**Description**

Each vial contains 13.25 mg of Na-DTPA i.e. sodium diethylenetriaminepentaacetate monohydrate (equivalent to 10 mg of pentetic acid), 800 micrograms (µg) of stannous chloride, di-hydrate and 7.10 mg of sodium chloride. The contents of the vial are lyophilised and sealed under nitrogen. The product is sterilised by membrane filtration.

As supplied, the product is sterile and pyrogen free.

The product contains no bactericide.

The product is designed for diagnostic use. Administration after reconstitution with sterile sodium pertechnetate 99mTc solution is by intravenous injection.

**Physical Characteristics of 99mTc**

Technetium-99m, with a physical half-life of six hours, decays by isomeric transition to technetium-99. Photons associated with this transition which are useful for detection and imaging studies are listed in Table 1. A decay chart is listed in Table 2.

**TABLE 1**

<table>
<thead>
<tr>
<th>Principal</th>
<th>Mean % per</th>
<th>Mean Energy (keV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Radiation</td>
<td>Disintegration</td>
<td></td>
</tr>
<tr>
<td>Gamma-2</td>
<td>87.2</td>
<td>140.5</td>
</tr>
</tbody>
</table>

**TABLE 2**

<table>
<thead>
<tr>
<th>Hours</th>
<th>Fraction Remaining</th>
<th>Hours</th>
<th>Fraction Remaining</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>1.000</td>
<td>5</td>
<td>0.562</td>
</tr>
<tr>
<td>1</td>
<td>0.891</td>
<td>6</td>
<td>0.501</td>
</tr>
<tr>
<td>2</td>
<td>0.794</td>
<td>7</td>
<td>0.447</td>
</tr>
<tr>
<td>3</td>
<td>0.708</td>
<td>8</td>
<td>0.398</td>
</tr>
<tr>
<td>4</td>
<td>0.631</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**External Radiation**

The specific gamma ray constant for 99mTc is 0.19mGy per MBq at 1cm. The first half-life thickness of lead for 99mTc is 0.2mm. A range of values for the relative attenuation of the radiation emitted by 99mTc resulting from the interposition of various thicknesses of lead is shown in Table 3.

**TABLE 3**

<table>
<thead>
<tr>
<th>Shield Thickness mm Pb</th>
<th>Coefficient of Attenuation (approx.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.2</td>
<td>0.5</td>
</tr>
<tr>
<td>0.95</td>
<td>0.1</td>
</tr>
<tr>
<td>1.8</td>
<td>0.01</td>
</tr>
<tr>
<td>2.7</td>
<td>0.001</td>
</tr>
<tr>
<td>3.6</td>
<td>0.0001</td>
</tr>
</tbody>
</table>
**Actions**

Following intravenous injection, $^{99m}$Tc-DTPA is rapidly distributed throughout the extracellular fluid space, where it is promptly cleared from the body by glomerular filtration. There is minimal binding to the renal parenchyma.

$^{99m}$Tc-DTPA tends to accumulate in intercranial lesions with excessive neovascularity or an altered blood-brain barrier. The agent does not accumulate in the choroid plexus.

Since the agent is excreted by glomerular filtration, the images of the kidneys obtained during the first few minutes after injection show the vascular pool within the kidney. Subsequent images represent radioactivity in both the collecting system and the renal pelvis.

**Indications**

$^{99m}$Tc-Pentastan may be used to perform kidney imaging, brain imaging, to assess renal perfusion and to estimate glomerular filtration rate.

**Contraindications**

None known.

**Warnings**

**Use During pregnancy:**

Adequate reproduction studies have not been performed in animals to determine whether this drug affects fertility in males or females, has teratogenic potential, or has other adverse affects on the foetus.

This radiopharmaceutical should not be administered to pregnant or nursing women or persons under 18 years of age unless the benefits to be gained outweigh the potential hazards. Ideally, examination of a woman of childbearing capabilities should be performed only during the 10 days following the onset of menses.

**Use During Lactation:**

Technetium ($^{99m}$Tc) is excreted in human milk. If administered to a nursing mother, formula feeding must be substituted.

**Precautions**

**General**

Radiopharmaceuticals should only be used by physicians who are qualified and licensed to handle radioactives. Contents of the kit are intended only for use in the preparation of technetium ($^{99m}$Tc) labelled DTPA. They should not be administered directly to the patient. Solutions containing Sodium Pertechnetate with antioxidants should not be used. At time of administration, the solution should be crystal clear.

**Dose Handling**

Radiation exposure to clinical personnel must be minimised. Care and appropriate safety measures should always be used. Pentastan multi-dose contains no bactericide. Aseptic procedures must be used at all times when handling the product.

**Patient Care**

Care should be taken to minimise unwanted radiation exposure to patients, consistent with proper patient management.

**Adverse Reactions**

No adverse reactions have been reported.
Dosage and Administration

Each vial contains Pentetic Acid intended for 2 doses.

Technetium (99mTc) Pentastan is prepared for clinical use as follows:

1. Using aseptic technique, add the required amount of sterile sodium pertechnetate (99mTc) solution. Use vent needle to release excess pressure.

2. Mix by shaking gently for 10 seconds.

3. Administer by I.V. injection within six hours.

It is recommended that reconstituted Pentastan be stored below 8°C, under refrigeration, and used within 6 hours of reconstitution.

The suggested dose for a normal (70kg) adult is:

- Brain Scan: 400-800 MBq
- Renal Scan: 200 MBq

For children, the dose should be reduced to that appropriate to the patient's weight.

The patient dose should be measured with a suitable radioactivity calibrator immediately before administration. Radiochemical purity should be checked prior to administration. Shielding should be used when preparing Technetium (99mTc) Pentastan.

Radiation Dosimetry

The estimated absorbed radiation dose per MBq to an average (70kg) patient from an intravenous injection of Technetium (99mTc) Pentastan is shown in Table 4.

TABLE 4
Radiation Dose Estimates (mGy. MBq⁻¹)

<table>
<thead>
<tr>
<th>Organ</th>
<th>Bladder Voided Regularly</th>
<th>Bladder not Voided</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kidneys</td>
<td>0.001</td>
<td>0.001</td>
</tr>
<tr>
<td>Bladder Wall</td>
<td>0.055</td>
<td>0.142</td>
</tr>
<tr>
<td>Whole Body</td>
<td>0.002</td>
<td>0.004</td>
</tr>
<tr>
<td>Ovaries</td>
<td>0.006</td>
<td>0.016</td>
</tr>
<tr>
<td>Testes</td>
<td>0.005</td>
<td>0.012</td>
</tr>
</tbody>
</table>

The radiation dose was calculated according to “MIRD” methods. Ref "S", Absorbed Dose per Unit Cumulated Activity for Selected Radionuclides and Organs, MIRD Pamphlet No.11, 1975.
Pentastan DTPA Reagent
Multi Dose Vials

Presentation

Pentastan Agent is supplied as a sterile, sealed under nitrogen, freeze-dried product, in 10 mL multidose vials. There are six vials to a pack.

Expiry

Expiry is 12 months from the date of manufacture. The expiry date is found on the pack.

Storage

Pentastan Agent should be stored between 2-8°C. (Refrigerate. Do not freeze).

Contact Details

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Facsimile: 02 9543 6511

ANSTO Health is a commercial enterprise of the Australian Nuclear Science and Technology Organisation (ANSTO), which is located at Lucas Heights, in Sydney, N.S.W.

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