Policy on Handling of Human Biological Samples

The Company's Commitment

The mission of Sight Sciences Inc. (the "Company") is to transform ophthalmology and optometry through the development and commercialization of proprietary devices that target the underlying causes of the world's most prevalent eye diseases. We are passionate about improving patients' lives. The Company's objective is to develop and market products for use in new treatment paradigms and to create an interventional mindset in eyecare whereby our products may be used in procedures which supplant conventional outdated approaches.

The Company operates under, and continuously strives to comply with, an array of global laws and regulations governing our business operations and practices, including the ethical and responsible handling of human biological samples. The purpose of this Policy on Handling of Human Biological Samples ("Policy") is to set forth the Company's requirements for the proper acquisition, use, storage, sharing, and disposal of human biological samples (the "Samples") by the Company. This Policy also outlines the Company's expectations of third-party researchers and suppliers ("Suppliers") who provide human biological samples to the Company to help ensure such Samples are acquired, used, stored, shared, and disposed of in an ethical and compliant manner.

Company Initiatives

The Company is committed to the ethical and responsible handling of human biological samples and expects all Suppliers to implement policies and procedures that address the responsible and ethical collection, use, handling, sharing, and disposal of Samples. The Company has a Standard Operating Procedure ("SOP") which addresses the process for handling, storing, and disposing of Samples to comply with applicable laws. The Company expects Suppliers who provide human biological samples to the Company to have appropriate ethical policies and procedures in place. Such policies and procedures should include the following:

Informed Consent: Suppliers should obtain the donor's (or their authorized representative) informed consent, as required by the U.S. Food and Drug Administration (or similar governing agency of the country of where the donor resides). The informed consent should include detailed information about the purpose of the research, the type of samples to be collected, the purpose for collecting the samples, how samples will be used, potential risks and benefits, and inform the donors of their right to withdraw their consent and how to withdraw their consent.

Privacy and Confidentiality: The Company expects Suppliers to protect and properly maintain donors' personal and Samples information in accordance with all applicable laws. All personal

information and Samples information should be anonymized or de-identified and should not be shared with Company.

Sample Handling and Storage: Suppliers should collect, label, and store Samples in accordance with relevant regulations, ethical guidelines, and standardized protocols to maintain the Samples' integrity. Suppliers should store Samples in facilities that comply with regulatory requirements for safety and security.

Use and Disposal of Samples: Samples should only be used, stored, and disposed of in accordance with the informed consent provided by the donor or their authorized representative. Suppliers must adhere to all relevant regulations, ethical guidelines, and standardized protocols governing the use of Samples. All Samples should be disposed of following established SOPs, protocols, and regulations to ensure environmental and personal safety.

Effective Date

This Policy is effective as of April 2, 2025