

Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

TechneScan HDP 3 mg, kit for radiopharmaceutical preparation

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Sodium Oxidronate 3.0 mg

The radionuclide is not part of the kit.

Excipients:

Sodium 12 mg

For the full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Kit for radiopharmaceutical preparation. Powder for solution for injection.

Grey-white lyophilized powder.

4 CLINICAL PARTICULARS

4.1 Therapeutic Indications

This medicinal product is for diagnostic use only.

After radiolabelling with sodium pertechnetate (^{99m}Tc) solution, the agent may be used for bone scintigraphy, where it delineates areas of altered osteogenesis.

4.2 Posology and method of administration

Posology

Adults

The average activity administered by single intravenous injection is 500 MBq (300 - 740 MBq) in a 70 kg adult. Other activities may be justifiable. It should be noted that in each country physicians should follow the Diagnostic Reference Levels and the rules set out by local law.

Elderly population

There is no special dosage regimen for the elderly patient.

Paediatric population

The use in paediatric children and adolescents has to be considered carefully, based upon clinical needs and assessing the risk/benefit ratio in this patient group. The activities to be administered to children and to adolescents were calculated according to the EANM dosage card (2014) by using the following Formula:

$A[\text{MBq}]_{\text{Administered}} = \text{Baseline Activity} \times \text{Multiple}$ (with a baseline activity of 35.0)

The resulting activities to be administered may be found in the following table:

Weight (kg)	Activity (MBq)	Weight (kg)	Activity (MBq)	Weight (kg)	Activity (MBq)
3	40	22	185	42	320
4	40	24	200	44	335
6	60	26	215	46	350
8	75	28	225	48	360
10	95	30	240	50	375
12	110	32	255	52 - 54	395
14	125	34	270	56 - 58	420
16	140	36	280	60 - 62	445
18	155	38	295	64 - 66	470

Weight (kg)	Activity (MBq)	Weight (kg)	Activity (MBq)	Weight (kg)	Activity (MBq)
20	170	40	310	68	490

In very young children (up to 1 year) a minimum dose of 40 MBq is necessary in order to obtain images of sufficient quality.

Method of administration

According to the followed reconstitution protocol the radiolabelled preparation is either for single use or multidose use. This medicinal product should be reconstituted before administration to the patient.

For instructions on reconstitution of the medicinal product before administration, see section 12.

For patient preparation, see section 4.4.

Image acquisition

The patient should void before scanning.

Images obtained shortly after injection (e.g. in the so-called 3-phase bone scan procedure) will only partly reflect metabolic activity.

Late phase static scintigraphy should be performed not earlier than 2 hours after injection.

Image acquisition should be performed according to clinical needs and/or current international guidelines.

4.3 Contraindications

Hypersensitivity to the active substance, to any of the excipients listed in section 6.1 or to any of the components of the labelled radiopharmaceutical.

4.4 Special warnings and precautions for use

Potential for hypersensitivity or anaphylactic reactions

If hypersensitivity or anaphylactic reactions occur, the administration of the medicinal product must be discontinued immediately and intravenous treatment initiated, if necessary. To enable immediate action in emergencies, the necessary medicinal products and equipment such as endotracheal tube and ventilator must be immediately available.

Individual benefit/risk justification

For each patient, the radiation exposure must be justifiable by the likely benefit. The activity administered should in every case be as low as reasonably achievable to obtain the required diagnostic information.

Renal impairment

Careful consideration of the benefit risk ratio in these patients is required since an increased radiation exposure is possible.

Generalised increased soft tissue uptake can be due to renal failure.

Paediatric population

For information on the use in paediatric population, see section 4.2.

In infants and children particular attention should be paid to the relatively higher radiation exposure of the epiphyses in growing bone. Careful consideration of the indication is required since the effective dose per MBq is higher than in adults (see section 11).

Patient preparation

The patient should be well hydrated before the start of the examination and urged to void as often as possible during the first hours after the study in order to reduce radiation.

Specific warnings

Inadvertent or accidental subcutaneous administration of technetium (^{99m}Tc) oxidronate should be avoided as perivascular inflammation has been described.

This medicinal product contains less than 1 mmol sodium (23 mg) per dose, i.e. essentially 'sodium-free'.

Precautions with respect to environmental hazard are in section 6.6.

4.5 Interaction with other medicinal products and other forms of interactions

An increased extraossal accumulation of the radioisotope was reported for iron-containing ingredients, administration of diphosphonates, various cytostatics (vincristine, cyclophosphamide, doxorubicin, methotrexate), immunosuppressive medicinal products (e.g. cortisone), antibiotics (gentamicin, amphotericin) and aluminium-containing drugs.

Regular medication with aluminium containing drugs may lead to abnormally high accumulation of ^{99m}Tc in the liver, presumably caused by the formation of labelled colloids.

In patients with hypercalcaemia soft-tissue uptake of bone-seeking radiopharmaceuticals may be observed.

4.6 Fertility, pregnancy and lactationWomen of childbearing potential

When an administration of radiopharmaceuticals to a woman of childbearing potential is intended, it is important to determine whether or not she is pregnant. Any woman who has missed a period should be assumed to be pregnant until proven otherwise. If in doubt about her potential pregnancy (if the woman has missed a period, if the period is very irregular, etc.), alternative techniques not using ionising radiation (if there are any) should be offered to the patient.

Pregnancy

Radionuclide procedures carried out on pregnant women also involve radiation doses to the foetus. Only essential investigations should be carried out during pregnancy when likely benefit exceeds the risk incurred by mother and foetus. Administration of 740 MBq technetium (^{99m}Tc) oxidronate to a patient results in an absorbed dose to the uterus of 4.7 mGy. Doses above 5 mGy would be regarded as a potential risk for the foetus.

Breast-feeding

^{99m}Tc will be excreted into breast milk.

Before administering a radioactive medicinal product to a mother who is breast-feeding consideration should be given to the possibility of delaying the administration of radionuclide until the mother has ceased breast-feeding and to what is the most appropriate choice of radiopharmaceutical, bearing in mind the secretion of activity in breast milk. If the administration is considered necessary, breast-feeding should be interrupted for 4 hours and the expressed feeds discarded.

There is no need for restriction of contact between the patient and infants/pregnant women after leaving the hospital.

Fertility

The effect of the administration of technetium (^{99m}Tc) oxidronate on pregnant women and fertility is unknown.

4.7 Effects on ability to drive and use machines

Technescan HDP has no or negligible influence on the ability to drive and use machines.

4.8 Undesirable effects

Very common ($\geq 1/10$)
Common ($\geq 1/100$ to $< 1/10$)
Uncommon ($\geq 1/1,000$ to $< 1/100$)
Rare ($\geq 1/10,000$ to $< 1/1,000$)
Very rare ($< 1/10,000$)
Not known (cannot be estimated from the available data)

Information on adverse reactions is available from spontaneous reporting. The reported reaction types are anaphylactoid reactions, psychovegetative reactions, as well as different kinds of injection site reactions and other general disorders. Onset of symptoms may be delayed 4 to 24 hours after administration.

Anaphylactoid reactions:

Anaphylactoid reactions were reported with a wide array of symptoms ranging from mild skin reactions to anaphylactic shock, which however was only reported in isolated cases.

Psychovegetative syndrome:

Single cases of severe psychovegetative reactions like circulatory collapse or syncope have been reported, however, most of the reported vegetative effects include gastrointestinal reactions like nausea or vomiting. Other reports include vasovagal reactions like headache or dizziness.

General disorders and administration site conditions

Injection site reactions are related to extravasation of the radioactive material during the injection, and the reported reactions rank from local swelling up to cellulitis. Extended extravasation may necessitate surgical treatment.

The following table subsumes the observed reaction types and symptoms. Due to the fact that only spontaneous reports could be analysed, no frequency indications could be provided.

Adverse Reactions sorted by System Organ Class

Immune system disorders

Frequency unknown*: Anaphylactic reactions (e.g. Anaphylactic shock, loss of consciousness, cardio-respiratory arrest, hypersensitivity, angioedema, tachycardia, hypertension, dyspnoea, conjunctivitis, rhinitis and nasal congestion, dermatitis, generalised pruritus, face oedema, laryngeal oedema, tongue oedema, and other types of oedema, urticaria, erythema, rash, dysgeusia, paraesthesia, sweating increased)

Nervous system disorders

Frequency unknown*: Psychovegetative syndrome (e.g. Syncope, circulatory collapse, dizziness, headache, tachycardia, bradycardia, hypotension, tremor, vision blurred, flushing)

Gastrointestinal disorders

Frequency unknown*: Vomiting, nausea, diarrhoea, abdominal pain

General disorders and administration site conditions

Frequency unknown*: Injection site reactions (e.g. Cellulitis, inflammation, pain, erythema, swelling), chest pain, chills

* Adverse reactions derived from spontaneous reporting

Exposure to ionising radiation is linked with cancer induction and a potential for development of hereditary defects. As the effective dose is 4.2 mSv (70 kg individual) when the maximal recommended activity of 740 MBq is administered these adverse reactions are expected to occur with a low probability.

Higher doses may be justified in some clinical circumstances. In all cases it is necessary to ensure that the risks of the radiation are less than from the disease itself.

This product contains no excipients that have a recognised action or effect, or knowledge of which is important for safe and effective use of the product.

Paediatric population

Frequency, type and severity of adverse reactions in children are expected to be the same as in adults.

Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via HPRC Pharmacovigilance; Earlsfort Terrace; IRL - Dublin 2; Tel: +353 1 6764971; Fax: +353 1 6762517; Website: www.hpra.ie; e-mail: medsafety@hpra.ie.

4.9 Overdose

In the event of administration of a radiation overdose with technetium (^{99m}Tc) oxidronate the absorbed dose to the patient should be reduced where possible by increasing the elimination of the radionuclide from the body by forced diuresis and frequent bladder voiding. It might be helpful to estimate the effective dose that was applied.

5 PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Diagnostic radiopharmaceuticals skeleton, ATC code: V09B A01

At chemical concentrations of radiopharmaceuticals and excipients used for diagnostic procedures technetium (^{99m}Tc) oxidronate does not appear to exert any pharmacodynamic effect.

5.2 Pharmacokinetic properties

Distribution

Intravenously administered technetium (^{99m}Tc) oxidronate is rapidly distributed throughout the extracellular space.

Organ uptake

Skeletal uptake begins almost immediately and proceeds rapidly. 30 minutes post injection 10 % of the initial dose is still present in whole blood. At 1 hour, 2 hours, 3 hours and 4 hours after injection these values are resp. 5 %, 3 %, 1.5 % and 1 %.

Elimination

Clearance from the body takes place via the kidneys. Of the administered activity approximately 30 % is cleared within the first hour, 48 % within two hours and 60 % within 6 hours.

5.3 Preclinical safety data

This agent is not intended for regular or continuous administration. Reproduction, mutagenicity studies and long-term carcinogenicity studies have not been carried out. Minimal liver abnormalities are seen at the level of 30 mg/kg in rats. In subacute toxicity studies rats do not react to the administration of 10 mg/kg/day for 14 days, dogs show histological changes in the liver (microgranuloma) after 3 and 10 mg/kg/day for 14 days. In dogs, which were treated for 14 consecutive days, long-lasting indurations at the site of injection were observed.

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Gentisic acid
Sodium chloride
Stannous chloride dihydrate
Hydrochloric acid
Sodium Hydroxide

6.2 Incompatibilities

This medicinal product must not be mixed with other medicinal products except those mentioned in section 12.

6.3 Shelf life

24 months

After reconstitution and labelling chemical and physical in-use stability has been demonstrated for 8 hours at 25°C. From a microbiological point of view, the product should be used immediately. If not used immediately, in use storage times and conditions prior to use are the responsibility of the user and would normally not be longer than 24 hours at 2 to 8°C, unless reconstitution / dilution (etc.) has taken place in controlled and validated aseptic conditions.

6.4 Special precautions for storage

Lyophilised powder: Do not store above 25°C.

Store in the original container in order to protect from light.

For storage conditions after radiolabelling of the medicinal product, see section 6.3.

Storage of radiopharmaceuticals should be in accordance with national regulation on radioactive materials.

6.5 Nature and contents of container

Colourless glass vial with grey stopper and blue crimp cap containing a grey white powder.

Techescan HDP is supplied as five vials in a carton box. 10 ml glass vial closed with a bromobutyl rubber stopper sealed with an aluminium crimp cap.

6.6 Special precautions for disposal and other handling

Any unused product or waste material should be disposed of in accordance with local requirements for radioactive materials

General warning

Radiopharmaceuticals should be received, used and administered only by authorised persons in designated clinical settings. Their receipt, storage, use, transfer and disposal are subject to the regulations and/or appropriate licences of the competent official organisation.

Radiopharmaceuticals should be prepared in a manner which satisfies both radiation safety and pharmaceutical quality requirements. Appropriate aseptic precautions should be taken.

Contents of the vial are intended only for use in the preparation of technetium (^{99m}Tc) oxidronate injection and are not to be administered directly to the patient without first undergoing the preparative procedure.

For instructions on reconstitution of the medicinal product before administration, see sections 12.

If at any time in the preparation of this product the integrity of this vial is compromised it should not be used.

Administration procedures should be carried out in a way to minimise risk of contamination of the medicinal product and irradiation of the operators. Adequate shielding is mandatory.

The content of the kit before extemporary preparation is not radioactive. However, after sodium pertechnetate (^{99m}Tc) is added, adequate shielding of the final preparation must be maintained.

The administration of radiopharmaceuticals creates risks for other persons from external radiation or contamination from spills of urine, vomit, etc. Radiation protection precautions in accordance with national regulations must therefore be taken.

Any unused medicinal product or waste material should be disposed of in accordance with local requirements for radioactive materials.

7 MARKETING AUTHORISATION HOLDER

Curium Netherlands B.V.
Westerduinweg 3
1755 LE Petten
The Netherlands

8 MARKETING AUTHORISATION NUMBER

PA0690/004/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 19 November 1993

Date of last renewal: 19 November 2008

10 DATE OF REVISION OF THE TEXT

August 2021

11 DOSIMETRY

Technetium (^{99m}Tc) is produced by means of a ($^{99}\text{Mo}/^{99m}\text{Tc}$) generator and decays with the emission of gamma radiation with an energy of 140 keV and a half-life of 6 hours to (^{99}Tc) Technetium which, in view of its long half-life of 2.13×10^5 years, can be regarded as quasi stable.

The dosimetry data listed below are from ICRP publication 80.

Absorbed doses: ^{99m}Tc-labelled phosphates and phosphonates (absorbed dose per unit activity administered [mGy/MBq]):

Organ	Adult	15 years	10 years	5 years	1 year
Adrenals	0.0021	0.0027	0.0039	0.0058	0.011
Bladder wall	0.048	0.06	0.088	0.073	0.13
Bone surface	0.063	0.082	0.13	0.22	0.53
Brain	0.0017	0.0021	0.0028	0.0043	0.0061
Breast	0.00071	0.00089	0.0014	0.0022	0.0042
Gall bladder	0.0014	0.0019	0.0035	0.0042	0.0067
Stomach wall	0.0012	0.0015	0.0025	0.0035	0.0066
Small intestine	0.0023	0.0029	0.0044	0.0053	0.0095
Colon	0.0027	0.0034	0.0053	0.0061	0.011
Upper large intestine	0.0019	0.0024	0.0039	0.0051	0.0089
Lower large intestine	0.0038	0.0047	0.0072	0.0075	0.013
Heart	0.0012	0.0016	0.0023	0.0034	0.006
Kidneys	0.0073	0.0088	0.012	0.018	0.032
Liver	0.0012	0.0016	0.0025	0.0036	0.0066
Lungs	0.0013	0.0016	0.0024	0.0036	0.0068
Muscles	0.0019	0.0023	0.0034	0.0044	0.0079
Oesophagus	0.001	0.0013	0.0019	0.003	0.0053
Ovaries	0.0036	0.0046	0.0066	0.007	0.012
Pancreas	0.0016	0.002	0.0031	0.0045	0.0082
Red marrow	0.0092	0.01	0.017	0.033	0.067
Skin	0.001	0.0013	0.002	0.0029	0.0055
Spleen	0.0014	0.0018	0.0028	0.0045	0.0079
Testes	0.0024	0.0033	0.0055	0.0058	0.011
Thymus	0.001	0.0013	0.0019	0.003	0.0053
Thyroid	0.0013	0.0016	0.0023	0.0035	0.0056
Uterus	0.0063	0.0076	0.012	0.011	0.018
Other tissue	0.0019	0.0023	0.0034	0.0045	0.0079
Effective dose [mSv/MBq]	0.0057	0.007	0.011	0.014	0.027

The effective dose resulting from the administration of 740 MBq for an adult weighing 70 kg is 4.2 mSv. For an administered activity of 740 MBq the typical radiation dose to the target organ (bone) is 47 mGy and the typical radiation dose to the critical organ (bladder wall) is 36 mGy.

12 INSTRUCTIONS FOR PREPARATION OF RADIOPHARMACEUTICALS

As with any pharmaceutical product, if at any time in the preparation of this product the integrity of this vial is compromised it should not be used.

Method of preparation

Preparation for multi-dose application

Aseptically add the required amount of sodium pertechnetate (^{99m}Tc) Injection (Fission or Non-Fission) with a maximum activity of 14 GBq, in a volume of 3-10 ml to one vial Technescan HDP.

Shake for 30 seconds to dissolve the contents. The preparation is then ready for injection.

Dilution should preferably be done with Sodium Chloride 0.9 % solution.

For a single patient at most 1 mg of HDP (1/3 of a vial) may be injected

Properties after labelling

After labelling the solution is colourless and clear to slightly opalescent.

Quality control

Examine by TLC on silica gel coated glass-fibre sheets.

1. Develop 5 to 10 microliter in 13.6% sodium acetate R. Technetium oxidronate complex and pertechnetate ion migrate near the solvent front, hydrolysed technetium and technetium in colloidal form remain at the start.
2. Develop 5 to 10 microliter in methyl ethyl ketone R. Pertechnetate ion migrates near the solvent front, technetium oxidronate complex and technetium in colloidal form remain at the start. For particulars consult the European Pharmacopoeia (Monograph 0641).

Individual and the total impurities may not be more than 5 %.

Radiochemical purity

A. Impurity A. Thin-layer chromatography.

Test solution. The preparation to be examined.

Plate: TLC silica gel plate R; use silica gel as the coating substance on a glass-fibre sheet.

Mobile phase: 136 g/L solution of sodium acetate R.

Application: 5-10 microliter.

Development: immediately, over a path of 10-15 cm in about 10 min.

Drying: in air.

Detection: suitable detector to determine the distribution of radioactivity.

Retardation factors: impurity A = 0.0 to 0.1; [^{99m}Tc]technetium oxidronate and impurity B = 0.9 to 1.0.

B. Impurity B. Thin-layer chromatography.

Test solution. The preparation to be examined.

Plate: TLC silica gel plate R; use silica gel as the coating substance on a glass-fibre sheet.

Mobile phase: methyl ethyl ketone R.

Application: 5-10 microliter; dry quickly.

Development: over a path of 10-15 cm in about 10 min.

Drying: in air.

Detection: suitable detector to determine the distribution of radioactivity.

Retardation factors: [^{99m}Tc]technetium oxidronate and impurity A = 0.0 to 0.1; impurity B = 0.9 to 1.0.

Limits:

sum of impurities A and B: maximum 5.0 per cent of the radioactivity due to technetium-99m in the chromatograms obtained in tests A and B.