to ≤21 yr
<b>COG ANBL00B1</b>	Neuroblastoma Biology Studies <a href="https://clinicaltrials.gov/ct2/show/NCT00904241">https://clinicaltrials.gov/ct2/show/NCT00904241</a>	Biology	<31 yr
<b>NANT 2004-05</b>	Neuroblastoma Biology Study (Any patient with high risk neuroblastoma who is not enrolled on a COG frontline therapeutic study is eligible if undergoing a disease evaluation.)	Biology	≥31 days
<b>COG ALTE03N1</b>	Key Adverse Events after Childhood Cancer (cardiac, stroke, secondary malign)(closed to AVN patients as of 11-26-08) <a href="https://clinicaltrials.gov/ct2/show/NCT00082745">https://clinicaltrials.gov/ct2/show/NCT00082745</a>	Non-therapeutic	≤21 yr at dx
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age
<b>COG ALTE15N2</b>	ALTE15N2: LEAHRN (Late Effects After High-Risk Neuroblastoma) Study <a href="https://clinicaltrials.gov/ct2/show/NCT03057626">https://clinicaltrials.gov/ct2/show/NCT03057626</a>	Non-therapeutic	≥5 yr
<b>AflacST17B1</b>	AflacST17B1: Immunophenotyping and Cytokine Profiling of Patients Receiving Therapeutic <sup>131</sup> I-MIBG for Relapsed/Refractory Neuroblastoma	Biology	≥1 to ≤30 yr

<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr
<b>Osteosarcoma treatment protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC#788607, IND#141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT0322003">https://clinicaltrials.gov/ct2/show/NCT0322003</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr
<b>Aflac ST1502 CHOANOME II</b>	AflacST1502: A Phase II Study of Sirolimus in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr
<b>AflacST1903</b>	A Maintenance Protocol of Sirolimus in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr



<b>COG ADVL1721</b>	A Non-randomized, Open-label, Multi-center, Phase I/II Study of PI3K Inhibitor Copanlisib in Pediatric Patients with Relapsed/ Refractory Solid Tumors or Lymphoma <a href="https://clinicaltrials.gov/ct2/show/NCT03458728">https://clinicaltrials.gov/ct2/show/NCT03458728</a>	I/II	≥6 mo to ≤21 yr
<b>Ignya RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr
<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>Colorado 18-2740</b>	A Phase I/Ib Study of Losartan in Combination with Sunitinib in the Treatment of Pediatric and Adult Patients with Relapsed or Refractory Osteosarcoma <a href="https://clinicaltrials.gov/ct2/show/NCT03900793">https://clinicaltrials.gov/ct2/show/NCT03900793</a>	I/Ib	10 to 40 yr
<b>COG ADVL1414</b>	ADVL1414, A Phase 1 Study of Selinexor (KPT-330, IND# 125052), A Selective XPO1 Inhibitor, in Recurrent and Refractory Pediatric Solid Tumors, including CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02323880">https://clinicaltrials.gov/ct2/show/NCT02323880</a>	I	≥12 mo to ≤21 yr
<b>COG ADVL1514</b>	ADVL1514, A Phase 1 Study of ABI-009 (Nab-Rapamycin) (IND# 133537) in Pediatric Patients with Recurrent or Refractory Solid Tumors, Including CNS Tumors as a Single Agent and in Combination with Temozolomide and Irinotecan <a href="https://clinicaltrials.gov/ct2/show/NCT02975882">https://clinicaltrials.gov/ct2/show/NCT02975882</a>	I	≥12 mo to ≤21 yr
<b>COG ADVL1921</b>	Phase 1 Study to Evaluate the Safety and Pharmacokinetics of Palbociclib (Ibrance®) in Combination with Irinotecan and Temozolomide in Pediatric Patients with Recurrent or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03709680">https://clinicaltrials.gov/ct2/show/NCT03709680</a>	I	≥2 to <21 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr
<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT038507491">https://clinicaltrials.gov/ct2/show/NCT038507491</a>	I	≥6 mo to ≤30 yr



<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr
<b>IMPACT</b>	Informational Meetings for Planning and Coordinating Treatment (IMPACT)	III	≥1 to <18 yr
<b>Osteosarcoma biology, supportive treatment, and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj
<b>NCICOVID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG APEC14B1</b>	APEC14B1: The Project:Everychild Protocol: A Registry, Eligibility Screening, Biology and Outcome Study <a href="https://clinicaltrials.gov/ct2/show/NCT02402244">https://clinicaltrials.gov/ct2/show/NCT02402244</a>	Non-therapeutic	≤25 yr
<b>COG APEC1621SC</b>	APEC1621SC: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) Screening Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT03155620">https://clinicaltrials.gov/ct2/show/NCT03155620</a>	NA	≥12 mo to ≤21 yr
<b>COG ALTE03N1</b>	Key Adverse Events after Childhood Cancer (cardiac, stroke, secondary malign)/(closed to AVN patients as of 11-26-08) <a href="https://clinicaltrials.gov/ct2/show/NCT00082745">https://clinicaltrials.gov/ct2/show/NCT00082745</a>	Non-therapeutic	≤21 yr at dx
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age
<b>DFHCC ctDNA</b>	Evaluation of ctDNA as a Prognostic Biomarker for Patients with Newly Diagnosed Localized Ewing Sarcoma or Osteosarcoma	Non-therapeutic	≥12 mo
<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr
<b>Retinoblastoma treatment protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC# 788607, IND# 141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr

<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT0322003">https://clinicaltrials.gov/ct2/show/NCT0322003</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr
<b>Aflac ST1502 CHOANOME II</b>	AflacST1502: A Phase II Study of Sirolimus in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr
<b>AflacST1903</b>	A Maintenance Protocol of Sirolimus in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr
<b>COG ADVL1721</b>	A Non-randomized, Open-label, Multi-center, Phase I/II Study of PI3K Inhibitor Copanlisib in Pediatric Patients with Relapsed/ Refractory Solid Tumors or Lymphoma <a href="https://clinicaltrials.gov/ct2/show/NCT03458728">https://clinicaltrials.gov/ct2/show/NCT03458728</a>	I/II	≥6 mo to ≤21 yr
<b>Ignitya RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr

<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>COG ADVL1921</b>	Phase 1 Study to Evaluate the Safety and Pharmacokinetics of Palbociclib (Ibrance®) in Combination with Irinotecan and Temozolomide in Pediatric Patients with Recurrent or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03709680">https://clinicaltrials.gov/ct2/show/NCT03709680</a>	I	≥2 to <21 yr
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr
<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT038507491">https://clinicaltrials.gov/ct2/show/NCT038507491</a>	I	≥6 mo to ≤30 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr
<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr
<b>Retinoblastoma biology, supportive treatment and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj
<b>NCICOVID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG APEC14B1</b>	APEC14B1: The Project:Everychild Protocol: A Registry, Eligibility Screening, Biology and Outcome Study <a href="https://clinicaltrials.gov/ct2/show/NCT02402244">https://clinicaltrials.gov/ct2/show/NCT02402244</a>	Non-therapeutic	≤25 yr
<b>COG APEC1621SC</b>	APEC1621SC: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) Screening Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT03155620">https://clinicaltrials.gov/ct2/show/NCT03155620</a>	NA	≥12 mo to ≤21 yr
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age
<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr

Rhabdo and non-rhabdo soft tissue sarcoma treatment protocols			
Study	Clinical trial name	Phase/type	Age
<b>COG ARST1431</b>	A Randomized Phase 3 Study of Vincristine, Dactinomycin, Cyclophosphamide (VAC) Alternating with Vincristine and Irinotecan (VI) Versus VAC/VI Plus Temsirolimus (TORI, Torisel, NSC# 683864, IND# 122782) in Patients with Intermediate Risk (IR) Rhabdomyosarcoma (RMS) <a href="https://clinicaltrials.gov/ct2/show/NCT02567435">https://clinicaltrials.gov/ct2/show/NCT02567435</a>	III	Feasibility Phase <21 yr  Efficacy Phase ≤40 yr
<b>Aflac ST1502 CHOANOME II</b>	AflacST1502: A Phase II Study of Sirolimus in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr
<b>AflacST1903</b>	A Maintenance Protocol of Sirolimus in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC# 788607, IND# 141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT0322003">https://clinicaltrials.gov/ct2/show/NCT0322003</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr

<b>COG ADVL1721</b>	A Non-randomized, Open-label, Multi-center, Phase I/II Study of PI3K Inhibitor Copanlisib in Pediatric Patients with Relapsed/ Refractory Solid Tumors or Lymphoma <a href="https://clinicaltrials.gov/ct2/show/NCT03458728">https://clinicaltrials.gov/ct2/show/NCT03458728</a>	I/II	≥6 mo to ≤21 yr
<b>Ignyta RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr
<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>COG ADVL1921</b>	Phase 1 Study to Evaluate the Safety and Pharmacokinetics of Palbociclib (Ibrance®) in Combination with Irinotecan and Temozolomide in Pediatric Patients with Recurrent or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03709680">https://clinicaltrials.gov/ct2/show/NCT03709680</a>	I	≥2 to <21 yr
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr
<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT038507491">https://clinicaltrials.gov/ct2/show/NCT038507491</a>	I	≥6 mo to ≤30 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr
<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr
<b>IMPACT</b>	Informational Meetings for Planning and Coordinating Treatment (IMPACT)	III	≥1 to <18 yr
<b>Rhabdo and non-rhabdo soft tissue sarcoma biology, supportive treatment and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj

<b>NCICOID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG APEC14B1</b>	APEC14B1: The Project:Everychild Protocol: A Registry, Eligibility Screening, Biology and Outcome Study <a href="https://clinicaltrials.gov/ct2/show/NCT02402244">https://clinicaltrials.gov/ct2/show/NCT02402244</a>	Non-therapeutic	≤25 yr
<b>COG APEC1621SC</b>	APEC1621SC: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) Screening Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT03155620">https://clinicaltrials.gov/ct2/show/NCT03155620</a>	NA	≥12 mo to ≤21 yr
<b>COG ALTE03N1</b>	Key Adverse Events after Childhood Cancer (cardiac, stroke, secondary malign)/closed to AVN patients as of 11-26-08 <a href="https://clinicaltrials.gov/ct2/show/NCT00082745">https://clinicaltrials.gov/ct2/show/NCT00082745</a>	Non-therapeutic	≤21 yr at dx
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age
<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr
<b>Wilms treatment protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>Aflac ST1502 CHOANOME II</b>	AflacST1502: A Phase II Study of Sunitinib in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr
<b>AflacST1903</b>	A Maintenance Protocol of Sunitinib in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC# 788607, IND# 141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr

<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT0322003">https://clinicaltrials.gov/ct2/show/NCT0322003</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr
<b>COG AREN1721</b>	AREN1721, A Randomized Phase 2 Trial of Axitinib/Nivolumab Combination Therapy vs. Single Agent Axitinib or Nivolumab for the Treatment of TFR/Translocation Renal Cell Carcinoma (tRCC) Across All Age Groups <a href="https://clinicaltrials.gov/ct2/show/NCT03595124">https://clinicaltrials.gov/ct2/show/NCT03595124</a>	II	≥12 mo
<b>COG AREN1921<sup>REQ</sup></b>	AREN1921, Treatment of Newly Diagnosed Diffuse Anaplastic Wilms Tumors (DAWT) and Relapsed Favorable Histology Wilms Tumors (FHWT) <a href="https://clinicaltrials.gov/ct2/show/NCT04322318">https://clinicaltrials.gov/ct2/show/NCT04322318</a>	II	≤30 yr
<b>COG ADVL1721</b>	A Non-randomized, Open-label, Multi-center, Phase I/II Study of PI3K Inhibitor Copanlisib in Pediatric Patients with Relapsed/Refractory Solid Tumors or Lymphoma <a href="https://clinicaltrials.gov/ct2/show/NCT03458728">https://clinicaltrials.gov/ct2/show/NCT03458728</a>	I/II	≥6 mo to ≤21 yr
<b>Ignya RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr
<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>COG ADVL1414</b>	ADVL1414, A Phase 1 Study of Selinexor (KPT-330, IND# 125052), A Selective XPO1 Inhibitor, in Recurrent and Refractory Pediatric Solid Tumors, including CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02323880">https://clinicaltrials.gov/ct2/show/NCT02323880</a>	I	≥12 mo to ≤21 yr
<b>COG ADVL1514</b>	ADVL1514, A Phase 1 Study of ABI-009 (Nab-Rapamycin) (IND# 133537) in Pediatric Patients with Recurrent or Refractory Solid Tumors, Including CNS Tumors as a Single Agent and in Combination with Temozolomide and Irinotecan <a href="https://clinicaltrials.gov/ct2/show/NCT02975882">https://clinicaltrials.gov/ct2/show/NCT02975882</a>	I	≥12 mo to ≤21 yr



<b>COG ADVL1921</b>	Phase 1 Study to Evaluate the Safety and Pharmacokinetics of Palbociclib (Ibrance®) in Combination with Irinotecan and Temozolomide in Pediatric Patients with Recurrent or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03709680">https://clinicaltrials.gov/ct2/show/NCT03709680</a>	I	≥2 to <21 yr
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr
<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT038507491">https://clinicaltrials.gov/ct2/show/NCT038507491</a>	I	≥6 mo to ≤30 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr
<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr
<b>Wilms biology, supportive treatment and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj
<b>NCICOVID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG APEC14B1</b>	APEC14B1: The Project:Everychild Protocol: A Registry, Eligibility Screening, Biology and Outcome Study <a href="https://clinicaltrials.gov/ct2/show/NCT02402244">https://clinicaltrials.gov/ct2/show/NCT02402244</a>	Non-therapeutic	≤25 yr
<b>COG APEC1621SC</b>	APEC1621SC: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) Screening Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT03155620">https://clinicaltrials.gov/ct2/show/NCT03155620</a>	NA	≥12 mo to ≤21 yr
<b>COG AREN03B2<sup>REQ</sup></b>	Renal Tumors Classification, Biology, and Banking Study <a href="https://clinicaltrials.gov/ct2/show/NCT00898365">https://clinicaltrials.gov/ct2/show/NCT00898365</a>	Biology	<30 yr
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age

<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr
<b>Other solid tumor and rare tumor treatment protocols</b>			
Study	Clinical trial name	Phase/type	Age
<b>COG AGCT1531</b>	AGCT1531: A Phase 3 Study of Active Surveillance for Low Risk and a Randomized Trial of Carboplatin vs. Cisplatin for Standard Risk Pediatric and Adult Patients with Germ Cell Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03067181">https://clinicaltrials.gov/ct2/show/NCT03067181</a>	III	<u>Low Risk</u> : <50 yr <u>Std</u> <u>Risk 1</u> : <11 yr <u>Std</u> <u>Risk 2</u> : ≥11 to <25 yr
<b>COG AGCT1532</b>	Phase 3 Accelerated BEP Trial: A Randomised Phase 3 Trial of Accelerated versus Standard BEP Chemotherapy for Patients with Intermediate and Poor-risk Metastatic Germ Cell Tumors	III	≥11 to ≤45 yr
<b>Alliance A031102</b>	A Randomized Phase III Trial Comparing Conventional-dose Chemotherapy using Paclitaxel, Ifosfamide, and Cisplatin (TIP) with High-dose Chemotherapy using Mobilizing Paclitaxel plus Ifosfamide followed by High-dose Carboplatin and Etoposide (TI-CE) as First Salvage Treatment in Relapsed or Refractory Germ Cell Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02375204">https://clinicaltrials.gov/ct2/show/NCT02375204</a>	III	≥14 yr males only
<b>Aflac ST1502 CHOANOME II</b>	AflacST1502: A Phase II Study of Sirolimus in Combination with Metronomic Chemotherapy in Children with Recurrent and/or Refractory Solid and CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02574728">https://clinicaltrials.gov/ct2/show/NCT02574728</a>	II	>12 mo to ≤30 yr
<b>AflacST1903</b>	A Maintenance Protocol of Sirolimus in Combination with Metronomic Chemotherapy in Children with High-Risk Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04469530">https://clinicaltrials.gov/ct2/show/NCT04469530</a>	II	≥12 mo to ≤30 yr
<b>COG ADVL1823</b>	ADVL1823, Larotrectinib (LOXO-101, NSC# 788607, IND# 141824) for Previously Untreated TRK Fusion Pediatric Solid Tumors and TRK Fusion Relapsed Pediatric Acute Leukemias <a href="https://clinicaltrials.gov/ct2/show/NCT03834961">https://clinicaltrials.gov/ct2/show/NCT03834961</a>	II	≤30 yr
<b>COG APEC1621A<sup>REQ</sup></b>	APEC1621A: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of LOXO-101 (Larotrectinib) in Patients with Tumors Harboring Actionable NTRK Fusions <a href="https://clinicaltrials.gov/ct2/show/NCT03213704">https://clinicaltrials.gov/ct2/show/NCT03213704</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621B<sup>REQ</sup></b>	APEC1621B: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) - Phase 2 Subprotocol of JNJ-42756493 (Erdafitinib) in Patients with Relapsed Pediatric Solid Tumors Harboring FGFR1/2/3/4 Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03210714">https://clinicaltrials.gov/ct2/show/NCT03210714</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621D<sup>REQ</sup></b>	APEC1621D NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of LY3023414 in Patients with Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03213678">https://clinicaltrials.gov/ct2/show/NCT03213678</a>	II	≥12 mo to ≤21 yr

<b>COG APEC1621F<sup>REQ</sup></b>	APEC1621F NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Ensartinib in Patients with Tumors Harboring ALK or ROS1 Genomic Alterations <a href="https://clinicaltrials.gov/ct2/show/NCT03213652">https://clinicaltrials.gov/ct2/show/NCT03213652</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621G<sup>REQ</sup></b>	APEC1621G NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) – Phase 2 Subprotocol of Vemurafenib in Patients with Tumors Harboring Actionable BRAF V600 Mutations <a href="https://clinicaltrials.gov/ct2/show/NCT0322003">https://clinicaltrials.gov/ct2/show/NCT0322003</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621H<sup>REQ</sup></b>	APEC1621H NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Olaparib in Patients with Tumors Harboring Defects in DNA Damage Repair Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03233204">https://clinicaltrials.gov/ct2/show/NCT03233204</a>	II	≥12 mo to ≤21 yr
<b>COG APEC1621I<sup>REQ</sup></b>	APEC1621I NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice)—Phase 2 Subprotocol of Palbociclib in Patients with Tumors Harboring Activating Alterations in Cell Cycle Genes <a href="https://clinicaltrials.gov/ct2/show/NCT03526250">https://clinicaltrials.gov/ct2/show/NCT03526250</a>	II	≥12 mo to ≤21 yr
<b>COG ARST1921</b>	ARST1921, A Safety, Pharmacokinetic and Efficacy Study of a $\gamma$ -Secretase Inhibitor, Nirogacestat (PF-03084014; IND# 146375), in Children and Adolescents with Progressive, Surgically Unresectable Desmoid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT04195399">https://clinicaltrials.gov/ct2/show/NCT04195399</a>	II	>12 mo to <18 yr
<b>LCH-IV (NACHO)</b>	LCH-IV International Collaborative Treatment Protocol for Children and Adolescents with Langerhans Cell Histiocytosis <a href="https://clinicaltrials.gov/ct2/show/NCT02205762">https://clinicaltrials.gov/ct2/show/NCT02205762</a>	II	≤18 yr
<b>Ignitya RXDX-101-03</b>	A Phase I/Ib, Open-Label, Dose-Escalation and Expansion Study of Entrectinib (RXDX-101) in Children and Adolescents with Recurrent or Refractory Solid Tumors and Primary CNS Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02650401">https://clinicaltrials.gov/ct2/show/NCT02650401</a>	I/Ib	≥2 to <22 yr
<b>LOXO-EXT-17005</b>	LOXO-EXT-17005: A Phase 1/2 Study of the TRK Inhibitor LOXO-195 in Adult and Pediatric Subjects with Previously Treated NTRK Fusion Cancers <a href="https://clinicaltrials.gov/ct2/show/NCT03215511">https://clinicaltrials.gov/ct2/show/NCT03215511</a>	I/II	≥1 mo
<b>AflacST1501</b>	Aflac ST1501, A Phase I Study of Abemaciclib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma and in Children with Recurrent and Refractory Solid Tumors Including Malignant Brain Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT02644460">https://clinicaltrials.gov/ct2/show/NCT02644460</a>	I	≥2 to ≤25 yr

<b>AflacST1603 GemAbrax</b>	AflacST1603: A Phase 1 Study Using Nab-paclitaxel (Abraxane®) in Combination with Gemcitabine for Pediatric Relapsed and Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT038507491">https://clinicaltrials.gov/ct2/show/NCT038507491</a>	I	≥6 mo to ≤30 yr
<b>EZH-102</b>	A Phase I Study of the EZH2 Inhibitor Tazemetostat in Pediatric Subjects with Relapsed or Refractory INI1-Negative Tumors or Synovial Sarcoma <a href="https://clinicaltrials.gov/ct2/show/NCT02601937">https://clinicaltrials.gov/ct2/show/NCT02601937</a>	I	≥6 mo to ≤21 yr
<b>AbbVie M13-833</b>	A Phase 1 Study of the Safety and Pharmacokinetics of Venetoclax in Pediatric and Young Adult Patients with Relapsed or Refractory Malignancies <a href="https://clinicaltrials.gov/ct2/show/NCT03236857">https://clinicaltrials.gov/ct2/show/NCT03236857</a>	I	<25 yr
<b>UCSD Cabozantinib cisRA</b>	A National Phase I Study of Cabozantinib in Combination with 13-cis-Retinoic Acid in Children with Relapsed or Refractory Solid Tumors <a href="https://clinicaltrials.gov/ct2/show/NCT03611595">https://clinicaltrials.gov/ct2/show/NCT03611595</a>	I	≥2 to ≤26 yr
<b>IMPACT</b>	Informational Meetings for Planning and Coordinating Treatment (IMPACT)	III	≥1 to <18 yr
<b>Other solid tumor and rare tumor biology, supportive treatment and non-therapeutic protocols</b>			
<b>Study</b>	<b>Clinical trial name</b>	<b>Phase/type</b>	<b>Age</b>
<b>AflacPM1702</b>	Aflac Precision Medicine Program (APMP) Study	Non-therapeutic	<30 yr
<b>CCPS</b>	Childhood Cancer Predisposition Study (CCPS) <a href="https://clinicaltrials.gov/ct2/show/NCT04511806">https://clinicaltrials.gov/ct2/show/NCT04511806</a>	Non-therapeutic	<21 yr primary subj
<b>NCICOVID</b>	NCI COVID-19 in Cancer Patients Study (NCCAPS): A Longitudinal Natural History Study <a href="https://clinicaltrials.gov/ct2/show/NCT04387656">https://clinicaltrials.gov/ct2/show/NCT04387656</a>	Non-therapeutic	Any age
<b>COG APEC14B1</b>	APEC14B1: The Project: Everychild Protocol: A Registry, Eligibility Screening, Biology and Outcome Study <a href="https://clinicaltrials.gov/ct2/show/NCT02402244">https://clinicaltrials.gov/ct2/show/NCT02402244</a>	Non-therapeutic	≤25 yr
<b>COG APEC1621SC</b>	APEC1621SC: NCI-COG Pediatric MATCH (Molecular Analysis for Therapy Choice) Screening Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT03155620">https://clinicaltrials.gov/ct2/show/NCT03155620</a>	NA	≥12 mo to ≤21 yr
<b>PPB DICER1</b>	International Pleuropulmonary Blastoma Registry for PPB, <i>DICER1</i> and Associated Conditions <a href="https://clinicaltrials.gov/ct2/show/NCT03382158">https://clinicaltrials.gov/ct2/show/NCT03382158</a>	Registry	Any age

<b>COG ALTE03N1</b>	Key Adverse Events after Childhood Cancer (cardiac, stroke, secondary malign)/closed to AVN patients as of 11-26-08 <a href="https://clinicaltrials.gov/ct2/show/NCT00082745">https://clinicaltrials.gov/ct2/show/NCT00082745</a>	Non-therapeutic	≤21 yr at dx
<b>COG ALTE05N1</b>	Umbrella Long-Term Follow-Up Protocol <a href="https://clinicaltrials.gov/ct2/show/NCT00736749">https://clinicaltrials.gov/ct2/show/NCT00736749</a>	Non-therapeutic	Any age
<b>QOL Thyroid</b>	The Quality of Life in Children and Adolescents with Thyroid Cancer	Non-therapeutic	2 to 21 yr
<b>NACHO BIO</b>	NACHO-BIO: A Translational Biology Platform to Advance Understanding of Pathogenesis and Improve Outcomes for Patients with Histiocytic Disorders	Non-therapeutic	Any age
<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr
<b>NACHO BIO</b>	NACHO-BIO: A Translational Biology Platform to Advance Understanding of Pathogenesis and Improve Outcomes for Patients with Histiocytic Disorders	Non-therapeutic	Any age
<b>Telehealth Home-based Transition</b>	Development and Pilot of a Telehealth Home-based Transition Intervention for Pediatric and Young Adults with Cancer	Non-therapeutic	≤29 yr