| Department                   | Project<br>Number | Project Title   | Principal Investigator | Source of Funding         | 23-24<br>Receipts |
|------------------------------|-------------------|---|------------------------|---------------------------|-------------------|
| Internal Medicine            |                   |   |                        |                           |                   |
|                              | 63891             | CorEvitas Inflammatory Bowel Disease (IBD) Drug Safety and Effectiveness Registry<br>(Corrona IBD-600 Registry)   | Dr Sonnier, W          | CorEvitas                 | 39,930.00         |
|                              | 63939             | Protocol No. 19767: A multicenter, international, randomized, active comparator-<br>controlled, double-blind, double dummy, parallel-group, 2-arm, Phase 3 study to<br>compare the efficacy and safety of the oral FXIa inhibitor asundexian (BAY 2433334)<br>with apixaban for the prevention of stroke or systemic embolism in male and<br>female participants aged 18 years and older with atrial fibrillation at risk for stroke<br>(BAY 19767) | Dr Malozzi, C          | Bayer Us LLC              | 16,257.00         |
|                              | 63949             | NN7533-4470- ASCENT A Multicentre Trial Evaluating the Efficacy and Safety of Oral<br>Decitabine-Tetrahydrouridine (NDec) in Patients with Sickle Cell Disease (NN7533-<br>4470 - ASCENT)   | Dr Hogue, A            | Novo Nordisk              | 2,800.00          |
|                              |                   |   |                        | Total Internal Medicine   | 58,987.00         |
| Mitchell Cancer<br>Institute |                   |   |                        |                           |                   |
|                              | 63799             | PALLAS: PALbociclib CoLlaborative Adjuvant Study: A randomized phase III trial of<br>Palbociclib with standard adjuvant endocrine therapy versus standard adjuvant<br>endocrine therapy alone for hormone receptor positive (HR+) / human epidermal<br>growth factor receptor 2 (HER2)-negative early breast cancer   | Dr Prodduturbar, P     | Alliance Foundation       | 2,100.00          |
|                              | 63937             | A Randomized, Double-blind, Double-dummy, Parallel Group Study to Assess<br>theEfficacy and Safety of Palonosetron HCl Buccal Film versus IV Palonosetron 0.25<br>mg for thePrevention of Chemotherapy-induced Nausea and Vomiting in Cancer<br>Patients ReceivingModerately Emetogenic Chemotherapy (LP-CT-PALO 202101)  | Dr Butler, T           | Xiamen LP Pharmaceuticals | 9,650.51          |
|                              | 63938             | BO44178- A Phase II, Randomized, Multicenter, Double-Blind, Controlled Study Of<br>RO7247669 Plus Platinum-Based Chemotherapy Versus Pembrolizumab Plus<br>Platinum-Based Chemotherapy In Patients With Previously Untreated Locally<br>Advanced Or Metastatic Non Small Cell Lung Cancer (Roche - BO44178)   | Dr Prodduturbar, P     | Roche - Genentech, Inc.   | 16,200.00         |

| Department | Project<br>Number | Project Title   | Principal Investigator | Source of Funding               | 23-24<br>Receipts |
|------------|-------------------|---|------------------------|---------------------------------|-------------------|
|            | 63946             | A Phase 3 Open-Label, Randomized Study of Fixed Duration Pirtobrutinib (LOXO-<br>305) plus Venetoclax and Rituximab versus Venetoclax and Rituximab in Previously<br>Treated Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma (BRUIN CLL-<br>322) (LOXO-BTX-20022) | Dr Alkharabsheh, O     | Loxo Oncology, Inc.             | 15,210.00         |
|            | 63953             | A Phase 4, Observational Study Evaluating the Efficacy and Safety of the Bruton<br>Tyrosine Kinase (BTK) Inhibitor Zanubrutinib in Patients With Waldenström<br>Macroglobulinemia (BeiGene - BGB-3111-402)  | Dr Alkharabsheh, O     | BeiGene, Ltd.                   | 10,200.00         |
|            | 63955             | A Phase IIIb, Single Arm, Open-label, Multicentre Study of Durvalumab in<br>Combination with Chemotherapy for the First Line Treatment for Patients with<br>Advanced Biliary Tract Cancers (TOURMALINE - D4191C00140)   | Dr Abdalla, A          | AstraZeneca Pharmaceuticals LP  | 34,202.80         |
|            | 63957             | A Randomized, Open-label, Phase 3 Study of Tarlatamab Compared With Standard<br>of Care in Subjects With Relapsed Small Cell Lung Cancer After Platinum-based First-<br>line Chemotherapy (DeLLphi 304) (AMG 757-20210004)  | Dr Prodduturbar, P     | Amgen, Inc.                     | 14,700.00         |
|            | 63960             | A Phase 3, Randomized Study to Compare Nemtabrutinib Versus Comparator<br>(Investigator's Choice of Ibrutinib or Acalabrutinib) in Participants With Untreated<br>Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma (BELLWAVE-011) (MK-<br>1026-011-0014)           | Dr Alkharabsheh, O     | Merck Sharp & Dohme LLC         | 10,500.00         |
|            |                   |   |                        | Total Mitchell Cancer Institute | 112,763.31        |

| Department<br>Neurology | Project<br>Number | Project Title   | Principal Investigator | Source of Funding          | 23-24<br>Receipts |
|-------------------------|-------------------|---|------------------------|----------------------------|-------------------|
|                         | 63867             | A Randomized, Double-blind, Placebo-controlled, Multicenter Study to Evaluate the safety, Tolerability and Efficacy of XEN1101 as Adjunctive Therapy in Focal-onset Epilepsy (Xenon XPF-008-201)                                      | Dr Naritoku, D         | Xenon Pharaceuticals, Inc. | 3,966.06          |
|                         | 63940             | A Randomized, Double-blind, Placebo-Controlled, Multicenter Phase 3 Study to<br>Evaluate the Safety, Tolerability, and Efficacy of XEN1101 as Adjunctive Therapy in<br>Focal-Onset Seizures (Xenon XPF-010-301)                       | Dr Naritoku, D         | Xenon Pharaceuticals, Inc. | 15,400.00         |
| ×                       | 63941             | A Randomized, Double-blind, Placebo-Controlled, Multicenter, Phase 3 Study to<br>Evaluate the Safety, Tolerability, and Efficacy of XEN1101 as Adjunctive Therapy in<br>Primary Generalized Tonic-Clonic Seizures (Xenon XPF-010-303) | Dr Naritoku, D         | Xenon Pharaceuticals, Inc. | 9,450.00          |
|                         |                   |   |                        | Total Neurology            | 28,816.06         |

| Department   | Project<br>Number | Project Title   | Principal Investigator | Source of Funding                     | 23-24<br>Receipts    |
|--------------|-------------------|---|------------------------|---------------------------------------|----------------------|
| Neurosurgery | 63952             | PAULA - Pelvic fixation and fusion during multilevel spinal surgery (SI-BONE -<br>PAULA)  | Dr Menger, R           | SI-BONE, Inc.                         | 26,857.50            |
|              |                   |   |                        | Total Neurosurgery                    | 26,857.50            |
| OB-GYN       | 63887             | A Phase 3, Randomized, Double- or Observer-Blinded, Placebo-Controlled Trial to<br>EvaluateThe Efficacy and Safety of a Respiratory Syncytial Virus (RSV) Perfusion<br>FSubunit Vacine in Infants Born to Women Vaccinated During Pregnancy<br>Investigational (Pfizer C3671008)                  | Dr Roth, T             | Pfizer<br>Total OB-GYN                | 2,329.27<br>2,329.27 |
| Pediatrics   | 63869             | Bayer Healthcare Pharmaceuticals, Inc. / "HEM-POWR: Observational Study<br>Evaluating Effectiveness and Safety of Real-World Treatment with Damoctocog alfa   | Dr Marri, P            | Bayer                                 | 2,860.00             |
|              | 03005             | pegol in Previously Treated Patients with Hemophilia A" (Bayer HEM-POWR)  | or worn, r             | Jayer                                 | 2,000.00             |
|              | 63913             | An Open-Label Extension Study to Evaluate the Long-Term Safety of Inclacumab<br>Administered to Participants with Sickle Cell Disease Who Have Participated in an<br>Inclacumab Clinical Trial (GBT2104-133)  | Dr Marri, P            | Global Blood Therapeutics, Inc.       | 8,560.60             |
|              | 63926             | A Phase 2, Double-Blind, 12-Week, Multicenter Study to Assess the Safety and<br>Effectiveness of Daily Oral Administration of Dexlansoprazole Delayed-Release<br>Capsules in Pediatric Subjects Aged 2 to 11 Years With Symptomatic Nonerosive<br>Gastroesophageal Reflux Disease (TAK-390MR_204) | Dr Gremse, D           | Takeda Development Ctr Americas, Inc. | 23,728.00            |

| Department  | Project<br>Number | Project Title  | Principal Investigator | Source of Funding                     | 23-24<br>Receipts |
|-------------|-------------------|--|------------------------|---------------------------------------|-------------------|
|             | 63927             | A Phase 2, Double-Blind, 36-Week, Multicenter Study to Assess the Safety and<br>Effectiveness of Daily Oral Administration of Dexlansoprazole Delayed-Release<br>Capsules for Healing of Erosive Esophagitis (EE) and Maintenance of Healed EE in<br>Pediatric Subjects Aged 2 to 11 Years With EE (TAK-390MR_205) | Dr Gremse, D           | Takeda Development Ctr Americas, Inc. | 33,399.00         |
|             | 63931             | Evaluation of a 7-Day Therapeutic Trial Dose of Commercial Sucraid® (sacrosidase)<br>Oral Solution for Alleviating Congenital Sucrase-Isomaltase Deficiency (CSID)<br>Symptoms in Pediatric Subjects with Low, Moderate, and Normal Sucrase Levels<br>(QOL - SSDXP-13)   | Dr Gremse, D           | QOL Medical, LLC                      | 23,702.60         |
|             | 63932             | A Randomized, Assessor-Blind, Parallel-Groups, Multicenter Trial Assessing the<br>Safety and Efficacy, Including Pharmacokinetic Assessments, of CLENPIQ in Children<br>Aged 2 Years to Less Than 9 Years (FIPCUS CLENPIQ 000359)  | Dr Gremse, D           | Ferring Pharmaceuticals               | 46,127.00         |
|             | 63951             | A Phase 1, Randomized, Parallel-group, Open-label, Multicenter Study to Evaluate<br>the Pharmacokinetics, Pharmacodynamics and Safety of Vonoprazan (10 or 20 mg<br>Once Daily) in Children Aged >= 6 to < 12 Years Who Have Symptomatic<br>Gastroesophageal Reflux Disease (VPED-103)                             | Dr Gremse, D           | Phathom Pharmaceuticals, Inc.         | 14,200.00         |
|             |                   |  |                        | Total Pediatrics                      | 152,577.20        |
| Sickle Cell |                   |  |                        |                                       |                   |
|             | 63945             | GBT 440-4R2 An Open Label, Observational, Prospective Registry of Participants<br>With Sickle Cell Disease (SCD) Treated With Oxbryta® (Voxelotor) (GBT-440-4R2)   | Dr Pack-Mabien, A      | Global Blood Therapeutics, Inc.       | 54,659.70         |
|             |                   |  |                        | Total Sickle Cell                     | 54,659.70         |

| e Principal Investigator   | Source of Funding   | 23-24<br>Receipts  |
|--|---|--|
| artial and Full Thickness Burn<br>Dr Bright, A                             | Mediwound   | 48,703.26  |
| ole of LINX for Reflux Disease (Torax Dr Richards, W                       | Torax Medical/Ethicon Endo-Surgery  | 9,355.00   |
| tor Prothrombin Complex Dr Butts, C<br>Survival in Patients with Traumatic | CSL Behring LLC   | 178,974.87   |
|  | Stimdia Medical, Inc.   | 13,460.00  |
|  | Total Surgery   | 250,493.13   |
|  |   |  |
| ency Urinary Incontinence: Post Dr Blanchard-Burch, K                      | Valencia Technologies   | 10,585.00  |
|  | Total Urology   | 10,585.00  |
|  | Total All Departments   | 698,068.17   |
|  | Partial and Full Thickness Burn Dr Bright, A<br>ole of LINX for Reflux Disease (Torax Dr Richards, W<br>le-Blind, Placebo-Controlled, Large<br>ctor Prothrombin Complex Dr Butts, C<br>Survival in Patients with Traumatic<br>s)<br>renic nerve to diaphragm<br>/entilated Patients (ReInvigorate Dr Mbaka, M | Partial and Full Thickness Burn Dr Bright, A Mediwound<br>ole of LINX for Reflux Disease (Torax Dr Richards, W Torax Medical/Ethicon Endo-Surgery<br>le Blind, Placebo-Controlled, Large<br>ctor Prothrombin Complex Dr Butts, C CSL Behring LLC<br>)<br>renic nerve to diaphragm<br>renic nerve to diaphragm<br>renic nerve to diaphragm<br>rent |

| Department | Project<br>Number | Project Title                         | Principal Investigator | Source of Funding        | 23-24<br>Receipts |
|------------|-------------------|---------------------------------------|------------------------|--------------------------|-------------------|
|            | 62648             | Royalty On Sales of Antibody Products | Dr Scammell, J         | Genovis / QED Bioscience | 2,261.00          |
|            |                   |                                       |                        | Total Other Research     | 2,261.00          |
|            |                   |                                       |                        | Total                    | 700,329.17        |