



CHAPTER 90
ARTICLE 23A - RIGHT TO TRY ACT

N.C.G.S. § 90-325.31.

Authorized access to and use of
individualized investigational drugs,
biological products, or devices.

SOURCE General Statutes of North Carolina, as published by the General Assembly	LAST AMENDMENT IN THIS TEXT No amendment history on record	RETRIEVED 21 September 2026, 05:52 UTC	SOURCE FINGERPRINT a3c908983ff9e3d46f8a47ee - c86335615e906adacbcef312 - f14a11c3b752bd3a
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§ 90-325.31(a) A manufacturer operating within an eligible facility and in accordance with all applicable federal law may make available to an eligible patient, and an eligible patient may request, the manufacturer's individualized investigational drug, biological product, or device from an eligible facility or manufacturer operating within an eligible facility. However, nothing in this Part shall be construed to require a manufacturer of an individualized investigational drug, biological product, or device to make such individualized investigational drug, biological product, or device available to an eligible patient.

§ 90-325.31(b) A manufacturer of an individualized investigational drug, biological product, or device may provide the individualized investigational drug, biological product, or device to an eligible patient without receiving compensation or may require the eligible patient to pay the costs of, or the costs associated with, the manufacture of the individualized investigational drug, biological product, or device. (2024-36, s. 1.)

APPARATUS III

Citation

FULL SECTION

N.C. Gen. Stat. § 90-325.31 (2026).

PINPOINT

N.C. Gen. Stat. § 90-325.31(a) (2026).

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