

**STANDARD TEST METHOD AND
PERFORMANCE SPECIFICATION
USED IN EVALUATING THE
PERFORMANCE CHARACTERISTICS OF
PROTECTORS FOR COMMOTIO CORDIS**

NOCSAE DOC (ND) 200-~~21~~ 22

Prepared By



**NATIONAL OPERATING COMMITTEE
ON STANDARDS FOR ATHLETIC EQUIPMENT**

This document contains **modifications in sections 5.2 and 16.4**. Modifications do not require Standards Committee approval, but discussion and review are appropriate. The Standards Committee or other committee appointed for the purpose may determine these modifications to be revisions that do require formal approval.

There is a **single requested revision in section 16.1.7** for consideration and voting by the Standards Committee.

Whether a modification or revision, new language is shown in **BLUE TEXT** and deletions are indicated in **Red Strike-Through**.

**No changes will be effective earlier
than February 2023**

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1. Scope

- 1.1 This standard test method describes laboratory equipment and basic requirements pertinent to projectile testing of protectors using the NOCSAE Thoracic Surrogate. It is believed that compliance with this test method will lead to improved protector performance, thus a reduced risk of commotio cordis. The protector shall be supplied as offered for sale with required hardware and fitting instructions along with any accessories required for the protector to function as designed. The requirements of this standard shall be subject to Equipment Level 2 compliance criteria, unless otherwise stated herein.
- 1.2 **All testing and requirements of this standard specification must be in accordance with NOCSAE DOC (ND) 001, except where modified herein.**
- 1.3 *This standard does not purport to address all of the safety problems, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices plus determine the applicability of regulatory limitations prior to use.*

2. Referenced Documents

- 2.1. NOCSAE DOC (ND) 001: *Standard Test Method and Equipment Used in Evaluating the Performance Characteristics of Headgear/Equipment*
- 2.2. NOCSAE DOC (ND) 021: *Standard Projectile Impact Test Method and Equipment Used in Evaluating the Performance Characteristics of Protective Headgear/Projectiles*

3. Significance and Use

- 3.1. The purpose of this test method is to provide reliable and repeatable measurements for the evaluation of various types of protectors. The pass/fail criteria are based on peak force (lbf, N) and within specified tolerances for other measures.

4. Terminology

- 4.1. NOCSAE Thoracic Surrogate (NTS): A device consisting of a damped loading surface, a load cell capable of measuring peak force, and a rigid back plate. The load cell is positioned in between the loading surface and the back plate and represents the cardiac silhouette.
- 4.2. Load Cell: A transducer that converts a mechanical force input to an electrical output signal.
- 4.3. Cardiac Load: The peak force measured by the load cell intended to represent the cardiac silhouette.
- 4.4. Extent of protective coverage: The minimum size of a chest protector's primary protective component, based on the torso length of the individual the model/size is intended to fit.

- 4.5. Linear Bearing Table: A sliding tabletop fitted with the necessary supports to position the thoracic surrogate at various positions in order to facilitate impacts at the designated locations. This table may be mounted to a separate device that allows for rapid elevation adjustments.
- 4.6. Primary Protective Component: That component of the protector/system that is the primary energy attenuation device, without which the protector or system will not meet the impact requirements.
- 4.7. Torso Length: The vertical distance from the base of the neck to the waistline when measuring on the front of the torso. More specifically, the measured vertical distance from the jugular notch to the navel.

5. Test Samples

- 5.1. See Sections 6 and 11, NOCSAE DOC 001, for QC/QA protocol testing.
- 5.2. For any standalone test report, a minimum of two samples per model in the smallest size offered for sale **in each of the size ranges specified in sections 13.2.1, 13.2.2, and 13.2.3 shall be submitted.** ~~must be submitted. Larger model/sizes may be requested if the submitted sample size does not adequately fit the surrogate.~~
 - 5.2.1. The submitted samples shall include the Torso Length of the individual the submitted model/size is intended to fit.
 - 5.2.2. In cases where the size of the primary protective component remains constant for all model/sizes, a minimum of two samples in the smallest size offered for sale shall be submitted. The samples shall include the longest Torso Length of the individual the model is intended to fit.

6. Conditioning Environments

- 6.1. Ambient Temperature: Expose protector, projectile, and equipment to testing environment for a minimum of four hours.

7. Sample Preparation

- 7.1. See section 10 of NOCSAE DOC 001.

8. Certification

- 8.1. See section 6 of NOCSAE DOC 001 for certification requirements.

9. Summary of Test Method

- 9.1. A protector is positioned on the NOCSAE Thoracic Surrogate, which is rigidly mounted to a linear bearing table to allow post impact kinematics. The projectile is aimed at the center of the cardiac silhouette and propelled at the surrogate such that the impact velocity is within 3% of the specified velocity. At impact, the peak force is measured by the cardiac load cell.

10. Test Instruments and Equipment

- 10.1. Projectile launching device capable of propelling the specified projectile to the specified velocity within 3% and having a targeting device that allows impacts within a $\frac{1}{4}$ inch radius within 24 inches from the muzzle or the point at which the projectile is first in "free flight."
- 10.2. Recommended Air Cannon Assembly Including Linear Bearing Table Assembly (See NOCSAE DOC 021, Figures 1 and 2).
- 10.3. NOCSAE Thoracic Surrogate comprising at least one 750 lbf maximum capacity single axis load cell.* Surrogate performance must be verified using the procedure outlined in Appendix 1.
- 10.4. A linear bearing table that is capable of providing some post impact motion with a weight not to exceed 12.5 lb (5.7 kg).
- 10.5. Data sampling rate shall be a minimum of 10,000 Hz per channel.
- 10.6. Data shall be filtered with a CFC120 low pass filter.
- 10.7. Impact shall be recorded on a device capable of capturing the load cell trace.

11. System Check

- 11.1. See section 18, NOCSAE DOC 001
- 11.2. Using a lacrosse ball or baseball that complies with the requirements of section 13, three impacts shall be performed on the unprotected NOCSAE Thoracic Surrogate at the cardiac load cell location. The impact velocity shall be 30 mph \pm 3% and shall impact the center ($\pm \frac{1}{4}$ inch) of the cardiac silhouette (Figure 1). The peak force measured by the load cell shall be no less than 130 lbf (578 N) and no more than 150 lbf (667 N) for each of the system check impacts. The projectile chosen shall be the same type of projectile designated by the manufacturer to be used during testing.

12. Chest Protector Positioning

- 12.1. Manufacturing fitting instructions shall be used to obtain a reasonable fit on the test surrogate. In the event that these instructions are unclear or result in a fit that is likely to yield erroneous test results, the technician shall fit the chest protector to the best of their ability.
 - 12.1.1. A chest protector manufacturer may further define the fit by supplying a protector positioning index. For example, the manufacturer could specify the location of the protector that is intended to line up with the center of the cardiac silhouette. If so specified, that resulting position shall be used instead of the fitting instructions provided the two do not provide conflicting information. In any case, the resulting fit must be reasonable as determined by the test technician.

* Note: NOCSAE compliant surrogate available from Southern Impact Research Center

12.2. Random Location: The technician shall fit the protector following Section 12.1 and then reposition the protector without altering or adjusting the protector's fitting system. In cases where repositioning the protector without adjusting the fitting system is likely to yield erroneous test results, the technician shall adjust the fitting system as needed to yield an appropriate test.

12.2.1. A chest protector manufacturer may provide guidance on fitting system adjustment for random impact locations. However, it is up to the test technician to decide whether the use of the additional guidance is reasonable.

13. Impact Locations

13.1. Cardiac Silhouette: Projectile shall impact the surrogate in the center of the cardiac silhouette ($\pm \frac{1}{4}$ inch) with the protector positioned according to Section 12.1. The surrogate shall be positioned so the base of the surrogate is perpendicular (± 2.5 degrees) to the line of travel of the projectile.

13.2. Random Location: Projectile shall impact the surrogate at the center of the cardiac silhouette ($\pm \frac{1}{4}$ inch), but the protector will be shifted from the position established in 12.1. The initial point of contact shall be a minimum of 1 inch from the initial point of contact of any other impact. The surrogate shall be positioned so the base of the surrogate is perpendicular (± 2.5 degrees) to the line of travel of the projectile.

Extent of Protective Coverage shall be established with the information provided in Section 5.2.1.

13.2.1. Protectors intended to fit Torso Lengths of 13.5 inches or less may be shifted up to 1.75 inches in any direction.

13.2.2. Protectors intended to fit Torso Lengths of more than 13.5 inches and less than 16.0 inches may be shifted up to 2.0 inches in any direction.

13.2.3. Protectors intended to fit Torso Lengths of 16.0 or greater may be shifted up to 2.25 inches in any direction.

14. Impact Attenuation Tests

14.1. The manufacturer shall designate the projectile, baseball or lacrosse ball, to be used during testing. Testing shall be conducted in accordance with TABLE 1.

14.1.1. Baseball: The baseball (s) used shall weigh 5 – 5.25 ounces (142 – 149 grams), have a circumference of 9 – 9.25 inches, and have a C-D at .25 inches of 200 – 300 lbs. and be of the construction specified and used by Major League Baseball.

14.1.2. Lacrosse Ball: The lacrosse ball(s) used shall weigh 5.0 to 5.25 ounces (141.8–148.8 grams), have a circumference of 7.75–8.0 inches and have a C-D at 25% of 115-150 lbs.

- 14.2. The NTS will be positioned with its impact site located within 24 inches (610 mm) from the end of the muzzle (or from the point at which the ball is released).
- 14.3. Protector repositioning during testing is anticipated.

TABLE 1
Impact Locations
Velocity – mph (m/s) \pm 3%

	Center of Cardiac Silhouette	Random Location #1	Random Location #2
Sample 1	30 (13.4)	30 (13.4)	30 (13.4)
Sample 2	50 (22.4)	50 (22.4)	50 (22.4)

15. Test Requirements

15.1. 30 mile per hour condition

- 15.1.1. For any impact, the peak force measured by the cardiac load cell shall not exceed 90 lbf (400N).

15.2. 50 mile per hour condition

- 15.2.1. For any impact, the peak force measured by the cardiac load cell shall not exceed 180 lbf (800N).

16. Labels and Warnings

- 16.1. Each protector shall be permanently and legibly labeled or marked in a manner such that the following information can be easily read and must not be obscured in any manner.

- 16.1.1. Name of manufacturer or a logo or recognizable mark that is registered and/or trademarked to the manufacturer.

- 16.1.2. Model designation

- 16.1.3. Size

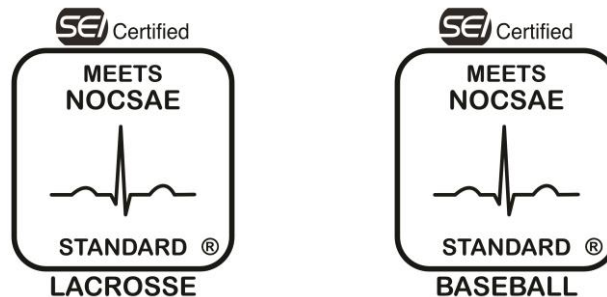
- 16.1.4. Month and year of manufacture

- 16.1.5. Notice that indicates additional information is available to the wearer. Such additional information may be on the packaging, insert, or removable tag.

- 16.1.6. A permanent, exact replica of the appropriate sport specific seal must appear legibly on the exterior of the protector's primary protective component.

- 16.1.7. **If the primary protective component is part of an assembly (such as inserted into a shirt like device) the supporting component shall be marked with a statement that communicates to the wearer that the**

product is not compliant with the stated performance standard unless assembled in accordance with the manufacturer's instructions.



NOTE: You must have an executed, valid license agreement with NOCSAE to use any of the NOCSAE logos at any time. NOCSAE, the NOCSAE seals/logos, and the National Operating Committee on Standards for Athletic Equipment are registered marks and the exclusive property of the Committee. Use of the marks in any manner is prohibited without prior written permission of the NOCSAE Board of Directors.

16.2. Packaging and/or instructional literature for protectors shall be permanently and legibly marked in a manner such that the following information can be easily read.

16.2.1. Information that warns the user that the protector can be damaged by contact with common substances (for example, certain solvents, cleaners, etc.) and that this damage may or may not be visible to the user. This information should also list any recommended cleaning agents and/or procedures.

16.2.2. Information that warns the user that for maximum performance the manufacturers fitting instructions must be followed.

16.3. Each protector shall have permanently affixed to the exterior of the protector a clearly legible statement which can be easily read without removal of any tape or other temporary or permanent part, which contains language which effectively communicates to the purchaser and user the following information, using the same or similar language:

WARNING:

**NO PROTECTOR CAN PREVENT ALL CARDIAC OR OTHER INJURY.
CATASTROPHIC INJURY AND DEATH MAY OCCUR TO THE WEARER
DESPITE USING THIS PROTECTOR.**

16.4. Additional information included with the protector shall describe proper fitting and use of the protector. **The fitting and use instructions should allow the wearer to determine if the wearer's cardiac silhouette is covered by the impact attenuating components that allow the protector to be certified.**

This standard is subject to revision at any time by the responsible technical authority and must be reviewed every five years and if not revised either reapproved or withdrawn. Your comments are invited either for revision, modification or creation of additional standards and should be addressed to NOCSAE's Executive Director. Check the web at www.nocsae.org to obtain the latest version of a standard.

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Appendix 1

Verification Procedure for NOCSAE Thoracic Surrogate [NTS]

This procedure is conducted to insure that the response of the NTS is correct when impacted with a projectile between the velocities of 29.1 to 30.9 MPH. Adjustments to the data acquisition system for each force transducer may be necessary. If the NTS response cannot be adjusted so that the response is correct, then service of the NTS is indicated prior to any product testing. Prior to commencing this procedure, carefully inspect all equipment to ensure your lab's mechanical system is in proper working order.

Validation Protocol:

Step 1: Setup the NTS and data acquisition system using the load cell output information given on the Interface Load Cell Calibration Certificate and according to NOCSAE DOC 200.

Step 2: Depending on the type of testing your lab will be doing, obtain a lacrosse ball or baseball with the specifications described in Section 14.1.1 [baseball] or 14.1.2. [lacrosse].

Step 3: The target response of the unprotected surrogate at 30 mph \pm 0.2 is 140 lbf \pm 2. Impact the NTS at the center of the cardiac silhouette at 30 mph \pm 0.2. If the projectile impact results in a response of 140 lbf \pm 2, no adjustments are needed at this time - proceed to Step 5.

Step 4: If the projectile impact at 30 mph \pm 0.2 mph does not give a response of 140 lbf \pm 2, adjust the system settings and repeat the impact. Repeat this step until the impact response is 140 lbf \pm 2.

Step 5: Impact the NTS at the center of the cardiac silhouette at 29.1 mph \pm 0.2. The response at this velocity must be greater than 130 lbf. If the response of your system is below 130 lbf, repeat Step 4 and target the higher end of 140 lbf \pm 2.

Step 6: Impact the NTS at the center of the cardiac silhouette at 30.9 mph \pm 0.2. The response at this velocity must be less than 150 lbf. If the response of your system is above 150 lbf, repeat Step 4 and target the lower end of 140 lbf \pm 2.

Step 7: Once you are satisfied with the response at the cardiac silhouette record the data acquisition settings for future use.

NOTE: If your laboratory will be testing with both baseball and lacrosse balls, your lab must validate the NTS with both projectiles to obtain settings for each projectile.

If you have any questions as to how to perform this procedure, contact Southern Impact Research Center at info@soimpact.com.

FEBRUARY 2017 MODIFICATIONS/REVISIONS

- Changed 18” to 24” in Section 10.1 to match the requirements in ND 021
- Changed referenced (ball characteristics) section from 12 to 13 in Section 11.2.
- Added “ALL” to example warning in section 15.3
- Added reference to ND 021 Figures 1 and 2 in section 10.2 and removed those figures from this document

JUNE 2017 MODIFICATIONS/REVISIONS

- REVISION: Reduced equipment level from 3 to 2 in Section 1.1.
- REVISION: Increased pass/fail criteria from 90 lb-f to 112 lb-f in Section 14.1.2.
- REVISION: Increased pass/fail criteria from 180 lb-f to 225 lb-f in Section 14.2.2.
- REVISION: Included a system check for the upper and lower load cells and added an upper range to the allowable load cell response (150 lb-f) in Section 11.2.
- Added a second figure depicting upper and lower load cell placement.
- Removed language requiring 200 ms data capture and time at which data capture begins.
- Added reference to Figure 2 in Section 12.2.
- Added Appendix 1
- Changed the word below to above in Step 6 of Appendix 1.

FEBRUARY 2018 MODIFICATIONS/REVISIONS

- REVISION: Removed COR requirement for baseball projectile Section 13.1.1

JULY 2019 MODIFICATIONS/REVISIONS

- REVISION: Added requirement to place logo on the primary protective component.
- Added a definition of Primary Protective Component.
- Changed “chest protector” to “protector”.

JANUARY 2021 MODIFICATIONS/REVISIONS

- REVISION: Eliminated use of the upper and lower load cells and their associated pass/fail criteria
- REVISION: Random impact locations updated to impact the cardiac silhouette while shifting the protector instead of moving the NTS.
- REVISION: Added definitions for Torso Length and Extent of Protective Coverage
- REVISION: Added requirement that manufacturers shall provide the Torso Length of the person the submitted sample is intended to fit

If approved: JANUARY 2022 MODIFICATIONS/REVISIONS

- REVISION: Added labeling requirement for protectors that have removable primary protective components.
- Added language to clarify test sample requirement in section 5.2
- Added language to clarify information that should be included with fit and use instructions