

**RADIOPAQUE DIAGNOSTIC AGENTS****Gaurav Patidar\*, Hemant Saini, Kuldeep Vinchurkar and Dr. Dinesh Kumar Mishra**

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**ABSTRACT**

This exploratory-descriptive, non-experimental quantitative studies aimed to study on the spot negative reactions to intravenous iodinated comparison media in the sufferers. The houses of a tissue extracellular area, specifically the microvascular thing of that space can be characterised by the usage of x-ray diagnostic modalities along with particulate assessment marketers. The gift invention affords a way for characterizing a assets of a tissue extravascular space using radiopaque particulate contrast. Moreover, the existing invention affords a novel elegance of particulate retailers suitable for use along side the methods provided herein. Iodinated radiopaque polymeric nanoparticles of sizes ranging among 30 and 350 nm were formed by way of emulsion polymerization of the monomer 2-methacryloyloxyethyl(2,3,5-

triiodobenzoate) in the presence of sodium dodecyl sulfate as surfactant and potassium persulfate as initiator. The impact of numerous polymerization parameters, e.G., monomer, initiator and surfactant concentrations on the molecular weight, polymerization yield, length and size distribution of the particles became elucidated. These polymeric nanoparticles are composed of ca. 58% with the aid of weight iodine, and are therefore predicted to own huge radiopaque nature. In vitro radiopacity of the iodinated nanoparticles of  $30.6 \pm 5$ . Zero nm diameter, dispersed in water and inside the dry kingdom, turned into confirmed with a CT scanner. In vivo CT-imaging carried out in a canine model by using intravenous administration of the uniform  $30.6 \pm 5$ . Zero nm diameter radiopaque nanoparticles dispersed in saline proven sizeable superior visibility of lymph nodes, liver, kidney and spleen. These outcomes indicate that those nanoparticles may be beneficial as new efficient evaluation agents for X-ray imaging programs.

**KEYWORDS:** Radiopaque nanoparticles, Emulsionpolymerization, Radiopacity, X-ray imaging.

## INTRODUCTION

Radiopaque retailers are tablets used to assist diagnose sure clinical problems. They contain iodine, which absorbs x-rays. Depending on how they may be given, radiopaque shops growth in a selected location of the body. The resulting immoderate degree of iodine lets in the x-rays to make a "image" of the place.

Radiopaque retailers are taken via mouth or given with the resource of enema or injection. X-rays are then used to check if there are any problems with the stomach, intestines, kidneys, or other elements of the frame.

Some radiopaque retailers, along with iohexol, iopamidol, and metrizamide are given by the use of injection into the spinal canal. X-rays are then used to assist diagnose problems or illnesses within the head, spinal canal, and nervous device.

The doses of radiopaque dealers can be unique for extraordinary patients and depend on the sort of take a look at. The energy of the solution is determined by how plenty iodine it incorporates. Different tests will require a selected energy and amount of solution depending on the age of the affected character, the evaluation wanted, and the x-ray tool used.

A catheter or syringe is used to location the answer of the radiopaque agent into the bladder or ureters to assist diagnose issues or diseases of the kidneys or other areas of the urinary tract. It can also be located into the uterus and fallopian tubes to help diagnose troubles or illness of those organs. After the take a look at is performed, the affected person expels most of the answer by means of urinating (after bladder or ureter research) or from the vagina (after uterine or fallopian tube research).<sup>[1]</sup>

## 1. Basic Principle

The use of radiographic contrast retailers dates almost from the invention of X-Rays. In 1896 Becheropacified the gastrointestinal tract of the guinea pig using lead subacetate.

The primary precept is that comparison marketers are administered in this type of manner as to adjust the absorption of x-rays with the aid of precise anatomic systems when it comes to

their surroundings. Contrast markers may be fantastic (iodine or barium compounds, for instance), or bad (gases).

Gaseous comparison markers take in fewer x-rays than tissues because of their low density, even though the effective atomic number can be higher. On the other hand, effective comparison markers take in more x-rays than tissues due to their excessive density and higher atomic range.<sup>[2]</sup>

## **2. Classification**

### **2.1 Iodine Compounds as Contrast Agents**

Almost all radiological examinations finished with injected comparison markers involve the administration of iodine-containing compounds.

The use of iodine compounds became first of all related to low toxicity and wonderful radio-opacity instead of physical considerations. However, it became also lucky that iodine compounds own bodily properties which lead them to higher evaluation agents than compounds with better atomic variety.

The K-fringe of iodine is 33.2 keV. At photon energies barely above this, iodine really has greater attenuating properties for x-rays than the same mass of lead.

The location of this K-area for iodine has sensible implications. Obviously the maximum contrast in radiographic studies using iodine compounds would be received through the use of a monochromatic beam of radiation of energy simply above 33.2 keV. This isn't always practicable, however, but what may be finished is to choose the height kilovoltage (kVp) for the exam on the way to provide a high proportion of the photons inside the 33-forty keV range. This is completed when a particularly low kVp is used.

Therefore, for contrast examinations the use of iodine compounds, top-quality approach requires a kVp inside the 60-eighty variety.<sup>[7]</sup>

#### **2.1.2 Ionic Water Soluble Agents Handled By Renal Excretion**

These are sodium or methylglucamine salts of 2,4,6 tri-iodobenzoic acid:

Positions three,5 are substituted with side chains which growth solubility and reduce toxicity.

Contrast media of this kind consist of:

- Diatrizoate (Urografin, Angiografín)

- Iothalamate (Conray)
- Metrizoate (Isopaque)

These compounds range best inside the aspect chains at positions three, 5. All of these dealers dissociate in method to produce ionic particles for every three iodine atoms. They are consequently called RATIO 1.5 AGENTS. Commercial preparations of those agents may be pure methylglucamine or pure sodium compounds, or a aggregate of both.

A exceedingly current addition to this organization of ionic retailers is a mixture of the sodium and methylglucamine salts of ioxaglic acid (Hexabrix).

### 2.1.2 Non-Ionic Contrast Agents

The look for evaluation agents with appropriate radio-opacity, low toxicity, low osmolality and suitability for intrathecal injection brought about metrizamide as the first agent of this group. Metrizamide is a glucosamidederivative of metrizoic acid. It is now not commercially available in Australia.

Non-ionic contrast dealers are now available; iopamidol, iohexol, ioversol, iopromide, iodixanol and iotrolan. Like Hexabrix, the primary four are RATIO 3 AGENTS however they do not dissociate in answer.

Iotrolan and iodixanol are non-ionic dimers and RATIO 6 AGENTS.

These compounds are appropriate for intrathecal use and have nearly entirely changed oily myelographicagents (observe that ioversol and iopromide have now not but been accepted for intrathecal use in Australia).

Another gain of those compounds is they seem to have much less toxicity than RATIO 1.5.

AGENTS, although it will take a few years of considerable utilization to verify this. Their essential downside is they value approximately 4-5 times as lots as RATIO 1.5 AGENTS.

Some hospitals have made the selection to apply non-ionic evaluation dealers most effective, because of their greater apparent safety. Other hospitals use these retailers selectively and the major indicators are:

- (i) Intrathecal examinations.

(ii) Examinations wherein higher osmolality markers may be painful or might also have a better chance of organ damage, e.g. Peripheral arteriography, cerebral and coronary angiography.

(iii) Patients at a high risk for damaging reactions from traditional media.(eight)

### 2.1.3 Intravenous Cholangiographic Agents

The best intravenous cholangiographic agent currently available in Australia is Biliscopin (meglumine iotroxate). This is a meglumine salt dimer of triiodobenzoic acid. An unsubstituted 5 role at the benzene ring mediates reversible binding to plasma proteins and guarantees biliary excretion without conjugation. In the blood, biliary assessment markers are preferentially sure to albumin and are consequently difficult to only restrain renal excretion.

The degree of bile duct opacification is to a massive extent determined with the aid of the plasma concentration of the contrast medium. A slow infusion allows to optimize the biliary excretion and additionally reduces adverse effects.

The common dose of Biliscopin is 100ml (105mg meglumine iotroxate/ml, 5.0g Iodine). This is administered slowly over a duration of 30 minutes to 60 mins through drip infusion or by infusion pump.

Optimal biliary opacification happens from 20 mins to ninety minutes following the end of the infusion.

Patients with mildly impaired liver characteristics may additionally excrete sufficient contrast to opacify the biliary duct device. When the bilirubin level exceeds 1.2mg%, the opacification effect is terrible.

Side outcomes are much like the ones of different iodinated contrast media but arise with more frequency.

This is thought to be because of the protein-binding characteristics required for biliary excretion.(nine).

#### 2.1.4 Oral Cholecystographic Agents

These are now not commercially available and are of historic interest most effective. They are derivatives of tri-iodobenzoic acid. The five position on the benzene ring isn't substituted. This is an essential structural characteristic determining biliary excretion. The molecules include hydrophilic and lipophilic corporations to permit biliary absorption and hepatic excretion.

#### 2.1.5 Oily Contrast Agents

These marketers are now not commercially available in Australia. They include Iophendylate (Myodil, Pantopaque), a myelographic agent and Lipiodrol Ultrafluide (Ethiodol), a lymphangiographic agent.

### 2.2 Barium Sulphate

Barium sulphate is derived from the mineral barytes. In aqueous suspension it has been the usual contrast agent for opacification of the gastro-intestinal tract, almost because the beginning of radiology. Barium has a K-part at 37.4keV. Therefore, a quite low kVp radiographic method could seem applicable however, in lots of cases, barium compounds are utilized in pretty high attention in the alimentary tract and the need for that is dubious. It could be of significance, however, in air contrast techniques in which only a thin coating of barium is used to define viscera. The many proprietary arrangements of barium sulphate which have appeared on the market over a few years differ of their components such as suspending dealers and many others.

### 2.3 Magnetic Resonance Imaging Contrast Agents

#### 2.3.1 Paramagnetic Contrast Agents

The magnetic dipole moments of paramagnetic materials are randomly aligned within the absence of an external magnetic area. When an outside magnetic area is gift, however, the magnetic moments align with the field and set off robust nearby magnetic fields that shorten-the T1 and T2 rest instances of adjacent protons. When the outside area is eliminated, the magnetic dipole moments of paramagnetic substances once more end up randomly aligned in order that there's no retained magnetization.

Paramagnetic substances have one or more debris (protons, neutrons or electrons) with a spin that is not cancelled via some other similar particle with an contrary spin. Magnetic dipole moments of unpaired electrons are very plenty larger than those of protons or neutrons, in

order that the neighborhood magnetic fields generated by way of unpaired electrons are very strong. Therefore, materials which have unpaired electrons, including the transitional factors, are very powerful paramagnetic contrast enhancers.

When paramagnetic ions are introduced to water, the relaxation of water molecules is more advantageous within the place of the paramagnetic substance. Both T1 and T2 relaxation instances are decreased.

It ought to be referred to that while iodine-containing contrast marketers in radiography immediately have an effect on film density with the aid of photon absorption, MRI comparison sellers have an indirect effect. It isn't always the actual contrast agent which alters the intensity of the picture but the presence of the comparison agent alters the rest characteristics of adjacent protons, for that reason in a roundabout way affecting the depth.

There are commercially to be had MRI evaluation dealers at present in Australia; viz, gadopentetatedimeglumine (Magnevist) and gadodiamide (Omniscan). In answer, the former substance dissociates.

Into two methylglucaminecations and one anion, containing a gadolinium atom chelated with diethylenetriaminepentaacetate.

On the opposite hand, gadodiamide is a non-ionic agent which does not dissociate in solution. In this molecule, gadolinium is chelated with diethylenetriaminepentaacetic acid bismethylamide.

### **2.3.2 Superparamagnetic Contrast Agents**

When placed in an external magnetic discipline, paramagnetic materials induce in addition magnetization that is immediately proportional to the electricity of the applied area. Increasing the external field increases the internet magnetization in a linear style. When there is no external magnetic subject, there may be no net magnetization.

Superparamagnetic materials, but, result in very sturdy magnetization in an outside magnetic subject, several orders of magnitude higher than paramagnetic marketers and in a non-linear style. There is, however, no magnetization within the absence of an outside area. It is this property which distinguishes superparamagnetic substances from ferromagnetic materials which preserve magnetization while there's no longer an external field. Particulate iron much

less than three hundred Angstrom devices in size is superparamagnetic; larger iron debris are ferromagnetic. In the latter, there's magnetic ordering of the unpaired electron spins of the iron atoms in areas which can be referred to as domains. When an external magnetic field is applied, those domain names, which have been previously randomly oriented, line up with the magnetic discipline and greatly beautify it. When the magnetic field is eliminated, this orientation stays. This is the assets of remanence that is the cardinal function of ferromagnetism. The particle size in superparamagnetic substances approximately equates to the size of a domain. When the external area is removed, the particles emerge as randomly orientated again and there may be no remanence.

Whereas paramagnetic substances are used diagnostically to increase proton sign, superparamagnetic or ferromagnetic substances spoil the signal, generating negative assessment.

Ferrites in particulate form were used as superparamagnetic evaluation dealers. Ferrites are iron oxides of the overall system  $\text{Fe}_2\text{O}_3\cdot\text{MO}$ , in which M is a divalent steel ion. Magnetite is a kind of ferrite, happening clearly, wherein the metal ion is  $\text{Fe}^{++}$ .

Particulate ferrites are taken up by way of the reticuloendothelial system. These assessment retailers gather in Kupffer cells in the liver and ruin the signal from normal liver while metastatic lesions maintain a signal as they do now not accumulate the contrast agent.

Particulate iron assessment sellers do have side consequences and aren't but to be had for scientific use.

## 2.4 Ultrasound Contrast Agents

Air bubbles were employed in echocardiography for over 30 years to transiently beautify the ventricular chambers and remarkable vessels. Initially "hand-made" ultrasound assessment retailers were created by way of agitating a saline answer previous to intravascular injection. More these days, a group of marketers comprising stabilized micro bubbles has been developed which behave as blood pool agents. These micro bubbles usually have a radius of 1 - 10  $\mu\text{m}$  and resonate while insolated with a 1 – 10MHz ultrasound pulse as is typically used in diagnostic imaging.

Coated by way of a variety of substances e.G. Galactose micro debris (Echovist, Levovist) or albumin (Infuson, Albunex), a number of the micro bubbles are sufficiently small and stable

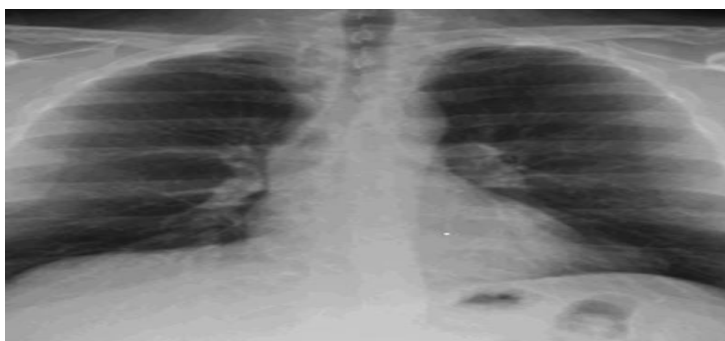


to traverse the pulmonary movement intact. The marketers flow into in the blood pool and result in a ten to twenty-five dB enhancement of echoes from flowing blood. The dealers commonly have an effect inside 20 – 30 seconds following intravenous administration and the consequences last for a couple of minutes. Side effects and toxicity are minimum with contemporary agents.

The best agent presently to be had in Australia is Levovist. This is an aqueous suspension of micro particles which include 99.9% galactose and 0.1% palmitic acid. After injection, the micro particles dissolve hastily, thereby releasing tiny air bubbles comparable in length to purple blood cells. The palmitic acid forms a defensive coating across the air bubbles, allowing a number of them to traverse the pulmonary flow intact. The most effective contraindication to Levovist is galactossaemia.(eleven).<sup>[3-7]</sup>

### 3. Opacity and Contrast

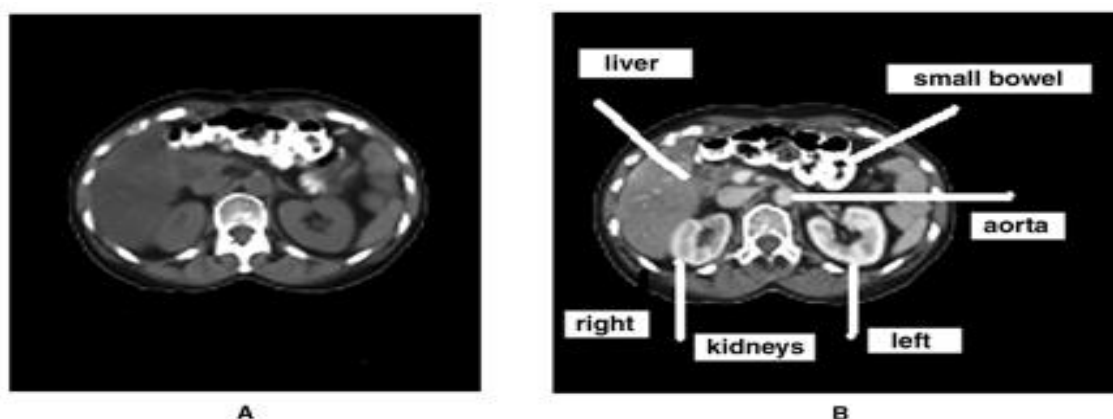
Darkening of the radiographic picture occurs at factors that correspond with interactions with x-rays. Radio density, a relative idea, describes the degree of film darkening. The photo seems lighter at points related to fewer collisions. Radiolucency and radiopacity correspond to relative densities of the object being imaged. Darker elements are radiolucent, and lighter aspects are radiopaque. For example, a dark-performing element of the picture is radiolucent relative to a mild-performing component (**Fig. 1**). Variations of radiolucency and radiopacity create photo assessment. Shading versions, which range from white to black (mild to dark), shape a seen analogue a photograph of anatomic points of interest. Without picture evaluation, there is no discernible structure.



**Fig 1: Chest radiograph.**

The bones, thoracic vascular structures, mediastinal areas (along with the coronary heart), and, within the decrease part of the photograph, the hemidiaphragms are particularly

radiopaque (light). The lungs and intercostal spaces are radiolucent (dark) The number one x-ray beam is directed from an outside supply to a material goal, which it penetrates in accordance with sure physical standards. The remnant beam describes the circulation of x-rays that emerge after interacting with the target material. It bureaucracy through the procedure of attenuation by using which quantities of the number one beam are removed. The resultant radiographic photo is an analogue of the attenuation sample of the remnant beam. In vivo, a radiopaque drug alters the attenuation sample as it blocks or slows the passage of x-rays and effects in a superb contrast study (expanded opacification of applicable portions of the picture) (**Fig.2**). The x-ray photograph is, in fact, a shadowgram, from time to time called a skiagram (from the Greek skia, that means shadow). Because the goal is improved radiographic visualization of tissues, radiopaque tablets are often known as radiocontrast tablets, marketers, or media. Negative contrast agents, consisting of air or carbon dioxide, are penetrated without problems by means of x-rays and selectively decorate picture radiolucency. Juxtaposition of superb and terrible image contrast is referred to as double assessment. For example, marked development of image formation can end result while the lumen of the colon is lined with a radiopaque drug, such as barium sulfate suspension (fantastic assessment), mixed with the creation of air (bad contrast). If a single mode of image assessment is hired, it's far a unmarried-contrast examine.



**Fig. 2: Shadowgram of Liver, Kidney and Aorta.**

Computed tomographic images of the abdomen showing opacification of the gastrointestinal tract by intraluminal presence of barium sulfate. Intravenous iodinated radiopaque drug has not been given in image A but has been given in image B and accounts for the relative opacification of the kidneys, aorta, and liver in the latter. Both radiopaque drugs elicit

positive contrast. Thus, the images do not illustrate a double-contrast study, in which positive and negative contrast occur. **Fig.3,4.**



**Fig. 3: Abdomen.**



**Fig. 4: Gastrointestinal Tract.**

#### 4. Image Formation

For conventional radiography, fluoroscopy, and CT, the remnant beam is first converted to seen mild. Intensifying displays use compounds referred to as phosphors (e.G., positive barium and gadolinium compounds) that take in higher-electricity x-rays and emit a more wide variety of lower-energy rays of seen light (fluorescence). For the image formed on a video display screen (CT and fluoroscopy), the significance of fluorescence appears obvious. Less apparent is its significance for the image formed on movie. Film is partially composed of an emulsion layer that uniformly contains many tiny silver halide crystals. The collision of photons of seen mild with any such crystals reduces cationic silver to metal silver. Many such photon–crystal reactions cause the blending of steel silver, growing a potential image. At this stage, the image continues to be invisible and should be rendered seen thru a multistep process of chemical reactions. Because the photon of visible light correctly activates the crystals, the multiplication of photons via fluorescence is nice.

The traditional radiographic photo is formed on movie; it's far a negative picture (i.E., radiolucent structures seem dark instead of light). With mild pictures, the bad image is transformed to a fine photograph, which corresponds to our notion of the object. It may appear paradoxical that the x-ray bad photograph depicts bone, as an example, as radiopaque, presenting it as a mild color as might a traditional photo (Figure 11). The reason behind this is that with x-ray imaging, bone tends to save you the x-ray photon from contributing to the picture, resulting in radiopacity. With light pictures, light is meditated from the item, which

renders the film radiolucent. Positive and poor pics should no longer be confused with tremendous and bad assessment.<sup>[14]</sup>

## 5. X-ray Generation

Not in contrast to a light bulb, the x-ray tube is a sealed, evacuated glass box with specialized electric circuitry. The tube houses a negatively charged electrode (the cathode) that features a filament. Facing the cathode is a positively charged electrode (the anode). The software of contemporary heats the steel filament to a white-warm nation (incandescence), and, via the process of thermionic emission, electrons are released from its floor. Functionally, this circuit submits a precise float of electrons to the cathode filament. Electron drift (modern) is the measure of electrons passing a given factor in step with unit of time. The ampere is the unit of cutting-edge, and the adopted unit for diagnostic radiography is the milliampere. By interface with every other circuit, a strong ability distinction (voltage) is carried out, which hurries up the cathode-generated electrons in the direction of excessive-speed collision with the tungsten- or molybdenum-containing target place of the definitely charged anode. A tube voltage of 100 kV, inside the variety of diagnostic radiography, propels electrons as much as 1/2 the rate of mild on the point of collision with the anode.

Braking radiation, additionally referred to as bremsstrahlung, is the main source of x-rays and is fashioned by the collision—with abrupt deceleration and deflection—of cathode-emitted electrons with positively charged anode atomic nuclei. X-ray photons are subsequently emitted with power equal to the diminution of kinetic power (speed) of the electron. Braking radiation bureaucracy a non-stop spectrum of energy. Another shape of x-ray era—characteristic radiation—is due to the interplay of cathode-generated electrons with inner-shell (on the whole K-shell) anode orbital electrons. It is discontinuous, forming spikes of photon emission in a specific, identifiable pattern related to the atomic range of the anode. Both forms of x-radiation generate the x-ray beam fashioned by the tube (primary beam), which is modified into the picture-forming beam (remnant beam) through interaction with the goal. A beam composed of a single strength (i.E., wavelength) is monochromatic. Current carried out to the filament is immediately proportional to the quantity of x-ray photons produced in line with unit of time (intensity) but has no impact on photon energy. On the alternative hand, photon electricity and depth range consistent with voltage input. In trendy, the number one beam spreads out over its path, and its intensity diminishes in inverse percentage to the square of distance traveled. Because the x-ray tube operates with alternating

modern-day, the height energy of cathode-generated electrons happens cyclically consistent with the frequency of the modern. Logically, the energy of a photon produced on the anode cannot exceed the peak strength of cathode-generated electrons.

### **6. Attenuation: Interaction of X-rays with Tissue and Radiopaque Drugs**

The primary x-ray beam is changed by means of interplay with a tissue goal. The following three consequences can occur for each photon:

1. No interplay. The photon completely penetrates the goal without deviation of strength or trajectory.
2. The photoelectric interaction. All the incident photon's strength is absorbed by using a goal (tissue) atom, in which case the photon ceases to exist. The excited atom ejects an electron (photoelectron) and is left in an ionized kingdom, however the photoelectron is not able to emerge from the target.
3. Compton scattering. A part of the incident photon's power is absorbed with the aid of an electron, a lower-energy photon maintains via a deflected direction (i.e., scatters), and the electron is ejected from the atom (ionization) however isn't able to emerge from the goal. With diagnostic radiography, scattered photons have a tendency to maintain within the forward route and reduce the first-rate of the photograph through darkening or fogging. The photoelectric interaction does now not produce scatter radiation due to the fact the incident photon is dissipated. Various fabric-precise coefficients of attenuation quantify the discount in intensity of the x-ray beam due to interactions with depend. Intensity is a measurement of the quantity of photons transmitted through the beam in step with unit of time and, within the case of visible light, connotes brightness. The significance of attenuation is inversely proportional to photon power. Stated some other way, a more energetic photon has a tendency to traverse farther the goal earlier than interacting with it. Beam quality describes the ability of the x-ray beam to penetrate count number.<sup>[8-12]</sup>

### **7. Adverse Drug Reaction**

Along with its wanted consequences, radiopaque marketers can purpose critical facet consequences including allergic reactions. These consequences may also occur almost straight away or a couple of minutes after the radiopaque agent are given. Although these severe aspect consequences appear most effective hardly ever, your health care professional can be prepared to present you on the spot clinical interest if wanted. If you have got any questions about this, test together with your medical doctor.<sup>[13]</sup>

Check with your doctor right away if any of the subsequent side effects occur:

**With injection into the spinal canal**

**Rare**

- Hallucinations (seeing hearing, or feeling matters that are not there)
- Paralysis of one facet of frame or of arms and legs

**For patients receiving gadolinium-based totally evaluation sellers (GBCAs)**

**Incidence no longer regarded**

- Burning or itching of the skin
- Joint stiffness
- Restrained variety of movement within the hands, fingers, legs, or ft
- Muscle weak point
- Pain deep within the hip bone or ribs
- Reddened or darkened patches at the skin
- Pores and skin swelling, hardening and/or tightening
- Yellow raised spots at the whites of the eyes

Some facet consequences may additionally occur that commonly do now not want medical attention. These aspect effects may work away all through remedy as your frame adjusts to the medicine. Also, your health care professional may be capable to inform you about approaches to save you or reduce some of those facet consequences. Check with your fitness care expert if any of the following side outcomes preserve or are bothersome or if you have any questions on them:<sup>[14-16]</sup>

**With oral or rectal use**

**Less commonplace**

- Diarrhea or laxative effect

**With injection into a vein or an artery**

**More commonplace**

- Unusual warmth and flushing of pores and skin

**Less not unusual**

- Chills
- Dizziness or lightheadedness

- Headache
- Nausea or vomiting
- Ache or burning at the place of injection
- Sweating
- Unusual or steel flavor
- Unusual thirst

### With injection into the spinal canal

#### More common

- Backache
- Dizziness
- Headache (mild to mild)
- Nausea and vomiting (slight to moderate)
- Stiffness of neck

#### Less not unusual or rare

- Difficult urination
- Drowsiness
- Headache (severe)
- Multiplied sensitivity of eyes to light
- Multiplied sweating
- Loss of appetite
- Ringing or buzzing in ears
- Unusual tiredness or weak point

### 8. Marketed Examples

Category	Generic name	Brand name	Manufacturer
<b>Ionic</b>	Diatrizoate	Gastrografin	Therapex
	Iothalamate	Conray	Liebel-Flarsheim Company
<b>Nonionic</b>	Iodixanol	Visipaque	GE Healthcare
	Iohexol	Omnipaque	GE Healthcare
	Iopamidol	Isovue	BIPSO GmbH
	Iopromide	Ultravist	Bayer Healthcare
	Ioversol	Optiray	Liebel-Flarsheim Company
	Ioxilan	Oxilan	Guerbet LLC

## CONCLUSION

Radiopaque drugs can interact with x-rays to enhance radiographic visualization of tissues and anatomical structures. The present invention offers a technique for characterizing a assets of a tissue extravascular area making use of radiopaque particulate evaluation. In vitro radiopacity of the iodinated nanoparticles of  $30.6 \pm 5$ . Zero nm diameter, dispersed in water and within the dry kingdom, was tested with a CT scanner. In vivo CT-imaging performed in a canine version via intravenous management of the uniform  $30.6 \pm 5.0$  nm diameter radiopaque nanoparticles dispersed in saline confirmed great enhanced visibility of lymph nodes, liver, kidney and spleen. These consequences indicate that these nanoparticles can be beneficial as new efficient evaluation dealers for X-ray imaging applications.

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