Gd-DTPA enhancement of posterior epidural scar: an experimental model.

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Gd-DTPA Enhancement of Posterior Epidural Scar: An Experimental Model

Because of the tremendous clinical and physiological importance of anterior epidural scar, an easily produced and reproducible model to assess potential pathways for lessening its formation is a necessity. We speculated whether posterior epidural scar (produced by the less complex surgery of laminectomy alone) could be considered equivalent to anterior scar from an imaging standpoint; that is, enhancement following Gd-DTPA irrespective of scar age. Posterior epidural scar in dogs showed the highest degree of enhancement 1 month after surgery, with a rapid decline thereafter out to 4 months postsurgery to a level equivalent to that of paraspinal muscle. Gd-153-DTPA time/activity curves paralleled the Gd-DTPA findings. Light microscopy showed granulation tissue after 1 month, and mature scar with large amounts of collagen 4 months after surgery. Electron microscopy showed tight capillary endothelial junctions.

An appropriate model for epidural scar, which has imaging characteristics similar to human anterior scar, necessitates an extensive lumbar laminectomy with anterior epidural dissection. A simple laminectomy, while easily performed, does not provide a physiologically correct time course of enhancement.

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A great deal of time and attention has been given to the control of epidural scar formation through placement of fat grafts or with various experimental agents [1–4]. These approaches have commonly used a simple laminectomy model to assess posterior epidural scar [5, 6]. This type of scar, while complicating the initial stage of reoperation, is otherwise not routinely involved in the failed back surgery syndrome, nor given a great deal of consideration in imaging studies [7]. This is in stark contrast to anterior or lateral epidural scar, which is associated with surgically remedial causes of recurrent pain (i.e., recurrent disk herniation) as well as the obviously important anatomy of the exiting nerve roots [8].

Because of the importance of anterior epidural scar, many techniques have been advocated for its diagnosis and distinction from disk material [9–13]. Recent work has demonstrated enhancement of epidural scar by MR following administration of Gd-DTPA in previously operated spines [14]. Anterior epidural scar enhances consistently. Anterior epidural scar has been seen to enhance in patients who had surgery as long as 30 years prior to the Gd-DTPA–enhanced studies. We have previously shown that marked enhancement of epidural scar in dogs (5 months after surgery) in a pattern similar to that seen in humans is secondary to scar vascularity and loose scar capillary endothelial junctions [15]. This model was surgically complex, since it necessitated an extensive laminectomy with dissection anterior to the thecal sac for placement of a plastic constrictive band.

Because of the tremendous clinical and physiologic importance of anterior epidural scar, an easily produced and reproducible model to assess potential pathways for lessening its formation is a necessity. We wondered whether posterior epidural scar produced by the less complex surgery of simple laminectomy could, in fact, be considered equivalent to anterior epidural scar. This equivalence, from
an imaging standpoint should be consistent enhancement irrespective of age, as seen in human anterior scar. To test
the hypothesis that posterior scar would enhance consistently despite the time since surgery, we undertook a four-part
study in dogs following simply laminectomy: (1) the time
course of enhancement of posterior scar with Gd-DTPA was
measured over 4 months, (2) the time/activity curves were
obtained for posterior epidural scar with Gd-153-DTPA, (3)
the histology of posterior scar was assessed with light
microscopy, and (4) the ultrastructure of scar, including the
endothelial morphology, was assessed with electron micros-
copy.

Materials and Methods
Animal Model

Ten adult female beagle dogs (12–14 kg) had laminectomy at the
L6–L7 level. Epidural scar has been shown to form consistently after
laminectomy in dogs [5]. Two dogs were sacrificed after 1 month for
histology. The other eight dogs had the imaging studies described
below.

Imaging Schemes

Imaging experiments were performed on a 1.5-T Siemens
superconducting magnet. Eight dogs were imaged at monthly intervals up
to 4 months postsurgery. The dogs were anesthetized with sodium
pentobarbital IV, intubated, and then placed supine on a 21-cm-
diameter resonator coil centered over the lower lumbar spine. Initially,
sagittal spin-echo images 400/13/4, (TR/TE/excitations) with a 4-mm
slice thickness and 50% gap were obtained for localization of the
axial images at the operative site. Following acquisition of a precon-
trast axial image at the surgical site, Gd-DTPA was injected intrave-
nously as a bolus (0.1 mmol/kg). Images were obtained every minute
for the first 10 min, and then every 5 min up to the end of the
examination (45 min). This procedure was repeated at monthly inter-
vals.

The dynamic MR studies were obtained by using a gradient-echo
technique (FLASH) with the following parameters: 20/10/8, 60° flip
angle, 10-mm slice thickness, and 256 phase-encoded cycles. With
the addition of recycling time, images could be obtained every 60
sec. The phase- and frequency-encoded directions were reversed,
so flow-related artifact would not project over the spine.

Data Analysis

Signal intensities (SI) were measured from regions of interest (ROI)
centered over the posterior epidural scar as well as paraspinal
musculature on each pre- and postcontrast image. ROIs included
approximately 0.1 cm², and were held constant in position and size
as each intensity measurement was taken. Hard-copy images were
obtained of the positioning of the axial images and ROIs on each dog
at month 1, which were subsequently used to position the imaging
sites and ROIs for the remaining 3 months.

Percent contrast enhancement was calculated by using the follow-
ing formula:

\[
\text{enhancement} = \frac{\text{post SI} - \text{pre SI}}{\text{pre SI}} \times 100
\]

Standards were not included because of the difficulty in maintaining
correct position of the standard on subsequent months, with the
inherent field drop-off associated with surface coils.

A sham experiment was performed on dog 256 at 3½ months
after surgery to assess signal variability over a 15-min period. A
precontrast axial image was obtained, and then saline was injected
into a forefoot vein and postinjection images repeated each minute
through 10 min postinjection, with a final image obtained at 15 min
postinjection. Signal intensities were measured for scar and muscle
as above.

Gd-153-DTPA

Two dogs, each at 1 month and 4 months after surgery, were
intubated and anesthetized. Surgical dissection was carried down to
the epidural scar. Gd-153-DTPA was then injected intravenously and
samples were obtained of venous blood, paraspinal musculature, and
posterior epidural scar at 1, 3, 5, and 10 min postinjection. At 1
month after surgery the dogs received 20 µCi Gd-153-DTPA, while
at 4 months after surgery they received 40 µCi. Sample activity was
subsequently counted, and the rate corrected to yield counts/gram
tissue for construction of a time/activity curve.

Histology

Two dogs were sacrificed at 1 month and four dogs sacrificed at
4 months after surgery, and samples of posterior epidural scar were
obtained. Samples for light microscopy were obtained from each of
the six dogs and fixed in 10% formaldehyde embedded conventionally
in paraffin. Samples for electron microscopy were obtained from
the four dogs sacrificed 4 months postsurgery and were fixed in 2.5%
glutaraldehyde buffered to pH 7.4 and 0.1 mol/l sodium cacodylate
and 7.5% sucrose for a minimum of 24 hr. Postfixation was performed
with 1% osmium tetroxide for 1 hr. Samples were embedded in Epon
812, and portions of interest were cut by using diamond knives and
were stained with uranyl acetate. Observations were made on a JEM
100cx electron microscope.

Fig. 1.—Sham experiment shows little signal variation over a 15-min
period.
Results

Dynamic MR

Of the 32 MR examinations performed in the eight dogs (four examinations each), 30 were without technical flaws. The month-4 (M4) examination in dog 256 did not have data beyond 20 min postinjection because of motion artifact. Similar problems were encountered in the M2 data in dog 254.

A sham examination was performed in dog 256 at 3½ months after surgery (Fig. 1). This showed minimal variation within the signal intensities over the examination time.

The signal intensities of the posterior epidural scar in the eight dogs showed a consistent trend over the 4-month period. A typical plot is shown in Figure 2A. The M1 enhancement curves showed the highest peaks during the 4-month period (X = 106%, R = 83–132%). Each succeeding month showed overall lower enhancement peaks (M2 X = 77%, R = 70–90%; M3 X = 52%, R = 35–85%; M4 X = 30%, R = 10–80%). There was generally a slow decline in enhancement from the time to peak out to the end of the study at 45 min after injection.

Paraspinal muscle, used as a control, showed consistently low levels of peak enhancement throughout the 4-month period (M1 X = 29%, R = 20–43%; M2 X = 20%, R = 14–25%; M3 X = 21%, R = 14–34%; M4 X = 21%, R = 11–34%). A typical plot is shown in Figure 2B. Times to peak enhancement for scar and muscle are shown in Table 1.

Gd-153-DTPA

The time/activity curves for blood at both 1 and 4 months after surgery showed the typical exponential decrease in activity as the gadolinium was cleared from the vascular system (Fig. 3). The time/activity curves for paraspinal muscle and scar paralleled the findings seen with the dynamic MR. That is, at month 1, the overall activity in epidural scar was higher than that of the low-level activity seen with paraspinal muscle. At month 4, paraspinal muscle and scar activity closely paralleled each other throughout the duration of the examination.

Histology

Light microscopy of the epidural scar at month 1 showed areas of granulation tissue with capillaries and plump fibroblasts, as well as additional areas showing more extensive collagen deposition (Fig. 4A). Histology at month 4 showed extensive mature scar composed principally of collagen, with a few intervening capillaries and fibrocytes (Fig. 4B). In two
Fig. 3.—A and B, Gd-153-DTPA time/activity curves at 1 month (A) and four months (B) after surgery for blood, posterior epidural scar, and paraspinal muscle. Scar shows a higher level of activity than muscle at 1 month, while scar and muscle have similar low levels of activity at 4 months, paralleling the Gd-DTPA MR experiment (Fig. 2).

Discussion

This study has demonstrated that a simple laminectomy model in dogs is not appropriate or applicable to the study of epidural scar in humans since it fails to show persistent enhancement beyond 1 month after surgery. After 1 month, the enhancement of scar decreased rapidly in this dog model and reached a level equivalent to that of paraspinal muscle. The dynamic MR findings were substantiated by the Gd-153-DTPA time/activity curves, which showed scar activity higher than muscle at month 1, but equivalent activity by month 4. In contrast, human anterior and lateral epidural scar shows persistent enhancement after Gd-DTPA administration even years after surgery [14]. In this model, posterior scar behaved more like typical peripheral scar; that is, it underwent rapid cicatrization with increasing amounts of collagen and decreasing amounts of water and vasculature [16]. Electron microscopy showed a “mature” endothelium at 4 months after surgery, with tight endothelial cell junctions. This differs from dogs, areas of metaplastic cartilage were seen at the laminectomy site.

Electron microscopy consistently demonstrated tight capillary endothelial junctions seen as electron-dense regions with close apposition of the outer leaflets of the cell membranes (Fig. 5). On only two sections were “loose” type endothelial junctions recognized (Fig. 6) where the outer leaflets of the adjacent cell membranes were less closely approximated to each other. Interspersed between the capillaries were occasional fibrocytes and a large amount of collagen (Fig. 7).
the leaky junctions that are present in immature scar in both our previous dog model and in humans [15].

Why the disparity in findings between scar in postoperative patients and this dog model? A major consideration is the nature and location of the scar itself. It is apparent that not all scar reacts in the same way over time, and it may be necessary to subdivide epidural scar in part on the basis of location. We postulate that anterior epidural scar cannot be considered to be in an environment that is equivalent to a pristine environment to which metaplastic cartilage was present intermixed with scar.

This study suggests that an appropriate model for epidural scar that has similar imaging characteristics to human epidural scar necessitates an extensive lumbar laminectomy with anterior epidural dissection. A simple laminectomy alone, while easily performed and commonly used, does not provide a physiologically correct time course of enhancement.

REFERENCES


Fig. 5.—Tight dog scar capillary endothelial junction shows close apposition of membrane leaflets (arrows) (electron micrograph x108,000). The junctions were commonly seen in the month-4 specimens.

Fig. 6.—“Loose” capillary endothelial junctions, where outer membrane leaflets show increased electron density (arrows) and do not as closely approximate each other. These junctions were rare in the specimens obtained 4 months after surgery (electron micrograph x81,000).

Fig. 7.—Interstice of mature scar shows abundant collagen, demonstrating its typical staining periodicity, and scattered fibrocytes (electron micrograph x6975).
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