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| Company/(Companies): | Period of Support | Name of Project | 1) A research grant or contract from this Company generates revenue for the University of Alberta | 2) A research grant or contract from this Company partially supports my research projects | 3) Educational activities or lectures for this Company generates revenue for the University of Alberta | | 4) Consulting or other services for this company generates personal income | | 5) I receive significant personal royalties from this Company | 6) I have equity in this Company | Comments |
|------------------------------------|-------------------|--|---|---|--|--------------------------|--|-------------------------------------|---|-------------------------------------|----------|
| | | | | | <\$10K | >\$10K | <\$10K | >\$10K | | | |
| Merck | 2016 to present | VeriCiguaT gIObal study in patients with heart failure and Reduced ejection frAction (VICTORIA) | <input checked="" type="checkbox"/> | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | |
| Boehringer Ingelheim | 2017 to present | STrategic Reperfusion in elderly patients Early After Myocardial Infaction (STREAM-II) | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | |
| Bayer | 2018 – 2020 | A Randomized Parellel-Group, Placebo Controlled, Double-Blind, Multi-Center Confirmatory Trial to Evaluate the Efficacy and Safety of the Oral sGC Stimulator on Physical Limitations in Patients with Heart Failure and Preserved Ejection Fraction (VITALITY) | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input checked="" type="checkbox"/> | |
| CSL Limited | 2018 – 2022 | A Phase 3, Multicenter, Double-blind, Randomized, Placebo-controlled, Parallel-group Study to Investigate the Efficacy and Safety of CSL112 in Subjects with Acute Coronary Syndrome (AEGIS II) | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | |
| Boehringer Ingelheim and Eli Lilly | 2020 - 2023 | A phase III, streamlined, multicentre, randomised, parallel group, double-blind, placebo-controlled, superiority trial to evaluate the effect of Empagliflozin on hospitalisation for heart failure and mortality in patients with acute myocardial infarction. (EMPACT-M) | <input checked="" type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> | |

Monetary amounts are on an annual basis.

** Please note this document should be placed in the context that Dr. Armstrong is working 0.5 FTE at the University.

In accordance with the American Heart Association conflict of interest policies.