

CT GUIDED NEUROLYTIC COELIAC PLEXUS BLOCK (NCPB) IN ABDOMINAL MALIGNANCIES: EXPERIENCE IN A CANCER HOSPITAL

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Abstract: The purpose of this study is to evaluate the efficacy of CT guided neurolytic coeliac plexus block (NCPB) in advanced pancreatic and other abdominal malignancies who become refractory to routine measures of pain relief. A total of nine patients were subjected to the procedure. A diagnostic block using local anaesthetic preceded the injection of alcohol to assess the expected response. Relief of pain was assessed objectively using a visual analog scale for pain (VAPS). There was significant relief of pain in 8 out of 9 patients. Thus it is concluded that CT guided neurolytic coeliac plexus block is an effective way to control pain in advanced abdominal malignancies and helps to improve terminal care.

Keywords: Coeliac plexus block, Neurolytic block, Abdominal malignancy, CT guidance

INTRODUCTION

Celiac plexus is the largest sympathetic innervation of abdominal viscera. It is located antero lateral to the origin of the celiac artery on both sides. Its involvement by disease, particularly in malignancies such as pancreatic and gastric leads to intractable visceral pain. Over time response to analgesics and opiates diminishes and celiac plexus block is an effective and durable way of managing such pain¹. CT guided NCPB offers accuracy of needle placement and is done with the patient in prone position. Patient should be off any analgesics on the day of the procedure and a diagnostic block using local anesthetics is first undertaken to evaluate efficacy. If there is significant relief in pain absolute alcohol is then injected for neurolysis. Best results are obtained with bilateral blocks and the effect lasts for 4 to 6 months before nerve regeneration occurs. Possible complications are hypotension, diarrhoea, puncture injuries and sexual dysfunction.

MATERIALS AND METHODS

Nine patients with abdominal malignancies were subjected to NCPB. All had inoperable or recurred intraabdominal malignancies & suffered from intractable upper abdominal pain and/or back pain. 7 patients had pancreatic carcinoma and 2 had gastric carcinoma. All were patients with pain severe enough to require frequent administration of intravenous analgesics and narcotic agents.

Pain was rated by the patient according to a visual analog scale for pain (VAPS) ranging from 0 (minimum) to 10 (maximum) before and after the procedure to objectively assess response. Patient was kept off all analgesics (oral and intravenous analgesics, transdermal patches) for period of 24 hrs prior to the procedure to eliminate the contribution of pain relief from

these analgesics. This allowed a more realistic assessment of pain relief solely from NCPB.

Informed consent was obtained with specific attention to complications associated with NCPB. The procedure was performed under supervision of qualified anaesthetist and under monitoring of vital parameters. The CT scans were performed on a spiral dual slice CT Scanner (Siemens Emotion Duo, Erlangen, Germany). Patient was placed prone on the CT table (Fig 1). After analyzing a previously acquired diagnostic CT scan to assess the overall extent of the disease and the approach, a limited guiding scan was performed with application of radio-opaque surface markers. The aorta and celiac artery origin were identified. Intravenous contrast was not considered necessary.

Under all aseptic precautions and after initial local instillation of anaesthetic agent in skin and subcutaneous tissue, a 22 gauge spinal needle was then introduced paraspinally at the chosen marker much like the procedure adopted for a CT guided biopsy. The needle tip was placed antero lateral to the aorta at

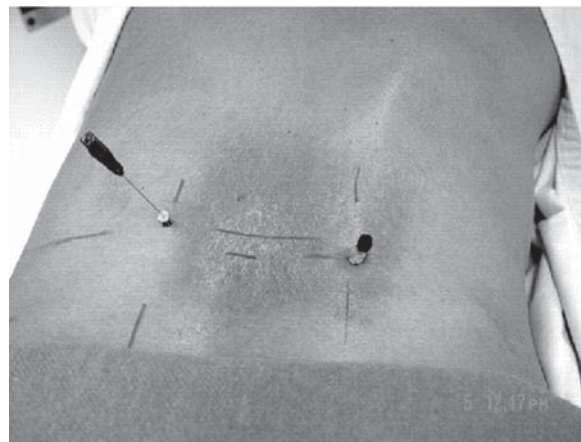


Fig 1. Patient position with needles

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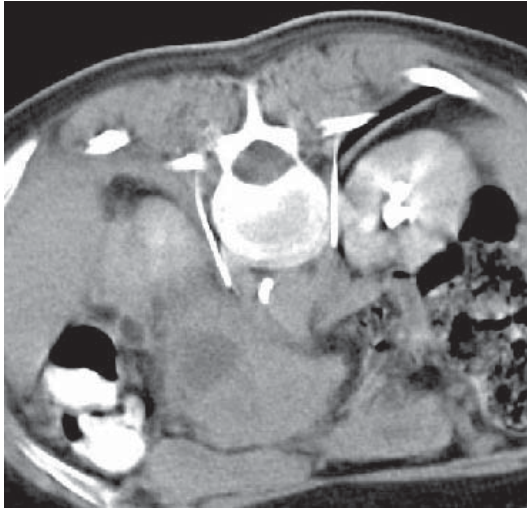


Fig 2. Final needle position

the level of the celiac artery (Fig 2). Needle trajectory was chosen to avoid the transverse process and medial margin of both kidneys. The right side entry point is usually somewhat more lateral than that on the left because optimal placement is between the vena cava and the aorta and requires an wider angle than that on the left.^[2] The needles are to be positioned on the right and left sides immediately adjacent to the area of the ganglion. Since the pleura is to be avoided, the needle entry point is more caudal than needle tip. Optimal placement on the right is between the aorta and vena cava, but a tip location adjacent to the posterior margin of the vena cava is acceptable. For the left side, it is to be adjacent to the superior mesenteric artery, but placement adjacent to the aorta is acceptable.² Needle position in either case was confirmed by limited guiding CT scans. 2ml of nonionic intravenous contrast medium (Omnipaque, Iohexol, GE healthcare) mixed with equal amount of local anaesthetic agent (Lignocaine 2% with adrenaline 1:200000). Various workers including Haaga et al also recommend use of air instillation instead of radiographic contrast.² However air will not permeate in tissue like alcohol would. After allowing few minutes for the spread of contrast, another limited CT scan confirmed the permeation of contrast around the aorta on either side (Fig 3 and 4). In case the pattern of contrast spread was not as desired, the needle position was accordingly modified. But total non permeation of contrast into the paraaortic region in spite of attempts at changing needle position pointed to the non feasibility of the procedure. As it happened in one of our patients who had substantial primary disease from carcinoma of pancreas encasing the abdominal aorta on all sides, providing no access for anaesthetic agent to the region of celiac plexus.

After having confirmed needle position, 13 to 15 ml local anaesthetic (bupivacaine 0.25%) mixed with 2ml nonionic contrast (Omnipaque, Iohexol, GE healthcare) was injected slowly on either side through the spinal needles. The VAPS score was then assessed after an interval of around 10 minutes for the effect of bupivacaine to set in. Relief in pain upwards of 50% was considered as measure of adequate response justifying further definitive alcohol ablation. Patient was also shifted to postoperative ward for



Fig 3. Spread of contrast

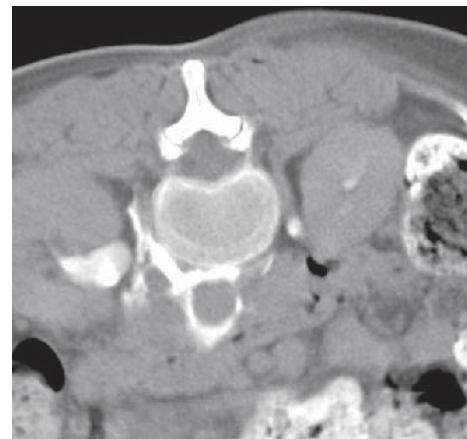


Fig 4. Note that the contrast outlines the aorta

careful observation of immediate and delayed complications if any of the procedure. Our experience in this regard was largely uneventful, with only complaint of pain around the needle insertion site and needle tract in 2 patients.

All patients with significant subjective pain relief and no evidence of complications (8 out of 9) were taken up for definitive alcohol ablation within 24 hrs. This involved placement of needles in the same manner as above. Then 50% v/v of 100% ethanol with 0.25% bupivacaine (total 13 to 15 ml) on either side was instilled to complete the procedure. Patients were again kept under observation overnight to assess pain relief and complications.

RESULTS

Pain relief

Out of 9 patients under review, definitive procedure with alcohol neurolysis was not performed in one due to failure of pain relief after diagnostic block using local anaesthetic. This was due to extensive primary disease encasing the aorta and celiac plexus completely, thereby disallowing permeation of the anaesthetic drug in the vicinity of the plexus. Other 8 patients successfully underwent the procedure and had significant drop

may be traumatized. We performed all our procedures using the posterior approach that possibly was the reason for lower complications.

The success of the neurolytic celiac plexus block, despite different approaches and methods used, depends on adequate spread of the injectate in the coeliac area. To evaluate CT patterns of neurolytic (mixed with contrast) spread, the coeliac area was divided on the frontal plane into four quadrants, upper right and left and lower right and left, as related to the coeliac artery. Results were expressed as the number of quadrants into which contrast spread, *i.e.*, four, three, two, or one quadrants with contrast. The patterns of contrast spread according to the number of quadrants with anatomic distortion were analyzed. Findings suggest that, using the single-needle anterior approach, the neurolytic spread in the coeliac area is highly hampered by the regional anatomic alterations. It also appears that only a complete (four quadrants) neurolytic spread in the coeliac area can guarantee long lasting analgesia, and that this picture may be obtained in a very limited fraction of patients with regional anatomic alterations¹¹. Regardless of the technique used to improve the spread of the injectate in the plexus area, failures are common due to regional infiltration by cancer tissue and anatomy distortion by either previous surgery or radiation therapy-induced fibrosis.

Workers have also gone further to compare conscious sedation with and without coeliac plexus block in patients undergoing thermal ablation of liver metastasis. (Alexander Beck, RSNA 2004). Results

reveal that percutaneous coeliac plexus block in addition to conscious sedation is a safe and reliable method to significantly decrease the need of pain medication and intervention time. So the future for CT guided coeliac plexus block holds a lot of promise.

REFERENCES

1. Oscar A. de Leon-Casasola. *Critical Evaluation of Chemical Neurolysis of the Sympathetic Axis for Cancer Pain. Cancer Control* 2000, 7(2):142-148
2. Haaga John R, Kori Shashidhar H, Eastwood DW, Borkowski GP. *Improved technique for CT guided celiac ganglia block. Am Journal of Radiology*:142, 1984:1201-1204
3. Boas RA. *Sympathetic blocks in clinical practice. Int Anesthesiol Clin* 1978;16:149-182
4. Busch Eric H, Kay Dennis, Branting Scott B. *Low Volume Neurolytic Celiac Plexus Block with Computed Tomography Guidance. Anesthesiology* 2003;99(5):1243-1244
5. Jabbal SS, Hunton J. *Reversible paraplegia following celiac plexus block. Anaesthesia*. 1992;47:857-858
6. Davies DD. *Incidence of major complications of neurolytic coeliac plexus block. Journal of the Royal Society of Medicine* 1993;86(5): 264-266
7. Sett SS, Taylor DC. *Aortic pseudoaneurysm secondary to celiac plexus block. Ann Vasc Surg* 1991;5:88-91
8. Kaplan R, Schiff Keren B, Alt E. *Aortic dissection as a complication of celiac plexus block. Anesthesiology* 1995;83:632-635
9. Lee JM. *J Korean Med Sci*. 2000;15(2):173-8]
10. Lee MJ, Mueller PR, van Sonnenberg E, Dawson SL, D'Agostino H, Saini S, Cats AM. *CT-guided celiac ganglion block with alcohol. Am J Roentgenol* 1993; 161: 633-6
11. De Cicco M, Matovic M, Bortolussi R, Coran F, Fantin, Dario, Fabiani F, Caserta M. *Celiac Plexus Block: Injectate Spread and Pain Relief in Patients with Regional Anatomic Distortions. Anesthesiology*: 2001; 94(4):561-565

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