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Non-invasive diagnostic imaging technology for mitochondria using radiolabeled lipophilic salts

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(54) Title: NON-INVASIVE DIAGNOSTIC IMAGING TECHNOLOGY FOR MITOCHONDRIA USING RADIOLABELED LIPOPHILIC SALTS

(57) Abstract: The invention provides a series of lipophilic phosphonium cations (PhCs) labeled with ¹⁸F for non-invasive assessment of $\Delta\Psi_m$, lipophilic ammonium cation analogs of the PhCs, and methods of using same for imaging and detection of mitochondrial-related pathologies in patients using PET or SPECT.

NON-INVASIVE DIAGNOSTIC IMAGING TECHNOLOGY FOR MITOCHONDRIA DYSFUNCTION USING RADIOLABELED LIPOPHILIC SALTS

5 This application claims the benefit of U.S. Provisional Application Serial No. 60/354,563
filed February 6, 2002, the teachings of which are incorporated herein by reference.

This invention was supported by National Institute of Health (NIH) Grant No. CA92871.
The United States government has certain rights to the invention.

10 BACKGROUND OF THE INVENTION

1. Field of the Invention.

The present invention provides novel radiolabeled lipophilic salts, particularly radiolabeled
lipophilic phosphonium and ammonium salts, which are capable of measuring mitochondrial
surface potential ($\Delta\Psi_m$). This invention also provides pharmaceutical compositions comprising
15 such radiolabeled lipophilic salts. Additionally this invention provides imaging methods for
identifying tissues or cells having aberrant levels of mitochondrial activity by selectively
localizing radiolabeled lipophilic salts of the invention into dysfunctional mitochondria. The
invention also provides on-invasive methods for an early and sensitive detection of tumor response
to chemotherapy agents. The invention further provides treatment methods comprising
20 administration of a high energy radiolabeled lipophilic salts to a patient, particularly patients
suffering from diseases or disorders associated with mitochondrial dysfunction.

2. Definition.

In the specification the term "comprising" shall be understood to have a broad meaning
25 similar to the term "including" and will be understood to imply the inclusion of a stated integer or
step or group of integers or steps but not the exclusion of any other integer or step or group of
integers or steps. This definition also applies to variations on the term "comprising" such as
"comprise" and "comprises".

30 3. Background.

Measurement of the mitochondrial membrane potential ($\Delta\Psi_m$) provides the single most
comprehensive reflection of mitochondrial bio-energetic function primarily because it directly
depends on the proper integration of diverse metabolic pathways that converge at

the mitochondria. Numerous diseases are associated with mitochondria dysfunction, including cancer, cardiovascular and liver diseases, degenerative and autoimmune disorders as well as aging and new pathologies related to mitochondria are identified each year.

5 Alterations in $\Delta\Psi_m$ is an important characteristic of a vast array of pathologies that either involve suppressed (e.g., cancer) or enhanced apoptosis (e.g., HIV, degenerative disease) as well as >100 diseases directly caused by mitochondrial dysfunction such as DNA mutations and oxidative stress (e.g., various types of myopathies).

10 There are SPECT imaging probes labeled with a technetium center which are capable of accumulation in the mitochondria and the technetium labeled probes have been used for mitochondria based imaging techniques. There are a number of commercially available imaging probes that detect a given pathology using imaging agents such as [^{99m}Tc]MBI, FDG.

15 [^{18}F]FDG detects malignant lesion due to enhanced glucose metabolism. Further, as mentioned above, [^{18}F]FDG is not able to differentiate neoplasm from inflammation. [^{18}F]FDG is most effective imaging probe for tumor detection but poorly distinguishes neoplasm from inflammation, posing a frequent diagnostic challenge. In
20 certain organs inflammation (e.g., tuberculosis) is a frequent pathologies among patients with suspected malignant lesion. For example, >10% of pulmonary hot spots indicated by [^{18}F]FDG PET are inflammatory process rather than neoplasm, as proven by surgery. In other words, about 10% of lung patients with [^{18}F]FDG PET indications may undergo unnecessary chest surgery, for a disease (inflammation) that otherwise can be treated in
25 non-surgical and less costly and morbid approaches.

 Current approaches for evaluation of efficacy of chemotherapy relies on alterations in tumor growth rate, a costly approach of limited sensitivity which involves months of follow up, repeated visits in clinic, multiple radiographic scans and frequently a number of
30 treatment cycles.

Technetium labeled mitochondria imaging agents are hampered by several limitations. More particularly, labeling a molecule with ^{99m}Tc requires a conjugating moiety to complex the technetium ion such that Tc-based imaging agents have a high molecular weight which reduces the permeability of the imaging agent in target areas.

- 5 Further, technetium imaging agents are imaged with SPECT which has relatively low spatial resolution and sensitivity when compared to comparable PET images.

There are technetium complexes, derivatives of [^{99m}Tc]annexin V, for apoptosis imaging by using SPECT. The novelty of the proposed [^{18}F]phosphonium cations (PhCs) is that they detect the apoptotic process via a change in $\Delta\Psi\text{m}$, whereas annexin V derivatives do so due to overexpression of specific membrane proteins.

- 10 [^{99m}Tc]annexin V detects apoptosis due to externalization of phosphatidylserine on the outer cytoplasm membrane. This event occurs at the end of the apoptosis process when the fragmented cell is transformed into clusters of molecules (apoptotic bodies). Shortly after the externalization of phosphatidylserine (termed "eat me" phospholipids) the apoptotic bodies are phagocytized by neighboring cells. Therefore, detection of overexpression of phosphatidylserine is limited to a narrow time window which may last a few days only. Furthermore, the time of appearance and the duration of this window may vary among different chemotherapy agents and subjects.

The collapse of $\Delta\Psi\text{m}$ is the point of no return of the apoptotic process. Therefore, the collapse of $\Delta\Psi\text{m}$ affords the earliest time point to detect apoptosis, rather the last event as in the case of annexin V, and the collapse persists independent of time.

25

Current approaches for the evaluation of myocardial perfusion and viability have several limitations, including masking of myocardial activity by high accumulation in the organs adjacent to the heart (Th-201, [^{99m}Tc]MIBI) and short half-life of the isotope ([^{13}N]-ammonia and $^{82}\text{Rubidium}$), thus limited to PET centers with an on-site cyclotron.

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It would be desirable to have a family of lipophilic salts which have an affinity for mitochondria, particularly mitochondria undergoing aberrant activity.

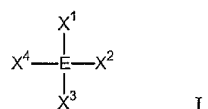
SUMMARY OF THE INVENTION

5 In view of the high incidence of cancer cases (~1.3 million per year in the USA), the high frequency of chemotherapy applications and the low frequency of successful chemotherapy, there is an urgent need for a non-invasive imaging probe of rapid and sensitive assessment of tumor response to treatment. The need for diagnostic means in oncology is best exemplified by the rapid transition of [¹⁸F]FDG PET from an
10 investigational to a preferred diagnostic tool for tumor detection within a few years.

There also exists a great need to diagnose and image cardiovascular diseases and disorders, many of which are associates with mitochondrial dysfunction. Thus there is also an urgent need for non-invasive imaging probes for rapid and sensitive measurement
15 of cardiac uptake of imaging agents having an affinity for dysfunctional mitochondria for the imaging of cardiovascular diseases such as myocardial perfusion.

The invention provides novel lipophilic salts, particularly lipophilic salts comprising a pharmaceutically acceptable anion and at least one phosphonium or
20 ammonium cations according to Formula I, and pharmaceutical compositions comprising cations of Formula I and at least one pharmaceutically acceptable carrier or excipient. Preferred lipophilic salts of the invention exhibit high affinity mitochondria, particularly dysfunctional mitochondria with enhanced or suppressed activity.

25 The present invention provides salts comprising at least one pharmaceutically acceptable anion and at least one cation according to Formula I



wherein

E is phosphorus or nitrogen; and

X^1 , X^2 , X^3 , and X^4 are independently selected from the group consisting of Ar and R, wherein at least one of X^1 , X^2 , X^3 , and X^4 is an Ar group;

Ar is optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

R is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope.

10

The salts of the invention which comprise a cation of Formula I are suitable for use in imaging or assessment, particularly PET or SPECT imaging, of mitochondrial dysfunction in a patient. Preferred salts of the invention are labeled with one or more radioisotopes, preferably including ^{11}C , ^{18}F , ^{76}Br , or ^{123}I and more preferably ^{18}F , ^{76}Br , or ^{123}I . The invention provides a phosphonium cation tracer labeled with ^{11}C -methyl group, e.g., [^{11}C]triphenylmethyl phosphonium (TPMP). Although suitable for use in medical centers situated at or near a cyclotron, the short half-life time of ^{11}C , e.g., about 20 minutes, limits the use of [^{11}C]TPMP at distant medical centers. Preferred salts of the invention comprise a ^{18}F , ^{76}Br , ^{123}I , or a combination thereof and are suitable for use in peripheral medical facilities and PET clinics.

20

The present invention provides lipophilic salts comprising a cation of Formula I or a subformula thereof which are preferentially taken up by dysfunctional mitochondria, e.g., mitochondria with suppressed or enhanced activity, and are suitable for use in imaging or radiotherapeutic applications. The invention provides imaging agents comprising a radiolabeled lipophilic cations, particularly lipophilic phosphonium or ammonium salts of the invention which has one or more radioisotopes which is capable of binding to dysfunctional mitochondria, e.g., mitochondria with suppressed or enhanced activity. More particularly, the radiolabeled lipophilic phosphonium or ammonium salts of the invention are suitable for use in measuring mitochondrial membrane potential ($\Delta\Psi\text{m}$) in vivo under a variety of conditions wherein the radiation

25

30

emitted by the radioisotope of the lipophilic phosphonium or ammonium salt is utilized to form the image. In preferred embodiments, radiolabeled lipophilic phosphonium or ammonium salts of the invention comprise one or more radioisotopes capable of emitting positron radiation and are suitable for use in positron emission tomography (PET).

5

According to yet another aspect, the present invention provides pharmaceutical compositions comprising radiolabeled salts of Formula I or the pharmaceutically acceptable salts or solvates thereof, which compositions are useful for the imaging variations in mitochondrial surface potential ($\Delta\Psi_m$), cells or tissues having dysfunctional mitochondria, and diseases or disorders associated with dysfunctional mitochondria. The invention further provides methods of imaging patients suffering from any of the above-recited diseases or disorders with an effective amount of a salt or composition of the invention.

10

15 Additionally this invention relates to the use of the salts of the invention (particularly labeled salts of this invention emitting high energy radiation) as therapeutic agents for the treatment of diseases and disorders associated with dysfunctional mitochondria for which the lipophilic phosphonium or ammonium salts of the invention have high affinity, e.g., disorders or diseases associated with dysfunctional mitochondria
20 activity. Typical disease and disorders include cancer, cardiovascular and liver diseases, degenerative disorders, autoimmune diseases and disorder, aging, DNA mutations, oxidative stress disorders, various myopathies, HIV, AIDS, and the like.

Preferred lipophilic cations, including phosphonium or ammonium salts, of the
25 invention preferentially localize to cells possessing mitochondria with elevated or suppressed levels of activity, e.g., dysfunctional mitochondrial activity.

BRIEF DESCRIPTION OF THE DRAWING

30 Figures 1 through 19 show results of Examples 9 through 19 which follow

DETAILED DESCRIPTION OF THE INVENTION

In addition to salts of Formula I, described above, the invention is further directed to lipophilic salts of Formula I (shown above) wherein the compounds provided by the invention are lipophilic salts of Formula I wherein

5

Ar is optionally substituted aryl having from 6 to 18 carbon atoms and between 1 and 3 rings, optionally substituted heteroaryl having from 3 to about 18 carbon atoms, between 1 and about 3 rings and between 1 and about 4 ring heteroatoms selected from N, O, and S, and optionally substituted aralkyl having between 7 and about 12 carbon

10

atoms; and
R is optionally substituted C₁₋₆alkyl, optionally substituted C₂₋₆alkenyl, optionally substituted C₂₋₆alkynyl, optionally substituted C₁₋₆haloalkyl having at least one F, Cl, Br, or I atom, optionally substituted cycloalkyl having between 3 and about 8 ring carbon atoms, optionally substituted aralkyl having between 7 and about 12 carbon atoms,

15

wherein at least one occurrence of R comprises at least one radioisotope.

Preferred salts of the invention include salts having at least one phosphonium cation of Formula I where E is phosphorus. Other preferred salts of the invention include those having at least one ammonium cation of Formula I where E is nitrogen. Other

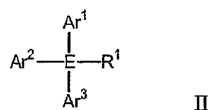
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preferred salts comprise a mixture of cations according to Formula I where each cation may be a phosphonium or ammonium cation.

Preferred salts of the invention comprise at least one R substituent which comprises a radioisotope capable of emitting positrons. Typically preferred positron emitting radioisotopes suitable for use in R substituents include ¹¹C, ¹⁸F, ¹²³I or any combination thereof.

25

Other preferred salts provided by the invention include salts comprising a cation of Formula II:



wherein

E is phosphorus or nitrogen; and

5 Ar¹, Ar², and Ar³ are independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

R¹ is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, 10 optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope.

More preferably, cations according to Formula II which are provided by the invention include those, wherein

15 Ar¹, Ar², and Ar³ are independently selected from the group consisting of optionally substituted aryl having from 6 to 18 carbon atoms and between 1 and 3 rings, optionally substituted heteroaryl having from 3 to about 18 carbon atoms, between 1 and about 3 rings and between 1 and about 4 ring heteroatoms selected from N, O, and S, and optionally substituted aralkyl having between 7 and about 12 carbon atoms; and

20 R¹ is optionally substituted C₁₋₆alkyl, optionally substituted C₂₋₆alkenyl, optionally substituted C₂₋₆alkynyl, optionally substituted C₁₋₆haloalkyl having at least one F, Cl, Br, or I atom, optionally substituted cycloalkyl having between 3 and about 8 ring carbon atoms, optionally substituted aralkyl having between 7 and about 12 carbon atoms, wherein at least one occurrence of R comprises at least one radioisotope.

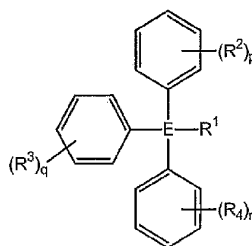
25 Particularly preferred cations of the invention according to Formula II comprise a R¹ group which is selected from the group consisting of ¹¹C-methyl, optionally substituted C₁₋₆alkyl, optionally substituted C₇₋₁₂aralkyl, optionally substituted C₆₋₁₂aryl, each of which is substituted with one or more ¹¹C-methyl, ¹¹C-methoxy, ¹⁸F, ⁷⁶Br, ¹²³I,

^{125}I , ^{131}I , or a combination thereof. More preferably, R^1 is ^{11}C -methyl, C_{2-6} alkyl substituted with one or more ^{18}F , or benzyl substituted with one or more ^{18}F , ^{76}Br , or ^{123}I .

The invention also provides salts comprising at least one cation according to
 5 Formula I or Formula II wherein R or R^1 comprises one or more radioisotopes suitable for use in radiation therapy.

The present invention further provides salts comprising a cation of Formula I which is represented by Formula III:

10



III

wherein

E is phosphorus or nitrogen;

15 R^1 is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope;

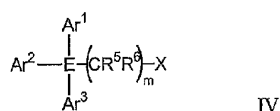
R^2 , R^3 , and R^4 are independently selected at each occurrence of R^2 , R^3 , and R^4
 20 from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide.

25

Preferred R^1 groups of Formula III include halo- C_{2-6} alkyl group or a halobenzyl group and more preferably R^1 of Formula III is selected from the group consisting of ω -fluoro- C_{2-6} alkyl, ω -iodo- C_{2-6} alkyl group, *ortho*, *meta* or *para*- fluorobenzyl group, or *ortho*, *meta* or *para*- iodobenzyl group.

5

Other preferred salts of the invention having a cation according to Formula I include those salts which comprise a cation according to Formula IV:



10

wherein

E is phosphorus or nitrogen;

Ar^1 , Ar^2 , and Ar^3 are independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted

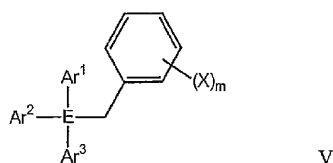
15 aralkyl; and

R^5 and R^6 are independently selected at each occurrence of R^5 and R^6 from the group consisting of hydrogen, halogen, hydroxy, amino, optionally substituted alkyl, optionally substituted haloalkyl, and optionally substituted alkoxy;

X is ^{11}C -methyl or a radioisotope of fluorine or iodine; and

20 m is a number from about 2 to about 6.

Yet other preferred salts of the invention having a cation according to Formula I include those salts which comprise a cation according to Formula V:



25

wherein

E is phosphorus or nitrogen;

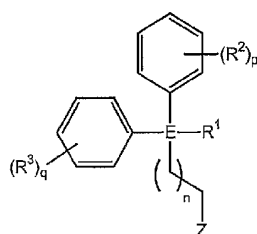
Ar¹, Ar², and Ar³ are independently selected from the group consisting of

- 5 optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

X is ¹¹C-methyl or a radioisotope of fluorine or iodine; and

m is a number from about 1 to about 5.

- 10 Still other preferred salts of the invention having a cation according to Formula I include those salts which comprise a cation according to Formula VI:



VI

- 15 wherein

E is phosphorus or nitrogen;

Z is chloro, fluoro, hydroxy, or methoxy;

n is a number from 1 to about 12;

p and q are independently selected numbers from zero to about 5;

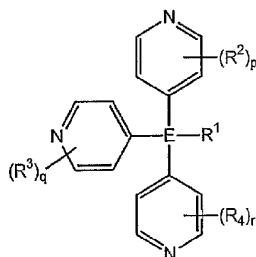
- 20 R¹ is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope; and

- 25 R² and R³ are independently selected at each occurrence of R² and R³ from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl,

optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide.

5

Other preferred salts of the invention having a cation according to Formula I include those salts which comprise a cation according to Formula VII:



VII

10

wherein

E is phosphorus or nitrogen;

p, q, and r are independently selected numbers from zero to about 4;

R¹ is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope;

R², R³, and R⁴ are independently selected at each occurrence of R², R³, and R⁴ from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide.

20

Particularly preferred salts comprising at least one pharmaceutically acceptable anion and at least one cation according to Formula I include those salts comprising at least one cation selected from the group consisting of:

- ¹¹C-methyl-triphenylphosphonium ion;
- 5 ¹¹C-methyl-tri-*ortho*-tolylphosphonium ion;
- ¹¹C-methyl-tri-*meta*-tolylphosphonium ion;
- ¹¹C-methyl-tri-*para*-tolylphosphonium ion;
- ¹⁸F-2-fluoroethyl-triphenylphosphonium ion;
- ¹⁸F-2-fluoroethyl -tri-*ortho*-tolylphosphonium ion;
- 10 ¹⁸F-2-fluoroethyl -tri-*meta*-tolylphosphonium ion;
- ¹⁸F-2-fluoroethyl -tri-*para*-tolylphosphonium ion;
- ¹⁸F-3-fluoropropyl-triphenylphosphonium ion;
- ¹⁸F-3- fluoropropyl -tri-*ortho*-tolylphosphonium ion;
- ¹⁸F-3- fluoropropyl -tri-*meta*-tolylphosphonium ion;
- 15 ¹⁸F-3- fluoropropyl -tri-*para*-tolylphosphonium ion;
- ¹⁸F-4-fluorobutyl-triphenylphosphonium ion;
- ¹⁸F-4- fluorobutyl-tri-*ortho*-tolylphosphonium ion;
- ¹⁸F-4- fluorobutyl -tri-*meta*-tolylphosphonium ion;
- ¹⁸F-4- fluorobutyl -tri-*para*-tolylphosphonium ion;
- 20 ¹⁸F-2-fluorobenzyl-triphenylphosphonium ion;
- ¹⁸F-2- fluorobenzyl -tri-*ortho*-tolylphosphonium ion;
- ¹⁸F-2- fluorobenzyl -tri-*meta*-tolylphosphonium ion;
- ¹⁸F-2- fluorobenzyl -tri-*para*-tolylphosphonium ion;
- ¹⁸F-3- fluorobenzyl -triphenylphosphonium ion;
- 25 ¹⁸F-3- fluorobenzyl -tri-*ortho*-tolylphosphonium ion;
- ¹⁸F-3- fluorobenzyl -tri-*meta*-tolylphosphonium ion;
- ¹⁸F-3- fluorobenzyl -tri-*para*-tolylphosphonium ion;
- ¹⁸F-4- fluorobenzyl -triphenylphosphonium ion;
- ¹⁸F-4- fluorobenzyl -tri-*ortho*-tolylphosphonium ion;
- 30 ¹⁸F-4- fluorobenzyl -tri-*meta*-tolylphosphonium ion;
- ¹⁸F-4- fluorobenzyl -tri-*para*-tolylphosphonium ion;

- ¹⁸F-3- fluoro-4-formyl-benzyl -triphenylphosphonium ion;
¹⁸F-3- fluoro-4-formyl-benzyl -tri-*ortho*-tolylphosphonium ion;
¹⁸F-3- fluoro-4-formyl-benzyl -tri-*meta*-tolylphosphonium ion;
¹⁸F-3- fluoro-4-formyl-benzyl -tri-*para*-tolylphosphonium ion;
5 (¹⁸F-4-fluorobenzyl)-(2-chloroethyl)-diphenylphosphonium ion;
(¹⁸F-4-fluorobenzyl)-(3-chloropropyl)-diphenylphosphonium ion;
(¹⁸F-4-fluorobenzyl)-(4-chlorobutyl)-diphenylphosphonium ion;
(¹⁸F-4-fluorobenzyl)-(6-chloropentyl)-diphenylphosphonium ion;
(¹⁸F-4-fluorobenzyl)-(5-chlorohexyl)-diphenylphosphonium ion;
10 ¹⁸F-2-fluoroethyl-tri(4-pyridyl)phosphonium ion;
¹⁸F-3-fluoropropyl- tri(4-pyridyl)phosphonium ion;
¹⁸F-4-fluorobutyl- tri(4-pyridyl)phosphonium ion;
¹⁸F-2-fluorobenzyl- tri(4-pyridyl)phosphonium ion;
¹⁸F-3- fluorobenzyl - tri(4-pyridyl)phosphonium ion;
15 ¹⁸F-4- fluorobenzyl - tri(4-pyridyl)phosphonium ion; and
¹⁸F-3- fluoro-4-formyl-benzyl - tri(4-pyridyl)phosphonium ion.

Preferred radiolabeled salts of the invention including those salts comprising a
 cation according to any one of Formula I, II, III, IV, V, VI, or VII, selectively localize to
 20 the mitochondria such that the ratio of radiation emitted from radiolabeled salts present in
 mitochondria to background radiation, c.g., radiolabeled salt not taken up in
 mitochondria is at least about 5:1. More preferably, salts of the invention are selectively
 taken up in dysfunctional mitochondria such that the ratio of to normal mitochondria is at
 least about 5:1.

25

The salts of the invention, particularly the lipophilic salts of the invention, have a
 distribution profile in the body which is a function of mitochondrial integrity and are
 suitable for use as diagnostic tools in the identification and imaging of various diseases
 and disorders associated with mitochondrial dysfunction. Moreover, the salts of the
 30 invention are useful diagnostic tools for assessing the efficacy of existing therapeutically
 drugs as well as the development of novel drugs. For example, the effectiveness of drugs

that trigger apoptosis (e.g., anticancer drugs) or suppress apoptosis (e.g., drugs that block the degenerative process in HIV) can be assessed by determining if the administration of said drugs can be monitored by observing the change (effective) or lack of change (not effective) to $\Delta\Psi_m$ by measuring $\Delta\Psi_m$ using the salts and imaging methods of the invention.

Preferred compounds of the invention, particularly compounds suitable for use in the imaging methods provided by the invention, include one or more radioisotopes capable of emitting one or more forms of radiation which are suitable for detection with any standard radiology equipment such as PET, SPECT, gamma cameras, MRI and the like. Preferred radioisotopes include tritium and isotopes of carbon, fluorine, technetium, iodine and other isotopes capable of emitting positrons. Particularly preferred radioisotopes include ^{11}C , ^{18}F , ^{76}Br , and ^{123}I .

The present invention further provides method of imaging which comprise the steps of:

- providing at least one radiolabeled salt comprising a pharmaceutically acceptable anion and at least one cation according to any one of Formula I, II, III, IV, V, VI, or VII;
- contacting cells or tissues with the radiolabeled salt; and
- making a radiographic image.

The imaging methods provided by the invention are suitable for assessing mitochondrial membrane potential ($\Delta\Psi_m$). More particularly, the imaging methods of the present invention are suitable for measuring change in mitochondrial membrane potential over time to assess the efficacy of therapeutic protocols or pharmaceutical treatments. Cells which exhibit suppressed or enhanced rates of apoptosis frequently also exhibit decreased or increased mitochondria activity. The salts provided by the present invention typically localize to cells in a concentration proportional to the level of mitochondria activity. Thus frequently when cells are experiencing reduced levels of apoptosis (e.g., cancer cells), a greater portion of the salt of the invention administered to the patient localizes to those cells, and vice versa, cells with enhanced levels of apoptosis

(e.g., auto immune disorders, tumor cells responsive to chemotherapy agents) will accumulate less salt of the invention than normal cells. Thus the imaging methods of the present invention are suitable for use in imaging of cells, tissues or other physiological targets which are experiencing suppressed or enhanced apoptosis.

5

The imaging methods of the present invention are generally suitable for imaging of any disease, disorder, or pathology which is related to mitochondria. Preferred diseases and disorders which are suitable for imaging include cancer (including neoplasms), cardiovascular diseases (including infraction and perfusion), liver diseases, 10 degenerative diseases or disorders, autoimmune disorders, aging, HIV infections, myopathies caused by oxidative stress or DNA mutation, or diseases and disorders associated with mitochondrial dysfunction.

The imaging methods of the invention are also suitable for use in assessing 15 efficacy of therapeutic drugs capable of triggering or suppressing apoptosis. The imaging methods of the invention may also be used to assess the efficacy of chemotherapy or radiation treatment protocols used to retard or destroy cancer and other malignant tumors.

20 The imaging methods of the invention which are suitable for assessing the efficacy of a therapeutic drug are also suitable in developing new therapeutic agents which are capable of disrupting mitochondrial function in target tissue.

The radiolabeled lipophilic salts of the invention and imaging methods using same 25 provide a non-invasive approach for early and sensitive assessment of treatment efficacy within a few days of starting a therapeutic protocol compared to current assessment methods which may require months. Most major anticancer drugs (e.g., taxol, cisplatin, vinblastine, and etoposide) induce their apoptotic effect via a cascade of events in which the collapse of $\Delta\Psi_m$ constitutes an early, obligatory and irreversible step of the apoptotic process. 30 Radiolabeled lipophilic salts of the invention accumulate mainly in the mitochondria and in direct correlation with $\Delta\Psi_m$. Cells affected by the treatment will accumulate

radiolabeled lipophilic salts of the invention much less than non-affected cells. Therefore, significant change between pre- and post-treatment scan will indicate tumor responding to treatment and lack of differences will indicate non-responding tumors. Collapse of $\Delta\Psi_m$ occurs within hours after treatment with most therapeutic agents.

5

The ability to monitor the first event of the irreversible phase of the apoptotic process affords a noninvasive method for early and sensitive detection of tumor response to treatment. In the clinical setting, the imaging methods provided by the present invention offer a powerful tool for tailoring of chemotherapy strategies that will most benefit the patient with reduced morbidity.

10

The radiolabeled lipophilic salts of the invention are also suitable for use in developing these new generations of chemotherapy agents. The radiolabeled lipophilic salts of the invention and imaging methods of using the same are suitable for use as non-invasive technique for an early and sensitive assessment at the molecular level of treatment efficacy in clinical studies. Moreover the imaging methods of the invention are suitable for use in selecting suitable malignant targets in test subjects, based on the functional integrity of mitochondria, upon which the novel drug can be tested.

15

20

The present invention further provides imaging methods suitable for use in the imaging of tumors with one or more salts having a cation according to Formula I or a subformula thereof. In preferred tumor imaging methods of the invention, the radiolabeled salt administered to a patient preferentially accumulates in mitochondria of malignant cells such that the concentration of radiolabeled cation of Formula I is greater in the mitochondria of the malignant cell than the concentration of the cation in adjacent normal cells.

25

The extent of cancerous disease (stage) is a major prognostic factor and non-invasive staging using imaging technologies has a key role in design of treatment strategies (e.g., surgery vs. radio-chemotherapy vs. adjuvant chemotherapy). The lipophilic salts of the present invention including salts having a cation according to Formula I accumulate in

30

malignant cells to a substantially greater extent than in normal cells. Administration of a radiolabeled salt of the present invention is suitable for the identification and imaging of malignant cells and tumors and is further suitable for measuring the stage of tumor development.

5

The tumor imaging methods of the invention are particularly suitable in certain embodiments for imaging of cancers, more particularly for imaging neoplasms including a variety of lung, breast, and prostate cancers. Moreover, the tumor imaging method of the invention may be used is capable of determining the extent of the cancerous disease

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(cancer stage).

The present invention provides methods of differentiating between malignant tumors, such as neoplasms, and tissue suffering from a variety of inflammation processes. The lipophilic salts of the invention including salts having a cation of Formula I accumulate

15

in malignant cells to a greater extent than in normal cells and accumulate in cellular components of inflammatory processes to a lesser extent than in normal cells such that a differential distinction can be made between malignant cells, normal cells and cells suffering from inflammation. Differential detection of malignancy will obviate the numerous unnecessary surgeries conducted each year and improve the cost-effectiveness of care

20

management in oncology.

The tumor imaging methods of the present invention are capable of distinguishing between tissue suffering from an inflammatory process and malignant lesions. While not wishing to be bound by theory, differential detection of malignant cells is possible

25

because malignant cells have a greater accumulation of the salts of the invention than normal cells and tissues or cellular components of inflammatory processes typically accumulate a lower concentration of the salts of the invention. Thus the concentration of any salt of the invention in the malignant lesion is significantly greater than normal tissue and tissue suffering from an inflammatory process.

30

The present invention further provides methods of imaging cardiovascular diseases, particularly methods of imaging the myocardia. The cardiovascular imaging

methods of the invention comprise the administration of at least one compound according to Formula I, or a subformula thereof to a patient suffering from or susceptible to a cardiovascular disease.

5 [18F]phosphonium cations of the invention are suitable for various cardiovascular diseases, particularly myocardial imaging. Myocytes contain the highest concentration of mitochondria and therefore the heart is by far the major organ target of phosphonium cations. In addition, phosphonium cation maintain excellent perfusion characteristics permitting high-contrast imaging of the heart. Infarct and heart failure involve apoptosis
10 followed by necrosis processes. [¹⁸F]Phosphonium cations are capable to accurately distinguish between myocardial segments in which the apoptotic process cannot be stopped by medication and revascularization, and myocardial area that can be salvaged by intervention.

15 Preferred imaging methods provided by the invention include the use of lipophilic salts according to any one of Formula I, II, III, IV, V, VI, or VII which are capable of generating at least a 2:1 target to background ratio of radiation intensity, or more preferably about a 5:1, about a 10:1 or about a 15:1 ratio of radiation intensity between target and background. In certain preferred methods the radiation intensity of the target
20 tissue is more intense than that of the background. In other embodiments, the invention provides methods where the radiation intensity of the target tissue is less intense than that of the background. Generally, any difference in radiation intensity between the target tissue and the background which is sufficient to allow for identification and visualization of the target tissue is sufficient for use in the methods of the present invention.

25 In preferred methods of the invention the compounds of the invention are excreted from tissues of the body quickly to prevent prolonged exposure to the radiation of the radiolabeled compound administered to the patient. Typically compounds according to Formula I or any subformula thereof are eliminated from the body in less than about 24
30 hours. More preferably, compounds of the invention are eliminated from the body in less than about 16 hours, 12 hours, 8 hours, 6 hours, 4 hours, 2 hours, 90 minutes, or 60

minutes. Typically preferred compounds are eliminated in between about 60 minutes and about 120 minutes.

Preferred compounds of the invention are stable in vivo such that substantially all,
5 e.g., more than about 50%, 60%, 70%, 80%, or more preferably 90% of the injected compound is not metabolized by the body prior to excretion.

Compounds and salts of the invention and imaging methods of the invention are useful in imaging a variety of conditions including cancer, cardiovascular and liver
10 diseases, HIV, AIDS, autoimmune disease, degenerative disorders, neoplasms, and the like.

Typical subjects to which compounds of the invention may be administered will be mammals, particularly primates, especially humans. For veterinary applications, a
15 wide variety of subjects will be suitable, e.g. livestock such as cattle, sheep, goats, cows, swine and the like; poultry such as chickens, ducks, geese, turkeys, and the like; and domesticated animals particularly pets such as dogs and cats. For diagnostic or research applications, a wide variety of mammals will be suitable subjects including rodents (e.g. mice, rats, hamsters), rabbits, primates, and swine such as inbred pigs and the like.
20 Additionally, for *in vitro* applications, such as *in vitro* diagnostic and research applications, body fluids and cell samples of the above subjects will be suitable for use such as mammalian, particularly primate such as human, blood, urine or tissue samples, or blood urine or tissue samples of the animals mentioned for veterinary applications.

25 The present invention also provide packaged pharmaceutical compositions comprising a pharmaceutical acceptable carrier and a salt comprising at least one pharmaceutically acceptable anion and a cation according to any one of Formula I, II, III, IV, V, VI, or VII. In certain embodiments the packaged pharmaceutical composition will comprise the reaction precursors necessary generate the compound or salt according to
30 Formula I or subformula thereof upon combination with a radiolabeled precursor. Other

packaged pharmaceutical compositions provided by the present invention further comprise indicia comprising at least one of:

- instructions for using the composition to image cells or tissues having increased or suppressed mitochondrial activity, or
- 5 instructions for using the composition to assess therapeutic effect of a drug protocol administered to a patient, or
- instructions for using the composition to selectively image malignant cells and tumors in the presence of inflammation, or
- instructions for using the composition to measure mitochondrial membrane
- 10 potential ($\Delta\Psi_m$).

In certain preferred embodiments, the invention provides a kit according to the invention contains from about 1 to about 30 mCi of the radionuclide-labeled imaging agent described above, in combination with a pharmaceutically acceptable carrier. The

15 imaging agent and carrier may be provided in solution or in lyophilized form. When the imaging agent and carrier of the kit are in lyophilized form, the kit may optionally contain a sterile and physiologically acceptable reconstitution medium such as water, saline, buffered saline, and the like.

20 The present invention further provides apparatus and synthetic protocols for the automated synthesis of ^{11}C , ^{18}F , ^{76}Br , or ^{123}I labeled salts of the invention, including salts comprising a cation according to any one of Formula I, II, III, VI, V, VII, and VIII, and preparation of pharmaceutical compositions comprising same. The half-life (120 min) of F-18 allows for distribution of cationic probes from central cyclotron to satellite PET

25 scanners, similarly to the rapidly evolving distribution system adopted for [^{18}F]FDG. Tagging the cationic probes with I-123 will allow for distribution from a manufacturing center to medical institutions equipped with SPECT.

Imaging agents of the invention may be used in accordance with the methods of

30 the invention by one of skill in the art, e.g., by specialists in nuclear medicine, to image sites having a dysfunctional mitochondria, e.g., mitochondria exhibiting aberrant activity,

in a subject or patient. Any site having a dysfunctional mitochondria, e.g., mitochondria exhibiting aberrant activity, may be imaged by the imaging methods and imaging agents of the present invention.

5 Images can be generated by virtue of differences in the spatial distribution of the imaging agents which accumulate at a site having a dysfunctional mitochondria, e.g., mitochondria exhibiting aberrant activity. The spatial distribution may be measured using any means suitable for the particular label, for example, a gamma camera, a PET
10 agent may be quantified using known methods for quantifying radioactive emissions. A particularly useful imaging approach employs more than one imaging agent to perform simultaneous studies. Alternatively, the imaging method may be carried out a plurality of times with increasing administered dose of the salt according to Formula I to perform successive studies using the split-dose image subtraction method.

15 Preferably, a detectably effective amount of the imaging agent of the invention is administered to a subject. In accordance with the invention, "a detectably effective amount" of the imaging agent of the invention is defined as an amount sufficient to yield an acceptable image using equipment which is available for clinical use. A detectably
20 effective amount of the imaging agent of the invention may be administered in more than one injection. The detectably effective amount of the imaging agent of the invention can vary according to factors such as the degree of susceptibility of the individual, the age, sex, and weight of the individual, idiosyncratic responses of the individual, the dosimetry. Detectably effective amounts of the imaging agent of the invention can also vary
25 according to instrument and film-related factors. Optimization of such factors is well within the level of skill in the art.

The amount of imaging agent used for diagnostic purposes and the duration of the imaging study will depend upon the radionuclide used to label the agent, the body mass
30 of the patient, the nature and severity of the condition being treated, the nature of therapeutic treatments which the patient has undergone, and on the idiosyncratic

responses of the patient. Ultimately, the attending physician will decide the amount of imaging agent to administer to each individual patient and the duration of the imaging study.

5 The compounds herein described may have one or more asymmetric centers or planes. Compounds of the present invention containing an asymmetrically substituted atom may be isolated in optically active or racemic forms. It is well known in the art how to prepare optically active forms, such as by resolution of racemic forms (racemates), by asymmetric synthesis, or by synthesis from optically active starting materials. Resolution
10 of the racemates can be accomplished, for example, by conventional methods such as crystallization in the presence of a resolving agent, or chromatography, using, for example a chiral HPLC column. Many geometric isomers of olefins, C=N double bonds, and the like can also be present in the compounds described herein, and all such stable isomers are contemplated in the present invention. *Cis* and *trans* geometric isomers of the
15 compounds of the present invention are described and may be isolated as a mixture of isomers or as separated isomeric forms. All chiral (enantiomeric and diastereomeric), and racemic forms, as well as all geometric isomeric forms of a structure are intended, unless the specific stereochemistry or isomeric form is specifically indicated.

20 When any variable occurs more than one time in any constituent or formula for a compound, its definition at each occurrence is independent of its definition at every other occurrence. Thus, for example, if a group is shown to be substituted with 0-2 R^{*}, then said group may optionally be substituted with up to two R^{*} groups and R^{*} at each occurrence is selected independently from the definition of R^{*}. Also, combinations of
25 substituents and/or variables are permissible only if such combinations result in stable compounds.

 As indicated above, various substituents of the various formulae (compounds of Formula I, II, III, IV, V, VI, or VII) are "optionally substituted", including Ar, Ar¹, Ar²,
30 Ar³, R, R¹, R², R³, R⁴, R⁵, R⁶, X, X¹, X², X³, X⁴, and Z of Formula I, II, III, IV, V, VI and VII, and such substituents as recited in the sub-formulae such as Formula I and

subformulae. The term "substituted," as used herein, means that any one or more hydrogens on the designated atom or group is replaced with a selection from the indicated group of substituents, provided that the designated atom's normal valence is not exceeded, and that the substitution results in a stable compound. When a substituent is oxo (keto, i.e., =O), then 2 hydrogens on an atom are replaced. The present invention is intended to include all isotopes (including radioisotopes) of atoms occurring in the present compounds.

When substituents such as Ar, Ar¹, Ar², Ar³, R, R¹, R², R³, R⁴, R⁵, R⁶, X, X¹, X², X³, X⁴, and Z of Formula I and subformulae thereof, and such substituents as recited in the sub-formulae are further substituted, they may be so substituted at one or more available positions, typically 1 to 3 or 4 positions, by one or more suitable groups such as those disclosed herein. Suitable groups that may be present on a "substituted" R¹, R², R or other group include *e.g.*, halogen; cyano; hydroxyl; nitro; azido; alkanoyl (such as a C₁₋₆ alkanoyl group such as acyl or the like); carboxamido; alkyl groups (including cycloalkyl groups, having 1 to about 8 carbon atoms, preferably 1, 2, 3, 4, 5, or 6 carbon atoms); alkenyl and alkynyl groups (including groups having one or more unsaturated linkages and from 2 to about 8, preferably 2, 3, 4, 5 or 6, carbon atoms); alkoxy groups having one or more oxygen linkages and from 1 to about 8, preferably 1, 2, 3, 4, 5 or 6 carbon atoms; aryloxy such as phenoxy; alkylthio groups including those having one or more thioether linkages and from 1 to about 8 carbon atoms, preferably 1, 2, 3, 4, 5 or 6 carbon atoms; alkylsulfinyl groups including those having one or more sulfinyl linkages and from 1 to about 8 carbon atoms, preferably 1, 2, 3, 4, 5, or 6 carbon atoms; alkylsulfonyl groups including those having one or more sulfonyl linkages and from 1 to about 8 carbon atoms, preferably 1, 2, 3, 4, 5, or 6 carbon atoms; aminoalkyl groups including groups having one or more N atoms and from 1 to about 8, preferably 1, 2, 3, 4, 5 or 6, carbon atoms; carbocyclic aryl having 6 or more carbons and one or more rings, (*e.g.*, phenyl, biphenyl, naphthyl, or the like, each ring either substituted or unsubstituted aromatic); arylalkyl having 1 to 3 separate or fused rings and from 6 to about 18 ring carbon atoms, with benzyl being a preferred arylalkyl group; arylalkoxy having 1 to 3 separate or fused rings and from 6 to about 18 ring carbon atoms, with O-benzyl being a

preferred arylalkoxy group, or a saturated, unsaturated, or aromatic heterocyclic group having 1 to 3 separate or fused rings with 3 to about 8 members per ring and one or more N, O or S atoms, e.g. coumarinyl, quinolinyl, isoquinolinyl, quinazolinyl, pyridyl, pyrazinyl, pyrimidyl, furanyl, pyrrolyl, thienyl, thiazolyl, triazinyl, oxazolyl, isoxazolyl, 5 imidazolyl, indolyl, benzofuranyl, benzothiazolyl, tetrahydrofuranyl, tetrahydropyranyl, piperidinyl, morpholinyl, piperazinyl, and pyrrolidinyl. Such heterocyclic groups may be further substituted, e.g. with hydroxy, alkyl, alkoxy, halogen and amino.

As used herein, "alkyl" is intended to include both branched and straight-chain 10 saturated aliphatic hydrocarbon groups, having the specified number of carbon atoms. Examples of alkyl include, but are not limited to, methyl, ethyl, *n*-propyl, *i*-propyl, *n*-butyl, *s*-butyl, *t*-butyl, *n*-pentyl, and *s*-pentyl. Preferred alkyl groups are C₁₋₆ alkyl groups. Especially preferred alkyl groups are methyl, ethyl, propyl, butyl, and 3-pentyl. The term C₁₋₄ alkyl as used herein includes alkyl groups consisting of 1 to 4 carbon 15 atoms, which may contain a cyclopropyl moiety. Suitable examples are methyl, ethyl, and cyclopropylmethyl.

"Cycloalkyl" is intended to include saturated ring groups, having the specified number of carbon atoms, such as cyclopropyl, cyclobutyl, cyclopentyl, or cyclohexyl. 20 Cycloalkyl groups typically will have 3 to about 8 ring members.

In the term "(C₃₋₈ cycloalkyl)C₁₋₄ alkyl", cycloalkyl, and alkyl are as defined above, and the point of attachment is on the alkyl group. This term encompasses, but is not limited to, cyclopropylmethyl, cyclohexylmethyl, and cyclohexylmethyl. 25

"Alkenyl" is intended to include hydrocarbon chains of either a straight or branched configuration comprising one or more unsaturated carbon-carbon bonds, which may occur in any stable point along the chain, such as ethenyl and propenyl. Alkenyl groups typically will have 2 to about 8 carbon atoms, more typically 2 to about 6 carbon 30 atoms.

"Alkynyl" is intended to include hydrocarbon chains of either a straight or branched configuration comprising one or more carbon-carbon triple bonds, which may occur in any stable point along the chain, such as ethynyl and propynyl. Alkynyl groups typically will have 2 to about 8 carbon atoms, more typically 2 to about 6 carbon atoms.

5

"Haloalkyl" is intended to include both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms, substituted with 1 or more halogen atoms. Examples of haloalkyl include, but are not limited to, mono-, di-, or tri-fluoromethyl, mono-, di-, or tri-chloromethyl, mono-, di-, tri-, tetra-, or penta-fluoroethyl, and mono-, di-, tri-, tetra-, or penta-chloroethyl. Typical haloalkyl groups will have 1 to about 8 carbon atoms, more typically 1 to about 6 carbon atoms.

10

"Alkoxy" represents an alkyl group as defined above with the indicated number of carbon atoms attached through an oxygen bridge. Examples of alkoxy include, but are not limited to, methoxy, ethoxy, *n*-propoxy, *i*-propoxy, *n*-butoxy, 2-butoxy, *t*-butoxy, *n*-pentoxy, 2-pentoxy, 3-pentoxy, isopentoxy, neopentoxy, *n*-hexoxy, 2-hexoxy, 3-hexoxy, and 3-methylpentoxy. Alkoxy groups typically have 1 to about 8 carbon atoms, more typically 1 to about 6 carbon atoms.

15

"Halolkoxy" represents a haloalkyl group as defined above with the indicated number of carbon atoms attached through an oxygen bridge.

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As used herein, the term "alkylthio" includes those groups having one or more thioether linkages and preferably from 1 to about 8 carbon atoms, more typically 1 to about 6 carbon atoms.

25

As used herein, the term "alkylsulfinyl" includes those groups having one or more sulfoxide (SO) linkage groups and typically from 1 to about 8 carbon atoms, more typically 1 to about 6 carbon atoms.

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As used herein, the term "alkylsulfonyl" includes those groups having one or more sulfonyl (SO₂) linkage groups and typically from 1 to about 8 carbon atoms, more typically 1 to about 6 carbon atoms.

5 As used herein, the term "alkylamino" includes those groups having one or more primary, secondary and/or tertiary amine groups and typically from 1 to about 8 carbon atoms, more typically 1 to about 6 carbon atoms.

"Halo" or "halogen" as used herein refers to fluoro, chloro, bromo, or iodo; and
10 "counter-ion" is used to represent a small, negatively charged species such as chloride, bromide, hydroxide, acetate, sulfate, and the like.

As used herein, "carbocyclic group" is intended to mean any stable 3- to 7-membered monocyclic or bicyclic or 7- to 13-membered bicyclic or tricyclic group, any
15 of which may be saturated, partially unsaturated, or aromatic. In addition to those exemplified elsewhere herein, examples of such carbocycles include, but are not limited to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, adamantyl, cyclooctyl, [3.3.0]bicyclooctanyl, [4.3.0]bicyclononanyl, [4.4.0]bicyclodecanyl, [2.2.2]bicyclooctanyl, fluorenyl, phenyl, naphthyl, indanyl, and tetrahydronaphthyl.

20 As used herein, the term "heterocyclic group" is intended to include saturated, partially unsaturated, or unsaturated (aromatic) groups having 1 to 3 (preferably fused) rings with 3 to about 8 members per ring at least one ring containing an atom selected from N, O or S. The nitrogen and sulfur heteroatoms may optionally be oxidized. The
25 term or "heterocycloalkyl" is used to refer to saturated heterocyclic groups.

The heterocyclic ring may be attached to its pendant group at any heteroatom or carbon atom that results in a stable structure. The heterocyclic rings described herein may be substituted on carbon or on a nitrogen atom if the resulting compound is stable. A
30 nitrogen in the heterocycle may optionally be quaternized. As used herein, the term "aromatic heterocyclic system" is intended to include any stable 5- to 7-membered

monocyclic or 10- to 14-membered bicyclic heterocyclic aromatic ring system which comprises carbon atoms and from 1 to 4 heteroatoms independently selected from the group consisting of N, O and S. It is preferred that the total number of S and O atoms in the aromatic heterocycle is not more than 2, more preferably not more than 1.

5

Examples of heterocycles include, but are not limited to, those exemplified elsewhere herein and further include acridinyl, azocinyl, benzimidazolyl, benzofuranyl, benzothiofuranyl, benzothiophenyl, benzoxazolyl, benzthiazolyl, benztriazolyl, benztetrazolyl, benzisoxazolyl, benzisothiazolyl, benzimidazolyl, carbazolyl, NH-carbazolyl, carbolinyl, chromanyl, chromenyl, cinnolinyl, decahydroquinolinyl, 2*H*,6*H*-1,5,2-dithiazinyl, dihydrofuro[2,3-*b*]tetrahydrofuran, furanyl, furazanyl, imidazolidinyl, imidazolyl, imidazolyl, 1*H*-indazolyl, indolenyl, indolyl, indolizyl, indolyl, 3*H*-indolyl, isobenzofuranyl, isochromanyl, isoindazolyl, isoindolyl, isoindolyl, isoquinolinyl, isothiazolyl, isoxazolyl, morpholinyl, naphthyridinyl, octahydroisoquinolinyl, oxadiazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl;- 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolidinyl, oxazolyl, oxazolidinyl, pyrimidinyl, phenanthridinyl, phenanthrolinyl, phenazinyl, phenothiazinyl, phenoxathiinyl, phenoxazinyl, phthalazinyl, piperazinyl, piperidinyl, pteridinyl, purinyl, pyranyl, pyrazinyl, pyrazolidinyl, pyrazolinyl, pyrazolyl, pyridazinyl, pyridoxazole, pyridoimidazole, pyridothiazole, pyridinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolinyl, 2*H*-pyrrolyl, pyrrolyl, quinazolinyl, quinolinyl, 4*H*-quinolizyl, quinoxalinyl, quinuclidinyl, tetrahydrofuranyl, tetrahydroisoquinolinyl, tetrahydroquinolinyl, 6*H*-1,2,5-thiadiazinyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,5-thiadiazolyl, 1,3,4-thiadiazolyl, thianthrenyl, thiazolyl, thienyl, thienothiazolyl, thienooxazolyl, thienoimidazolyl, thiophenyl, triazinyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl, 1,3,4-triazolyl, and xanthenyl.

Preferred heterocyclic groups include, but are not limited to, pyridinyl, pyrimidinyl, furanyl, thienyl, pyrrolyl, pyrazolyl, pyrrolidinyl, morpholinyl, piperidinyl, piperazinyl, and imidazolyl. Also included are fused ring and spiro compounds containing, for example, the above heterocycles.

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As used herein, the term "carbocyclic aryl" includes groups that contain 1 to 3 separate or fused rings and from 6 to about 18 ring atoms, without hetero atoms as ring members. Specifically preferred carbocyclic aryl groups include phenyl, and naphthyl including 1-naphthyl and 2-naphthyl.

A "pharmaceutically acceptable carrier" refers to a biocompatible solution, having due regard to sterility, pH, isotonicity, stability, and the like and can include any and all solvents, diluents (including sterile saline, Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, Lactated Ringer's Injection and other aqueous buffer solutions), dispersion media, coatings, antibacterial and antifungal agents, isotonic agents, and the like. The pharmaceutically acceptable carrier may also contain stabilizers, preservatives, antioxidants, or other additives, which are well known to one of skill in the art, or other vehicle as known in the art.

As used herein, "pharmaceutically acceptable salts" refer to derivatives of the disclosed compounds wherein the parent compound is modified by making non-toxic acid or base salts thereof. Examples of pharmaceutically acceptable salts include, but are not limited to, mineral or organic acid salts of basic residues such as amines; alkali or organic salts of acidic residues such as carboxylic acids; and the like. The pharmaceutically acceptable salts include the conventional non-toxic salts or the quaternary ammonium salts of the parent compound formed, for example, from non-toxic inorganic or organic acids. For example, conventional non-toxic acid salts include those derived from inorganic acids such as hydrochloric, hydrobromic, sulfuric, sulfamic, phosphoric, nitric and the like; and the salts prepared from organic acids such as acetic, propionic, succinic, glycolic, stearic, lactic, malic, tartaric, citric, ascorbic, pamoic, malefic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, mesylic, sulfanilic, 2-acetoxybenzoic, fumaric, toluenesulfonic, methanesulfonic, ethane disulfonic, oxalic, isethionic, $\text{HOOC}-(\text{CH}_2)_n-\text{COOH}$ where n is 0-4, and the like. The pharmaceutically acceptable salts of the present invention can be synthesized from a parent compound that contains a basic or acidic moiety by conventional chemical methods. Generally, such

salts can be prepared by reacting free acid forms of these compounds with a stoichiometric amount of the appropriate base (such as Na, Ca, Mg, or K hydroxide, carbonate, bicarbonate, or the like), or by reacting free base forms of these compounds with a stoichiometric amount of the appropriate acid. Such reactions are typically carried out in water or in an organic solvent, or in a mixture of the two. Generally, non-aqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred, where practicable. Lists of additional suitable salts may be found, e.g., in *Remington's Pharmaceutical Sciences*, 17th ed., Mack Publishing Company, Easton, PA, p. 1418 (1985).

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Because of the correlation between elevated or suppressed mitochondria activity and a variety of diseases and disorders, an imaging agent and methods of imaging using same that can assessing changes in mitochondria surface potential is an effective diagnostic tool for testing for the presence of a variety of disease states associated with triggering or suppressing apoptosis in cells. Moreover, imaging agents suitable for use in imaging or assessing changes in mitochondria surface potential are suitable for use in studying a variety of diseases including cancer, cardiovascular or liver diseases, HIV, AIDS, autoimmune disease, degenerative disorders, neoplasms, and the like.

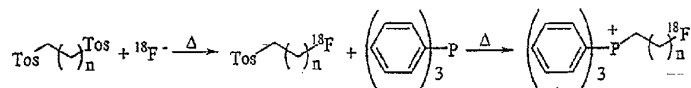
20 EXAMPLES

The present invention is further illustrated by the following examples which should not be construed as limiting in any way. The contents of all cited references (including literature references, issued patents, published patent applications) as cited throughout this application are hereby expressly incorporated by reference. The practice of the present invention will employ, unless otherwise indicated, conventional techniques, which are within the skill of the art. Such techniques are explained fully in the literature.

EXAMPLE 1: [^{18}F]3-fluoropropyltriphenyl-phosphonium ion ([^{18}F]FPTP)

30 The synthesis starts with the [^{18}F]fluoride from the cyclotron target transferred

onto an anion exchange column (Trap and Release column (DW-TRC) D and W, Inc., Oakdale, TN, USA). The column is eluted with aqueous potassium carbonate (2.3 mg dissolved in 0.3 mL) into a 5 cc v-vial containing Kryptofix. The Kryptofix, potassium carbonate, [^{18}F]fluoride mixture is dried at 120 °C; and, 7 mg of propyl ditosylate (Aldrich) is added in 0.5 mL acetonitrile. After heating at 80°C for 5 minutes, 21 mg of triphenylphosphine (Aldrich) in 0.5 mL of toluene is added. The acetonitrile is evaporated away and the toluene mixture heated to boiling for 3 - 5 minutes. After evaporating the toluene and cooling the vial, 0.5 mL of high-pressure liquid chromatography (HPLC) solvent [35:65 acetonitrile:water (0.1 M ammonium formate)] is added to the vial. The mixture is filtered through a 0.45 μm Teflon HPLC filter (Alltech 13 mm) and injected onto a preparative HPLC column (Waters Novapak C-18 6 μm , 7.8 x 300 mm) at 7 mL/min for purification. The product is collected on a rotary evaporator modified to allow addition and removal of solvents, the HPLC solvent evaporated and the radiolabeled phosphonium salt dissolved in sterile normal saline. The overall decay corrected radiochemical yield of [^{18}F]FPTP calculated from starting [^{18}F]fluoride is 12 percent. After sterile filtration (PALL-Gelman 0.2 μm Tuffryn) into a sterile vial, the solution is checked for radiochemical, chemical purity and specific activity by analytical HPLC [40:60 acetonitrile:water (0.1 M ammonium formate), Waters Novapak C-18 60 Å 4 μm , 3.9 x 150 mm] at 3 mL/min with a known concentration of cold standard characterized solution of 3-fluoropropyltriphenyl-phosphonium bromide [physical data: mp 313-316 °C; ^1H NMR (CDCl_3 , δ) 1.81-2.17 (m, 2H), 4.01-4.11 (m, 2H), 4.72-4.75 (m, 1H), 4.87-4.90 (m, 1H), 7.69-7.88 (m, 15H)]. The synthesis is summarized in Scheme 1.



Scheme 1: Synthesis of [^{18}F]alkyl phosphonium ions.

EXAMPLE 2: [^{18}F]2-fluoroethyltriphenyl-phosphonium ion ([^{18}F]FETP)

[^{18}F]FETP was prepared according to the method used to prepare [^{18}F]FPTP as described in Example 1 *supra*.

EXAMPLE 3: [^{18}F]2-fluorobutyltriphenyl-phosphonium ion ([^{18}F]FBTP)

[^{18}F]FBTP was prepared according to the method used to prepare [^{18}F]FPTP as described in Example 1 *supra*.

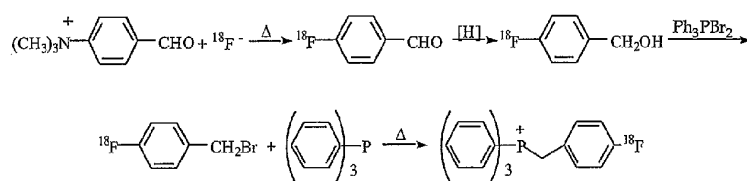
5

EXAMPLE 4: [^{18}F]fluorobenzyltriphenylphosphonium ion

After collecting and drying the [^{18}F]fluoride in the potassium carbonate/Kryptofix (as described in Example 1), trimethylammoniumbenzaldehyde triflate salt (7 mg) in 0.2 mL of dimethylsulfoxide (vacuum distilled from barium oxide prior to use) is added to the
10 mixture. After heating at 120°C for 5 minutes, the aldehyde is diluted with 5 mL of HPLC water and collected on a C-18 solid phase extraction cartridge (Waters C-18 Plus Sep-Pak) then washed with 10 mL of HPLC water and dried by inert gas flow through the cartridge for 3 minutes. The aldehyde is eluted from the cartridge with 2 mL of diethylether (Aldrich) and passed through 10% sodium borohydride on basic alumina
15 (Aldrich- 200 -400 mg) to reduce it to the alcohol. The alcohol is subsequently converted to [^{18}F]fluorobenzyl bromide by mixing with triphenylphosphine dibromide (75 - 100 mg) in 1 mL of methylene chloride for 5 minutes. After passing through a silica solid phase extraction cartridge (Waters Silica Classic Sep-Pak) and washing with 1 mL of methylene chloride, the [^{18}F]fluorinated benzyl bromide is added to 21 mg of
20 triphenylphosphine (or its analog) dissolved in 0.5 mL toluene in a 5 mL vial. The methylene chloride/ether solvent is evaporated away at low heat with inert gas flow, the vial capped and heated to boiling for 3-5 minutes. After evaporating the toluene and cooling the vial, 0.5 mL of high-pressure liquid chromatography (HPLC) solvent [50:50 acetonitrile:water (0.1 M ammonium formate)] is added to the vial. The mixture is filtered
25 through a 0.45 μm Teflon HPLC filter (Alltech 13 mm) and injected onto a preparative HPLC column (Waters Novapak C-18 6 μm , 7.8 x 300 mm) at 7 mL/min for purification. The product is collected on a rotary evaporator modified to allow addition and removal of solvents, the HPLC solvent evaporated and the radiolabeled phosphonium salt dissolved in sterile normal saline. After sterile filtration (PALL-

Gelman 0.2 μm Tuffryn) into a sterile vial, the solution is checked for radiochemical, chemical purity and specific activity by analytical HPLC [40:60 acetonitrile:water (0.1 M ammonium formate), Waters Novapak C-18 60 \AA 4 μm , 3.9 x 150 mm] at 3 ml/min and compared to a known standard of p-fluorobenzyltriphenylphosphonium ion as in example

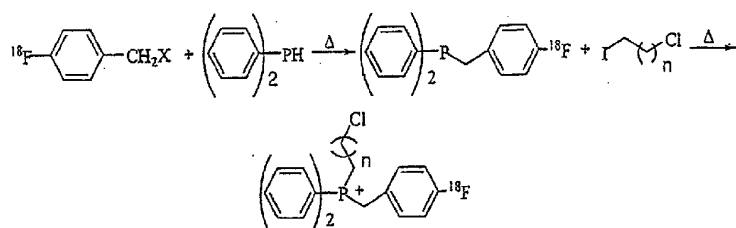
- 5 1. The overall decay corrected radiochemical yield of [^{18}F]fluorobenzyltriphenyl phosphonium ion calculated from starting [^{18}F]fluoride is 14 percent. [Physical Data: mp 313-316 $^{\circ}\text{C}$; ^1H NMR (D^6 -dmsO, δ) 5.17-5.21 (d, 2H), 6.99-7.08 (m, 4H), 7.67-7.94 (m, 15H)]. The synthesis is summarized in Scheme 2..



10 Scheme 2: Synthesis of [^{18}F]aryl phosphonium ions.

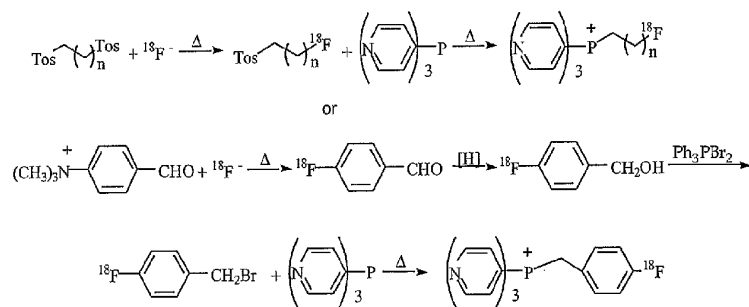
EXAMPLE 5: Radiolabeled diphenyl(haloalkyl)([^{18}F]fluorobenzyl)phosphonium ion

- Radiolabeled haloalkylphosphonium ion derivatives are prepared by the synthesis of a ^{18}F -fluorobenzyl halide as described in Example 4. The radiolabeled benzyl halide
 15 will be attached to diphenylphosphine to create a radiolabeled phosphine: Next the radiolabeled phosphine will be converted to the haloalkylphosphonium ion by reaction with the appropriate haloalkyl iodide. The synthesis of the radiolabeled chloroalkyl phosphonium ion is summarized in Scheme 3. Other haloalkyl species including chloro, bromo and iodo attached to alkyl chains of varying length, branching
 20 and site of halide substitution can also be prepared.

Scheme 3: synthesis of the [¹⁸F]-benzyl-chloroalkyl phosphonium ion.

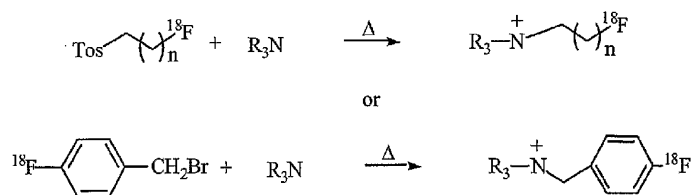
5 EXAMPLE 6: radiolabeled [¹⁸F]fluoroalkyl-tripyriddyphosphonium ion and radiolabeled [¹⁸F]fluorobenzyl-tripyriddyphosphonium ion

Radiolabeled phosphonium ions containing pyridyl rings are prepared according to the generalized reaction scheme provided in Scheme 4. The
10 radiosynthesis involves the reaction of ¹⁸F-fluoroalkyl (Example 1) and benzyl (Example 4) moieties as described above with tripyridylphosphine.

Scheme 4: synthesis of [¹⁸F]-alkyl or aryl tripyridylphosphonium ion

15 EXAMPLE 7: Radiolabeled ammonium ions

The procedures described in Examples 1-6 for preparation of radiolabeled phosphonium ions are also applicable for the preparation of quaternary ammonium ions as illustrated in Scheme 5. Quaternary ammonium ions have comparable biodistribution to phosphonium ions.



R = phenyl and/or alkyl moieties

Scheme 5: synthesis of radiolabeled ammonium ions

EXAMPLE 8: Whole body PET/SPECT Imaging

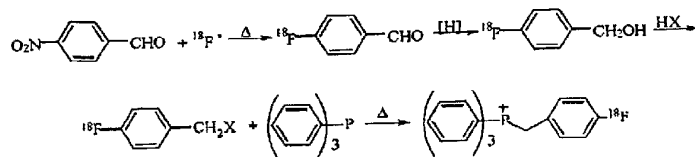
- 5 One imaging protocol suitable for delivery of a salt of the invention involves intravenous administration of the salt and acquisition of static scan for several minutes per bed position. The exact scan duration may be varied depending upon patient size, salt dosage, and the nature of the tissue to be imaged. However imaging parameters for imaging primates, particularly humans, can be modified as necessary by one skilled in the
- 10 radiological arts and familiar with PET and/or SPECT imaging with other radiopharmaceutical agents.

Example 9: Radiolabeling of F-18-FBnTP and stability in plasma after injection of the agent into mice

- 15 Radiochemical purity of F-18-FBnTP and its stability in vivo was measured by chromatography.

Methods:

- Preparation of F-18-fluorobenzyl triphenyl phosphonium (F-18-FBnTP): The synthesis of
- 20 labeling FBnTP with F-18 in the ortho position is described schematically below. After collecting the F-18-fluoride in Kryptofix (as with the fluoroalkyl derivatives), a nitrobenzaldehyde in acetonitrile is added to the mixture. After heating, the aldehyde is reduced to the alcohol and subsequently converted to a radiolabeled benzyl halide. The fluorinated benzyl halide is reacted with triphenylphosphine or its analog in toluene. The
- 25 mixture is purified and quality control performed, as discussed above.



5

Synthesis of o-F-18-fluorobenzyl triphenyl phosphonium ions

Samples of mouse plasma were obtained from heparinized whole blood collected 5, 15 and 30 min p.i. of the tracer. To eliminate binding to plasma proteins, the plasma was added to solid urea to give a final concentration of 8M urea. The plasma-urea was loaded into a column switch HPLC system (Hilton 2000), in which the plasma passes through a small capture column (Oasis Sorbent, Waters Corp.), which retains lipophilic solutes, while polar species fail to bind and are detected as they pass through a positron detector. After four minutes, the capture column is free of plasma proteins and polar species. Then, the contents of the capture column was swept onto an analytical column (Prodigy ODS-3, Phenomenex) by 40% acetonitrile, 60% triethylamine acetate buffer pH 4.1 at 1mL/min, where separation of the parent compound and lipophilic metabolites occurs. The effluent from the analytical column also passes through the positron flow detector. The proportion of each species is determined from the area under each chromatographic peak. Results: Radiochemical purity of the ¹⁸F-labeled FBnTP was more than 95%. Chromatography of plasma revealed a single radio-peak, comprising 97% of total activity with no other peak observed. Figure 1 depicts the chromatogram of plasma collected 30 min p.i., and the parental compound incubated for a similar time period (30 min) in buffered saline. Plasma activity and parental compound peaked at same time (~6 min).

These radiochromatograms are good evidence of the F-18-FBnTP can be labeled with high radiochemical purity and stable more than 30min after intravenous injection of the tracer in mice in vivo.

30

Example 10: Stability of FBnTP fluorination

Methods: Mice were injected via the tail vein with 25 μ Ci of F-18-PhC; 5 and 30 min p.i., the left or right femur bone was removed, and bone radioactivity together with standard (1:100 of injected dose) were counted in a gamma counter. A parallel group of mice was injected with free fluorides (F-18). Three mice of each group were studied at each time point. Radioactivity is represented as percentage of injected dosed (% ID) and total bone uptake was calculated as activity in femur bone x 20.

Results: Figure 2 depicts the total bone uptake at 5 and 30 min p.i. The marginal bone uptake of F-18-PhC, compared to the bone uptake in mice injected with F-18 only, indicates the stability of fluorination of the phosphonium compound. The minimal bone uptake of F-18-FBnTP in bone indicate the lack of free fluorides, meaning the stability of fluorination

Example 11: Mitochondria Membrane Potential (MMP) -Dependent Uptake

The extent of MMP-dependent cellular uptake of F-18-FBnTP was assessed using CCCP, a known protonophore that selectively abolishes the MMP.

Methods: Human lung carcinoma H549 cells (10/ml) were incubated with 0.1 μ Ci/ml F-18-FBnTP for 30 min. Samples of the suspension (1 mL) were transferred to Eppendorf's vials and placed in a 37°C bath. Varying concentrations (30, 60, 90, 120 μ M) of CCCP were added to the suspension. After 30 min of incubation with CCCP, the Eppendorf vials were centrifuged for 1 min, and activity in pellet and supernatant was immediately counted.

Results: Figure 3 depicts the cellular uptake of F-18-FBnTP in the presence of varying concentrations of CCCP. CCCP induced a dose-dependent decrease of the F-18-FBnTP cellular uptake. The large majority (86%) of F-18-FBnTP cellular uptake is MMP-dependent.

Example 12:. Biodistribution of novel F-18-fluorophosphonium cations in comparison with various tracers.

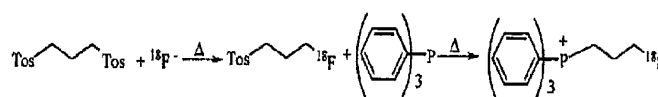
- Methods: The biodistribution of novel phosphonium compounds F-18-fluorbenzyl triphenyl phosphonium (FBnTP) and F-18-fluoropropyl triphenyl phosphonium cation (FPTP) was studied in adult mice and compared with the tracers C-11-triphenylmethyl phosphonium (TPMP), tetraphenyl phosphonium (TPP) and Tc-99m-sestamibi (MIBI). F-18-FBnTP was prepared as described above. Preparation of FPTP is described below. TPMP was prepared as previously described (Madar I, J Nucl Med. 1999;40:1180-5).
- TPP and MIBI were purchased from (NEN and Dupont, respectively). Three to five mice were used for the biodistribution study of each tracer. Nonanesthetized animals were injected i.v. with tracer solution (FBnTP or FPTP, TPMP 25 μ Ci; MIBI 40 μ Ci; TPP 2 μ Ci, all in a volume of 0.2 mL saline) then killed by neck dislocation at 60 min after injection. The organs and tissues of interest were removed and counted by a radioactivity counter along with standards (1:1000) (LKB Wallac, 1282 Compugamma CS).

Preparation of F-18-fluoropropyl triphenyl phosphonium ion (FPTP):

Schematic drawing of F-18-FPTP synthesis is given below. F-18-FPTP was

- first prepared as the nonradioactive compound, fluoropropyltriphenyl phosphonium bromide, from 1-fluoro-3-bromopropane and triphenylphosphine. The cold compound was characterized by proton and carbon-13 NMR, as well as HPLC. This standard was used for comparison during the purification, quality control and determination of specific activity of the radiolabeled F-18-FPTP. The F-18-FPTP was synthesized as described in figure 3. Briefly, 1,3-ditosylpropane in acetonitrile was added to a dried vial containing F-18-fluoride, Kryptofix and potassium carbonate. After heating at 80°C for 4 minutes on a heat gun, triphenylphosphine in toluene was added to the vial. After 5 minutes, the volume of the solution decreased to a few microliters. After cooling to room temperature, HPLC solvent was added to the vial. The mixture was

filtered through a 0.45 micron filter and injected onto a semi-preparative HPLC column. The product, F-18-FPTP, was collected, the solvent evaporated and the remaining dry F-18-FPTP redissolved in sterile normal saline. The solution was filtered through a sterile 0.22 micron filter into a sterile evacuated vial. An aliquot was removed to determine chemical and radiochemical purity by analytical HPLC. The specific activity was also determined at this time.



Synthesis of F-18-fluoropropyltriphenyl phosphonium ion

15

Results: The biodistribution of the tracers is presented in Table 1. Our comparative biodistribution studies in mice indicate that F-18-FBnTP act even better than C-11-TPMP as a PET perfusion agent. F-18-FBnTP uptake in heart is significantly greater than the other tracers, even that of C-11-TPMP, whereas clearance from blood is as good as that of the C-11-TPMP. Many investigators have reported that apoptosis plays a significant role in acute myocardial infarction and the pathogenesis of other forms of heart failure. These data suggest that F-18-FBnTP has a potential utility to assess myocardial diseases including apoptosis in connection with the MMP function as well as the myocardial perfusion tracer.

30

Table 1: Biodistribution of fluorophosphonium cations in mice 60 min after iv. injection in comparing with various tracers.

	[¹⁸ F]FBnTP	[¹⁸ F]FPTP	[¹¹ C]TPMP	[³ H]TPP	[^{99m} Tc]MIBI
blood	0.02 ± 0.00	0.01 ± 0.00	0.03 ± 0.00	0.09 ± 0.02	0.26 ± 0.05
brain	0.07 ± 0.01	0.03 ± 0.00	0.06 ± 0.02	0.06 ± 0.02	0.08 ± 0.04
heart	35.39 ± 5.02	9.86 ± 0.82	13.4 ± 0.95	20.7 ± 1.92	5.08 ± 0.25
lung	7.38 ± 1.55	2.02 ± 0.34	2.20 ± 0.21	2.81 ± 0.41	0.98 ± 0.19
liver	3.00 ± 1.12	4.22 ± 1.79	5.68 ± 0.60	6.12 ± 1.08	5.55 ± 1.08
spleen	1.79 ± 0.57	1.23 ± 0.26	1.43 ± 0.18	1.93 ± 0.47	1.27 ± 0.33
kidney	4.89 ± 0.88	5.74 ± 0.54	3.79 ± 0.83	4.37 ± 0.66	12.6 ± 2.57
muscle	4.31 ± 1.30	2.06 ± 0.53	2.49 ± 0.35	2.22 ± 0.41	1.56 ± 0.31

* = %ID/g

- 10 Example 13. Differentiation of tumor from inflammation using F-18-Phosphonium cations .

- The carcinogen nitrosomethyl urea (NMU) induces carcinoma tumors solely in the mammary gland of female rats. Therefore, the orthotopic NMU
- 15 mammary tumor is an excellent model for evaluating tracer tumor selectivity by contrasting radioactivity accumulating in mammary gland infested with carcinoma cells and healthy mammary gland. Freund Complete Adjuvant (FCA) is a well studied inflammation agent in rats Induction of FCA inflammation in NMU-bearing rats allows for direct quantitation of
- 20 tracer capability to differentiate carcinoma from inflammation.

Methods:

- 0.1 ml of nitrosomethyl urea (NMU) was introduced i.p in female rats (150 g). When tumor reached an approximate size of 1 to 1.5 cm, FCA was
- 25 injected (0.15 ml) to the hind limb footpad. Uptake assays were carried out 3 days thereafter. F-18-FBnTP (0.25 mCi) was injected via tail vein. Sixty minutes later tumor, inflamed tissue and healthy muscle tissue of the opposite limb (control) were collected on ice, weighed and counted in a gamma counter together with standards..

30

Results: Figure 3 depicts F-18-FBnTP activity in malignant mammary gland (Tumor), healthy mammary gland (Control gland), inflammation site (Inflammation), normalized to muscle. F-18-FBnTP uptake in tumor is 4-times greater than that in healthy gland, and 3-times greater than in inflammation. F-18-FBnTP differential uptake in mammary gland carcinoma versus healthy gland and inflammation muscle is a good evidence for the efficacy of the invention.

- The accumulation of the novel F-18-phosphonium cations, F-18-FBnTP and F-18-fluoropropyl triphenyl phosphonium (F-18-FPTP) in comparison with F18-FDG in FCA-induced inflammation and healthy tissue is presented in Table 2. Induction of inflammation and activity counting were carried out as described above.
- Fluorophosphonium compounds accumulate in inflammation much less than FDG (Table 2). These data provide strong evidence to our claim that F-18-phosphonium compound are suitable for differentiation of tumor from inflammation and may resolve a major drawback of F-18-FDG.
- Table 2: Accumulation of fluorophosphonium compounds (FBnTP and FPTP) FDG in inflammation tissue and healthy muscle 3 days after administration of FCA.

Table 2: Accumulation of fluorophosphonium compounds (FBnTP and FPTP) FDG in inflammation tissue and healthy muscle 3 days after administration of FCA. and

	Blood	Inflammation	Control	Inf/Control
¹⁸ F FPTP	*0.037 ± 0.01	0.07 ± 0.01	0.29 ± 0.06	0.25 ± 0.04
¹⁸ F FDG	0.17 ± 0.06	1.20 ± 0.16	1.05 ± 0.35	1.20 ± 0.29
¹⁸ F FBnTP	0.10 ± 0.01	0.27 ± 0.22	0.76 ± 0.21	0.42 ± 0.40
¹⁸ F FDG	0.40 ± 0.14	1.42 ± 0.53	0.68 ± 0.14	2.12 ± 0.73

*= %ID/g tissue

Example 14: Detection of response of lung carcinoma tumor to the chemotherapy agent taxeter _ In vivo

Methods: 2x10⁶ human lung carcinoma A549 cells were inoculated s.q. in 12 nude mice. When tumor size reached an approximate size of 5 _ 10 mm, six mice were injected i.v. with taxeter and six mice served as control. Uptake assays were carried out 48 hours thereafter. 25 μ Ci of F-18-FBnTP were injected i.v., and tumor and muscle tissue was dissected after 60 min.

Results: Figure 5 depicts tumor activity normalized to muscle. Taxeter produced a nearly 50% decrease in F-18-FBnTP accumulation in tumor, compared to non-treated mice.

Example 15: Detection of apoptosis induced by androgen depletion

Methods: Male rats were castrated and uptake assays were conducted 4 days thereafter. The ventral (VP), anterior (AP) and the dorsolateral (DLP) lobes of the prostate were dissected together with hind limb muscle. After counting tracer activity in the tissue samples, TUNEL staining was carried out and the fractions of apoptotic cells in the ventral and anterior lobes were measured.

Results: Data in Figure 6 are the mean of 9 treated and 8 control rats, normalized to muscle. Castration induced a lobe-specific decrease in the ventral aspects, but not in the anterior and the dorsolateral lobes. These findings are in line with fraction of apoptotic cells, as counted in the stained histological section. In the VP lobe, 12.4% \pm 3.8% of cells demonstrated DNA laddering (figure 8), compared to only 3.3% \pm 1.7% of cells in the AP lobe (figure 9).

Example 16: F-18-FBnTP selectivity for prostate carcinoma

The data presented above show that F-18-PhC is capable of detecting the apoptotic process in the whole animal. In the prostate, moreover,

5 alterations in F-18-PhC accumulation correlate with extent of apoptosis in the target tissue. These data suggest suitability of F-18-PhC as a PET tracer for measuring the efficacy of chemotherapy in prostate carcinoma and most probably in other type of carcinomas as well.

10 However, the capability of a tracer to accurately report extent of apoptosis may depend on the selectivity of the tracer to the tumor. To address this question, tumor selectivity of F-18-FBnTP was studied in orthotopic model of prostate carcinoma.

15 Methods: 2x10⁶ cells were injected, under anesthesia, into the prostate epithelial tissue of nude mice. When tumor reached an approximate size of 5 mm, uptake assays were performed as described above. Tumor selectivity of F-18-FBnTP and F-18-FDG was compared (3 mice in each group)

20 Results: Figure 9 depicts the accumulation of tracer in the normal and malignant prostate, normalized to muscle. The uptake ratio of malignant-to-normal prostate tissue was 2.5 for F-18-PhC and 1.25 for FDG. This data provides further support for the suitability of F-18-PhC to detect prostate carcinoma and to measure response to treatment.

25 Example 17: C-11-TPMP uptake kinetics in the myocardium

We have examined the performance of C-11-triphenyl phosphonium cation (TPMP) for assessing regional myocardial flow in dog using PET (Kraus, 1994).

30

Methods: Four mCi of C-11-TPMP were introduced i.v., and dynamic images of increasing duration (15 sec to 20 min) were acquired over a total time of 85 min. Images were acquired on the GE 4096+ PET scanner (15 slices, 6.5 mm slice thickness). Images were reconstructed using
5 backprojection, and corrected for attenuation. A detailed description of methods appears in Kraus, 1994.

Results: Axial sections of the heart at 5, 30 and 60 min after injection are shown in Figure 10. These images show excellent visualization of the
10 myocardium with a high contrast to the surrounding lung tissue. During the plateau time period heart/lung ratio was >14:1, and heart/blood >100:1.

The extraction fraction of C-11-TPMP in the dog heart as a function of myocardial blood flow is shown in Figure 11. Under baseline conditions
15 (flow = 69 ml/min/100g), the extraction fraction is very high (91%). A five-fold increase (by adenosine) in flow resulted in a 39% decrease of the extraction fraction.

To investigate the relationship between myocardial blood flow and
20 C-11-TPMP uptake in the heart, the LAD was occluded and the C-11-TPMP accumulation in tissue samples compared to microsphere determined regional myocardial blood flow. A significant correlation ($r=0.93$, $p < 0.01$) was found at 5 min after LAD occlusion (Figure 12). The non-infracted/infracted myocardium ratio was 12.1 ± 2.4 .

25 These data point out the excellent features of the phosphonium cation as a perfusion agent for assessing myocardial blood flow, compared to other currently used SPECT perfusion agents. Thallium 201 extraction under baseline flow is about 80%, decreasing to about 60% for five-fold
30 increases in flow rate. MIBI extraction for a normal flow is about 60%, decreasing to 40% for high flow rates. The advantages of C-11-TPMP PET

technology over these SPECT agents are: (1) an overall higher myocardial extraction of C-11-TPMP; (2) a prolonged retention of C-11-TPMP in the myocardium; and (3) the better temporal and spatial resolution of the PET scanner for better documentation of ischemic regions in the myocardium.

5

Example 18: F-18-FBnTP uptake kinetics in the myocardium

Methods: Mongrel dogs (BW = 35 kg) were injected with 3-4 mCi of F-18-FBnTP. Images were acquired on a GE 4096+ scanner (15 slices, 6.5 mm slice thickness). PET scans of increasing duration (15 sec to 20 min) were acquired over a total time of 85 min post-injection. Arterial blood samples (0.5 ml in volume) were collected every few seconds for the first 3 minutes and at gradually increasing intervals (1 to 10 min) for the remaining time of the imaging study. The vascular and myocardial kinetics of F-18-FBnTP was analyzed using the ROI method.

10

15

Results: F-18-FBnTP demonstrated a rapid washout from the blood pool (fig. 13). F-18-FBnTP accumulated rapidly in the myocardium, reached equilibrium within a few minutes which persisted for the remaining scanning time (fig. 14). Rapid uptake was seen in both the left and right ventricle (fig. 15). In contrast, F-18-FBnTP demonstrated rapid clearance from the atrium as well as from the adjacent lungs (fig 15). At 60 - 85 min post-injection period the ratio of myocardium to atrium and to lung was >15:1. Consequently, F-18-FBnTP afforded high-contrast cardiac images of an excellent visual clarity (fig.16).

20

25

Example 19: F-18-FBnTP myocardial accumulation in heart failure.

Pacing of the mongrel dog heart at high rate (210bpm) for a four weeks is a well-established model of heart failure. The advantage of this model is that the cardiomyopathy solely involve apoptosis without stenosis of coronary artery or related ischemia. Therefore, in this model the affect of apoptosis on FBnTP uptake can be dissected.

30

Methods: Preparation of dog and data acquisition was performed as described above. Mongrel dog was installed with a pacemaker in the rib case and underwent FBnTP PET scan (presented above). Following the baseline scan, the dog's heart was paced at a rate of 210 bpm for 4 weeks and a second scan was acquired.

Results: Figure 17 depicts the short-axis images of FBnTP before and after 4-wks pacing. Pacing produced a significant decrease in FBnTP uptake of 40-60% throughout most of the inferior wall (Figures 18-19, Table 3). Despite that the PET scan was not gated, but due to the superb perfusion capacities of FBnTP and consequently excellent clarity of the myocardium left ventricular wall, pacing-induced remodeling of the myocardium typical to heart failure, including thinning of the left ventricular wall and dilation of the left ventricular chamber can be seen clearly.

Moreover, pacing induced a significant decrease in the accumulation of F-18-FBnTP throughout the entire inferior wall, indicating an enhanced process of heart failure mediating apoptosis of myocytes in this segment. Figure 18 depicts co-registered images before and after pacing. A significant decrease in F-18FBnTP uptake is seen in the inferior wall.

Quantitation of FBnTP in the myocardium was performed using region of interest placed on co-registered myocardial images, whose activity was normalized to the injected dose, before and after pacing. An example for ROI placement is illustrated in figure 18.

Myocardial pacing induced a significant ($p < 0.001$) decrease of 40 to 60% in the inferior wall (see table 3)

Table 3: F-18-FBnTP myocardial uptake before and after pacing. For positioning of ROI see fig. 18. Data were derived as illustrated in Fig. 19

	Baseline	4-Wks Pacing	Ratio P/B
Slices 46-49			
ROI1	*108.62	60.70	0.56
ROI2	118.89	90.93	0.76
ROI3	119.11	97.86	0.82
Slices 36-37			
ROI1	72.22	44.46	0.62
ROI2	89.40	81.06	0.91
ROI3	102.46	86.93	0.85

* average activity (% of injected dose) accumulated over the over 36 to 85 min per slice, mean of activity of denoted coronal slices.
Position of ROIs as depicted in the above upper left image.
Same ROIs template was used to retrieve all data.

15

20 The disclosures of all articles and references mentioned in this application, including patents, are incorporated herein by reference.

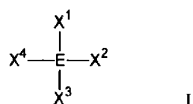
The invention and the manner and process of making and using it, are now described in such full, clear, concise and exact terms as to enable any person skilled in the art to which it pertains, to make and use the same. It is to be understood that the foregoing describes preferred embodiments of the present invention and that modifications may be made therein without departing from the spirit or scope of the present invention as set forth in the claims. To particularly point out and distinctly claim the subject matter regarded as invention, the following claims conclude this specification.

30

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A salt comprising at least one pharmaceutically acceptable anion and at least one cation according to Formula I

5



wherein

E is phosphorus or nitrogen; and

10 X^1 , X^2 , X^3 , and X^4 are independently selected from the group consisting of Ar and R, wherein at least one of X^1 , X^2 , X^3 , and X^4 is an Ar group;

Ar is optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

15 R is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope.

2. The salt of claim 1, wherein

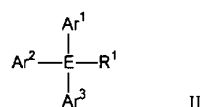
20 Ar is optionally substituted aryl having from 6 to 18 carbon atoms and between 1 and 3 rings, optionally substituted heteroaryl having from 3 to about 18 carbon atoms, between 1 and about 3 rings and between 1 and about 4 ring heteroatoms selected from N, O, and S, and optionally substituted aralkyl having between 7 and about 12 carbon atoms; and

25 R is optionally substituted C_{1-6} alkyl, optionally substituted C_{2-6} alkenyl, optionally substituted C_{2-6} alkynyl, optionally substituted C_{1-6} haloalkyl having at least one F, Cl, Br, or I atom, optionally substituted cycloalkyl having between 3 and about 8 ring carbon atoms, optionally substituted aralkyl having between 7 and about 12 carbon atoms, wherein at least one occurrence of R comprises at least one radioisotope.

3. The salt of claim 1 or claim 2, wherein the R group comprises one or more positron emitting radioactive isotopes.

4. The salt of any one of claims 1 to 3, wherein the R group comprises one or more isotopes selected from ^{11}C , ^{18}F , ^{76}Br , ^{123}I or any combination thereof

5. The salt of claim 1, wherein the cation is according to Formula II:



wherein

E is phosphorus or nitrogen; and

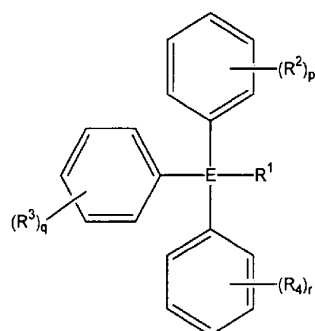
Ar^1 , Ar^2 , and Ar^3 are independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

R^1 is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope.

6. The salt of claim 5, wherein

Ar^1 , Ar^2 , and Ar^3 are independently selected from the group consisting of optionally substituted aryl having from 6 to 18 carbon atoms and between 1 and 3 rings, optionally substituted heteroaryl having from 3 to about 18 carbon atoms, between 1 and about 3 rings and between 1 and about 4 ring heteroatoms selected from N, O, and S, and optionally substituted aralkyl having between 7 and about 12 carbon atoms; and

- R¹ is optionally substituted C₁₋₆alkyl, optionally substituted C₂₋₆alkenyl, optionally substituted C₂₋₆alkynyl, optionally substituted C₁₋₆haloalkyl having at least one F, Cl, Br, or I atom, optionally substituted cycloalkyl having between 3 and about 8 ring carbon atoms, optionally substituted aralkyl having between 7 and about 12 carbon atoms, wherein at least one occurrence of R comprises at least one radioisotope.
- 5
7. The salt of any one of claims 1 to 6, wherein E is phosphorus.
8. The salt of any one of claims 1 to 6, wherein E is nitrogen.
- 10
9. The salt of any one of claims 6 to 8, wherein R¹ is selected from the group consisting of ¹¹C-methyl, optionally substituted C₁₋₆alkyl, optionally substituted C₇₋₁₂aralkyl, optionally substituted C₆₋₁₂aryl, each of which is substituted with one or more ¹¹C-methyl, ¹¹C-methoxy, ¹⁸F, ⁷⁶Br, ¹²³I, ¹²⁵I, ¹³¹I, or a combination thereof.
- 15
10. The salt of any one of claims 6 to 8, wherein R¹ is ¹¹C-methyl, C₂₋₆alkyl substituted with one or more ¹⁸F, or benzyl substituted with one or more ¹⁸F.
11. The salt of any one of claims 5 to 10, wherein R¹ comprises one or more radioisotope suitable for use in radiation therapy.
- 20
12. The salt of claim 1, wherein the cation is according to Formula III:



III

wherein

E is phosphorus or nitrogen;

5 p, q, and r are independently selected numbers from zero to about 5;

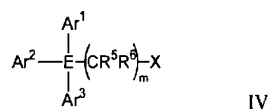
R¹ is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope;

10 R², R³, and R⁴ are independently selected at each occurrence of R², R³, and R⁴ from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide.

13. The salt of claim 12, wherein R¹ is a halo-C₂₋₆alkyl group or a halobenzyl group.

20 14. The salt of claim 13, wherein R¹ is ω-fluoro-C₂₋₆alkyl, ω-iodo-C₂₋₆alkyl group, *ortho*, *meta* or *para*- fluorobenzyl group, or *ortho*, *meta* or *para*- iodobenzyl group.

15. The salt of claim 1, wherein the cation is according to Formula IV:



5 wherein

E is phosphorus or nitrogen;

Ar¹, Ar², and Ar³ are independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted alkyl; and

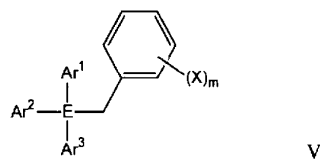
10 R⁵ and R⁶ are independently selected at each occurrence of R⁵ and R⁶ from the group consisting of hydrogen, halogen, hydroxy, amino, optionally substituted alkyl, optionally substituted haloalkyl, and optionally substituted alkoxy;

X is ¹¹C-methyl or a radioisotope of fluorine or iodine; and

m is a number from about 2 to about 6.

15

16. The salt of claim 1, wherein the cation is according to Formula V:



20 wherein

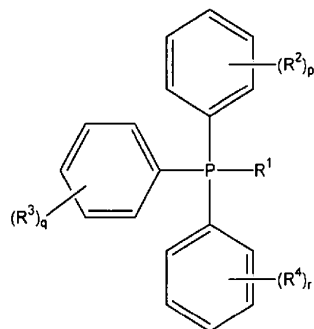
E is phosphorus or nitrogen;

Ar¹, Ar², and Ar³ are independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted alkyl; and

25 X is ¹¹C-methyl or a radioisotope of fluorine or iodine; and

m is a number from about 1 to about 5.

17. A salt comprising at least one pharmaceutically acceptable anion and a cation represented by the following formula:



5

wherein

R¹ is alkyl, alkenyl, alkynyl, haloalkyl, cycloalkyl, or aralkyl, with R¹ being substituted by one or more groups comprising ¹⁸F and optionally substituted by one or more additional groups;

10 R², R³, and R⁴ are independently selected at each occurrence of R², R³, and R⁴ from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide; and

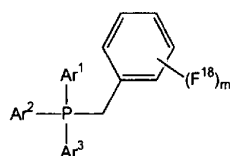
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p, q, and r are independently selected numbers from zero to 5.

18. The salt of claim 17, wherein R¹ is aralkyl.

20

19. A salt comprising at least one pharmaceutically acceptable anion and a cation represented by the following formula:



wherein

- Ar¹, Ar², and Ar³ are independently selected from the group consisting of
 5 optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted
 aralkyl; and

m is an integer of to 1 to 5.

20. The salt of claim 19 wherein at least one of Ar¹, Ar², and Ar³ is aryl or
 10 phenyl.

21. The salt of claim 19 wherein m is 1.

22. A compound which comprises:
 15 ¹⁸F-4-fluorobenzyl-triphenylphosphonium;
¹⁸F-2-fluorobenzyl-triphenylphosphonium;
¹⁸F-3-fluorobenzyl-triphenylphosphonium
¹⁸F-2-fluoroethyl-triphenylphosphonium;
¹⁸F-2-fluoroethyl-tri-*ortho*-tolylphosphonium;
 20 ¹⁸F-2-fluoroethyl-tri-*meta*-tolylphosphonium;
¹⁸F-2-fluoroethyl-tri-*para*-tolylphosphonium;
¹⁸F-3-fluoropropyl-triphenylphosphonium;
¹⁸F-3-fluoropropyl-tri-*ortho*-tolylphosphonium;
¹⁸F-3-fluoropropyl-tri-*meta*-tolylphosphonium;
 25 ¹⁸F-3-fluoropropyl-tri-*para*-tolylphosphonium;
¹⁸F-4-fluorobutyl-triphenylphosphonium;
¹⁸F-4-fluorobutyl-tri-*ortho*-tolylphosphonium;
¹⁸F-4-fluorobutyl-tri-*meta*-tolylphosphonium;

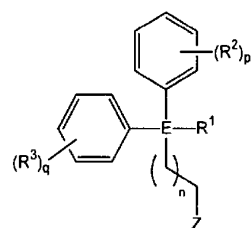
- 54 -

- 5 ¹⁸F-4- fluorobutyl-tri-*para*-tolylphosphonium;
¹⁸F-2- fluorobenzyl-tri-*ortho*-tolylphosphonium;
¹⁸F-2- fluorobenzyl-tri-*meta*-tolylphosphonium;
¹⁸F-2- fluorobenzyl-tri-*para*-tolylphosphonium;
¹⁸F-3- fluorobenzyl-tri-*ortho*-tolylphosphonium;
¹⁸F-3- fluorobenzyl-tri-*meta*-tolylphosphonium;
¹⁸F-3- fluorobenzyl-tri-*para*-tolylphosphonium;
¹⁸F-4- fluorobenzyl-tri-*ortho*-tolylphosphonium;
¹⁸F-4- fluorobenzyl-tri-*meta*-tolylphosphonium;
10 ¹⁸F-4- fluorobenzyl-tri-*para*-tolylphosphonium;
¹⁸F-3- fluoro-4-formyl-benzyl-triphenylphosphonium;
¹⁸F-3- fluoro-4-formyl-benzyl-tri-*ortho*-tolylphosphonium;
¹⁸F-3- fluoro-4-formyl-benzyl-tri-*meta*-tolylphosphonium;
¹⁸F-3- fluoro-4-formyl-benzyl-tri-*para*-tolylphosphonium;
15 (¹⁸F-4-fluorobenzyl)-(2-chloroethyl)-diphenylphosphonium;
(¹⁸F-4-fluorobenzyl)-(3-chloropropyl)-diphenylphosphonium;
(¹⁸F-4-fluorobenzyl)-(4-chlorobutyl)-diphenylphosphonium;
(¹⁸F-4-fluorobenzyl)-(6-chloropentyl)-diphenylphosphonium;
(¹⁸F-4-fluorobenzyl)-(5-chlorohexyl)-diphenylphosphonium;
20 ¹⁸F-2-fluoroethyl-tri(4-pyridyl)phosphonium;
¹⁸F-3-fluoropropyl-tri(4-pyridyl)phosphonium;
¹⁸F-4-fluorobutyl-tri(4-pyridyl)phosphonium;
¹⁸F-2-fluorobenzyl-tri(4-pyridyl)phosphonium;
¹⁸F-3-fluorobenzyl-tri(4-pyridyl)phosphonium;
25 ¹⁸F-4-fluorobenzyl-tri(4-pyridyl)phosphonium; or
¹⁸F-3-fluoro-4-formyl-benzyl-tri(4-pyridyl)phosphonium.
- 30 23. The salt of claim 1 which comprises a pharmaceutically acceptable anion
and a cation selected from the group consisting of:

	¹¹ C-methyl-triphenylphosphonium ion
	¹¹ C-methyl-tri- <i>ortho</i> -tolylphosphonium ion
	¹¹ C-methyl-tri- <i>meta</i> -tolylphosphonium ion
	¹¹ C-methyl-tri- <i>para</i> -tolylphosphonium ion
5	¹⁸ F-2-fluoroethyl-triphenylphosphonium ion
	¹⁸ F-2-fluoroethyl -tri- <i>ortho</i> -tolylphosphonium ion
	¹⁸ F-2-fluoroethyl -tri- <i>meta</i> -tolylphosphonium ion
	¹⁸ F-2-fluoroethyl -tri- <i>para</i> -tolylphosphonium ion
	¹⁸ F-3-fluoropropyl-triphenylphosphonium ion
10	¹⁸ F-3- fluoropropyl -tri- <i>ortho</i> -tolylphosphonium ion
	¹⁸ F-3- fluoropropyl -tri- <i>meta</i> -tolylphosphonium ion
	¹⁸ F-3- fluoropropyl -tri- <i>para</i> -tolylphosphonium ion
	¹⁸ F-4-fluorobutyl-triphenylphosphonium ion
	¹⁸ F-4- fluorobutyl-tri- <i>ortho</i> -tolylphosphonium ion
15	¹⁸ F-4- fluorobutyl -tri- <i>meta</i> -tolylphosphonium ion
	¹⁸ F-4- fluorobutyl -tri- <i>para</i> -tolylphosphonium ion
	¹⁸ F-2-fluorobenzyl-triphenylphosphonium ion
	¹⁸ F-2- fluorobenzyl -tri- <i>ortho</i> -tolylphosphonium ion
	¹⁸ F-2- fluorobenzyl -tri- <i>meta</i> -tolylphosphonium ion
20	¹⁸ F-2- fluorobenzyl -tri- <i>para</i> -tolylphosphonium ion
	¹⁸ F-3- fluorobenzyl -triphenylphosphonium ion
	¹⁸ F-3- fluorobenzyl -tri- <i>ortho</i> -tolylphosphonium ion
	¹⁸ F-3- fluorobenzyl -tri- <i>meta</i> -tolylphosphonium ion
	¹⁸ F-3- fluorobenzyl -tri- <i>para</i> -tolylphosphonium ion
25	¹⁸ F-4- fluorobenzyl -triphenylphosphonium ion
	¹⁸ F-4- fluorobenzyl -tri- <i>ortho</i> -tolylphosphonium ion
	¹⁸ F-4- fluorobenzyl -tri- <i>meta</i> -tolylphosphonium ion
	¹⁸ F-4- fluorobenzyl -tri- <i>para</i> -tolylphosphonium ion
	¹⁸ F-3- fluoro-4-formyl-benzyl -triphenylphosphonium ion
30	¹⁸ F-3- fluoro-4-formyl-benzyl -tri- <i>ortho</i> -tolylphosphonium ion
	¹⁸ F-3- fluoro-4-formyl-benzyl -tri- <i>meta</i> -tolylphosphonium ion

¹⁸F-3- fluoro-4-formyl-benzyl -tri-*para*-tolylphosphonium ion
(¹⁸F-4-fluorobenzyl)-(2-chloroethyl)-diphenylphosphonium ion
(¹⁸F-4-fluorobenzyl)-(3-chloropropyl)-diphenylphosphonium ion
(¹⁸F-4-fluorobenzyl)-(4-chlorobutyl)-diphenylphosphonium ion
5 (¹⁸F-4-fluorobenzyl)-(6-chloropentyl)-diphenylphosphonium ion
(¹⁸F-4-fluorobenzyl)-(5-chlorohexyl)-diphenylphosphonium ion
¹⁸F-2-fluoroethyl-tri(4-pyridyl)phosphonium ion
¹⁸F-3-fluoropropyl- tri(4-pyridyl)phosphonium ion
¹⁸F-4-fluorobutyl- tri(4-pyridyl)phosphonium ion
10 ¹⁸F-2-fluorobenzyl- tri(4-pyridyl)phosphonium ion
¹⁸F-3- fluorobenzyl - tri(4-pyridyl)phosphonium ion
¹⁸F-4- fluorobenzyl - tri(4-pyridyl)phosphonium ion
¹⁸F-3- fluoro-4-formyl-benzyl - tri(4-pyridyl)phosphonium ion

24. The salt of claim 1, wherein the cation is according to Formula VI:



VI

wherein

E is phosphorus or nitrogen;

Z is chloro, fluoro, hydroxy, or methoxy;

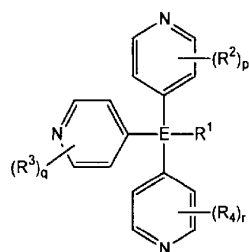
n is a number from 1 to about 12;

p and q are independently selected numbers from zero to about 5;

R^1 is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope; and

- 5 R^2 and R^3 are independently selected at each occurrence of R^2 and R^3 from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally
- 10 substituted mono or dialkylcarboxamide.

25. The salt of claim 1, wherein the cation is according to Formula VII:



VII

15

wherein

E is phosphorus or nitrogen;

p, q, and r are independently selected numbers from zero to about 4;

- 20 R^1 is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope;

- R^2 , R^3 , and R^4 are independently selected at each occurrence of R^2 , R^3 , and R^4 from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted
- 25

alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide.

5 26. The salt of any one of claims 1 to 25, wherein the salt exhibits a mitochondria to non-mitochondria ratio of at least about 5:1.

 27. The salt of claim 26, wherein the salt exhibits a selectivity ratio for dysfunctional mitochondria to normal mitochondria of at least about 5:1.

10

 28. The salt of any one of claims 1 to 27 which is suitable for use in a positron emission tomography (PET) or single photon emission computed tomography (SPECT).

 29. The salt of any one of claims 1 to 28 which has a bio-distribution
15 correlated to mitochondrial integrity.

 30. A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a salt or compound according to any one of claims 1 to 29.

20 31. A package comprising a pharmaceutical composition of claim 30 in a container and further comprising indicia comprising at least one of:

 instructions for using the composition to image cells or tissues having increased or suppressed mitochondrial activity, or

 instructions for using the composition to assess therapeutic effect of a drug
25 protocol administered to a patient, or

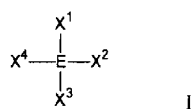
 instructions for using the composition to selectively image malignant cells and tumors in the presence of inflammation, or

 instructions for using the composition to measure mitochondrial membrane potential ($\Delta\Psi_m$).

30

 32. An imaging method comprising the steps of:

providing a radiolabeled salt comprising at least one pharmaceutically acceptable anion and at least one cation according to Formula I



5

wherein

E is phosphorus or nitrogen; and

X^1 , X^2 , X^3 , and X^4 are independently selected from the group consisting of Ar and R, wherein at least one of X^1 , X^2 , X^3 , and X^4 is an Ar group;

10

Ar is optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

R is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope;

15

contacting cells or tissues with the radiolabeled salt; and
making a radiographic image.

20

33. An imaging method comprising:

contacting cells or tissues with a salt or compound according to any one of claims 2 to 29; and
making a radiographic image.

25

34. The method of claim 32 or claim 33, wherein the imaging method is suitable for measuring mitochondrial membrane potential ($\Delta\Psi_m$).

35. The method of any one of claims 32 to 34, wherein the imaging method is suitable for measuring suppressed or enhanced apoptosis.

36. The method of any one of claims 32 to 34, wherein the imaging method is suitable for imaging a pathology that involves either enhanced or suppressed apoptosis.
- 5 37. The method of claim 36, wherein the imaging method is suitable for imaging a disease associated with suppressed apoptosis.
38. The method of claim 36, wherein the imaging method is suitable for imaging a disease associated with enhanced apoptosis.
- 10 39. The method of any one of claims 32 to 36, wherein the imaging method is suitable for imaging of cancer, degenerative diseases or disorders, autoimmune disorders, aging, HIV infections, myopathies caused by oxidative stress or DNA mutation, or diseases and disorders associated with mitochondrial dysfunction.
- 15 40. The method of any one of claims 32 to 34, wherein the imaging method is suitable for use in imaging mitochondrial dysfunction.
41. The method of any one of claims 32 to 34, wherein the imaging method is suitable for assessing efficacy of therapeutic drugs capable of triggering or suppressing apoptosis.
- 20 42. The method of claim 41, wherein the imaging method is suitable for assessing the efficacy of therapeutic protocol.
- 25 43. The method of claim 42, wherein the therapeutic protocol is selected from drug, radiation or a combination thereof.
44. The method of any one of claims 32 to 34, wherein the imaging method is suitable for imaging of tumors.
- 30

45. The method of claim 44, wherein radiolabeled salt preferentially accumulates in mitochondria of tumour cells.
46. The method of claim 44 or 45 wherein the tumor is a neoplasm.
- 5 47. The method of claim 44 or 45 wherein the tumor is a lung tumor, a breast tumor or a prostate tumor.
- 10 48. The method of claim 44 wherein the imaging method is capable of distinguishing between tissue inflammation and tumors.
- 15 49. The method of any one of claims 32 to 34, wherein the imaging method is suitable for imaging cancer.
50. The method of claim 49, wherein the imaging method is capable of determining the extent of the cancerous disease (cancer stage).
- 20 51. The method of claim 49 or claim 50, wherein radiolabeled salt preferentially accumulates in mitochondria of cancer cells.
52. The method of any one of claims 49 to 51, wherein the cancer is a neoplasm.
- 25 53. The method of any one of claims 49 to 51, wherein the cancer is lung cancer.
- 30 54. The method of claim 49, wherein the imaging method is capable of distinguishing between tissue inflammation and cancer.

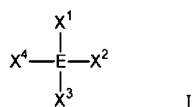
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55. The method of any one of claims 32 to 34, wherein the imaging method is suitable in developing new therapeutic agents which are capable of disrupting mitochondrial function in target tissue.
- 5 56. The method of any one of claims 32 to 55, wherein the radiolabeled salt exhibits a target to non-target ratio of at least about 5:1.
57. The method of any one of claims 32 to 56, wherein the radiolabeled salt is stable in vivo.
- 10 58. The method of any one of claims 32 to 34, wherein myocardial cells or tissue is imaged.
59. The method of any one of claims 32 to 58, wherein the radiolabeled salt substantially localizes to a site or sites having dysfunctional mitochondria within about 15 120 minutes after administration.
60. The method of claims 32 to 58, wherein the radiolabeled salt substantially localizes to a site or sites dysfunctional mitochondria within about 60 minutes after administration.
- 20 61. The method of claims 32 to 58, wherein the radiolabeled salt substantially localizes to a site or sites dysfunctional mitochondria within about 30 minutes after administration.
- 25 62. The method of any one of claims 32 to 61, wherein the radiolabeled salt is detected by a gamma camera, positron emission tomography (PET) or single photon emission tomography (SPECT).
- 30 63. The method of any one of claims 32 to 62, wherein the subject is a human, rat, mouse, cat, dog, horse, sheep, cow, monkey, avian, or amphibian.

64. An imaging method comprising:

- (a) administering a salt comprising at least one pharmaceutically acceptable anion and at least one cation according to the following Formula I to a patient suffering or susceptible to cancer, liver disease, degenerative disease or disorder, autoimmune disorder, HIV infection, or myopathy caused by oxidative stress or DNA mutation:



wherein

E is phosphorus or nitrogen; and

- X^1 , X^2 , X^3 , and X^4 are independently selected from the group consisting of Ar and R, wherein at least one of X^1 , X^2 , X^3 , and X^4 is an Ar group;

Ar is optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

- R is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted haloalkyl, optionally substituted cycloalkyl, optionally substituted aralkyl, wherein at least one occurrence of R comprises at least one radioisotope; and

(b) making a radiographic image.

65. The method of claim 64 wherein the patient has been selected for diagnosis or treatment of cancer, liver disease, degenerative disease or disorder, autoimmune disorder, HIV infection, or myopathy caused by oxidative stress or DNA mutation.

66. The method of claim 64 wherein the patient is selected for diagnosis or treatment of cancer and the compound is administered to the selected patient.

67. The method of claim 64 wherein the patient is selected for diagnosis or treatment of lung cancer, breast cancer, prostate cancer, liver disease, degenerative

disease or disorder, HIV infection, or myopathy caused by oxidative stress or DNA mutation and the compound is administered to the selected patient.

68. The method of any one of claims 64 to 67 wherein:

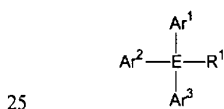
5 Ar is optionally substituted aryl having from 6 to 18 carbon atoms and between 1 and 3 rings, optionally substituted heteroaryl having from 3 to about 18 carbon atoms, between 1 and about 3 rings and between 1 and about 4 ring heteroatoms selected from N, O, and S, and optionally substituted aralkyl having between 7 and about 12 carbon atoms; and

10 R is optionally substituted C₁₋₆alkyl, optionally substituted C₂₋₆alkenyl, optionally substituted C₂₋₆alkynyl, optionally substituted C₁₋₆haloalkyl having at least one F, Cl, Br, or I atom, optionally substituted cycloalkyl having between 3 and about 8 ring carbon atoms, optionally substituted aralkyl having between 7 and about 12 carbon atoms, wherein at least one occurrence of R comprises at least one radioisotope.

15 69. The method of any one of claims 64 to 68 wherein E is phosphorus or nitrogen.

20 70. The method of any one of claims 64 to 69 wherein at least one R group comprises one or more ¹⁸F.

71. The method of any one of claims 64 to 67, wherein the cation is represented by the following formula:



wherein

E is phosphorus or nitrogen; and

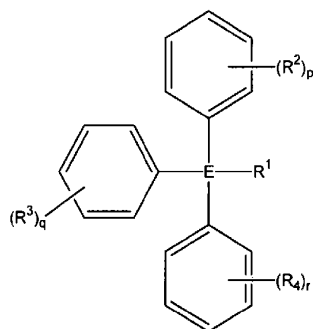
Ar^1 , Ar^2 , and Ar^3 are each independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

R^1 is alkyl, alkenyl, alkynyl, haloalkyl, cycloalkyl, or aralkyl, with R^1 being substituted by one or more groups comprising ^{18}F and optionally substituted by one or more additional groups.

72. The method of claim 71 wherein E is phosphorus.

73. The method of claim 71 or 72 wherein R^1 is selected from the group consisting of optionally substituted C_{1-6} alkyl, optionally substituted C_{7-12} aralkyl, optionally substituted C_{6-12} aryl, each of which is substituted with one or more ^{18}F .

74. The method of any one of claims 64 to 67, wherein the cation is represented by the following formula:



wherein

E is phosphorus or nitrogen;

p, q, and r are independently selected numbers from zero to about 5;

R¹ is alkyl, alkenyl, alkynyl, haloalkyl, cycloalkyl, or aralkyl, with R¹ being substituted by one or more ¹⁸F and optionally substituted by one or more additional groups;

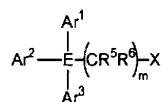
- 5 R², R³, and R⁴ are independently selected at each occurrence of R², R³, and R⁴ from the group consisting of hydrogen, halogen, cyano, nitro, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted alkoxy, optionally substituted (cycloalkyl)alkyl, optionally substituted alkylthio, optionally substituted alkylsulfinyl, or optionally substituted alkylsulfonyl, and optionally substituted mono or dialkylcarboxamide.

10

75. The method of claim 74 wherein E is phosphorus.

76. The method of any one of claims 64 to 67, wherein the cation is represented by the following formula:

15



wherein

E is phosphorus or nitrogen;

- 20 Ar¹, Ar², and Ar³ are independently selected from the group consisting of optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted aralkyl; and

R⁵ and R⁶ are independently selected at each occurrence of R⁵ and R⁶ from the group consisting of hydrogen, halogen, hydroxy, amino, optionally substituted alkyl, optionally substituted haloalkyl, and optionally substituted alkoxy;

- 25 X is ¹¹C-methyl or a radioisotope of fluorine or iodine; and
m is a number from about 2 to about 6.

77. The method of claim 76 wherein E is phosphorus.

78. The method of any one of claims 64 to 67 wherein the salt comprises:
¹¹C-methyl-triphenylphosphonium;
¹¹C-methyl-tri-*ortho*-tolylphosphonium;
¹¹C-methyl-tri-*meta*-tolylphosphonium ion; or
5 ¹¹C-methyl-tri-*para*-tolylphosphonium ion.
79. The method of any one of claims 64 to 78, wherein the salt is detected by positron emission tomography.
- 10 80. An imaging agent comprising at least one radiolabeled salt or compound of any one of claims 1 to 29 or a pharmaceutically composition of claim 30.
81. A salt comprising at least one pharmaceutically acceptable anion and at least one cation substantially as hereinbefore described with reference to any one of
15 Examples 1 to 7.
82. An imaging method substantially as hereinbefore described with reference to any one of Examples 8 and 13 to 19.
- 20 83. An imaging agent substantially as hereinbefore described with reference to any one of Examples 9 to 13.

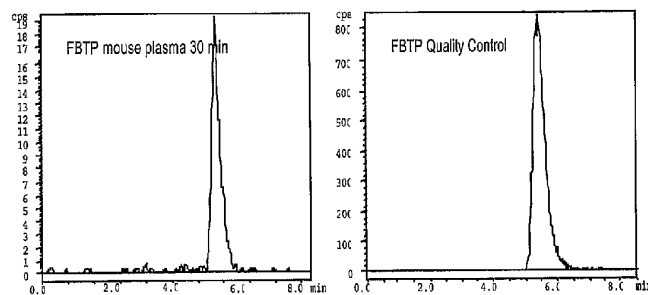


Figure 1 Radiochromatograms of ^{18}F -FBnTP (quality control) and plasma 30 min p.i. of the tracer into mice.

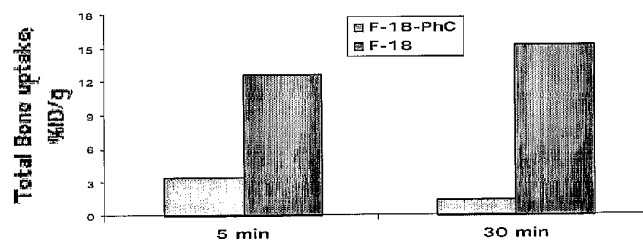


Figure 2: Bone uptake of F-18-FBnTP and F-18 in mice

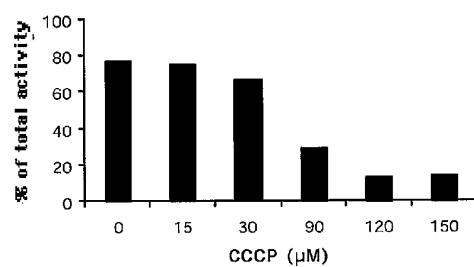


Figure 3. Mitochondrial membrane potential-dependent uptake of ^{18}F -FBnTP in vitro

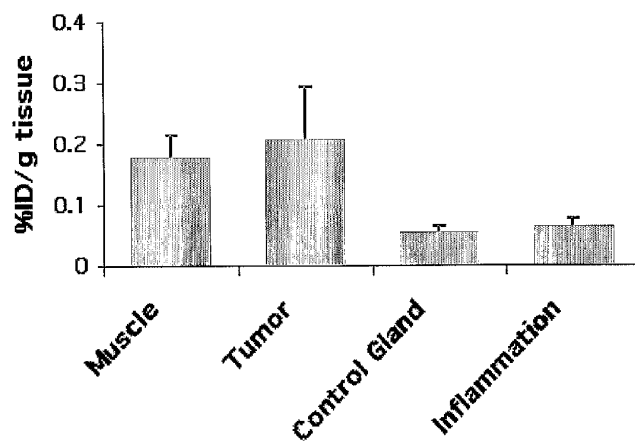


Figure 4. Accumulation of ^{18}F -FBnTP in malignant (Tumor) and healthy (control) mammary gland and in healthy (Muscle) and inflammation muscle, in rats.

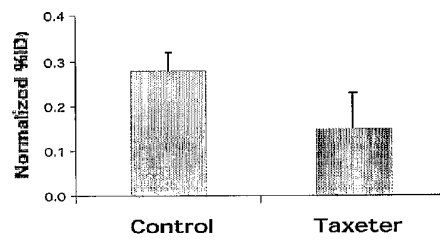


Figure 5: Cellular uptake of F-18-FBnTP in lung carcinoma A549 tumor.

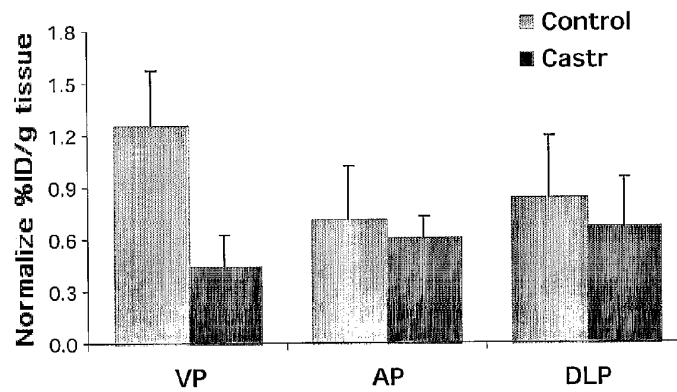


Figure 6: Accumulation of F-18-FBnTP in the prostate lobes 4 days after castration. Castration-produced apoptosis induced significant lobe-specific decrease in the ventral aspect, compared to control

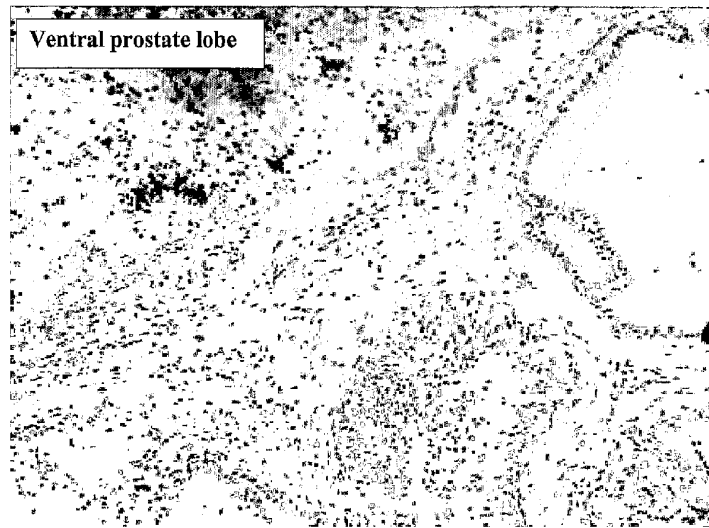


Figure 7

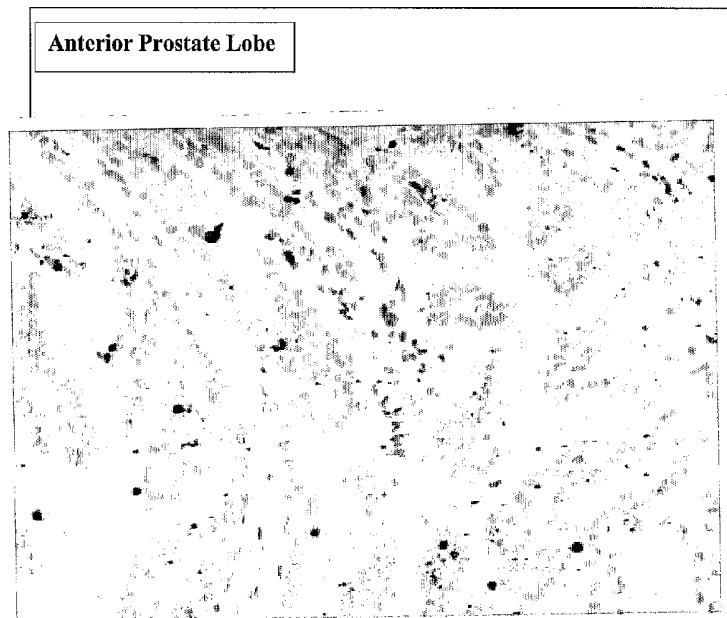


FIG 8

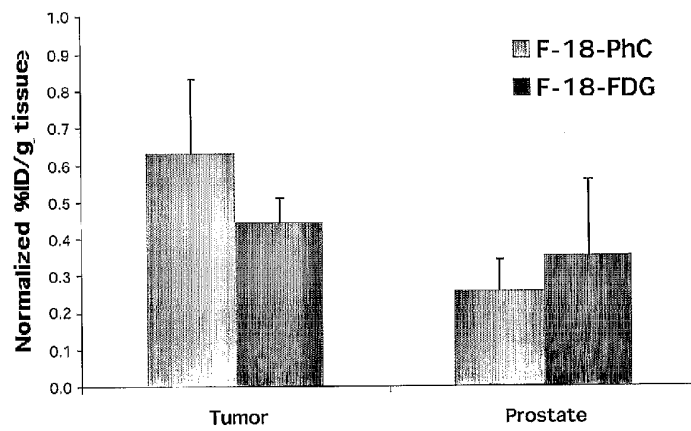


Figure 9: Accumulation of F-18FbnTP and F-18-FDG in prostate tumor in mice.

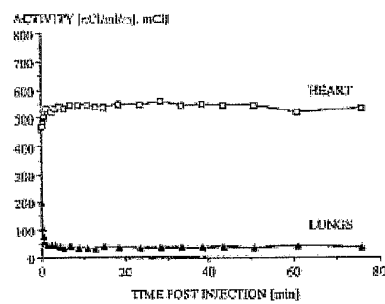


Figure 10: C-11TPMP myocardial uptake

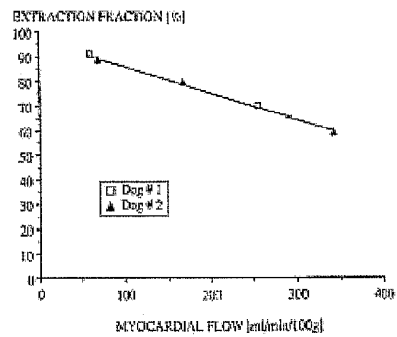


Figure 11: C-11-TPMP extraction fraction

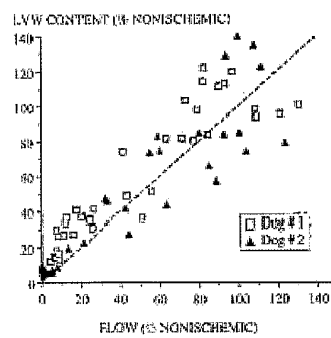


Figure 12: C-11-TPMP myocardial uptake as function of coronary blood flow

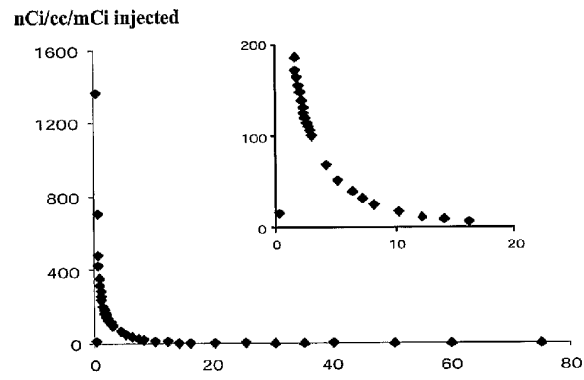


Figure 13: F-18-FBnTP rapid clearance from blood

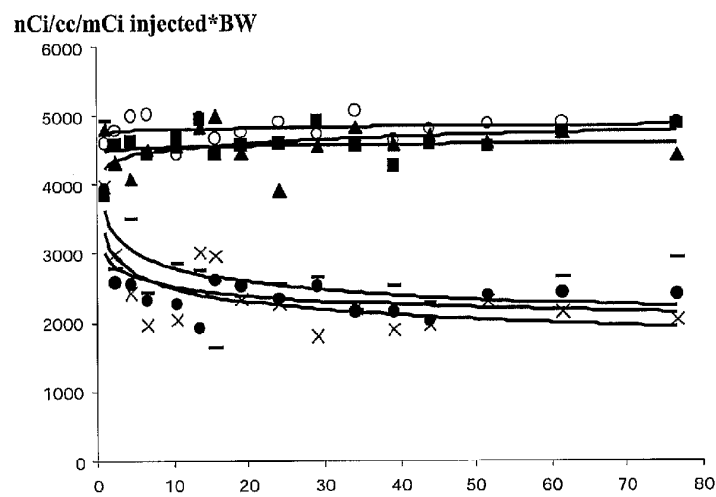
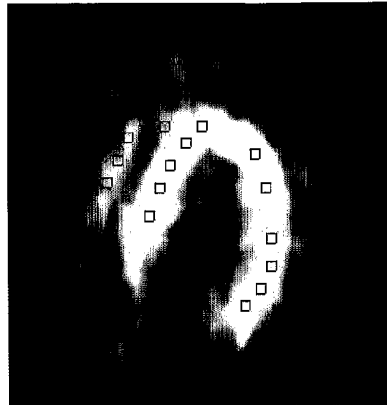
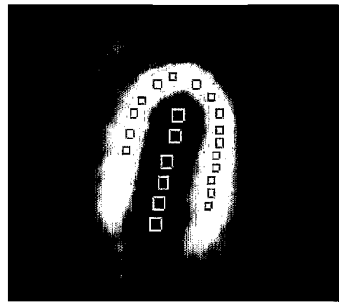


Figure 14: F-18-FBnTP accumulation kinetics in the left and right ventricles in 3 consecutive axial slices (8 to 10). Each trace is the mean of ROIs as above demonstrated. This figure is a good example for the rapid fact kinetics of FBnTP and uniform distribution in the myocardium.



nCi/cc/mCi injected * BW

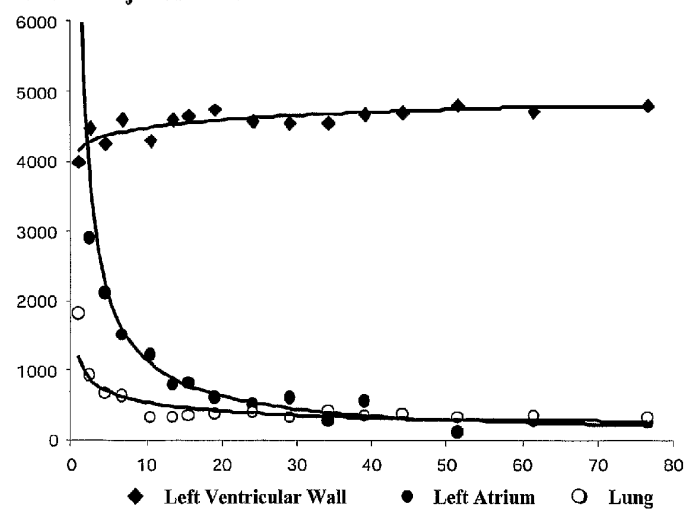


Figure 15: F-18-FBnTP uptake kinetics in the left ventricle and lung. ROIs for generating the left ventricle time-activity curve are presented on the corresponding slice. This figure is an excellent example of the preferential accumulation of FBnTP in the myocardium, compared to the adjacent lungs, resulting in the very high-quality images of the myocardium as presented in the figure below.

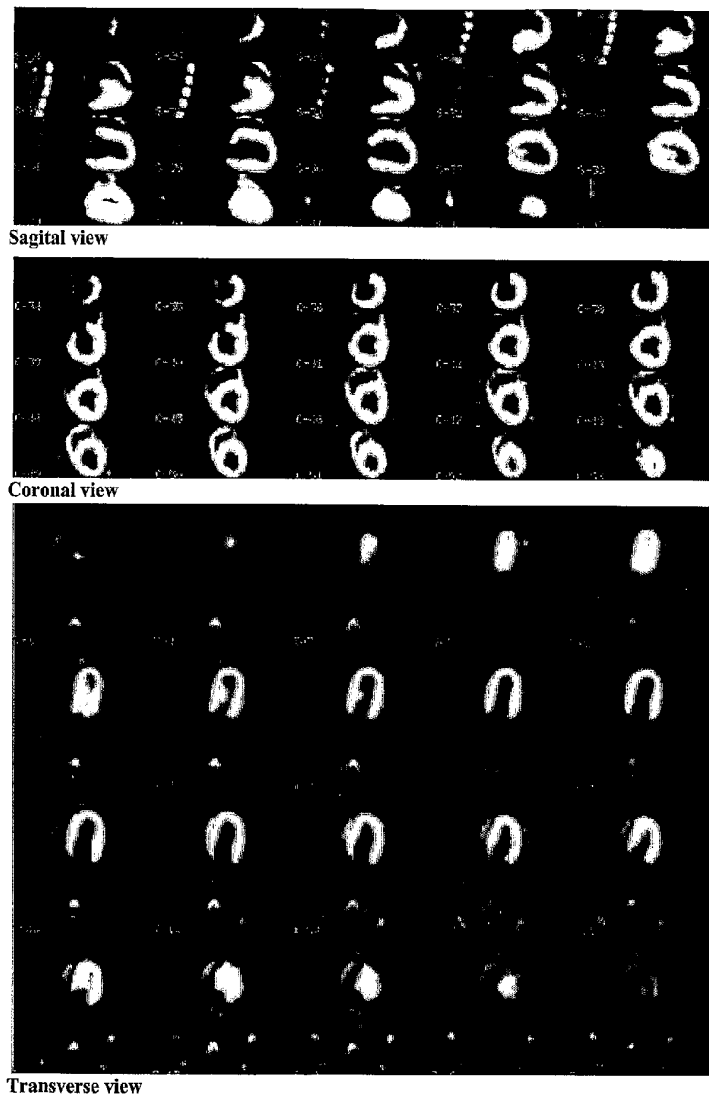


Figure 16: F-18-FBnTP myocardial accumulation in heart failure.

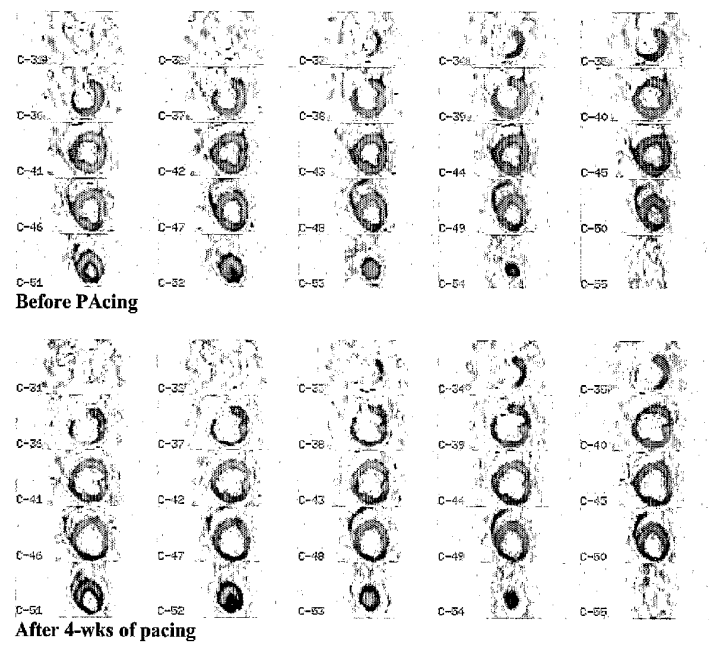


Figure 17: F-18-FBNTP PET short-axis images of the myocardium before and after pacing

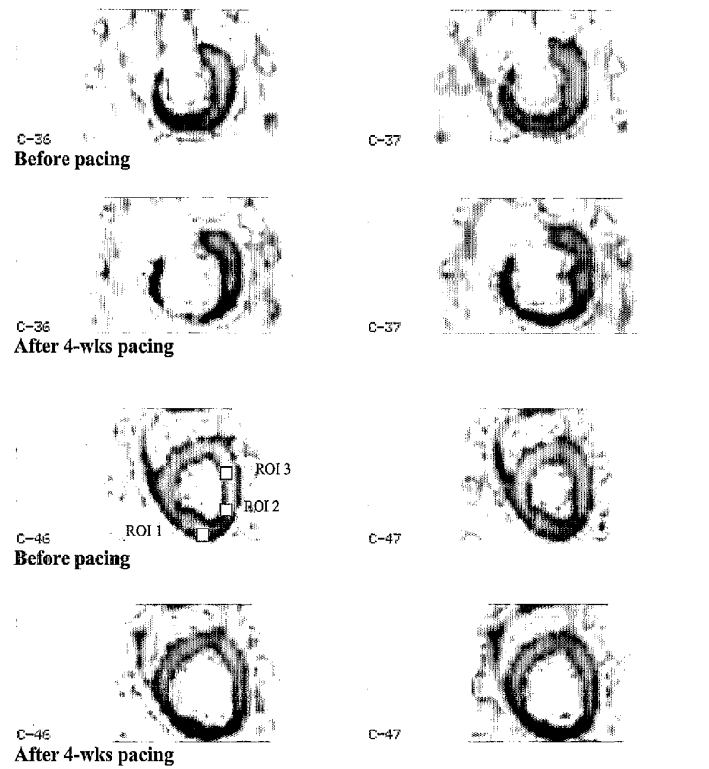


Figure 18: A decrease in F-18-FBnTp uptake in the inferior wall following pacing-induced heart failure.

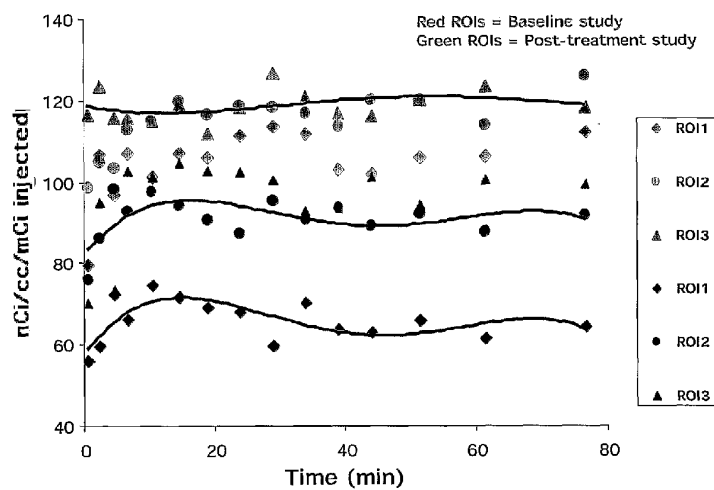


Figure 19: Time-activity course of [^{18}F]FBnTP myocardial uptake before (red) and after 4 weeks of pacing. Each data point represents the mean activity of four slices (slice 46 to 49). Positioning of region of interest (ROI) is shown in the left upper cardiac image. The same ROIs template was used to retrieve data in all cardiac sections.