RISK OF TRANSFORAMINAL EPIDURAL INJECTIONS

To the Editor:

There have been recent reports of serious complications arising from the injection of steroids into the spinal neural foramina especially in the cervical region (1-5). These complications have followed the relatively routine interventional pain clinic procedure of therapeutic injection to treat radicular pain. The procedures typically involve fluoroscopically-guided needle placement into the neural foramen with injection of water soluble contrast to document needle position. Disastrous events including paraplegia, quadriplegia and death have occurred shortly after the subsequent injection of particulate steroid into the needle. At least some of these procedures have been performed by experienced practitioners presumably using well-accepted injection technique. As interventional pain specialists, it behooves us to objectively re-examine the risks and benefits of neural foraminal injection so that we may maintain safe and efficacious medical practice for the benefit of our patients with persistent pain.

Cervical transforaminal steroid injections have been used increasingly over the past decade to diagnose and treat neck and upper extremity pain syndromes. From review and extrapolation of insurance data, it is estimated that over 100,000 cervical transforaminal steroid injections have been performed in the United States in the past 10 years.

Review of the recent complications has revealed the following commonalities:

1. A sharp-tipped needle was placed into the neural foramen under fluoroscopic guidance.
2. Contrast was injected and apparently documented good needle position.
3. A particulate steroid was injected subsequent to injection of contrast.
4. The complication occurred within minutes of the steroid injection.
5. Patient evaluation after the event revealed extensive spinal cord infarction as the pathological event.

Most interventional pain specialists agree that the transforaminal approach to the epidural space has advantages in the diagnosis and treatment of certain pain syndromes (6-24). When irritation or inflammation of a specific spinal nerve root or dorsal root ganglion is suspected as the cause of intractable pain, transforaminal injection is the most direct route to apply medication to the pain generator. Furthermore, transforaminal injections are more likely to distribute therapeutic steroid to the anterior epidural space, which is the region most likely affected by irritation from disrupted or herniated intervertebral discs.

Spinal Cord Blood Supply

Certainly, injection into the foramen poses unique risks because of the presence of foraminal arteries that are not present in the dorsal epidural space. These foraminal arteries are the beginning of a rather tenuous anastomotic arterial network of feeder vessels that deliver blood to the spinal cord. The arteries that supply the spinal cord are end arteries and collateral blood supply is lacking throughout much of the cord. The spinal radicular arteries vary in number and location and travel with the nerve root through the foramen en route to the cord. In the thoracolumbar region the artery of Adamkiewicz exits the aorta and typically enters the spinal canal through a lower thoracic or upper lumbar foramen most commonly from the left side (2). An ascending sacral radicular artery and a second thoracic radicular artery may also contribute to thoracolumbar cord blood supply. In the cervical region, the vertebral arteries give rise to the anterior spinal artery at the upper cervical cord. The anterior spinal artery then descends through the spinal canal supplying the upper cord. The anterior spinal artery receives contributions from one or more radicular arteries, which originate from the more proximal portions of the vertebral artery and course through one or more of the cervical neural foramina with the nerve root. The most common scenario involves a single radicular feeder artery exiting the right proximal vertebral artery and following the C6 nerve root into the spinal canal through the right C5-6 neural foramen.

Minimizing Risk

It is most likely that the spinal cord and/or brainstem infarction documented as the pathologic basis for injury after cervical transforaminal injection is the sequelae of embolism of particulate steroid into the downstream spinal cord arteriolar system. This is the only mechanism that would explain the size and distribution of infarction (anterol cord, Brown-Sequard lesion) associated with these events. Arterial spasm from needle penetration is an unlikely cause of injury and the long history of angiography has demonstrated that needles can safely be passed into arteries without causing spasm. External vascular compression from the injectate would also be unlikely to cause abrupt and extensive spinal cord infarction.

In order for injection into the neural foramen to cause spinal cord embolization with subsequent cord infarction, two conditions must first be met:

1. The tip of the needle must lie within the lumen of radicular artery that is supplying the arterial feeder system of the cord.
2. The injected material must be capable of causing downstream interruption of blood flow.

Those of us who routinely use particulate steroids such as methylprednisolone and triamcinolone in pain clinical practice know that these compounds are capable of clogging the lumen of a 30 gauge needle. They are also capable of occluding blood flow as they are distributed to the arborized arteriolar network within the substance of the spinal cord. The fact that relatively large sections of the spinal cord even including portions of the brainstem have been infarcted by single transforaminal injections attests to the fact that the radicular arteries arborize into a vast interconnected network of end-artery vessels.

Yet it seems clear that if we can avoid injecting particulate steroid into the radicular artery, transforaminal injection is a relatively safe procedure. It is likely that the disastrous complications described above occurred either because the contrast flow pattern was intravascular but was incorrectly interpreted or because the needle tip migrated into the artery after the contrast was injected. No other explanation seems feasible. Meticulous injection technique should therefore minimize
the risk of intravascular injection and the following points should be considered:

1. Correct interpretation of the results of contrast injection is crucial to make absolutely certain that the nerve root and/or epidural space is outlined with contrast prior to injecting steroid. If contrast flows out of the needle tip and outlines the exiting nerve root lateral to the neuroforamen, then by definition the needle tip cannot be intravascular.

a. Contrast should outline the exiting nerve root for a clear and definable distance laterally.

b. It is not necessarily a problem if contrast moves retrograde through the neuroforamen and into the epidural space but if the nerve root is not simultaneously visualized then it is sometimes difficult to determine whether the contrast moving medially is intravascular or epidural.

2. Once contrast injection has confirmed safe needle position, great pains must be taken to keep the needle completely stationary for the remainder of the procedure.

a. Use of a pigtail extension attached to the needle hub will facilitate exchange of contrast and medication syringes without moving the needle itself.

b. Injection of the active medication should be done incrementally with frequent fluoroscopic visualization to make sure the needle remains in proper position.

c. Subtle movement of the perineural contrast pool visualized by repeat fluoroscopy during incremental injection of steroid indicates that the needle tip is remaining extra vascular.

In addition, some experts have recommended using blunt tip needles for transforaminal injection although these needles are 22 gauge and are currently not available in 25 gauge diameter making them less desirable for some practitioners. Blunt tip needles have been demonstrated to be less likely to penetrate vascular structures. Clear steroid preparations such as betamethasone may also decrease risk although it is unclear whether non-particulate steroid preparations will prove as efficacious as the particulate medications.

As evidenced by the long history of safe and efficacious transforaminal injection in the experience of many practitioners, it is certainly possible to perform this procedure with good result. Nonetheless, recent events demonstrate the potential for disaster with injection into the neuroforamen and each interventional pain specialist should reexamine the indications and techniques for this procedure. In addition, informed consent for transforaminal injection should include an explanation of the risk of spinal cord injury.

REFERENCES


