VALIDITY OF A SET OF CLINICAL CRITERIA TO RULE OUT INJURY TO THE CERVICAL SPINE IN PATIENTS WITH BLUNT TRAUMA

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ABSTRACT

Background Because clinicians fear missing occult cervical-spine injuries, they obtain cervical radiographs for nearly all patients who present with blunt trauma. Previous research suggests that a set of clinical criteria (decision instrument) can identify patients who have an extremely low probability of injury and who consequently have no need for imaging studies.

Methods We conducted a prospective, observational study of such a decision instrument at 21 centers across the United States. The decision instrument required patients to meet five criteria in order to be classified as having a low probability of injury: no midline cervical tenderness, no focal neurologic deficit, normal alertness, no intoxication, and no painful, distracting injury. We examined the performance of the decision instrument in 34,069 patients who underwent radiography of the cervical spine after blunt trauma.

Results The decision instrument identified all but 8 of the 818 patients who had cervical-spine injury (sensitivity, 99.0 percent [95 percent confidence interval, 98.0 to 99.6 percent]). The negative predictive value was 99.8 percent (95 percent confidence interval, 99.6 to 100 percent), which is too low to justify its widespread use. Eight of the 818 patients who had cervical-spine injury had a low probability of injury according to the decision instrument, and only one of these two patients received surgical treatment. According to the results of assessment with the decision instrument, radiographic imaging could have been avoided in the cases of 4309 (12.6 percent) of the 34,069 evaluated patients.

Conclusions A simple decision instrument based on clinical criteria can help physicians to identify reliably the patients who need radiography of the cervical spine after blunt trauma. Application of this instrument could reduce the use of imaging in such patients. (N Engl J Med 2000;343:94-9.)

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BECAUSE unrecognized injury to the cervical spine can produce catastrophic neurologic disability, clinicians liberally order radiographs of the cervical spine, and as a result the majority of the radiographs are normal. Eliminating even a small proportion of the approximately 800,000 cervical-spine radiographs ordered annually in the United States for patients with blunt trauma could lead to substantial savings and decrease patients’ exposure to ionizing radiation.

Several small studies have suggested that patients with blunt trauma have a low probability of injury to the cervical spine if they meet all five of the following criteria: they do not have tenderness at the posterior midline of the cervical spine, they have no focal neurologic deficit, they have a normal level of alertness, they have no evidence of intoxication, and they do not have a clinically apparent, painful injury that might distract them from the pain of a cervical-spine injury.

Although the combination of these five criteria was reported to have a sensitivity of 100 percent for ruling out cervical-spine injury, the lower confidence limit for the sensitivity of the instrument was only 89 percent, which is too low to justify its widespread use. We organized the National Emergency X-Radiography Utilization Study (NEXUS) to validate this set of criteria and to test the hypothesis that patients with blunt trauma who meet all five of the above criteria have a very low probability of clinically significant injury to the cervical spine.

METHODS

Participating Centers

Twenty-one centers across the United States participated in this prospective, observational study. Among them were university and community hospitals, hospitals with and without residency programs, and public and private hospitals; they varied in size, in the level of activity in the emergency department, and in the level of trauma care they provided. The study was designed to assess the validity of the following five criteria (the decision instrument)

*The centers and investigators participating in the National Emergency X-Radiography Utilization Study (NEXUS) are listed in the Appendix.
VALIDITY OF CLINICAL CRITERIA TO RULE OUT INJURY TO THE CERVICAL SPINE IN PATIENTS WITH BLUNT TRAUMA

in ruling out cervical-spine injury in patients with blunt trauma: the absence of tenderness at the posterior midline of the cervical spine, the absence of a focal neurologic deficit, a normal level of alertness, no evidence of intoxication, and absence of clinically apparent pain that might distract the patient from the pain of a cervical-spine injury. Patients who met all five criteria were considered to have a low probability of injury and not to require radiographic or other imaging.

At each center, a physician in the emergency department served as a liaison to the study investigators, and a designated radiologist ensured that the collection of radiologic data was carried out completely and correctly. The liaison physician at each center attended a one-hour training session led by the regional study coordinator, at which the overall study design was presented and the decision instrument and each of the five criteria were explained. The liaison physicians were then responsible for training the participating clinicians in their emergency departments, in some cases through similar brief, formal training sessions and in others, informally. Searchable information and guidelines ("help" screens) were available on computer to assist all the participating clinicians.

Patients

All the patients with blunt trauma who underwent radiography of the cervical spine in a participating emergency department were included in the study. Patients with penetrating trauma and those who underwent cervical-spine imaging for any other reason, unrelated to trauma, were not eligible for inclusion. The participating clinicians were reminded at training sessions that although use of clinical criteria for risk assessment ("clinical clearance") of patients with trauma had come into wider practice, there was no definitive evidence regarding the safety of the clinical decision instrument on which the study focused. They were also cautioned against using the set of criteria being tested as the sole determinant of whether patients needed imaging. The ultimate decision whether to order radiography was made at the discretion of the treating physician, according to the criteria he or she ordinarily used, and was not determined in any way by participation in the study.

The study was prospective and observational. The protocol neither required nor directed any element of the care of enrolled patients and thus posed no risk to the patients. For purposes of confidentiality, the data on the patients were transformed with the use of unique identifying numbers before this information was downloaded into the central data bank, such that it was impossible for individual patients to be identified on the basis of study data. A waiver of informed consent was granted by each institution participating in the study.

Cervical-Spine Radiography

A standard series of three views of the cervical spine (cross-table lateral view, anteroposterior view, and open-mouth view of the odontoid) was obtained in all patients, unless computed tomography (CT) or magnetic resonance imaging of the entire spine was performed because plain-film radiography was impractical or impossible. Other imaging studies (oblique views, flexion-extension radiographs, or CT images) could be ordered in addition to the three-view series of radiographs at the discretion of the treating physician.

Collection of Data

The clinicians prospectively recorded demographic data for each study patient and noted whether each of the study criteria was present, was absent, or could not be assessed. For criteria that could not be assessed (for example, tenderness in a comatose patient), the patient was considered not to have met that criterion.

The results of all the evaluations were recorded on data forms before imaging of the cervical spine. Every patient with a completed data form underwent imaging. All the study sites agreed to obtain the radiographs only after a voucher attesting that the data form had been completed had been issued. If an attending physician believed that even the minimal delay associated with completing the brief data form might be harmful to the patient, the physician could obtain the study voucher before imaging by indicating that the patient was “unstable.” In such cases, the clinicians were encouraged to complete an assessment of the patient with respect to the five criteria as soon as possible, preferably before the results of radiography were known. Designating a patient as “unstable” was considered equivalent to identifying a clinically significant injury (defined below) and was a reason for considering the patient not to have a low probability of injury.

The five criteria for a low probability of injury were not explicitly defined, but possible interpretations of the criteria were reviewed during the training sessions at each center. In addition, information provided to the liaison physician and clinicians at each site included descriptions of possible characteristics that could exclude patients from being classified as having a low risk. This information was included in the guidelines available to the clinicians on the computer (and is available on request from the authors).

Assessment of Injuries

All the radiographs were formally interpreted by the designated radiologists at the study sites. Diagnoses of cervical-spine injury and determination of the type of any fracture were made according to the final interpretation of all the imaging studies. When the results they reported were ambiguous, the radiologists reviewed both their reports and the original radiographs to make the final determination of the type of any fracture. Neither the formal interpretation by the radiologists nor the classification of injuries was done with knowledge of findings recorded on study data forms.

A list of potential cervical-spine injuries was created before data were collected, and each injury on this list was categorized as clinically significant or not clinically significant (Table 1). Injuries that were not clinically significant were those that typically require no specific treatment and those that, if not identified, would be expected to result in no harm. Radiographically documented cervical-spine injuries were categorized as not clinically significant only if they were isolated and there was no evidence of other bony injury or ligamentous or spinal cord injury. All other cervical-spine injuries were considered clinically significant. In any case in which the radiologists’ report was unclear as to the exact nature of the injury, the injury was classified as clinically significant. During the study, the cervical-spine injuries were reviewed on an ongoing basis to identify any patient with a clinically significant injury in whom use of the decision instrument had failed (i.e., had indicated that the patient had a low probability of injury). A rule for stopping the study was in place, to be activated if five such cases were identified.

Statistical Analysis

To determine the sensitivity of the decision instrument to within 0.5 percent, we needed to enroll at least 737 patients with cervical-spine injury.9 Patients were considered to have a low probability of cervical-spine injury if they were clinically stable and met all five of the clinical criteria. For patients who did not meet all five of the criteria and who were reported to have a radiographically documented cervical-spine injury, the decision instrument was considered to have yielded a true positive result. For patients who did not meet one or more of the five criteria and who had a radiographically documented injury, the result was considered false negative. The result was true negative for patients who met all the criteria for a low probability of injury and who had no evidence of cervical-spine injury on radiography, whereas it was false positive for those who did not meet all the criteria but who had no injury.

Because we enrolled only patients who underwent imaging, we reviewed the neurosurgical records and quality-assurance logs of each participating site three months after the completion of the study, in order to identify any cases in which cervical-spine injury was missed because of an initial failure to obtain imaging studies.
The fracture was not accompanied by anterior soft-teroinferior portion of the second cervical vertebra. But whose plain films showed a fracture of the an-

jury, one was a 54-year-old man with a history of

who did have radiographically documented cervical-

identified as having a low probability of injury but

The instrument yielded a false negative result for 8 of

results of assessment with the clinical decision instru-

The study population consisted of 34,069 patients
evaluated by imaging of the cervical spine after blunt

trauma. Of these 34,069 patients, 818 (2.4 percent)

were 8 years old or younger.

The distribution of the patients according to the

presence or absence of cervical-spine injury and the

results of assessment with the clinical decision instru-

ment is shown in Table 2. The resulting perform-

ance of the decision instrument is shown in Table 3. The instrument yielded a false negative result for 8 of

the 818 patients with radiographically documented cervical-spine injury; 2 of these 8 patients were among

578 patients who met the predefined criteria for clin-

cally significant injury.

The eight patients whom the decision instrument

identified as having a low probability of injury but

who did have radiographically documented cervical-

spine injury are described more fully in Table 4. Of

the two patients who had a clinically significant in-

jury, one was a 54-year-old man with a history of

multiple motorcycle accidents who had no symptoms

but whose plain films showed a fracture of the an-
teriorinferior portion of the second cervical vertebra.

The fracture was not accompanied by anterior soft-
tissue swelling or any other abnormal finding on ei-

ther plain films or CT scans; the injury was described

in most of the radiology reports as an avulsion of the

end plate of this vertebra. However, it was also
described in one report as an “extension-teardrop”
fracture and thus met our criteria for clinically sig-

nificant injury. The patient remained asymptomatic

during a 24-hour hospitalization and refused any

treatment other than a soft cervical collar, which he

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<table>
<thead>
<tr>
<th>Patient's Sex/Age (yr)</th>
<th>Cervical-Spine Injury</th>
<th>Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>M/53 C6–C7</td>
<td>Chipped osteophyte</td>
<td></td>
</tr>
<tr>
<td>M/54 C2</td>
<td>Extension (teardrop)  fracture; normal alignment without soft-tissue swelling</td>
<td></td>
</tr>
<tr>
<td>M/20 C7</td>
<td>Anterosuperior end-plate avulsion, without soft-tissue swelling</td>
<td>Treatment with soft collar only; no sequelae</td>
</tr>
<tr>
<td>F/18 C5</td>
<td>Wedge compression fracture</td>
<td>Minimal loss of body height</td>
</tr>
<tr>
<td>F/81 C2</td>
<td>Isolated lateral-mass avulsion</td>
<td>Treatment with soft collar</td>
</tr>
<tr>
<td>M/84 C2</td>
<td>Isolated lateral-mass avulsion</td>
<td>Treatment with hard collar for 2 days, followed by soft collar</td>
</tr>
<tr>
<td>M/57 C6</td>
<td>Laminal fracture</td>
<td></td>
</tr>
</tbody>
</table>

* A negative result indicated that the patient was considered to have such a low probability of cervical-spine injury that imaging was not necessary.

Although there have been several case reports of occult injury to the cervical spine, most of these cases involved patients who either were inadequately evaluated or did not meet at least one of our criteria for low risk. Nevertheless, fear of missing a clinically occult injury has prompted physicians to order images of the cervical spine for virtually all patients who have blunt trauma. As a result, for each injury detected, a large number of films with negative findings are ordered. Because of the consequent human and economic losses, combined with medicolegal issues and concern about quality assurance, validation of selective criteria for ruling out probable cervical-spine injury requiring radiography in patients with blunt trauma is an important priority.

This study is the culmination of a series of investigations designed to derive and validate to a high level of confidence a set of clinical criteria that identify cervical-spine injury requiring radiography. Although it had already been demonstrated that an instrument based on the criteria we used has very high negative predictive value, a study of this size (with more than 737 patients with a fracture) was required for reliable estimation of its sensitivity.

In rare instances, this decision instrument will undoubtedly miss individual cases of cervical-spine injury. In this study, the overall rate of missed cervical-spine injuries was less than 1 in 4000 patients. To place the characteristics of the decision instrument in perspective, we can consider that every full-time emergency physician orders cervical-spine imaging in approximately 32 patients annually (given that approximately 25,000 full-time–equivalent emergency physicians in the United States order about 800,000 films each year); physicians can therefore expect to encounter a case of occult cervical-spine injury (which occurs less than once for every 4000 radiographs obtained) perhaps once in every 125 years of clinical practice. Missed cases of clinically significant cervical-spine injury (which occurred in only two patients in our study) and those requiring specific therapy (one patient) appear to be even more rare.

Two of our patients did have clinically significant injury, according to our formal definition, that was missed, although one of them seemed clearly not to have had an acute injury and had no clinical sequelae even though he essentially refused treatment. Only one patient with a false negative result underwent spe-
cific treatment for the injury; this case may actually represent misapplication, rather than failure, of the decision instrument, since the patient had loss of consciousness, a clavicular fracture, and neurologic symptoms (paresthesias).

Nevertheless, on extremely rare occasions, a missed injury may lead to profound consequences for an individual patient. We believe that although clinicians can generally adhere to the clinical criteria in the decision instrument, they should be free to make exceptions for individual patients on clinical grounds. In any case, no decision instrument is ever likely to be 100 percent sensitive, and the medical and economic costs of a quixotic search for absolute diagnostic certainty can lead to more harm than good.38

In this study, application of the decision instrument would have decreased the overall ordering of radiographs by only 12.6 percent. This decrease is far smaller than the reduction of almost one third in the use of radiography that would have been predicted by the results of our previous study, conducted in a single hospital,8 and may reflect an influence of the previous study on the ordering of radiographs at institutions participating in this study. In emergency departments with more liberal use of imaging, the effect of the adoption of the decision instrument could be greater than that seen here. In any case, even a reduction of one eighth in the ordering of radiographs, as our study indicates is possible, would translate into a substantial decrease (by about 100,000) in the number of cervical-spine radiographs obtained each year in the United States.

For a decision instrument to be valuable, it must be shown to be reliable when used by different practitioners.39,40 Each of the criteria of our decision instrument has been shown to have good-to-excellent interobserver reliability (kappa, 0.58 to 0.86), and interobserver agreement for the decision instrument as a whole is excellent (kappa, 0.73).36 In this study, the decision instrument had an extremely high sensitivity when applied by a very large number of clinicians at various sites. Although physicians who participate in a multicenter trial are not perfectly representative of all physicians, the participation of physicians at all levels of training and in many different environments in this study provides powerful evidence of the external validity of the clinical criteria. Furthermore, the ability of such clinicians to apply the criteria with high sensitivity, despite minimal formal training, suggests that the use of the decision instrument can be widely taught, without undue expenditure of human or economic resources.

We chose not to define the individual criteria of the decision instrument explicitly, for two reasons. First, we do not believe such criteria can be precisely defined in a clinically meaningful way. An attempt to define a “distracting” injury, for example, with a long list of various injuries that could distract a patient from a cervical-spine injury would be extremely misleading. Some contusions, for example, may be associated with extreme pain, whereas not all long-bone fractures are particularly painful. Therefore, we allowed the clinicians to judge whether the patients had an injury that could produce distracting pain and thus required cervical-spine imaging. Similarly, we believe that evidence of intoxication and the level of alertness are best evaluated on the basis of clinical judgment, rather than laboratory tests or uniform criteria.

Second, if physicians were required to consult precise definitions for each criterion whenever evaluating a patient with blunt trauma — for example, to check the exact length a laceration needed to be to qualify as a distracting injury — the decision instrument would rapidly fall into disuse. We believe our limited number of criteria are straightforward, logical, and easy to remember, and that they thus can be readily applied at the bedside.

Because this study was strictly observational, it is possible that there were patients with cervical-spine injury at the study sites who met the decision-instrument criteria but did not undergo radiography and were thus not included in the study. It is impossible to identify every patient with a potential illness or injury, since some patients may have symptoms so minor that they do not seek medical care or their presentation does not suggest the possibility of the disease or injury in question. Such patients cannot be identified by this or any other decision instrument. The question we wished to address was whether some of the many patients who are currently considered candidates for cervical-spine imaging can be safely classified as having such a low probability of injury on clinical grounds that radiography need not be performed, with consequent cost savings and medical benefits. Our methods have allowed us to answer this question.

In summary, this prospective, multicenter study confirms the validity of a decision instrument based on five clinical criteria for identifying, with a high degree of confidence, patients with blunt trauma who have an extremely low probability of having sustained injury to the cervical spine. The sensitivity of this set of criteria approaches 100 percent for clinically important injuries, and its general application should result in both clinical and economic benefit. As with any other clinical tool, it should be applied with great care and should not replace clinical judgment in the care of individual patients. There may be compelling reasons to order cervical-spine images in individual cases, even if all the criteria for a low probability of injury are met.

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VALIDITY OF CLINICAL CRITERIA TO RULE OUT INJURY TO THE CERVICAL SPINE IN PATIENTS WITH BLUNT TRAUMA

We are indebted to Guy Merchant, project coordinator, for his outstanding contributions to the project, as well as to the house officers and attending physicians at the participating sites, without whose cooperation and hard work this study would not have been possible.

APPENDIX

The following centers and investigators collaborated in the NEXUS study: Principal investigator — W. Mower; Co-investigator — J. Hoffman; Steering Committee — J. Hoffman, W. Mower, K. Todd, A. Wolfson, and M. Zucker; Site investigators — Antelope Valley Medical Center (Los Angeles): M. Brown and R. Sisson; Bellevue Hospital (New York): W. Goldberg and R. Siegmann; Cedars-Sinai Medical Center (Los Angeles): J. Geiderman and B. Pressman; Crawford Long Hospital (Atlanta): S. Pitts and W. Davis; Egleston Children’s Hospital (Atlanta): H. Simon and T. Ball; Emory University Medical Center (Atlanta): D. Lowery and S. Tiggies; Grady Hospital (Atlanta): C. Finney and S. Tiggies; Hennepin County Medical Center (Minneapolis): B. Mahoney and J. Hollerman; Jacobs Medical Center (Bronx, N.Y.): M. Touger, P. Gennis, and N. Nathanson; Maricopa Medical Center (Phoenix, Ariz.): C. Pollack and M.Connell; Mercy Hospital of Pittsburgh (Pittsburgh): M. Turturro and B. Carlin; Midway Hospital (Los Angeles): D. Kalmanson and G. Berman; Ohio State University Medical Center (Columbus): D. Martin and C. Mueller; Southern Regional Hospital (Decatur, Ga.): W. Watkins and E. Hadley; State University of New York at Stony Brook (Stony Brook): P. Viccillo and S. Fuchs; University of California, Davis, Medical Center (Sacramento): E. Panacek and J. Holmes; University of California, Los Angeles, Center for the Health Sciences (Los Angeles); J. Hoffman and M. Zucker; University of California, San Francisco, Fresno University Medical Center (Fresno, Calif.): G. Henley and R. Lesperance; University of Maryland Medical Center (Baltimore): B. Browne and S. Marris; University of Pittsburgh Medical Center (Pittsburgh): A. Wolfson and J. Towers; Hermann Hospital, University of Texas Health Science Center (Houston): N. Adame, Jr., and J. Harris, Jr.

REFERENCES