GASTRIC OUTLET OBSTRUCTION DURING BONE SCAN.

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ABSTRACT

A female patient of 40 years was diagnosed to have gastric outlet obstruction during radionuclide bone scintigraphy (Fig.1)

INTRODUCTION

Noninvasive measurement of gastric accomodation and emptying with radionuclide technetium 99 metastable (99m Tc) is feasible.¹ At present, a barostat study is the gold standard, but it is invasive and possibly induces artifacts as a result of positive intraluminal balloon pressure.

CASE REPORT

A female patient of 40 years came for whole body bone scan after left-sided mastectomy due to duct cell carcinoma and desmoplasia. Intravenous injection of 20 milli-Curies 99m Tc MDP (methylene diphosphonate) was given. Blood pool images under Siemens gamma camera were normal, but static views after 3.5 hours showed only non-osseous uptakes in right breast nodule and unusually distended stomach. Ultrasound scan by sonoline SL-2 showed a 9.9 mm hyperechoic nodule in right breast, pockets of small ascites, slow gastric emptying and lower part of esophagus was inflamed.² The patient vomitted during gamma camera scan (static view). The patient complained of anorexia and nausea, but never vomitted before.



Fig. 1A 99m Tc MDP uptake in stomach (S) and right breast (RB)

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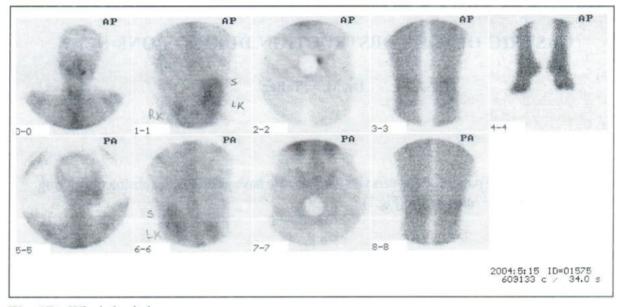


Fig. 1B Whole body bone scan

DISCUSSION

The current gold standard for the measurement of accomodation is the gastric barostat, involving the introduction of a balloon into the gastric fundus.3 In addition to the discomfort associated with this invasive and time-consuming procedure, the presence of a balloon in the stomach has been shown to cause dilatation of the antrum as a result of meal displacement and induction of exaggerated proximal gastric relaxation.4 Radionuclide studies using pertechnetate, sucralfate, DTPA and sulfur colloid particles are well- known in gastric disorders,5,6 however, MDP was not used for this purpose yet, although non-osseous uptake of MDP during threephases bone scan is a recognized fact.7 The stomach uptake must be distinguished from splenic or renal accumulation.8 The mechanism for localization of bone-scanning agents in noncalcified soft tissue, particularly damaged muscle, may be related to movement of calcium from plasma into damaged muscle cells through abnormally permeable sarcolemma.9 Other possible mechanisms of soft-tissue uptake include binding to immature collagen and atypical binding of the Tc-99m phosphate to phosphatase enzymes.10,11

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