The Joint Commission has approved the following revisions for prepublication. While revised requirements are published in the semiannual updates to the print manuals (as well as in the online E-dition®), accredited organizations and paid subscribers can also view them in the monthly periodical The Joint Commission Perspectives®. To begin your subscription, call 800-746-6578 or visit http://www.jcrinc.com.

Please note: Where applicable, this report shows current standards and EPs first, with deleted language struck-through. Then, the revised requirement follows in bold text, with new language underlined.

APPLICABLE TO THE LABORATORY ACCREDITATION PROGRAM
Effective July 1, 2022

Environment of Care (EC) Chapter

EC.02.04.01

The laboratory manages laboratory equipment risks.

Element(s) of Performance for EC.02.04.01

5. The laboratory monitors and reports all incidents in which laboratory equipment is suspected in or attributed to the death, serious injury, or serious illness of any individual, as required by the Safe Medical Devices Act of 1990.
   Note: Laboratory equipment includes instruments and machines intended for use in diagnosing disease or other conditions.

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Key: □ indicates that documentation is required; ▲ indicates an identified risk area;